

Medicare Part D: 5 Tier Closed Venture Formulary 2022

Please click here.

For Medicare Part D: Prior Authorization Criteria

Please click here.

For Medicare Part D: Step Therapy Criteria

Please click here.

For more recent information or other questions, please contact:

Freedom Blue PPO (PA) Customer Service at 1-800-550-8722

Security Blue HMO-POS Customer Service at 1-800-935-2583

Community Blue Medicare HMO Customer Service at 1-888-234-5397

Blue Rx PDP Customer Service at 1-800-290-3914

For TTY users, *711 National Relay Service*, Monday through Sunday, 8:00 a.m. to 8:00 p.m.

Visit medicare.highmark.com.

Formulary ID: 22037 Version: 25

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Note to existing members: This formulary has changed since last year. Please review this document to make sure that it still contains the drugs you take.

When this drug list (formulary) refers to “we,” “us,” or “our,” it means Highmark Senior Health Company, Highmark Choice Company, Highmark Senior Solutions Company or HM Health Insurance Company.

When it refers to “plan” or “our plan,” it means Freedom Blue PPO, Security Blue HMO-POS, Community Blue Medicare HMO, or Blue Rx PDP.

This document includes a list of the drugs (formulary) for our plan which is current as of July 1, 2022. For an updated formulary, please contact us. Our contact information, along with the date we last updated the formulary, appears on the front and back cover pages.

You must generally use network pharmacies to use your prescription drug benefit. Benefits, formulary, pharmacy network, and/or copayments/coinsurance may change on January 1, 2022, and from time to time during the year.

What is the Freedom Blue PPO, Security Blue HMO-POS, Community Blue Medicare HMO, and Blue Rx PDP Formulary?

A formulary is a list of covered drugs selected by our plans in consultation with a team of health care providers, which represents the prescription therapies believed to be a necessary part of a quality treatment program. Our plans will generally cover the drugs listed in our formulary as long as the drug is medically necessary, the prescription is filled at one of our plan’s network pharmacies, and other plan rules are followed. For more information on how to fill your prescriptions, please review your Evidence of Coverage.

Can the Freedom Blue PPO, Security Blue HMO-POS, Community Blue Medicare HMO, and Blue Rx PDP Formulary (drug list) change?

Most changes in drug coverage happen on January 1, but we may add or remove drugs on the Drug List during the year, move them to different cost-sharing tiers, or add new restrictions. We must follow Medicare rules in making these changes.

Changes that can affect you this year: In the below cases, you will be affected by coverage changes during the year:

- **New generic drugs.** We may immediately remove a brand name drug on our Drug List if we are replacing it with a new generic drug that will appear on the same or lower cost sharing tier and with the same or fewer restrictions. Also, when adding the new generic drug, we may decide to keep the brand name drug on our Drug List, but immediately move it to a different cost-sharing tier or add new restrictions. If you are currently taking that brand name drug, we may not tell you in advance before we make that change, but we will later provide you with information about the specific change(s) we have made.

- o If we make such a change, you or your prescriber can ask us to make an exception and continue to cover the brand name drug for you. The notice we provide you will also include information on how to request an exception, and you can also find information in the section below entitled “How do I request an exception to the Freedom Blue PPO, Security Blue HMO-POS, Community Blue Medicare HMO, or Blue Rx PDP Formulary?”
- **Drugs removed from the market.** If the Food and Drug Administration deems a drug on our formulary to be unsafe or the drug’s manufacturer removes the drug from the market, we will immediately remove the drug from our formulary and provide notice to members who take the drug.
- **Other changes.** We may make other changes that affect members currently taking a drug. For instance, we may add a generic drug that is not new to market to replace a brand name drug currently on the formulary or add new restrictions to the brand name drug or move it to different cost sharing tier or both. Or we may make changes based on new clinical guidelines. If we remove drugs from our formulary, add prior authorization, quantity limits and/or step therapy restrictions on a drug, or move a drug to a higher cost-sharing tier, we must notify affected members of the change at least 30 days before the change becomes effective, or at the time the member requests a refill of the drug, at which time the member will receive a 31-day supply of the drug.
 - o If we make these other changes, you or your prescriber can ask us to make an exception and continue to cover the brand name drug for you. The notice we provide you will also include information on how to request an exception, and you can also find information in the section below entitled “How do I request an exception to the Freedom Blue PPO, Security Blue HMO-POS, Community Blue Medicare HMO, or Blue Rx PDP Formulary?”

Changes that will not affect you if you are currently taking the drug. Generally, if you are taking a drug on our 2022 formulary that was covered at the beginning of the year, we will not discontinue or reduce coverage of the drug during the 2022 coverage year except as described above. This means these drugs will remain available at the same cost-sharing and with no new restrictions for those members taking them for the remainder of the coverage year. You will not get direct notice this year about changes that do not affect you. However, on January 1 of the next year, such changes would affect you, and it is important to check the Drug List for the new benefit year for any changes to drugs.

The enclosed formulary is current as of July 1, 2022. To get updated information about the drugs covered by our plans, please contact us. Our contact information appears on the front and back cover pages. In the event of mid-year non-maintenance formulary changes, members will be notified by mail and prospective members will receive an update with this formulary. The most up-to-date formulary is available on our website, [medicare.highmark.com](https://www.medicare.highmark.com).

How do I use the Freedom Blue PPO, Security Blue HMO-POS, Community Blue Medicare HMO, and Blue Rx PDP Formulary?

There are two ways to find your drug within the formulary:

Medical Condition

The formulary begins on page 8. The drugs in this formulary are grouped into categories depending on the type of medical conditions that they are used to treat. For example, drugs used to treat a heart condition are listed under the category, “Cardiovascular – Hypertension & Lipids.” If you know what your drug is used for, look for the category name in the list that begins on page number 8. Then look under the category name for your drug.

Alphabetical Listing

If you are not sure what category to look under, you should look for your drug in the Index that begins at the end of this document. The Index provides an alphabetical list of all of the drugs included in this document. Both brand name drugs and generic drugs are listed in the Index. Look in the Index and find your drug. Next to your drug, you will see the page number where you can find coverage information. Turn to the page listed in the Index and find the name of your drug in the first column of the list.

What are generic drugs?

Our plans cover both brand name drugs and generic drugs. A generic drug is approved by the FDA as having the same active ingredient as the brand name drug. Generally, generic drugs cost less than brand name drugs.

Are there any restrictions on my coverage?

Some covered drugs may have additional requirements or limits on coverage. These requirements and limits may include:

- **Prior Authorization:** Our plan requires you or your physician to get prior authorization for certain drugs. This means that you will need to get approval from our plans before you fill your prescriptions. If you don't get approval, our plans may not cover the drug.
- **Quantity Limits:** For certain drugs, our plans limit the amount of the drug covered. For example, our plans provide tablets per prescription for 100mg Imitrex. This may be in addition to a standard one-month or three-month supply.
- **Step Therapy:** In some cases, our plans require you to first try certain drugs to treat your medical condition before we will cover another drug for that condition. For example, if Drug A and Drug B both treat your medical condition, our plans may not cover Drug B unless you try Drug A first. If Drug A does not work for you, our plans will then cover Drug B.

You can find out if your drug has any additional requirements or limits by looking in the formulary that begins on page 8. You can also get more information about the restrictions applied to specific covered drugs by visiting our website. We have posted online document(s) that explain(s) our prior authorization and step therapy restrictions. You may also ask us to send you a copy. Our contact information, along with the date we last updated the formulary, appears on the front and back cover pages.

You can ask our plans to make an exception to these restrictions or limits or for a list of other, similar drugs that may treat your health condition. See the section, “How do I request an exception to the Freedom Blue PPO, Security Blue HMO-POS, Community Blue Medicare HMO, or Blue Rx PDP Formulary?” on page 5 for information about how to request an exception.

What if my drug is not on the Freedom Blue PPO, Security Blue HMO-POS, Community Blue Medicare HMO, and Blue Rx PDP Formulary?

If your drug is not included in this formulary (list of covered drugs), you should first contact Customer Service and ask if your drug is covered.

If you learn that our plan does not cover your drug, you have two options:

- You can ask Customer Service for a list of similar drugs that are covered by our plans. When you receive the list, show it to your doctor and ask your doctor to prescribe a similar drug that is covered by our plan.
- You can ask our plan to make an exception and cover your drug. See below for information about how to request an exception.

How do I request an exception to the Freedom Blue PPO, Security Blue HMO-POS, Community Blue Medicare HMO, or Blue Rx PDP Formulary?

You can ask our plan to make an exception to our coverage rules. There are several types of exceptions that you can ask us to make.

- You can ask us to cover a drug even if it is not on our formulary. If approved, this drug will be covered at a pre-determined cost-sharing level, and you would not be able to ask us to provide the drug at a lower cost-sharing level.
- You can ask us to cover a formulary drug at a lower cost-sharing level if this drug is not on the specialty tier. If approved, this would lower the amount you must pay for your drug.
- You can ask us to waive coverage restrictions or limits on your drug. For example, for certain drugs, our plans limit the amount of the drug that we will cover. If your drug has a quantity limit, you can ask us to waive the limit and cover a greater amount.

Generally, our plan will only approve your request for an exception if the alternative drugs included on the plan's formulary, the lower cost-sharing drug or additional utilization restrictions, would not be as effective in treating your condition and/or would cause you to have adverse medical effects.

You should contact us to ask us for an initial coverage decision for a formulary, or utilization restriction exception. **When you request a formulary or utilization restriction exception, you should submit a statement from your prescriber or physician supporting your request.** Generally, we must make our decision within 72 hours of getting your prescriber's supporting statement. You can request an expedited (fast) exception if you or your doctor believe that your health could be seriously harmed by waiting up to 72 hours for a decision. If your request to expedite is granted, we must give you a decision no later than 24 hours after we get a supporting statement from your doctor or other prescriber.

What do I do before I can talk to my doctor about changing my drugs or requesting an exception?

As a new or continuing member in our plan, you may be taking drugs that are not on our formulary. Or you may be taking a drug that is on our formulary but your ability to get it is limited. For example, you may need a prior authorization from us before you can fill your prescription. You should talk to your doctor to decide if you should switch to an appropriate drug that we cover or request a formulary exception so that we will cover the drug you take. While you talk to your doctor to determine the right course of action for you, we may cover your drug in certain cases during the first 90 days you are a member of our plan.

For each of your drugs that is not on our formulary or if your ability to get your drugs is limited, we will cover a temporary 31-day supply. If your prescription is written for fewer days, we'll allow refills to provide up to a maximum 31-day supply of medication. After your first 31-day supply, we will not pay for these drugs, even if you have been a member of the plan less than 90 days.

If you are a resident of a long-term care facility and you need a drug that is not on our formulary or if your ability to get your drugs is limited, but you are past the first 90 days of membership in our plan, we will cover a 31-day emergency supply of that drug while you pursue a formulary exception.

The above transition process will be implemented to accommodate you if you have an immediate need for a non-formulary drug or a drug that requires prior authorization due to a change in your level of care while you are waiting for an exception request to be processed.

For more information

For more detailed information about your plan's prescription drug coverage, please review your Evidence of Coverage and other plan materials.

If you have questions about your plan, please contact us. Our contact information, along with the date we last updated the formulary, appears on the front and back cover pages.

If you have general questions about Medicare prescription drug coverage, please call Medicare at 1-800-MEDICARE (1-800-633-4227) 24 hours a day / 7 days a week. TTY users should call 1-877-486-2048. Or visit <http://www.medicare.gov>.

Freedom Blue PPO, Security Blue HMO-POS, Community Blue Medicare HMO, or Blue Rx PDP Formulary

The formulary that begins on the next page provides coverage information about the drugs covered by your plan. If you have trouble finding your drug in the list, turn to the Index that begins at the end of this document.

The first column of the chart lists the drug name. Brand name drugs are capitalized (e.g., ABELCET) and generic drugs are listed in lowercase italics (e.g., *abacavir*).

The information in the Requirements/Limits column tells you if Freedom Blue PPO, Security Blue HMO-POS, Community Blue Medicare HMO, or Blue Rx PDP has any special requirements for coverage of your drug.

The following is a Formulary Format Example Only:

Drug Name	Venture Drug Tier	Requirements/ Limits
Anti - Infectives		
<i>XYZ DRUG</i>	T5	QL- 28

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List of Abbreviations

T1: Cost-Sharing Tier 1 includes preferred generic drugs. This is the lowest cost-sharing tier.

T2: Cost-Sharing Tier 2 includes generic drugs.

T3: Cost-Sharing Tier 3 includes preferred brand name drugs and may include some single-sourced drugs (those generic drugs made by a single manufacturer).

T4: Cost-Sharing Tier 4 includes non-preferred brand name drugs and may include some single-sourced generic drugs (those generic drugs made by a single manufacturer).

T5: Cost-Sharing Tier 5 includes specialty drugs. This is the highest cost-sharing tier.

LA: Limited access

PA: Prior authorization required

PA-BvD: This drug may be covered under Medicare part B or D depending on the circumstance. Information may need to be submitted describing the use and setting of the drug to make the determination.

PA-NS: Prior authorization required for new starts only

QL: Quantity limit applies. The quantity limit is noted for each drug. For example, if the quantity limit is QL (90 EA per 180 days), the quantity limit would be 90 units per 180-day supply.

QL: Quantity limit applies. The quantity limit is noted for each drug. For example, if the quantity limit is QL (90 EA per 180 days), the quantity limit would be 90 units per 180-day supply.

SI: Select Insulin Drug under the Part D Senior Savings Model. For complete coverage details on these preferred select insulins see Chapter 6 of your Evidence of Coverage.

ST: Step therapy applies

ST-NS: Step therapy applies to new starts only

Below is a list of drug name formatting patterns that may appear in the following pages.

List of Patterns

lowercase italics: Generic drugs

UPPERCASE BOLD: Brand name drugs

Drug Name	Drug Tier	Requirements/Limits
Anti - Infectives		
<i>abacavir oral solution</i>	T4	
<i>abacavir oral tablet</i>	T3	
<i>abacavir-lamivudine</i>	T4	
ABELCET	T4	PA-BvD
<i>acyclovir oral capsule</i>	T2	
<i>acyclovir oral suspension 200 mg/5 ml</i>	T2	
<i>acyclovir oral tablet</i>	T2	
<i>acyclovir sodium intravenous solution</i>	T2	PA-BvD
<i>adefovir</i>	T4	
<i>albendazole</i>	T4	
<i>amantadine hcl oral capsule</i>	T2	QL (124 EA per 31 days)
<i>amantadine hcl oral solution</i>	T2	
<i>amantadine hcl oral tablet</i>	T2	
AMBISOME	T5	PA-BvD
<i>amikacin injection solution 500 mg/2 ml</i>	T2	
<i>amoxicillin oral capsule</i>	T1	
<i>amoxicillin oral suspension for reconstitution</i>	T1	
<i>amoxicillin oral tablet</i>	T1	
<i>amoxicillin oral tablet, chewable 125 mg, 250 mg</i>	T1	
<i>amoxicillin-pot clavulanate oral suspension for reconstitution</i>	T2	
<i>amoxicillin-pot clavulanate oral tablet</i>	T2	
<i>amoxicillin-pot clavulanate oral tablet, chewable</i>	T2	
<i>amphotericin b</i>	T4	PA-BvD
<i>ampicillin oral capsule 500 mg</i>	T2	
<i>ampicillin sodium injection recon soln 1 gram, 10 gram, 125 mg</i>	T2	
<i>ampicillin-sulbactam injection</i>	T2	
APTIVUS	T5	
ARIKAYCE	T5	PA
<i>atazanavir</i>	T4	
<i>atovaquone</i>	T4	
<i>atovaquone-proguanil</i>	T2	
AVYCAZ	T5	
<i>azithromycin intravenous</i>	T2	
<i>azithromycin oral packet</i>	T2	
<i>azithromycin oral tablet</i>	T2	
<i>aztreonam</i>	T4	

Drug Name	Drug Tier	Requirements/Limits
BARACLUDE ORAL SOLUTION	T3	
BETHKIS	T4	PA
BICILLIN C-R	T3	
BICILLIN L-A INTRAMUSCULAR SYRINGE 600,000 UNIT/ML	T3	
BIKTARVY	T5	QL (31 EA per 31 days)
<i>caspofungin intravenous recon soln 50 mg</i>	T5	
<i>caspofungin intravenous recon soln 70 mg</i>	T4	
CAYSTON	T5	
<i>cefaclor oral capsule 500 mg</i>	T2	
<i>cefaclor oral tablet extended release 12 hr</i>	T2	
<i>cefadroxil oral capsule</i>	T2	
<i>cefadroxil oral suspension for reconstitution 250 mg/5 ml</i>	T2	
<i>cefadroxil oral suspension for reconstitution 500 mg/5 ml</i>	T3	
<i>cefadroxil oral tablet</i>	T3	
<i>cefazolin injection recon soln 1 gram, 10 gram, 500 mg</i>	T2	
<i>cefdinir oral capsule</i>	T2	
<i>cefepime injection</i>	T4	
<i>cefixime</i>	T2	
<i>cefotetan injection</i>	T2	
<i>cefoxitin</i>	T2	
<i>cefpodoxime</i>	T2	
<i>cefprozil</i>	T2	
<i>ceftazidime</i>	T4	
<i>ceftriaxone injection recon soln 1 gram, 10 gram, 2 gram, 250 mg, 500 mg</i>	T2	
<i>cefuroxime axetil oral tablet</i>	T2	
<i>cefuroxime sodium injection recon soln 750 mg</i>	T2	
<i>cefuroxime sodium intravenous recon soln 1.5 gram</i>	T2	
<i>cephalexin</i>	T2	
<i>chloroquine phosphate oral tablet 250 mg</i>	T2	PA; QL (50 EA per 30 days)
<i>chloroquine phosphate oral tablet 500 mg</i>	T2	PA; QL (25 EA per 30 days)
CIMDUO	T5	QL (31 EA per 31 days)
<i>ciprofloxacin hcl oral</i>	T1	
<i>ciprofloxacin in 5 % dextrose intravenous piggyback 200 mg/100 ml</i>	T2	

Drug Name	Drug Tier	Requirements/Limits
<i>clarithromycin</i>	T2	
<i>clindamycin hcl</i>	T2	
<i>clindamycin in 5 % dextrose</i>	T2	
CLINDAMYCIN PEDIATRIC	T2	
<i>clindamycin phosphate injection</i>	T2	
<i>clindamycin phosphate intravenous</i>	T2	
<i>clotrimazole mucous membrane</i>	T2	
COARTEM	T4	
<i>colistin (colistimethate na)</i>	T4	
COMPLERA	T5	
CRESEMBA ORAL	T5	
DALVANCE	T5	
<i>dapsone oral</i>	T3	
<i>daptomycin</i>	T5	
DELSTRIGO	T5	QL (31 EA per 31 days)
<i>demeclocycline</i>	T2	
DESCOVY ORAL TABLET 200-25 MG	T5	QL (31 EA per 31 days)
<i>dicloxacillin</i>	T2	
DIFICID ORAL SUSPENSION FOR RECONSTITUTION	T5	QL (136 ML per 12 days)
DIFICID ORAL TABLET	T5	QL (20 EA per 10 days)
DOVATO	T5	QL (31 EA per 31 days)
DOXY-100	T2	
<i>doxycycline hyclate oral capsule</i>	T2	
<i>doxycycline hyclate oral tablet 100 mg</i>	T2	
<i>doxycycline hyclate oral tablet 150 mg, 50 mg, 75 mg</i>	T4	
<i>doxycycline hyclate oral tablet 20 mg</i>	T1	
<i>doxycycline hyclate oral tablet, delayed release (dr/ec) 100 mg, 200 mg, 50 mg</i>	T2	
<i>doxycycline hyclate oral tablet, delayed release (dr/ec) 150 mg, 75 mg</i>	T1	
<i>doxycycline hyclate oral tablet, delayed release (dr/ec) 80 mg</i>	T4	
<i>doxycycline monohydrate oral capsule 100 mg, 50 mg</i>	T2	
<i>doxycycline monohydrate oral capsule 150 mg, 75 mg</i>	T4	
<i>doxycycline monohydrate oral suspension for reconstitution</i>	T2	

Drug Name	Drug Tier	Requirements/Limits
<i>doxycycline monohydrate oral tablet</i>	T2	
E.E.S. 400 ORAL TABLET	T4	
EDURANT	T5	
<i>efavirenz</i>	T3	
<i>efavirenz-emtricitabin-tenofof</i>	T5	
<i>efavirenz-lamivu-tenofof disop</i>	T5	QL (31 EA per 31 days)
<i>emtricitabine</i>	T3	
<i>emtricitabine-tenofovir (tdf)</i>	T5	
EMTRIVA ORAL SOLUTION	T3	
EMVERM	T5	
<i>entecavir</i>	T4	
EPCLUSA ORAL PELLETS IN PACKET 150-37.5 MG	T5	PA; QL (28 EA per 28 days)
EPCLUSA ORAL PELLETS IN PACKET 200-50 MG	T5	PA; QL (56 EA per 28 days)
EPCLUSA ORAL TABLET	T5	PA; QL (28 EA per 28 days)
EPIVIR HBV ORAL SOLUTION	T3	
ERAXIS(WATER DILUENT)	T4	
<i>ertapenem</i>	T4	
ERYPED 200	T4	
ERY-TAB ORAL TABLET,DELAYED RELEASE (DR/EC) 250 MG, 333 MG	T2	
ERYTHROCIN (AS STEARATE) ORAL TABLET 250 MG	T2	
ERYTHROCIN INTRAVENOUS RECON SOLN 500 MG	T3	
<i>erythromycin ethylsuccinate oral suspension for reconstitution 200 mg/5 ml</i>	T4	
<i>erythromycin ethylsuccinate oral tablet</i>	T4	
<i>erythromycin oral tablet</i>	T2	
<i>ethambutol</i>	T2	
<i>etravirine oral tablet 100 mg</i>	T4	
<i>etravirine oral tablet 200 mg</i>	T5	
EVOTAZ	T3	
<i>famciclovir</i>	T2	
FIRVANQ	T4	
<i>fluconazole</i>	T2	
<i>fluconazole in nacl (iso-osm) intravenous piggyback 200 mg/100 ml, 400 mg/200 ml</i>	T2	
<i>flucytosine</i>	T4	

Drug Name	Drug Tier	Requirements/Limits
<i>fosamprenavir</i>	T4	
<i>fosfomycin tromethamine</i>	T4	
FUZEON SUBCUTANEOUS RECON SOLN	T5	
<i>gentamicin in nacl (iso-osm) intravenous piggyback 100 mg/100 ml, 60 mg/50 ml, 80 mg/100 ml, 80 mg/50 ml</i>	T2	
<i>gentamicin injection solution 40 mg/ml</i>	T1	
GENVOYA	T5	
<i>griseofulvin microsize oral suspension</i>	T2	
<i>griseofulvin microsize oral tablet</i>	T4	
<i>griseofulvin ultramicrosize</i>	T4	
HARVONI ORAL PELLETS IN PACKET	T5	PA; QL (28 EA per 28 days)
HARVONI ORAL TABLET 90-400 MG	T5	PA; QL (28 EA per 28 days)
<i>hydroxychloroquine oral tablet 100 mg</i>	T4	QL (93 EA per 31 days)
<i>hydroxychloroquine oral tablet 200 mg</i>	T2	QL (93 EA per 31 days)
<i>hydroxychloroquine oral tablet 300 mg</i>	T4	QL (62 EA per 31 days)
<i>hydroxychloroquine oral tablet 400 mg</i>	T4	QL (31 EA per 31 days)
<i>imipenem-cilastatin</i>	T2	
IMPAVIDO	T5	
INTELENCE ORAL TABLET 25 MG	T4	
ISENTRESS	T3	
ISENTRESS HD	T5	
<i>isoniazid oral solution</i>	T2	
<i>isoniazid oral tablet</i>	T1	
<i>itraconazole oral capsule</i>	T2	PA
<i>itraconazole oral solution</i>	T4	PA
<i>ivermectin oral</i>	T2	
JULUCA	T5	
<i>ketoconazole oral</i>	T2	
<i>lamivudine</i>	T2	
<i>lamivudine-zidovudine</i>	T2	
<i>ledipasvir-sofosbuvir</i>	T5	PA; QL (28 EA per 28 days)
<i>levofloxacin in d5w intravenous piggyback 500 mg/100 ml, 750 mg/150 ml</i>	T2	
<i>levofloxacin intravenous</i>	T3	
<i>levofloxacin oral solution</i>	T3	
<i>levofloxacin oral tablet</i>	T2	
LEXIVA ORAL SUSPENSION	T3	
<i>linezolid</i>	T4	

Drug Name	Drug Tier	Requirements/Limits
<i>linezolid in dextrose 5%</i>	T4	
LIVTENCITY	T5	PA; QL (124 EA per 31 days)
<i>lopinavir-ritonavir oral solution</i>	T4	
<i>lopinavir-ritonavir oral tablet 100-25 mg</i>	T3	
<i>lopinavir-ritonavir oral tablet 200-50 mg</i>	T5	
<i>maraviroc</i>	T5	
MAVYRET ORAL PELLETS IN PACKET	T5	PA; QL (140 EA per 28 days)
MAVYRET ORAL TABLET	T5	PA; QL (84 EA per 28 days)
<i>mefloquine</i>	T2	
<i>meropenem</i>	T4	
<i>methenamine hippurate</i>	T2	
<i>metronidazole in nacl (iso-os)</i>	T2	
<i>metronidazole oral tablet</i>	T1	
<i>micafungin</i>	T5	
<i>minocycline oral capsule</i>	T2	
<i>minocycline oral tablet</i>	T2	
<i>minocycline oral tablet extended release 24 hr 105 mg, 115 mg, 135 mg, 45 mg, 65 mg, 80 mg, 90 mg</i>	T4	
<i>moxifloxacin oral</i>	T3	
<i>moxifloxacin-sod.chloride(iso)</i>	T4	
<i>nafcillin injection</i>	T2	
<i>neomycin</i>	T2	
<i>nevirapine</i>	T2	
<i>nitazoxanide</i>	T4	
<i>nitrofurantoin macrocrystal oral capsule 100 mg</i>	T2	QL (90 EA per 365 days)
<i>nitrofurantoin macrocrystal oral capsule 25 mg</i>	T2	QL (360 EA per 365 days)
<i>nitrofurantoin macrocrystal oral capsule 50 mg</i>	T2	QL (180 EA per 365 days)
<i>nitrofurantoin monohyd/m-cryst</i>	T2	QL (90 EA per 365 days)
NORVIR ORAL POWDER IN PACKET	T4	
NORVIR ORAL SOLUTION	T4	
NUZYRA	T5	
<i>nystatin oral</i>	T2	
ODEFSEY	T5	QL (31 EA per 31 days)
<i>ofloxacin oral tablet 300 mg, 400 mg</i>	T2	
<i>oseltamivir oral capsule 30 mg</i>	T2	QL (170 EA per 365 days)
<i>oseltamivir oral capsule 45 mg, 75 mg</i>	T2	QL (90 EA per 365 days)
<i>oseltamivir oral suspension for reconstitution</i>	T3	QL (1080 ML per 365 days)

Drug Name	Drug Tier	Requirements/Limits
<i>oxacillin in dextrose(iso-osm) intravenous piggyback 1 gram/50 ml</i>	T2	
<i>oxacillin injection recon soln 1 gram, 2 gram</i>	T2	
<i>oxacillin injection recon soln 10 gram</i>	T4	
<i>paromomycin</i>	T2	
PASER	T4	
<i>penicillin g pot in dextrose intravenous piggyback 2 million unit/50 ml, 3 million unit/50 ml</i>	T4	
<i>penicillin g potassium injection recon soln 20 million unit</i>	T2	
<i>penicillin g procaine intramuscular syringe 1.2 million unit/2 ml</i>	T2	
<i>penicillin g sodium</i>	T4	
<i>penicillin v potassium</i>	T1	
<i>pentamidine inhalation</i>	T4	PA-BvD; QL (1 EA per 28 days)
<i>pentamidine injection</i>	T4	
PIFELTRO	T5	QL (62 EA per 31 days)
<i>piperacillin-tazobactam intravenous recon soln 2.25 gram, 3.375 gram, 4.5 gram, 40.5 gram</i>	T4	
<i>polymyxin b sulfate</i>	T2	
<i>praziquantel</i>	T4	
PREVYMIS ORAL TABLET 240 MG	T4	
PREVYMIS ORAL TABLET 480 MG	T5	
PREZCOBIX	T5	
PREZISTA ORAL SUSPENSION	T5	
PREZISTA ORAL TABLET 150 MG, 600 MG, 800 MG	T5	
PREZISTA ORAL TABLET 75 MG	T3	
PRIFTIN	T4	
<i>primaquine</i>	T3	
<i>pyrazinamide</i>	T4	
<i>pyrimethamine</i>	T5	PA
<i>quinine sulfate</i>	T3	PA; QL (42 EA per 28 days)
RELENZA DISKHALER	T3	
REYATAZ ORAL POWDER IN PACKET	T4	
<i>ribavirin oral capsule</i>	T2	
<i>ribavirin oral tablet 200 mg</i>	T2	
<i>rifabutin</i>	T4	
<i>rifampin</i>	T2	
<i>rimantadine</i>	T2	

Drug Name	Drug Tier	Requirements/Limits
<i>ritonavir</i>	T3	
RUKOBIA	T5	QL (62 EA per 31 days)
SELZENTRY ORAL SOLUTION	T5	
SELZENTRY ORAL TABLET 150 MG, 300 MG, 75 MG	T5	
SELZENTRY ORAL TABLET 25 MG	T4	
SIRTURO	T5	
SIVEXTRO INTRAVENOUS	T5	
SIVEXTRO ORAL	T5	QL (6 EA per 31 days)
<i>sofosbuvir-velpatasvir</i>	T5	PA; QL (28 EA per 28 days)
SOVALDI ORAL PELLETS IN PACKET	T5	PA; QL (28 EA per 28 days)
SOVALDI ORAL TABLET 400 MG	T5	PA; QL (28 EA per 28 days)
<i>streptomycin</i>	T3	
STRIBILD	T5	
<i>sulfadiazine</i>	T2	
<i>sulfamethoxazole-trimethoprim oral</i>	T1	
SUPRAX ORAL SUSPENSION FOR RECONSTITUTION 500 MG/5 ML	T3	
SYMFI	T5	QL (31 EA per 31 days)
SYMFI LO	T5	QL (31 EA per 31 days)
SYMTUZA	T5	QL (31 EA per 31 days)
TEFLARO	T5	
<i>tenofovir disoproxil fumarate</i>	T3	
<i>terbinafine hcl oral</i>	T1	QL (90 EA per 180 days)
<i>tetracycline</i>	T2	
<i>tigecycline</i>	T5	
TIVICAY ORAL TABLET 10 MG	T4	
TIVICAY ORAL TABLET 25 MG, 50 MG	T5	
TIVICAY PD	T4	
TOBI PODHALER INHALATION CAPSULE, W/INHALATION DEVICE	T3	PA; QL (224 EA per 56 days)
<i>tobramycin in 0.225 % nacl</i>	T5	PA
<i>tobramycin inhalation</i>	T4	PA
<i>tobramycin sulfate injection solution</i>	T1	
TOLSURA	T5	PA; QL (130 EA per 31 days)
TRECATOR	T4	
<i>trimethoprim</i>	T2	
TRIUMEQ	T5	
TRIUMEQ PD	T5	QL (186 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
TRIZIVIR	T5	
TYBOST	T3	
VABOMERE	T4	
<i>valacyclovir</i>	T2	
<i>valganciclovir oral recon soln</i>	T5	
<i>valganciclovir oral tablet</i>	T3	
<i>vancomycin intravenous recon soln 1,000 mg, 10 gram, 500 mg, 750 mg</i>	T4	
<i>vancomycin oral capsule 125 mg</i>	T4	QL (124 EA per 31 days)
<i>vancomycin oral capsule 250 mg</i>	T4	QL (248 EA per 31 days)
<i>vancomycin oral recon soln</i>	T4	
VEMLIDY	T5	QL (31 EA per 31 days)
VIRACEPT ORAL TABLET	T5	
VIREAD ORAL POWDER	T3	
VIREAD ORAL TABLET 150 MG, 200 MG, 250 MG	T3	
VIVJOA	T4	PA; QL (18 EA per 84 days)
<i>voriconazole intravenous</i>	T5	PA
<i>voriconazole oral suspension for reconstitution</i>	T5	
<i>voriconazole oral tablet</i>	T4	
VOSEVI	T5	PA; QL (28 EA per 28 days)
XENLETA ORAL	T5	
XIFAXAN ORAL TABLET 200 MG	T4	QL (9 EA per 3 days)
XIFAXAN ORAL TABLET 550 MG	T5	PA; QL (62 EA per 31 days)
XOFLUZA ORAL TABLET 40 MG, 80 MG	T3	QL (9 EA per 365 days)
ZEMDRI	T5	
ZEPATIER	T5	PA; QL (28 EA per 28 days)
<i>zidovudine</i>	T2	
ZOSYN IN DEXTROSE (ISO-OSM) INTRAVENOUS PIGGYBACK 2.25 GRAM/50 ML, 3.375 GRAM/50 ML	T3	
ZYVOX INTRAVENOUS PIGGYBACK 600 MG/300 ML	T4	
Antineoplastic / Immunosuppressant Drugs		
<i>abiraterone oral tablet 250 mg</i>	T5	PA-NS; QL (124 EA per 31 days)
<i>abiraterone oral tablet 500 mg</i>	T5	PA-NS; QL (62 EA per 31 days)
ALECENSA	T5	PA-NS; QL (248 EA per 31 days)
ALUNBRIG ORAL TABLET 180 MG, 90 MG	T5	PA-NS; QL (31 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
ALUNBRIG ORAL TABLET 30 MG	T5	PA-NS; QL (186 EA per 31 days)
ALUNBRIG ORAL TABLETS,DOSE PACK	T5	PA-NS; QL (30 EA per 365 days)
<i>anastrozole</i>	T2	
ASTAGRAF XL ORAL CAPSULE,EXTENDED RELEASE 24HR 0.5 MG, 1 MG	T3	PA-BvD
ASTAGRAF XL ORAL CAPSULE,EXTENDED RELEASE 24HR 5 MG	T4	PA-BvD
AYVAKIT	T5	PA-NS; QL (31 EA per 31 days)
AZASAN	T4	PA-BvD
<i>azathioprine oral tablet 100 mg, 75 mg</i>	T4	PA-BvD
<i>azathioprine oral tablet 50 mg</i>	T2	PA-BvD
BALVERSA	T5	PA-NS
<i>bexarotene oral</i>	T5	PA-NS
<i>bexarotene topical</i>	T5	PA-NS; QL (60 GM per 28 days)
<i>bicalutamide</i>	T2	
BOSULIF	T5	PA-NS
BRAFTOVI ORAL CAPSULE 75 MG	T5	PA-NS; QL (186 EA per 31 days)
BRUKINSA	T5	PA-NS; QL (124 EA per 31 days)
CABOMETYX	T5	PA-NS; QL (31 EA per 31 days)
CALQUENCE	T5	PA-NS; QL (62 EA per 31 days)
CALQUENCE (ACALABRUTINIB MAL)	T5	PA-NS; QL (62 EA per 31 days)
CAPRELSA	T5	PA-NS
COMETRIQ	T5	PA-NS
COPIKTRA	T5	PA-NS; QL (62 EA per 31 days)
COTELIC	T5	PA-NS; LA
<i>cyclophosphamide oral</i>	T3	PA-BvD
<i>cyclosporine modified</i>	T2	PA-BvD
<i>cyclosporine oral capsule</i>	T2	PA-BvD
DAURISMO ORAL TABLET 100 MG	T5	PA-NS; QL (31 EA per 31 days)
DAURISMO ORAL TABLET 25 MG	T5	PA-NS; QL (62 EA per 31 days)
DROXIA	T4	
ELIGARD	T4	
ELIGARD (3 MONTH)	T4	
ELIGARD (4 MONTH)	T4	
ELIGARD (6 MONTH)	T4	
EMCYT	T3	
ENSPRYNG	T5	PA; QL (1 ML per 28 days)

Drug Name	Drug Tier	Requirements/Limits
ENVARBUS XR	T4	PA-BvD
ERIVEDGE	T5	PA-NS; QL (31 EA per 31 days)
ERLEADA	T5	PA-NS; QL (124 EA per 31 days)
<i>erlotinib</i>	T5	PA-NS; QL (31 EA per 31 days)
<i>everolimus (antineoplastic) oral tablet 10 mg, 2.5 mg, 7.5 mg</i>	T5	PA-NS; QL (31 EA per 31 days)
<i>everolimus (antineoplastic) oral tablet 5 mg</i>	T5	PA-NS; QL (62 EA per 31 days)
<i>everolimus (antineoplastic) oral tablet for suspension 2 mg, 5 mg</i>	T5	PA-NS; QL (62 EA per 31 days)
<i>everolimus (antineoplastic) oral tablet for suspension 3 mg</i>	T5	PA-NS; QL (93 EA per 31 days)
<i>everolimus (immunosuppressive)</i>	T5	PA-BvD
<i>exemestane</i>	T2	
EXKIVITY	T5	PA-NS; QL (124 EA per 31 days)
FIRMAGON KIT W DILUENT SYRINGE SUBCUTANEOUS RECON SOLN 120 MG	T5	
FIRMAGON KIT W DILUENT SYRINGE SUBCUTANEOUS RECON SOLN 80 MG	T4	
FOTIVDA	T5	PA-NS; QL (21 EA per 28 days)
GAVRETO	T5	PA-NS; QL (124 EA per 31 days)
GENGRAF	T2	PA-BvD
GILOTRIF	T5	PA-NS; QL (31 EA per 31 days)
<i>hydroxyurea</i>	T2	
IBRANCE	T5	PA-NS; QL (21 EA per 28 days)
ICLUSIG	T5	PA-NS; QL (31 EA per 31 days)
IDHIFA ORAL TABLET 100 MG	T5	PA-NS; QL (31 EA per 31 days)
IDHIFA ORAL TABLET 50 MG	T5	PA-NS; QL (62 EA per 31 days)
<i>imatinib oral tablet 100 mg</i>	T5	PA-NS; QL (93 EA per 31 days)
<i>imatinib oral tablet 400 mg</i>	T5	PA-NS; QL (62 EA per 31 days)
IMBRUVICA ORAL CAPSULE 140 MG	T5	PA-NS; QL (124 EA per 31 days)
IMBRUVICA ORAL CAPSULE 70 MG	T5	PA-NS; QL (31 EA per 31 days)
IMBRUVICA ORAL SUSPENSION	T5	PA-NS; QL (216 ML per 25 days)
IMBRUVICA ORAL TABLET	T5	PA-NS; QL (31 EA per 31 days)
INLYTA	T5	PA-NS; QL (124 EA per 31 days)
INQOVI	T5	PA-NS; QL (5 EA per 28 days)
INREBIC	T5	PA-NS; QL (124 EA per 31 days)
IRESSA	T5	PA-NS
JAKAFI	T5	PA-NS; QL (62 EA per 31 days)
KISQALI FEMARA CO-PACK ORAL TABLET 200 MG/DAY(200 MG X 1)-2.5 MG	T5	PA-NS; QL (49 EA per 28 days)

Drug Name	Drug Tier	Requirements/Limits
KISQALI FEMARA CO-PACK ORAL TABLET 400 MG/DAY(200 MG X 2)-2.5 MG	T5	PA-NS; QL (70 EA per 28 days)
KISQALI FEMARA CO-PACK ORAL TABLET 600 MG/DAY(200 MG X 3)-2.5 MG	T5	PA-NS; QL (91 EA per 28 days)
KISQALI ORAL TABLET 200 MG/DAY (200 MG X 1)	T5	PA-NS; QL (21 EA per 28 days)
KISQALI ORAL TABLET 400 MG/DAY (200 MG X 2)	T5	PA-NS; QL (42 EA per 28 days)
KISQALI ORAL TABLET 600 MG/DAY (200 MG X 3)	T5	PA-NS; QL (63 EA per 28 days)
KLISYRI	T4	PA
KOSELUGO ORAL CAPSULE 10 MG	T5	PA-NS; QL (279 EA per 31 days)
KOSELUGO ORAL CAPSULE 25 MG	T5	PA-NS; QL (124 EA per 31 days)
<i>lapatinib</i>	T5	PA-NS
<i>lenalidomide</i>	T5	PA-NS; QL (21 EA per 28 days)
LENVIMA	T5	PA-NS
<i>letrozole</i>	T2	
<i>leucovorin calcium oral</i>	T2	
LEUKERAN	T4	
<i>leuprolide subcutaneous kit</i>	T2	
LONSURF	T5	PA-NS
LORBRENA ORAL TABLET 100 MG	T5	PA-NS; QL (31 EA per 31 days)
LORBRENA ORAL TABLET 25 MG	T5	PA-NS; QL (93 EA per 31 days)
LUMAKRAS	T5	PA-NS; QL (248 EA per 31 days)
LUPKYNIS	T5	PA; QL (186 EA per 31 days)
LUPRON DEPOT	T5	ST
LUPRON DEPOT (3 MONTH)	T5	ST
LUPRON DEPOT (4 MONTH)	T5	ST
LUPRON DEPOT (6 MONTH)	T5	ST
LYNPARZA	T5	PA-NS; QL (124 EA per 31 days)
LYSODREN	T3	
MATULANE	T5	
<i>megestrol oral suspension 400 mg/10 ml (40 mg/ml)</i>	T2	PA
<i>megestrol oral suspension 625 mg/5 ml (125 mg/ml)</i>	T4	PA
<i>megestrol oral tablet</i>	T2	PA-NS
MEKINIST	T5	PA-NS
MEKTOVI	T5	PA-NS; QL (186 EA per 31 days)
<i>mercaptopurine</i>	T2	

Drug Name	Drug Tier	Requirements/Limits
MESNEX ORAL	T3	
<i>methotrexate sodium (pf) injection solution</i>	T2	PA-BvD
<i>methotrexate sodium injection</i>	T2	PA-BvD
<i>methotrexate sodium oral</i>	T1	PA-BvD
MYCAPSSA	T5	PA; QL (124 EA per 31 days)
<i>mycophenolate mofetil</i>	T2	PA-BvD
<i>mycophenolate sodium</i>	T2	PA-BvD
NEORAL	T3	PA-BvD
NERLYNX	T5	PA-NS; QL (186 EA per 31 days)
<i>nilutamide</i>	T5	
NINLARO	T5	PA-NS
NUBEQA	T5	PA-NS; QL (124 EA per 31 days)
<i>octreotide acetate injection solution 1,000 mcg/ml, 200 mcg/ml</i>	T3	PA
<i>octreotide acetate injection solution 100 mcg/ml, 50 mcg/ml</i>	T2	PA
<i>octreotide acetate injection solution 500 mcg/ml</i>	T5	PA
ODOMZO	T5	PA-NS; LA
ONUREG	T5	PA-NS; QL (14 EA per 28 days)
ORGOVYX	T5	PA-NS; QL (31 EA per 31 days)
PEMAZYRE	T5	PA-NS; QL (14 EA per 21 days)
PIQRAY ORAL TABLET 200 MG/DAY (200 MG X 1)	T5	PA-NS; QL (28 EA per 28 days)
PIQRAY ORAL TABLET 250 MG/DAY (200 MG X1-50 MG X1), 300 MG/DAY (150 MG X 2)	T5	PA-NS; QL (56 EA per 28 days)
POMALYST	T5	PA-NS; QL (21 EA per 28 days)
PROGRAF ORAL GRANULES IN PACKET	T4	PA-BvD
PURIXAN	T4	
QINLOCK	T5	PA-NS; QL (93 EA per 31 days)
RETEVMO ORAL CAPSULE 40 MG	T5	PA-NS; QL (186 EA per 31 days)
RETEVMO ORAL CAPSULE 80 MG	T5	PA-NS; QL (124 EA per 31 days)
ROZLYTREK ORAL CAPSULE 100 MG	T5	PA-NS; QL (155 EA per 31 days)
ROZLYTREK ORAL CAPSULE 200 MG	T5	PA-NS; QL (93 EA per 31 days)
RUBRACA	T5	PA-NS; QL (124 EA per 31 days)
RYDAPT	T5	PA-NS; QL (248 EA per 31 days)
SANDIMMUNE ORAL SOLUTION	T3	PA-BvD
SCEMBLIX ORAL TABLET 20 MG	T5	PA-NS; QL (62 EA per 31 days)
SCEMBLIX ORAL TABLET 40 MG	T5	PA-NS; QL (310 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
SIGNIFOR	T5	PA
<i>sirolimus</i>	T2	PA-BvD
SOLTAMOX	T4	
<i>sorafenib</i>	T5	PA-NS; QL (124 EA per 31 days)
SPRYCEL	T5	PA-NS; QL (31 EA per 31 days)
STIVARGA	T5	PA-NS; QL (84 EA per 28 days)
<i>sunitinib</i>	T5	PA-NS
SYNRIBO	T5	
TABLOID	T3	
TABRECTA	T5	PA-NS; QL (124 EA per 31 days)
<i>tacrolimus oral</i>	T2	PA-BvD
TAFINLAR	T5	PA-NS
TAGRISSO	T5	PA-NS; LA; QL (31 EA per 31 days)
TALZENNA ORAL CAPSULE 0.25 MG	T5	PA-NS; QL (93 EA per 31 days)
TALZENNA ORAL CAPSULE 0.5 MG	T5	PA-NS; QL (62 EA per 31 days)
TALZENNA ORAL CAPSULE 0.75 MG, 1 MG	T5	PA-NS; QL (31 EA per 31 days)
<i>tamoxifen</i>	T1	
TASIGNA	T5	PA-NS; QL (124 EA per 31 days)
TAZVERIK	T5	PA-NS; QL (248 EA per 31 days)
TEPMETKO	T5	PA-NS; QL (62 EA per 31 days)
THALOMID ORAL CAPSULE 100 MG, 150 MG, 50 MG	T5	PA-NS; QL (28 EA per 28 days)
THALOMID ORAL CAPSULE 200 MG	T5	PA-NS; QL (56 EA per 28 days)
TIBSOVO	T5	PA-NS; QL (62 EA per 31 days)
<i>toremifene</i>	T3	
TRELSTAR INTRAMUSCULAR SUSPENSION FOR RECONSTITUTION 11.25 MG, 22.5 MG	T3	PA
TRELSTAR INTRAMUSCULAR SUSPENSION FOR RECONSTITUTION 3.75 MG	T5	PA
<i>tretinoin (antineoplastic)</i>	T5	
TREXALL	T3	PA-BvD
TRUSELTIQ ORAL CAPSULE 100 MG/DAY (100 MG X 1)	T5	PA-NS; QL (21 EA per 28 days)
TRUSELTIQ ORAL CAPSULE 125 MG/DAY(100 MG X1-25MG X1), 50 MG/DAY (25 MG X 2)	T5	PA-NS; QL (42 EA per 28 days)

Drug Name	Drug Tier	Requirements/Limits
TRUSELTIQ ORAL CAPSULE 75 MG/DAY (25 MG X 3)	T5	PA-NS; QL (63 EA per 28 days)
TUKYSA ORAL TABLET 150 MG	T5	PA-NS; QL (124 EA per 31 days)
TUKYSA ORAL TABLET 50 MG	T5	PA-NS; QL (248 EA per 31 days)
TURALIO	T5	PA-NS; QL (124 EA per 31 days)
VENCLEXTA ORAL TABLET 10 MG	T4	PA-NS
VENCLEXTA ORAL TABLET 100 MG, 50 MG	T5	PA-NS
VENCLEXTA STARTING PACK	T5	PA-NS
VERZENIO	T5	PA-NS; QL (62 EA per 31 days)
VIJOICE ORAL TABLET 125 MG, 50 MG	T5	PA-NS; QL (31 EA per 31 days)
VIJOICE ORAL TABLET 250 MG/DAY (200 MG X1-50 MG X1)	T5	PA-NS; QL (56 EA per 28 days)
VITRAKVI ORAL CAPSULE 100 MG	T5	PA-NS; QL (62 EA per 31 days)
VITRAKVI ORAL CAPSULE 25 MG	T5	PA-NS; QL (186 EA per 31 days)
VITRAKVI ORAL SOLUTION	T5	PA-NS; QL (310 ML per 31 days)
VIZIMPRO	T5	PA-NS; QL (31 EA per 31 days)
VONJO	T5	PA-NS; QL (124 EA per 31 days)
VOTRIENT	T5	PA-NS; QL (124 EA per 31 days)
WELIREG	T5	PA-NS; QL (93 EA per 31 days)
XALKORI	T5	PA-NS; QL (62 EA per 31 days)
XATMEP	T4	PA-BvD
XERMELO	T5	PA; QL (93 EA per 31 days)
XGEVA	T5	PA-NS
XOSPATA	T5	PA-NS; QL (124 EA per 31 days)
XPOVIO ORAL TABLET 100 MG/WEEK (50 MG X 2), 40MG TWICE WEEK (40 MG X 2), 80 MG/WEEK (40 MG X 2)	T5	PA-NS; QL (8 EA per 28 days)
XPOVIO ORAL TABLET 40 MG/WEEK (40 MG X 1), 60 MG/WEEK (60 MG X 1)	T5	PA-NS; QL (4 EA per 28 days)
XPOVIO ORAL TABLET 60MG TWICE WEEK (120 MG/WEEK)	T5	PA-NS; QL (24 EA per 28 days)
XPOVIO ORAL TABLET 80MG TWICE WEEK (160 MG/WEEK)	T5	PA-NS; QL (32 EA per 28 days)
XTANDI ORAL CAPSULE	T5	PA-NS; QL (124 EA per 31 days)
XTANDI ORAL TABLET 40 MG	T5	PA-NS; QL (124 EA per 31 days)
XTANDI ORAL TABLET 80 MG	T5	PA-NS; QL (62 EA per 31 days)
YONSA	T5	PA-NS; QL (124 EA per 31 days)
ZEJULA	T5	PA-NS; QL (93 EA per 31 days)
ZELBORAF	T5	PA-NS

Drug Name	Drug Tier	Requirements/Limits
ZOLINZA	T5	PA-NS
ZORTRESS ORAL TABLET 1 MG	T5	PA-BvD
ZYDELIG	T5	PA-NS; QL (62 EA per 31 days)
ZYKADIA	T5	PA-NS; QL (93 EA per 31 days)
Autonomic / Cns Drugs, Neurology / Psych		
ABILIFY MAINTENA	T5	QL (1 EA per 28 days)
<i>acetaminophen-codeine oral solution 120-12 mg/5 ml</i>	T2	PA; QL (5167 ML per 31 days)
<i>acetaminophen-codeine oral tablet</i>	T2	PA; QL (403 EA per 31 days)
AIMOVIG AUTOINJECTOR SUBCUTANEOUS AUTO-INJECTOR 140 MG/ML	T3	PA; QL (1 ML per 28 days)
AIMOVIG AUTOINJECTOR SUBCUTANEOUS AUTO-INJECTOR 70 MG/ML	T3	PA; QL (2 ML per 28 days)
AJOVY AUTOINJECTOR	T3	PA; QL (1.5 ML per 28 days)
AJOVY SYRINGE	T3	PA; QL (1.5 ML per 28 days)
<i>almotriptan malate oral tablet 12.5 mg</i>	T2	QL (8 EA per 28 days)
<i>almotriptan malate oral tablet 6.25 mg</i>	T2	QL (16 EA per 28 days)
ALPRAZOLAM INTENSOL	T4	PA
<i>alprazolam oral tablet 0.25 mg, 0.5 mg</i>	T2	PA; QL (93 EA per 31 days)
<i>alprazolam oral tablet 1 mg, 2 mg</i>	T2	PA; QL (155 EA per 31 days)
<i>alprazolam oral tablet extended release 24 hr 0.5 mg, 1 mg</i>	T2	PA; QL (31 EA per 31 days)
<i>alprazolam oral tablet extended release 24 hr 2 mg</i>	T2	PA; QL (155 EA per 31 days)
<i>alprazolam oral tablet extended release 24 hr 3 mg</i>	T2	PA; QL (93 EA per 31 days)
<i>amitriptyline</i>	T2	PA-NS
<i>amitriptyline-chlordiazepoxide</i>	T4	PA-NS
<i>amoxapine</i>	T1	
<i>apomorphine</i>	T5	PA; QL (60 ML per 30 days)
APTENSIO XR	T4	ST; QL (31 EA per 31 days)
APTIOM ORAL TABLET 200 MG, 800 MG	T4	
APTIOM ORAL TABLET 400 MG, 600 MG	T5	
<i>aripiprazole oral solution</i>	T4	PA-NS
<i>aripiprazole oral tablet 10 mg, 15 mg, 2 mg, 5 mg</i>	T3	PA-NS
<i>aripiprazole oral tablet 20 mg, 30 mg</i>	T4	PA-NS
<i>aripiprazole oral tablet, disintegrating</i>	T4	PA-NS

Drug Name	Drug Tier	Requirements/Limits
ARISTADA INITIO	T5	QL (4.8 ML per 365 days)
ARISTADA INTRAMUSCULAR SUSPENSION,EXTENDED REL SYRING 1,064 MG/3.9 ML	T5	QL (3.9 ML per 28 days)
ARISTADA INTRAMUSCULAR SUSPENSION,EXTENDED REL SYRING 441 MG/1.6 ML	T5	QL (1.6 ML per 28 days)
ARISTADA INTRAMUSCULAR SUSPENSION,EXTENDED REL SYRING 662 MG/2.4 ML	T5	QL (2.4 ML per 28 days)
ARISTADA INTRAMUSCULAR SUSPENSION,EXTENDED REL SYRING 882 MG/3.2 ML	T5	QL (3.2 ML per 28 days)
<i>armodafinil</i>	T4	PA; QL (31 EA per 31 days)
<i>asenapine maleate</i>	T4	QL (62 EA per 31 days)
<i>atomoxetine oral capsule 10 mg, 25 mg, 40 mg</i>	T4	QL (62 EA per 31 days)
<i>atomoxetine oral capsule 100 mg, 60 mg, 80 mg</i>	T4	QL (31 EA per 31 days)
<i>atomoxetine oral capsule 18 mg</i>	T4	QL (124 EA per 31 days)
AUBAGIO	T5	PA; QL (31 EA per 31 days)
<i>baclofen oral tablet 10 mg</i>	T1	
<i>baclofen oral tablet 20 mg</i>	T2	
BAFIERTAM	T5	PA; QL (124 EA per 31 days)
BANZEL ORAL SUSPENSION	T5	PA-NS
<i>benztropine oral</i>	T2	PA
BRIVIACT ORAL	T5	
<i>bromocriptine</i>	T4	
<i>buprenorphine</i>	T4	PA; QL (4 EA per 28 days)
<i>buprenorphine hcl sublingual tablet 2 mg</i>	T3	QL (93 EA per 31 days)
<i>buprenorphine hcl sublingual tablet 8 mg</i>	T3	QL (62 EA per 31 days)
<i>buprenorphine-naloxone sublingual film 12-3 mg, 4-1 mg, 8-2 mg</i>	T2	QL (62 EA per 31 days)
<i>buprenorphine-naloxone sublingual film 2-0.5 mg</i>	T2	QL (93 EA per 31 days)
<i>buprenorphine-naloxone sublingual tablet</i>	T4	ST; QL (93 EA per 31 days)
<i>bupropion hcl oral tablet</i>	T2	
<i>bupropion hcl oral tablet extended release 24 hr 150 mg</i>	T2	QL (93 EA per 31 days)
<i>bupropion hcl oral tablet extended release 24 hr 300 mg</i>	T2	QL (31 EA per 31 days)
<i>bupropion hcl oral tablet extended release 24 hr 450 mg</i>	T4	
<i>bupropion hcl oral tablet sustained-release 12 hr</i>	T2	QL (62 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
<i>buspirone</i>	T2	
<i>butalbital-acetaminop-caf-cod oral capsule 50-300-40-30 mg</i>	T4	PA; QL (403 EA per 31 days)
<i>butalbital-acetaminop-caf-cod oral capsule 50-325-40-30 mg</i>	T4	PA; QL (372 EA per 31 days)
<i>butalbital-acetaminophen oral tablet 50-325 mg</i>	T2	QL (372 EA per 31 days)
<i>butalbital-acetaminophen-caff oral capsule 50-300-40 mg</i>	T4	QL (403 EA per 31 days)
<i>butalbital-acetaminophen-caff oral capsule 50-325-40 mg</i>	T4	QL (372 EA per 31 days)
<i>butalbital-acetaminophen-caff oral tablet</i>	T4	QL (372 EA per 31 days)
<i>butalbital-aspirin-caffeine oral capsule</i>	T2	
<i>butorphanol nasal</i>	T2	QL (5 ML per 28 days)
BUTRANS	T4	PA; QL (4 EA per 28 days)
CAMBIA	T4	
CAPLYTA	T5	PA-NS; QL (31 EA per 31 days)
<i>carbamazepine oral capsule, er multiphase 12 hr</i>	T2	
<i>carbamazepine oral suspension 100 mg/5 ml</i>	T1	
<i>carbamazepine oral tablet</i>	T1	
<i>carbamazepine oral tablet extended release 12 hr</i>	T2	
<i>carbamazepine oral tablet, chewable</i>	T1	
<i>carbidopa-levodopa</i>	T2	
<i>carbidopa-levodopa-entacapone</i>	T2	
<i>celecoxib</i>	T2	ST; QL (62 EA per 31 days)
CELONTIN ORAL CAPSULE 300 MG	T4	
<i>chlorpromazine oral</i>	T4	
<i>citalopram oral solution</i>	T3	
<i>citalopram oral tablet</i>	T1	
<i>clobazam oral suspension</i>	T4	PA-NS; QL (496 ML per 31 days)
<i>clobazam oral tablet</i>	T4	PA-NS; QL (62 EA per 31 days)
<i>clomipramine</i>	T4	PA-NS
<i>clonazepam oral tablet 0.5 mg</i>	T2	QL (93 EA per 31 days)
<i>clonazepam oral tablet 1 mg</i>	T2	QL (124 EA per 31 days)
<i>clonazepam oral tablet 2 mg</i>	T2	QL (310 EA per 31 days)
<i>clonazepam oral tablet, disintegrating 0.125 mg, 0.25 mg, 0.5 mg</i>	T2	QL (93 EA per 31 days)
<i>clonazepam oral tablet, disintegrating 1 mg</i>	T2	QL (124 EA per 31 days)
<i>clonazepam oral tablet, disintegrating 2 mg</i>	T2	QL (310 EA per 31 days)
<i>clonidine hcl oral tablet extended release 12 hr</i>	T4	PA

Drug Name	Drug Tier	Requirements/Limits
<i>clorazepate dipotassium oral tablet 15 mg</i>	T2	QL (186 EA per 31 days)
<i>clorazepate dipotassium oral tablet 3.75 mg, 7.5 mg</i>	T2	QL (93 EA per 31 days)
<i>clozapine oral tablet 100 mg, 25 mg</i>	T2	QL (279 EA per 31 days)
<i>clozapine oral tablet 200 mg</i>	T2	QL (124 EA per 31 days)
<i>clozapine oral tablet 50 mg</i>	T2	QL (93 EA per 31 days)
<i>clozapine oral tablet, disintegrating 100 mg, 25 mg</i>	T2	QL (279 EA per 31 days)
<i>clozapine oral tablet, disintegrating 12.5 mg</i>	T2	QL (93 EA per 31 days)
<i>clozapine oral tablet, disintegrating 150 mg</i>	T4	QL (186 EA per 31 days)
<i>clozapine oral tablet, disintegrating 200 mg</i>	T4	QL (124 EA per 31 days)
COPAXONE SUBCUTANEOUS SYRINGE 20 MG/ML	T5	ST; QL (31 ML per 31 days)
COPAXONE SUBCUTANEOUS SYRINGE 40 MG/ML	T5	ST; QL (12 ML per 28 days)
<i>cyclobenzaprine oral tablet 10 mg, 7.5 mg</i>	T2	PA; QL (93 EA per 31 days)
<i>cyclobenzaprine oral tablet 5 mg</i>	T2	PA; QL (155 EA per 31 days)
<i>dalfampridine</i>	T5	PA; QL (62 EA per 31 days)
<i>dantrolene oral</i>	T4	
DAYTRANA	T4	PA; QL (30 EA per 30 days)
<i>desipramine oral tablet 10 mg, 25 mg, 50 mg</i>	T3	
<i>desipramine oral tablet 100 mg, 150 mg, 75 mg</i>	T4	
<i>desvenlafaxine</i>	T4	
<i>desvenlafaxine succinate</i>	T4	QL (31 EA per 31 days)
<i>dexmethylphenidate oral capsule, er biphasic 50-50</i>	T2	QL (31 EA per 31 days)
<i>dexmethylphenidate oral tablet 10 mg</i>	T2	QL (62 EA per 31 days)
<i>dexmethylphenidate oral tablet 2.5 mg, 5 mg</i>	T2	QL (93 EA per 31 days)
<i>dextroamphetamine sulfate oral solution</i>	T3	
<i>dextroamphetamine sulfate oral tablet 10 mg</i>	T4	QL (186 EA per 31 days)
<i>dextroamphetamine sulfate oral tablet 15 mg, 20 mg, 30 mg</i>	T4	QL (62 EA per 31 days)
<i>dextroamphetamine sulfate oral tablet 5 mg</i>	T4	QL (341 EA per 31 days)
<i>dextroamphetamine-amphetamine oral capsule, extended release 24hr</i>	T3	QL (31 EA per 31 days)
<i>dextroamphetamine-amphetamine oral tablet 10 mg, 30 mg</i>	T2	QL (62 EA per 31 days)
<i>dextroamphetamine-amphetamine oral tablet 12.5 mg, 15 mg, 5 mg, 7.5 mg</i>	T1	QL (62 EA per 31 days)
<i>dextroamphetamine-amphetamine oral tablet 20 mg</i>	T2	QL (93 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
DIACOMIT ORAL CAPSULE 250 MG	T5	PA-NS; QL (341 EA per 31 days)
DIACOMIT ORAL CAPSULE 500 MG	T5	PA-NS; QL (186 EA per 31 days)
DIACOMIT ORAL POWDER IN PACKET 250 MG	T5	PA-NS; QL (341 EA per 31 days)
DIACOMIT ORAL POWDER IN PACKET 500 MG	T5	PA-NS; QL (186 EA per 31 days)
DIAZEPAM INTENSOL	T2	QL (248 ML per 31 days)
<i>diazepam oral solution 5 mg/5 ml (1 mg/ml)</i>	T2	QL (1500 ML per 31 days)
<i>diazepam oral tablet</i>	T2	QL (124 EA per 31 days)
<i>diazepam rectal</i>	T4	
<i>diclofenac epolamine</i>	T4	PA; QL (62 EA per 31 days)
<i>diclofenac potassium oral tablet 50 mg</i>	T1	
<i>diclofenac sodium oral</i>	T1	
<i>diclofenac sodium topical drops</i>	T2	QL (450 ML per 28 days)
<i>diclofenac sodium topical gel 1 %</i>	T3	QL (900 GM per 28 days)
<i>diclofenac-misoprostol</i>	T2	
<i>diflunisal</i>	T2	
<i>dihydroergotamine nasal</i>	T4	PA; QL (8 ML per 31 days)
DILANTIN	T4	
DILANTIN EXTENDED	T4	
DILANTIN INFATABS	T4	
DILANTIN-125	T4	
<i>dimethyl fumarate oral capsule, delayed release(dr/ec) 120 mg (14)- 240 mg (46)</i>	T5	PA; QL (120 EA per 365 days)
<i>dimethyl fumarate oral capsule, delayed release(dr/ec) 120 mg, 240 mg</i>	T5	PA; QL (62 EA per 31 days)
<i>divalproex oral capsule, delayed rel sprinkle</i>	T2	
<i>divalproex oral tablet extended release 24 hr 250 mg</i>	T2	
<i>divalproex oral tablet extended release 24 hr 500 mg</i>	T3	
<i>divalproex oral tablet, delayed release (dr/ec)</i>	T2	
<i>donepezil oral tablet 10 mg, 5 mg</i>	T2	
<i>donepezil oral tablet 23 mg</i>	T3	QL (31 EA per 31 days)
<i>donepezil oral tablet, disintegrating</i>	T2	
<i>doxepin oral capsule</i>	T2	PA-NS
<i>doxepin oral concentrate</i>	T2	PA-NS
<i>doxepin oral tablet</i>	T2	PA
DRIZALMA SPRINKLE ORAL CAPSULE, DELAYED REL SPRINKLE 20 MG	T4	PA-NS; QL (93 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
DRIZALMA SPRINKLE ORAL CAPSULE, DELAYED REL SPRINKLE 30 MG, 60 MG	T4	PA-NS; QL (62 EA per 31 days)
DRIZALMA SPRINKLE ORAL CAPSULE, DELAYED REL SPRINKLE 40 MG	T4	PA-NS; QL (31 EA per 31 days)
<i>duloxetine oral capsule, delayed release(dr/ec) 20 mg, 60 mg</i>	T2	QL (62 EA per 31 days)
<i>duloxetine oral capsule, delayed release(dr/ec) 30 mg</i>	T2	QL (31 EA per 31 days)
<i>duloxetine oral capsule, delayed release(dr/ec) 40 mg</i>	T3	QL (31 EA per 31 days)
DUOPA	T5	PA-BvD
<i>eletriptan oral tablet 20 mg</i>	T4	QL (12 EA per 28 days)
<i>eletriptan oral tablet 40 mg</i>	T4	QL (6 EA per 28 days)
EMGALITY PEN	T3	PA; QL (1 ML per 28 days)
EMGALITY SYRINGE SUBCUTANEOUS SYRINGE 120 MG/ML	T3	PA; QL (1 ML per 28 days)
EMGALITY SYRINGE SUBCUTANEOUS SYRINGE 300 MG/3 ML (100 MG/ML X 3)	T3	PA; QL (3 ML per 28 days)
EMSAM	T5	QL (30 EA per 30 days)
ENDOCET ORAL TABLET 10-325 MG	T3	PA; QL (372 EA per 31 days)
ENDOCET ORAL TABLET 5-325 MG, 7.5-325 MG	T2	PA; QL (372 EA per 31 days)
<i>entacapone</i>	T2	
EPIDIOLEX	T5	PA-NS
EPITOL	T1	
EPRONTIA	T4	PA-NS; QL (496 ML per 31 days)
EQUETRO	T4	
<i>ergoloid</i>	T2	PA
<i>ergotamine-caffeine</i>	T2	PA
<i>escitalopram oxalate oral solution</i>	T4	QL (620 ML per 31 days)
<i>escitalopram oxalate oral tablet 10 mg</i>	T2	QL (45 EA per 30 days)
<i>escitalopram oxalate oral tablet 20 mg, 5 mg</i>	T2	QL (30 EA per 30 days)
<i>estazolam</i>	T2	
<i>eszopiclone</i>	T2	PA; QL (31 EA per 31 days)
<i>ethosuximide oral capsule</i>	T3	
<i>ethosuximide oral solution</i>	T2	
<i>etodolac oral capsule</i>	T3	
<i>etodolac oral tablet</i>	T2	
<i>etodolac oral tablet extended release 24 hr</i>	T3	
EVRYSDI	T5	PA; QL (240 ML per 31 days)

Drug Name	Drug Tier	Requirements/Limits
FANAPT ORAL TABLET	T4	QL (62 EA per 31 days)
FANAPT ORAL TABLETS,DOSE PACK	T4	QL (16 EA per 365 days)
<i>felbamate oral suspension</i>	T2	
<i>felbamate oral tablet</i>	T4	
<i>fenoprofen oral capsule 400 mg</i>	T4	
<i>fentanyl citrate buccal lozenge on a handle 1,200 mcg</i>	T5	PA; QL (40 EA per 31 days)
<i>fentanyl citrate buccal lozenge on a handle 1,600 mcg</i>	T5	PA; QL (30 EA per 31 days)
<i>fentanyl citrate buccal lozenge on a handle 200 mcg</i>	T4	PA; QL (124 EA per 31 days)
<i>fentanyl citrate buccal lozenge on a handle 400 mcg</i>	T5	PA; QL (119 EA per 31 days)
<i>fentanyl citrate buccal lozenge on a handle 600 mcg</i>	T5	PA; QL (79 EA per 31 days)
<i>fentanyl citrate buccal lozenge on a handle 800 mcg</i>	T5	PA; QL (59 EA per 31 days)
<i>fentanyl citrate buccal tablet, effervescent 100 mcg, 200 mcg</i>	T5	PA; QL (124 EA per 31 days)
<i>fentanyl citrate buccal tablet, effervescent 400 mcg</i>	T5	PA; QL (119 EA per 31 days)
<i>fentanyl citrate buccal tablet, effervescent 600 mcg</i>	T5	PA; QL (79 EA per 31 days)
<i>fentanyl citrate buccal tablet, effervescent 800 mcg</i>	T5	PA; QL (59 EA per 31 days)
<i>fentanyl transdermal patch 72 hour 100 mcg/hr</i>	T4	PA; QL (10 EA per 30 days)
<i>fentanyl transdermal patch 72 hour 12 mcg/hr</i>	T4	PA; QL (20 EA per 30 days)
<i>fentanyl transdermal patch 72 hour 25 mcg/hr</i>	T2	PA; QL (20 EA per 30 days)
<i>fentanyl transdermal patch 72 hour 50 mcg/hr</i>	T2	PA; QL (17 EA per 30 days)
<i>fentanyl transdermal patch 72 hour 75 mcg/hr</i>	T4	PA; QL (12 EA per 30 days)
FENTORA BUCCAL TABLET, EFFERVESCENT 100 MCG, 200 MCG	T5	PA; QL (124 EA per 31 days)
FENTORA BUCCAL TABLET, EFFERVESCENT 400 MCG	T5	PA; QL (119 EA per 31 days)
FENTORA BUCCAL TABLET, EFFERVESCENT 600 MCG	T5	PA; QL (79 EA per 31 days)
FENTORA BUCCAL TABLET, EFFERVESCENT 800 MCG	T5	PA; QL (59 EA per 31 days)
FETZIMA ORAL CAPSULE,EXT REL 24HR DOSE PACK	T4	PA-NS; QL (56 EA per 365 days)
FETZIMA ORAL CAPSULE,EXTENDED RELEASE 24 HR 120 MG, 40 MG, 80 MG	T4	PA-NS; QL (31 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
FETZIMA ORAL CAPSULE,EXTENDED RELEASE 24 HR 20 MG	T4	PA-NS; QL (93 EA per 31 days)
<i> fingolimod </i>	T5	PA; QL (31 EA per 31 days)
FINTEPLA	T5	PA-NS; QL (360 ML per 30 days)
FIRDAPSE	T5	PA; QL (248 EA per 31 days)
FLECTOR	T4	PA; QL (62 EA per 31 days)
<i> fluoxetine (pmd) </i>	T1	
<i> fluoxetine oral capsule </i>	T1	
<i> fluoxetine oral solution </i>	T1	
<i> fluoxetine oral tablet 10 mg, 20 mg </i>	T1	
<i> fluphenazine decanoate </i>	T2	
<i> fluphenazine hcl injection </i>	T2	
<i> fluphenazine hcl oral concentrate </i>	T2	
<i> fluphenazine hcl oral tablet </i>	T1	
<i> flurazepam </i>	T2	
<i> flurbiprofen oral tablet 100 mg </i>	T2	
<i> fluvoxamine </i>	T2	
<i> frovatriptan </i>	T4	QL (12 EA per 28 days)
FYCOMPA ORAL SUSPENSION	T5	
FYCOMPA ORAL TABLET 10 MG, 12 MG, 4 MG, 6 MG, 8 MG	T5	
FYCOMPA ORAL TABLET 2 MG	T4	
<i> gabapentin oral capsule 100 mg, 400 mg </i>	T2	PA-NS; QL (270 EA per 30 days)
<i> gabapentin oral capsule 300 mg </i>	T2	PA-NS; QL (360 EA per 30 days)
<i> gabapentin oral solution 250 mg/5 ml </i>	T3	PA-NS; QL (2160 ML per 30 days)
<i> gabapentin oral tablet 600 mg </i>	T2	PA-NS; QL (180 EA per 30 days)
<i> gabapentin oral tablet 800 mg </i>	T2	PA-NS; QL (120 EA per 30 days)
<i> galantamine oral capsule,ext rel. pellets 24 hr </i>	T3	
<i> galantamine oral solution </i>	T3	
<i> galantamine oral tablet </i>	T2	
<i> glatiramer subcutaneous syringe 20 mg/ml </i>	T5	QL (31 ML per 31 days)
<i> glatiramer subcutaneous syringe 40 mg/ml </i>	T5	QL (12 ML per 28 days)
GLATOPA SUBCUTANEOUS SYRINGE 20 MG/ML	T5	QL (31 ML per 31 days)
GLATOPA SUBCUTANEOUS SYRINGE 40 MG/ML	T5	QL (12 ML per 28 days)
<i> guanfacine oral tablet extended release 24 hr </i>	T2	PA
<i> haloperidol decanoate </i>	T2	

Drug Name	Drug Tier	Requirements/Limits
<i>haloperidol lactate injection</i>	T1	
<i>haloperidol lactate oral</i>	T2	
<i>haloperidol oral tablet 0.5 mg, 1 mg, 10 mg, 2 mg, 5 mg</i>	T1	
<i>haloperidol oral tablet 20 mg</i>	T2	
HETLIOZ	T5	PA; QL (31 EA per 31 days)
HETLIOZ LQ	T5	PA; QL (158 ML per 31 days)
HORIZANT ORAL TABLET EXTENDED RELEASE 300 MG	T4	PA; QL (90 EA per 30 days)
HORIZANT ORAL TABLET EXTENDED RELEASE 600 MG	T4	PA; QL (60 EA per 30 days)
<i>hydrocodone-acetaminophen oral tablet 10-325 mg, 5-325 mg, 7.5-325 mg</i>	T2	PA; QL (372 EA per 31 days)
<i>hydrocodone-ibuprofen oral tablet 10-200 mg, 5-200 mg</i>	T3	PA; QL (155 EA per 31 days)
<i>hydromorphone oral liquid</i>	T4	PA; QL (1550 ML per 31 days)
<i>hydromorphone oral tablet</i>	T2	PA; QL (186 EA per 31 days)
IBU ORAL TABLET 600 MG, 800 MG	T1	
<i>ibuprofen oral suspension</i>	T1	
<i>ibuprofen oral tablet 400 mg, 600 mg, 800 mg</i>	T1	
<i>imipramine hcl</i>	T2	PA-NS
<i>imipramine pamoate</i>	T4	PA-NS
INBRIJA INHALATION CAPSULE, W/INHALATION DEVICE	T5	PA; QL (300 EA per 30 days)
INDOCIN	T4	
<i>indomethacin oral</i>	T1	
INGREZZA INITIATION PACK	T5	PA; QL (56 EA per 365 days)
INGREZZA ORAL CAPSULE 40 MG	T5	PA; QL (62 EA per 31 days)
INGREZZA ORAL CAPSULE 60 MG, 80 MG	T5	PA; QL (31 EA per 31 days)
INVEGA HAFYERA INTRAMUSCULAR SYRINGE 1,092 MG/3.5 ML	T4	QL (3.5 ML per 180 days)
INVEGA HAFYERA INTRAMUSCULAR SYRINGE 1,560 MG/5 ML	T4	QL (5 ML per 180 days)
INVEGA SUSTENNA INTRAMUSCULAR SYRINGE 117 MG/0.75 ML	T5	QL (0.75 ML per 28 days)
INVEGA SUSTENNA INTRAMUSCULAR SYRINGE 156 MG/ML	T5	QL (1 ML per 28 days)
INVEGA SUSTENNA INTRAMUSCULAR SYRINGE 234 MG/1.5 ML	T5	QL (1.5 ML per 28 days)
INVEGA SUSTENNA INTRAMUSCULAR SYRINGE 39 MG/0.25 ML	T4	QL (0.25 ML per 28 days)

Drug Name	Drug Tier	Requirements/Limits
INVEGA SUSTENNA INTRAMUSCULAR SYRINGE 78 MG/0.5 ML	T5	QL (0.5 ML per 28 days)
INVEGA TRINZA INTRAMUSCULAR SYRINGE 273 MG/0.88 ML	T5	QL (0.875 ML per 84 days)
INVEGA TRINZA INTRAMUSCULAR SYRINGE 410 MG/1.32 ML	T5	QL (1.315 ML per 84 days)
INVEGA TRINZA INTRAMUSCULAR SYRINGE 546 MG/1.75 ML	T5	QL (1.75 ML per 84 days)
INVEGA TRINZA INTRAMUSCULAR SYRINGE 819 MG/2.63 ML	T5	QL (2.625 ML per 84 days)
KESIMPTA PEN	T5	PA; QL (0.4 ML per 28 days)
<i>ketoprofen oral capsule 25 mg</i>	T2	
<i>ketoprofen oral capsule,ext rel. pellets 24 hr 200 mg</i>	T2	
<i>ketorolac oral</i>	T2	
KLOXXADO	T3	
KYNMOBI SUBLINGUAL FILM 10 MG, 15 MG, 20 MG, 25 MG, 30 MG	T5	PA; QL (155 EA per 31 days)
<i>lacosamide oral</i>	T4	
<i>lamotrigine oral tablet</i>	T2	
<i>lamotrigine oral tablet extended release 24hr</i>	T4	
<i>lamotrigine oral tablet, chewable dispersible</i>	T2	
<i>lamotrigine oral tablet,disintegrating</i>	T2	
LATUDA ORAL TABLET 120 MG, 20 MG, 40 MG, 60 MG	T5	PA-NS; QL (31 EA per 31 days)
LATUDA ORAL TABLET 80 MG	T5	PA-NS; QL (62 EA per 31 days)
<i>levetiracetam oral solution 100 mg/ml</i>	T2	
<i>levetiracetam oral tablet</i>	T2	
<i>levetiracetam oral tablet extended release 24 hr</i>	T2	
<i>lithium carbonate</i>	T1	
LORAZEPAM INTENSOL	T2	QL (155 ML per 31 days)
<i>lorazepam oral tablet 0.5 mg</i>	T2	QL (124 EA per 31 days)
<i>lorazepam oral tablet 1 mg</i>	T2	QL (186 EA per 31 days)
<i>lorazepam oral tablet 2 mg</i>	T2	QL (155 EA per 31 days)
<i>loxapine succinate</i>	T2	
LUCEMYRA	T5	
LYBALVI	T5	PA-NS; QL (31 EA per 31 days)
MARPLAN	T3	
MAVENCLAD (10 TABLET PACK)	T5	PA; QL (40 EA per 365 days)
MAVENCLAD (4 TABLET PACK)	T5	PA; QL (40 EA per 365 days)

Drug Name	Drug Tier	Requirements/Limits
MAVENCLAD (5 TABLET PACK)	T5	PA; QL (40 EA per 365 days)
MAVENCLAD (6 TABLET PACK)	T5	PA; QL (40 EA per 365 days)
MAVENCLAD (7 TABLET PACK)	T5	PA; QL (40 EA per 365 days)
MAVENCLAD (8 TABLET PACK)	T5	PA; QL (40 EA per 365 days)
MAVENCLAD (9 TABLET PACK)	T5	PA; QL (40 EA per 365 days)
MAYZENT ORAL TABLET 0.25 MG	T5	PA; QL (155 EA per 31 days)
MAYZENT ORAL TABLET 1 MG, 2 MG	T5	PA; QL (31 EA per 31 days)
MAYZENT STARTER(FOR 1MG MAINT)	T4	PA; QL (14 EA per 365 days)
MAYZENT STARTER(FOR 2MG MAINT)	T5	PA; QL (24 EA per 365 days)
<i>mefenamic acid</i>	T4	
<i>meloxicam oral tablet</i>	T1	
<i>memantine oral capsule,sprinkle,er 24hr</i>	T3	
<i>memantine oral solution</i>	T3	
<i>memantine oral tablet</i>	T2	
<i>memantine oral tablets,dose pack</i>	T4	
<i>metaxalone</i>	T2	PA
<i>methadone oral solution 10 mg/5 ml</i>	T2	PA; QL (1033 ML per 31 days)
<i>methadone oral solution 5 mg/5 ml</i>	T2	PA; QL (2066 ML per 31 days)
<i>methadone oral tablet 10 mg</i>	T2	PA; QL (206 EA per 31 days)
<i>methadone oral tablet 5 mg</i>	T2	PA; QL (248 EA per 31 days)
<i>methamphetamine</i>	T4	PA
<i>methylphenidate hcl oral cap,er sprinkle,biphasic 40-60</i>	T4	QL (31 EA per 31 days)
<i>methylphenidate hcl oral capsule, er biphasic 30-70 10 mg, 20 mg, 40 mg, 50 mg, 60 mg</i>	T4	QL (31 EA per 31 days)
<i>methylphenidate hcl oral capsule, er biphasic 30-70 30 mg</i>	T2	QL (31 EA per 31 days)
<i>methylphenidate hcl oral capsule,er biphasic 50-50 10 mg</i>	T2	QL (186 EA per 31 days)
<i>methylphenidate hcl oral capsule,er biphasic 50-50 20 mg</i>	T4	QL (93 EA per 31 days)
<i>methylphenidate hcl oral capsule,er biphasic 50-50 30 mg</i>	T4	QL (62 EA per 31 days)
<i>methylphenidate hcl oral capsule,er biphasic 50-50 40 mg</i>	T2	QL (62 EA per 31 days)
<i>methylphenidate hcl oral capsule,er biphasic 50-50 60 mg</i>	T4	QL (31 EA per 31 days)
<i>methylphenidate hcl oral tablet</i>	T2	QL (93 EA per 31 days)
<i>methylphenidate hcl oral tablet extended release 10 mg</i>	T2	QL (31 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
<i>methylphenidate hcl oral tablet extended release 20 mg</i>	T2	QL (93 EA per 31 days)
<i>methylphenidate hcl oral tablet extended release 24hr 18 mg, 18 mg (bx rating), 27 mg, 27 mg (bx rating), 36 mg, 36 mg (bx rating), 54 mg, 54 mg (bx rating)</i>	T4	QL (31 EA per 31 days)
<i>methylphenidate hcl oral tablet, chewable 10 mg</i>	T2	QL (186 EA per 31 days)
<i>methylphenidate hcl oral tablet, chewable 2.5 mg, 5 mg</i>	T2	QL (93 EA per 31 days)
<i>mirtazapine oral tablet 15 mg, 30 mg, 45 mg</i>	T2	
<i>mirtazapine oral tablet 7.5 mg</i>	T3	
<i>mirtazapine oral tablet, disintegrating</i>	T3	
<i>modafinil</i>	T2	PA; QL (31 EA per 31 days)
<i>molindone</i>	T4	
<i>morphine concentrate oral solution</i>	T2	PA; QL (310 ML per 31 days)
<i>morphine oral capsule, er multiphase 24 hr 120 mg</i>	T3	PA; QL (51 EA per 31 days)
<i>morphine oral capsule, er multiphase 24 hr 30 mg, 45 mg, 60 mg, 75 mg, 90 mg</i>	T3	PA; QL (62 EA per 31 days)
<i>morphine oral solution 10 mg/5 ml</i>	T2	PA; QL (2800 ML per 31 days)
<i>morphine oral solution 20 mg/5 ml (4 mg/ml)</i>	T2	PA; QL (1400 ML per 31 days)
<i>morphine oral tablet</i>	T2	PA; QL (186 EA per 31 days)
<i>morphine oral tablet extended release 100 mg</i>	T3	PA; QL (62 EA per 31 days)
<i>morphine oral tablet extended release 15 mg, 30 mg, 60 mg</i>	T3	PA; QL (100 EA per 31 days)
<i>morphine oral tablet extended release 200 mg</i>	T3	PA; QL (31 EA per 31 days)
<i>nabumetone</i>	T1	
<i>naloxone injection solution</i>	T2	
<i>naloxone injection syringe</i>	T2	
<i>naloxone nasal</i>	T3	
<i>naltrexone</i>	T2	
NAMENDA TITRATION PAK	T4	PA
NAMZARIC	T4	PA
NAPRELAN CR ORAL TABLET, ER MULTIPHASE 24 HR 750 MG	T4	
<i>naproxen oral suspension</i>	T1	
<i>naproxen oral tablet</i>	T1	
<i>naproxen oral tablet, delayed release (dr/ec)</i>	T2	
<i>naproxen sodium oral tablet 275 mg, 550 mg</i>	T1	
<i>naratriptan oral tablet 1 mg</i>	T2	QL (20 EA per 28 days)

Drug Name	Drug Tier	Requirements/Limits
<i>naratriptan oral tablet 2.5 mg</i>	T2	QL (9 EA per 28 days)
NARCAN	T3	
NAYZILAM	T4	PA-NS; QL (10 EA per 30 days)
<i>nefazodone</i>	T2	
NEUPRO	T4	
<i>nortriptyline</i>	T2	
NOURIANZ	T5	PA; QL (31 EA per 31 days)
NUCYNTA	T4	PA; QL (186 EA per 31 days)
NUEDEXTA	T3	PA; QL (62 EA per 31 days)
NUPLAZID	T5	PA-NS; QL (31 EA per 31 days)
NURTEC ODT	T5	QL (18 EA per 28 days)
<i>olanzapine intramuscular</i>	T2	
<i>olanzapine oral</i>	T2	QL (31 EA per 31 days)
<i>olanzapine-fluoxetine</i>	T2	
ONGENTYS	T4	PA; QL (31 EA per 31 days)
ONZETRA XSAIL	T5	QL (16 EA per 28 days)
<i>oxaprozin</i>	T2	
<i>oxazepam</i>	T4	
<i>oxcarbazepine</i>	T2	
OXTELLAR XR	T4	
<i>oxycodone oral capsule</i>	T2	PA; QL (186 EA per 31 days)
<i>oxycodone oral concentrate</i>	T2	PA; QL (180 ML per 31 days)
<i>oxycodone oral solution</i>	T2	PA; QL (4133 ML per 31 days)
<i>oxycodone oral tablet 10 mg, 15 mg, 20 mg, 5 mg</i>	T2	PA; QL (186 EA per 31 days)
<i>oxycodone oral tablet 30 mg</i>	T3	PA; QL (138 EA per 31 days)
<i>oxycodone-acetaminophen oral tablet 10-325 mg, 2.5-325 mg</i>	T3	PA; QL (372 EA per 31 days)
<i>oxycodone-acetaminophen oral tablet 5-325 mg, 7.5-325 mg</i>	T2	PA; QL (372 EA per 31 days)
<i>paliperidone oral tablet extended release 24hr 1.5 mg, 3 mg, 9 mg</i>	T4	QL (31 EA per 31 days)
<i>paliperidone oral tablet extended release 24hr 6 mg</i>	T4	QL (62 EA per 31 days)
<i>paroxetine hcl oral suspension</i>	T4	
<i>paroxetine hcl oral tablet</i>	T1	
<i>paroxetine hcl oral tablet extended release 24 hr</i>	T3	
<i>perphenazine</i>	T2	
<i>perphenazine-amitriptyline</i>	T2	PA-NS
PERSERIS	T5	QL (1 EA per 28 days)

Drug Name	Drug Tier	Requirements/Limits
<i>phenelzine</i>	T2	
<i>phenobarbital</i>	T2	PA-NS
PHENYTEK	T4	
<i>phenytoin oral suspension 125 mg/5 ml</i>	T2	
<i>phenytoin oral tablet, chewable</i>	T2	
<i>phenytoin sodium extended</i>	T2	
<i>pimozide</i>	T4	
<i>piroxicam</i>	T2	
PONVORY	T5	PA; QL (31 EA per 31 days)
PONVORY 14-DAY STARTER PACK	T5	PA; QL (28 EA per 365 days)
<i>pramipexole oral tablet</i>	T2	
<i>pregabalin oral capsule 100 mg, 150 mg, 200 mg, 25 mg, 50 mg, 75 mg</i>	T2	PA-NS; QL (93 EA per 31 days)
<i>pregabalin oral capsule 225 mg, 300 mg</i>	T2	PA-NS; QL (62 EA per 31 days)
<i>pregabalin oral solution</i>	T2	PA-NS; QL (930 ML per 31 days)
<i>pregabalin oral tablet extended release 24 hr</i>	T4	PA; QL (31 EA per 31 days)
<i>primidone</i>	T2	
PROCENTRA	T3	
<i>protriptyline</i>	T4	
<i>pyridostigmine bromide oral syrup</i>	T2	
<i>pyridostigmine bromide oral tablet 30 mg</i>	T3	
<i>pyridostigmine bromide oral tablet 60 mg</i>	T2	
<i>pyridostigmine bromide oral tablet extended release</i>	T2	
<i>quetiapine oral tablet 100 mg, 200 mg, 300 mg, 400 mg, 50 mg</i>	T2	QL (62 EA per 31 days)
<i>quetiapine oral tablet 25 mg</i>	T1	QL (62 EA per 31 days)
<i>quetiapine oral tablet extended release 24 hr</i>	T3	QL (62 EA per 31 days)
QUILLIVANT XR	T4	ST; QL (360 ML per 30 days)
RADICAVA ORS STARTER KIT SUSP	T5	PA; QL (70 ML per 28 days)
<i>ramelteon</i>	T4	QL (31 EA per 31 days)
<i>rasagiline</i>	T3	
REXULTI	T5	PA-NS; QL (31 EA per 31 days)
REYVOW ORAL TABLET 100 MG	T4	QL (8 EA per 28 days)
REYVOW ORAL TABLET 50 MG	T4	QL (4 EA per 28 days)
RISPERDAL CONSTA INTRAMUSCULAR SUSPENSION, EXTENDED REL RECON 12.5 MG/2 ML, 25 MG/2 ML, 37.5 MG/2 ML	T4	QL (2 EA per 28 days)

Drug Name	Drug Tier	Requirements/Limits
RISPERDAL CONSTA INTRAMUSCULAR SUSPENSION,EXTENDED REL RECON 50 MG/2 ML	T5	QL (2 EA per 28 days)
<i>risperidone oral solution</i>	T2	QL (496 ML per 31 days)
<i>risperidone oral tablet 0.25 mg, 0.5 mg, 1 mg, 2 mg</i>	T1	QL (31 EA per 31 days)
<i>risperidone oral tablet 3 mg</i>	T1	QL (93 EA per 31 days)
<i>risperidone oral tablet 4 mg</i>	T1	QL (124 EA per 31 days)
<i>risperidone oral tablet,disintegrating 0.25 mg, 0.5 mg, 1 mg, 2 mg</i>	T4	QL (31 EA per 31 days)
<i>risperidone oral tablet,disintegrating 3 mg</i>	T4	QL (93 EA per 31 days)
<i>risperidone oral tablet,disintegrating 4 mg</i>	T4	QL (124 EA per 31 days)
<i>rivastigmine</i>	T2	QL (30 EA per 30 days)
<i>rivastigmine tartrate</i>	T2	
<i>rizatriptan oral tablet 10 mg</i>	T2	QL (12 EA per 28 days)
<i>rizatriptan oral tablet 5 mg</i>	T2	QL (24 EA per 28 days)
<i>rizatriptan oral tablet,disintegrating 10 mg</i>	T2	QL (12 EA per 28 days)
<i>rizatriptan oral tablet,disintegrating 5 mg</i>	T2	QL (24 EA per 28 days)
<i>ropinirole oral tablet</i>	T2	
<i>ropinirole oral tablet extended release 24 hr</i>	T4	
ROWEEPRA ORAL TABLET 500 MG	T2	
<i>rufinamide</i>	T5	PA-NS
RYTARY	T3	ST
SECUADO	T5	QL (31 EA per 31 days)
<i>selegiline hcl</i>	T2	
<i>sertraline oral concentrate</i>	T1	
<i>sertraline oral tablet</i>	T1	
SPRITAM	T4	
<i>sulindac</i>	T2	
<i>sumatriptan nasal spray,non-aerosol 20 mg/actuation</i>	T2	QL (8 EA per 28 days)
<i>sumatriptan nasal spray,non-aerosol 5 mg/actuation</i>	T2	QL (32 EA per 28 days)
<i>sumatriptan succinate oral tablet 100 mg</i>	T2	QL (9 EA per 28 days)
<i>sumatriptan succinate oral tablet 25 mg</i>	T2	QL (36 EA per 28 days)
<i>sumatriptan succinate oral tablet 50 mg</i>	T2	QL (18 EA per 28 days)
<i>sumatriptan succinate subcutaneous cartridge 4 mg/0.5 ml</i>	T2	QL (6 ML per 28 days)
<i>sumatriptan succinate subcutaneous cartridge 6 mg/0.5 ml</i>	T4	QL (4 ML per 28 days)

Drug Name	Drug Tier	Requirements/Limits
<i>sumatriptan succinate subcutaneous pen injector 4 mg/0.5 ml</i>	T4	QL (6 ML per 28 days)
<i>sumatriptan succinate subcutaneous pen injector 6 mg/0.5 ml</i>	T4	QL (4 ML per 28 days)
<i>sumatriptan succinate subcutaneous solution</i>	T2	QL (4 ML per 28 days)
<i>sumatriptan-naproxen</i>	T4	QL (9 EA per 28 days)
SUNOSI	T4	PA; QL (31 EA per 31 days)
SYMPAZAN ORAL FILM 10 MG, 20 MG	T5	PA-NS; QL (62 EA per 31 days)
SYMPAZAN ORAL FILM 5 MG	T4	PA-NS; QL (62 EA per 31 days)
TEGRETOL ORAL SUSPENSION	T4	
TEGRETOL ORAL TABLET	T4	
TEGRETOL XR	T4	
TEGSEDI	T5	PA; QL (6 ML per 28 days)
<i>temazepam</i>	T2	QL (31 EA per 31 days)
TENCON	T2	QL (372 EA per 31 days)
<i>tetrabenazine oral tablet 12.5 mg</i>	T5	PA; QL (93 EA per 31 days)
<i>tetrabenazine oral tablet 25 mg</i>	T5	PA; QL (124 EA per 31 days)
<i>thioridazine</i>	T2	
<i>thiothixene</i>	T1	
<i>tiagabine oral tablet 12 mg, 16 mg</i>	T3	
<i>tiagabine oral tablet 2 mg, 4 mg</i>	T4	
<i>tizanidine</i>	T2	
<i>tolcapone</i>	T5	
<i>topiramate oral capsule, sprinkle</i>	T2	
<i>topiramate oral tablet</i>	T2	
<i>tramadol oral tablet 100 mg</i>	T4	PA; QL (124 EA per 31 days)
<i>tramadol oral tablet 50 mg</i>	T2	PA; QL (240 EA per 30 days)
<i>tramadol-acetaminophen</i>	T2	PA; QL (372 EA per 31 days)
<i>tranylcypromine</i>	T2	
<i>trazodone oral tablet 100 mg, 150 mg, 50 mg</i>	T1	
<i>trazodone oral tablet 300 mg</i>	T2	
<i>triazolam</i>	T4	PA
<i>trifluoperazine</i>	T2	
<i>trihexyphenidyl</i>	T2	
<i>trimipramine</i>	T3	PA-NS
TRINTELLIX	T3	PA-NS
TROKENDI XR	T4	
UBRELVY ORAL TABLET 100 MG	T5	QL (17 EA per 28 days)
UBRELVY ORAL TABLET 50 MG	T5	QL (34 EA per 28 days)

Drug Name	Drug Tier	Requirements/Limits
<i>valproic acid</i>	T2	
<i>valproic acid (as sodium salt) oral solution 250 mg/5 ml</i>	T2	
VALTOCO	T4	PA-NS; QL (10 EA per 30 days)
<i>venlafaxine oral capsule,extended release 24hr 150 mg, 37.5 mg</i>	T2	QL (31 EA per 31 days)
<i>venlafaxine oral capsule,extended release 24hr 75 mg</i>	T2	QL (93 EA per 31 days)
<i>venlafaxine oral tablet</i>	T2	
<i>venlafaxine oral tablet extended release 24hr</i>	T4	QL (31 EA per 31 days)
VERSACLOZ	T4	QL (558 ML per 31 days)
<i>vigabatrin</i>	T5	PA-NS
VIGADRONE	T5	PA-NS
VIIBRYD ORAL TABLETS,DOSE PACK 10 MG (7)- 20 MG (23)	T3	PA-NS; QL (60 EA per 365 days)
<i>vilazodone</i>	T3	PA-NS; QL (31 EA per 31 days)
VIVITROL	T5	
VRAYLAR ORAL CAPSULE	T5	PA-NS; QL (31 EA per 31 days)
VRAYLAR ORAL CAPSULE,DOSE PACK	T4	PA-NS; QL (14 EA per 365 days)
VTOL LQ	T4	QL (5723 ML per 31 days)
VUMERITY	T5	PA; QL (124 EA per 31 days)
VYVANSE ORAL CAPSULE	T4	ST; QL (31 EA per 31 days)
WAKIX	T5	PA; QL (62 EA per 31 days)
XCOPRI	T5	PA-NS
XCOPRI MAINTENANCE PACK ORAL TABLET 250MG/DAY(150 MG X1-100MG X1), 350 MG/DAY (200 MG X1-150MG X1)	T5	PA-NS
XCOPRI TITRATION PACK ORAL TABLETS,DOSE PACK 12.5 MG (14)- 25 MG (14)	T4	PA-NS
XCOPRI TITRATION PACK ORAL TABLETS,DOSE PACK 150 MG (14)- 200 MG (14), 50 MG (14)- 100 MG (14)	T5	PA-NS
XYREM	T5	PA; QL (540 ML per 30 days)
XYWAV	T5	PA; QL (540 ML per 30 days)
<i>zaleplon oral capsule 10 mg</i>	T2	PA; QL (62 EA per 31 days)
<i>zaleplon oral capsule 5 mg</i>	T2	PA; QL (93 EA per 31 days)
ZEBUTAL	T2	QL (372 EA per 31 days)
ZELAPAR	T5	
ZENZEDI ORAL TABLET 10 MG, 5 MG	T2	QL (62 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
ZENZEDI ORAL TABLET 15 MG, 2.5 MG, 20 MG, 30 MG, 7.5 MG	T4	QL (62 EA per 31 days)
ZEPOSIA	T5	PA; QL (31 EA per 31 days)
ZEPOSIA STARTER KIT	T5	PA; QL (74 EA per 365 days)
ZEPOSIA STARTER PACK	T5	PA; QL (14 EA per 365 days)
<i>ziprasidone hcl</i>	T3	QL (62 EA per 31 days)
<i>ziprasidone mesylate</i>	T3	
<i>zolmitriptan nasal spray,non-aerosol 5 mg</i>	T4	QL (8 EA per 28 days)
<i>zolmitriptan oral tablet 2.5 mg</i>	T2	QL (16 EA per 28 days)
<i>zolmitriptan oral tablet 5 mg</i>	T2	QL (8 EA per 28 days)
<i>zolmitriptan oral tablet,disintegrating 2.5 mg</i>	T2	QL (16 EA per 28 days)
<i>zolmitriptan oral tablet,disintegrating 5 mg</i>	T2	QL (8 EA per 28 days)
<i>zolpidem oral</i>	T2	PA; QL (31 EA per 31 days)
<i>zolpidem sublingual</i>	T3	PA; QL (31 EA per 31 days)
ZOMIG NASAL SPRAY,NON-AEROSOL 2.5 MG	T4	QL (16 EA per 28 days)
<i>zonisamide</i>	T2	
ZUBSOLV SUBLINGUAL TABLET 0.7-0.18 MG, 2.9-0.71 MG, 8.6-2.1 MG	T3	QL (62 EA per 31 days)
ZUBSOLV SUBLINGUAL TABLET 1.4-0.36 MG	T3	QL (93 EA per 31 days)
ZUBSOLV SUBLINGUAL TABLET 11.4-2.9 MG, 5.7-1.4 MG	T3	QL (31 EA per 31 days)
ZYPREXA RELPREVV INTRAMUSCULAR SUSPENSION FOR RECONSTITUTION 210 MG	T4	QL (2 EA per 28 days)
Cardiovascular, Hypertension / Lipids		
<i>acebutolol</i>	T1	
<i>aliskiren</i>	T4	
ALTOPREV ORAL TABLET EXTENDED RELEASE 24 HR 20 MG, 60 MG	T4	
ALTOPREV ORAL TABLET EXTENDED RELEASE 24 HR 40 MG	T5	
<i>amiloride</i>	T1	
<i>amiloride-hydrochlorothiazide</i>	T1	
<i>amiodarone oral</i>	T2	
<i>amlodipine</i>	T1	
<i>amlodipine-atorvastatin</i>	T2	
<i>amlodipine-benazepril</i>	T1	
<i>amlodipine-olmesartan</i>	T3	QL (31 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
<i>amlodipine-valsartan</i>	T2	
ANTARA ORAL CAPSULE 30 MG, 90 MG	T4	
<i>aspirin-dipyridamole</i>	T2	
<i>atenolol</i>	T1	
<i>atenolol-chlorthalidone</i>	T1	
<i>atorvastatin</i>	T1	
<i>benazepril</i>	T1	
<i>benazepril-hydrochlorothiazide</i>	T1	
<i>betaxolol oral</i>	T1	
<i>bisoprolol fumarate</i>	T1	
<i>bisoprolol-hydrochlorothiazide</i>	T1	
BRILINTA	T3	
<i>bumetanide</i>	T1	
CABLIVI INJECTION KIT	T5	PA; QL (31 EA per 31 days)
CAMZYOS	T5	PA; QL (31 EA per 31 days)
<i>candesartan</i>	T1	
<i>candesartan-hydrochlorothiazid</i>	T1	
<i>captopril</i>	T1	
CARDIZEM LA ORAL TABLET EXTENDED RELEASE 24 HR 120 MG	T4	
CARTIA XT	T1	
<i>carvedilol</i>	T1	
<i>carvedilol phosphate</i>	T4	
<i>chlorthalidone oral tablet 25 mg, 50 mg</i>	T1	
<i>cholestyramine (with sugar) oral powder in packet</i>	T2	
CHOLESTYRAMINE LIGHT ORAL POWDER IN PACKET	T2	
<i>cilostazol</i>	T2	
<i>clonidine</i>	T2	
<i>clonidine hcl oral tablet</i>	T1	
<i>clopidogrel oral tablet 75 mg</i>	T1	
<i>colesevelam</i>	T3	
<i>colestipol oral packet</i>	T2	
<i>colestipol oral tablet</i>	T2	
CORLANOR ORAL SOLUTION	T4	PA; QL (420 ML per 28 days)
CORLANOR ORAL TABLET 5 MG	T4	PA; QL (93 EA per 31 days)
CORLANOR ORAL TABLET 7.5 MG	T4	PA; QL (62 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
DIGITEK ORAL TABLET 125 MCG (0.125 MG)	T1	QL (62 EA per 31 days)
DIGITEK ORAL TABLET 250 MCG (0.25 MG)	T2	QL (31 EA per 31 days)
<i>digoxin oral solution</i>	T3	QL (155 ML per 31 days)
<i>digoxin oral tablet 125 mcg (0.125 mg)</i>	T1	QL (62 EA per 31 days)
<i>digoxin oral tablet 250 mcg (0.25 mg)</i>	T2	QL (31 EA per 31 days)
<i>digoxin oral tablet 62.5 mcg (0.0625 mg)</i>	T4	QL (124 EA per 31 days)
<i>diltiazem hcl oral capsule,extended release 12 hr</i>	T1	
<i>diltiazem hcl oral capsule,extended release 24 hr 360 mg, 420 mg</i>	T1	
<i>diltiazem hcl oral capsule,extended release 24hr 120 mg, 180 mg, 240 mg, 300 mg</i>	T1	
<i>diltiazem hcl oral tablet</i>	T1	
<i>diltiazem hcl oral tablet extended release 24 hr 180 mg, 240 mg, 300 mg, 360 mg</i>	T1	
DILT-XR	T1	
<i>dofetilide</i>	T3	
DOPTELET (10 TAB PACK)	T5	PA
DOPTELET (15 TAB PACK)	T5	PA
DOPTELET (30 TAB PACK)	T5	PA
<i>doxazosin</i>	T1	
EDARBYCLOR	T4	
ELIQUIS DVT-PE TREAT 30D START	T3	QL (74 EA per 30 days)
ELIQUIS ORAL TABLET 2.5 MG	T3	QL (60 EA per 30 days)
ELIQUIS ORAL TABLET 5 MG	T3	QL (74 EA per 30 days)
<i>enalapril maleate oral solution</i>	T4	
<i>enalapril maleate oral tablet</i>	T1	
<i>enalapril-hydrochlorothiazide</i>	T1	
<i>enoxaparin subcutaneous syringe 100 mg/ml, 120 mg/0.8 ml, 150 mg/ml</i>	T4	
<i>enoxaparin subcutaneous syringe 30 mg/0.3 ml, 40 mg/0.4 ml, 60 mg/0.6 ml, 80 mg/0.8 ml</i>	T2	
ENTRESTO ORAL TABLET 24-26 MG	T3	QL (186 EA per 31 days)
ENTRESTO ORAL TABLET 49-51 MG	T3	QL (93 EA per 31 days)
ENTRESTO ORAL TABLET 97-103 MG	T3	QL (62 EA per 31 days)
<i>eplerenone</i>	T2	
<i>ethacrynic acid</i>	T4	
<i>ezetimibe</i>	T2	
<i>ezetimibe-simvastatin</i>	T3	

Drug Name	Drug Tier	Requirements/Limits
<i>felodipine</i>	T2	
<i>fenofibrate micronized oral capsule 134 mg, 200 mg, 43 mg, 67 mg</i>	T2	
<i>fenofibrate nanocrystallized</i>	T2	
<i>fenofibrate oral tablet 120 mg, 40 mg</i>	T4	
<i>fenofibrate oral tablet 160 mg, 54 mg</i>	T2	
<i>fenofibric acid (choline)</i>	T3	
<i>flecainide</i>	T2	
<i>fluvastatin oral capsule</i>	T1	
<i>fluvastatin oral tablet extended release 24 hr</i>	T3	
<i>fondaparinux subcutaneous syringe 10 mg/0.8 ml, 5 mg/0.4 ml, 7.5 mg/0.6 ml</i>	T5	
<i>fondaparinux subcutaneous syringe 2.5 mg/0.5 ml</i>	T4	
<i>fosinopril</i>	T1	
<i>fosinopril-hydrochlorothiazide</i>	T1	
FRAGMIN SUBCUTANEOUS SOLUTION 25,000 ANTI-XA UNIT/ML	T5	
FRAGMIN SUBCUTANEOUS SYRINGE 10,000 ANTI-XA UNIT/ML, 12,500 ANTI-XA UNIT/0.5 ML, 18,000 ANTI-XA UNIT/0.72 ML, 7,500 ANTI-XA UNIT/0.3 ML	T5	
FRAGMIN SUBCUTANEOUS SYRINGE 15,000 ANTI-XA UNIT/0.6 ML, 2,500 ANTI-XA UNIT/0.2 ML, 5,000 ANTI-XA UNIT/0.2 ML	T3	
<i>furosemide injection</i>	T2	
<i>furosemide oral solution 10 mg/ml, 40 mg/5 ml (8 mg/ml)</i>	T2	
<i>furosemide oral tablet</i>	T1	
<i>gemfibrozil</i>	T1	
<i>heparin (porcine) injection solution</i>	T2	
<i>hydralazine oral</i>	T1	
<i>hydrochlorothiazide</i>	T1	
<i>icosapent ethyl oral capsule 0.5 gram</i>	T4	QL (248 EA per 31 days)
<i>icosapent ethyl oral capsule 1 gram</i>	T4	QL (124 EA per 31 days)
<i>indapamide</i>	T1	
<i>irbesartan</i>	T1	QL (31 EA per 31 days)
<i>irbesartan-hydrochlorothiazide</i>	T2	QL (31 EA per 31 days)
ISORDIL	T4	
<i>isosorbide dinitrate oral tablet</i>	T2	
<i>isosorbide mononitrate</i>	T1	

Drug Name	Drug Tier	Requirements/Limits
<i>isosorbide-hydralazine</i>	T4	
<i>isradipine</i>	T2	
JANTOVEN	T1	
JUXTAPID ORAL CAPSULE 10 MG, 20 MG, 30 MG, 5 MG	T5	PA
<i>labetalol oral</i>	T1	
LIPOFEN	T4	
<i>lisinopril</i>	T1	
<i>lisinopril-hydrochlorothiazide</i>	T1	
LIVALO	T4	
<i>losartan oral tablet 100 mg</i>	T1	QL (31 EA per 31 days)
<i>losartan oral tablet 25 mg</i>	T1	QL (93 EA per 31 days)
<i>losartan oral tablet 50 mg</i>	T1	QL (62 EA per 31 days)
<i>losartan-hydrochlorothiazide</i>	T1	
<i>lovastatin</i>	T1	
MATZIM LA	T2	
<i>metolazone</i>	T2	
<i>metoprolol succinate</i>	T1	
<i>metoprolol ta-hydrochlorothiaz</i>	T1	
<i>metoprolol tartrate oral</i>	T1	
<i>metyrosine</i>	T3	
<i>mexiletine</i>	T3	
<i>minoxidil oral</i>	T2	
<i>moexipril</i>	T1	
MULPLETA	T5	PA
MULTAQ	T4	
<i>nadolol</i>	T1	
<i>nebivolol oral tablet 10 mg, 2.5 mg</i>	T4	QL (93 EA per 31 days)
<i>nebivolol oral tablet 20 mg</i>	T4	QL (62 EA per 31 days)
<i>nebivolol oral tablet 5 mg</i>	T4	QL (217 EA per 31 days)
NEXLETOL	T4	PA; QL (31 EA per 31 days)
NEXLIZET	T4	PA; QL (31 EA per 31 days)
<i>niacin oral tablet 500 mg</i>	T5	
<i>niacin oral tablet extended release 24 hr 1,000 mg, 750 mg</i>	T3	
<i>niacin oral tablet extended release 24 hr 500 mg</i>	T3	QL (31 EA per 31 days)
NIACOR	T4	
<i>nicardipine oral</i>	T2	
<i>nifedipine oral tablet extended release</i>	T2	

Drug Name	Drug Tier	Requirements/Limits
<i>nifedipine oral tablet extended release 24hr</i>	T2	
<i>nimodipine</i>	T2	
<i>nisoldipine</i>	T2	
NITRO-BID	T2	
<i>nitroglycerin sublingual</i>	T2	
<i>nitroglycerin transdermal patch 24 hour</i>	T2	
<i>nitroglycerin translingual</i>	T2	
NITROSTAT	T4	
NYMALIZE ORAL SYRINGE 60 MG/10 ML	T4	
<i>olmesartan oral tablet 20 mg, 40 mg</i>	T2	QL (31 EA per 31 days)
<i>olmesartan oral tablet 5 mg</i>	T2	QL (93 EA per 31 days)
<i>olmesartan-amlodipin-hcthiazid</i>	T3	
<i>olmesartan-hydrochlorothiazide</i>	T2	QL (31 EA per 31 days)
<i>omega-3 acid ethyl esters</i>	T3	QL (124 EA per 31 days)
ORENITRAM ORAL TABLET EXTENDED RELEASE 0.125 MG	T4	PA; QL (93 EA per 31 days)
ORENITRAM ORAL TABLET EXTENDED RELEASE 0.25 MG	T4	PA; QL (186 EA per 31 days)
ORENITRAM ORAL TABLET EXTENDED RELEASE 1 MG	T5	PA; QL (186 EA per 31 days)
ORENITRAM ORAL TABLET EXTENDED RELEASE 2.5 MG	T5	PA; QL (521 EA per 31 days)
ORENITRAM ORAL TABLET EXTENDED RELEASE 5 MG	T5	PA; QL (261 EA per 31 days)
PACERONE ORAL TABLET 100 MG, 200 MG, 400 MG	T2	
<i>pentoxifylline</i>	T2	
<i>perindopril erbumine</i>	T1	
<i>phenoxybenzamine</i>	T5	PA
<i>pindolol</i>	T1	
PRALUENT PEN	T4	PA; QL (2 ML per 28 days)
<i>prasugrel</i>	T3	
<i>pravastatin</i>	T1	
<i>prazosin</i>	T1	
PREVALITE ORAL POWDER IN PACKET	T2	
PROMACTA ORAL POWDER IN PACKET 12.5 MG	T5	PA; QL (372 EA per 31 days)
PROMACTA ORAL POWDER IN PACKET 25 MG	T5	PA; QL (31 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
PROMACTA ORAL TABLET 12.5 MG, 25 MG	T5	PA; QL (31 EA per 31 days)
PROMACTA ORAL TABLET 50 MG, 75 MG	T5	PA; QL (62 EA per 31 days)
<i>propafenone</i>	T2	
<i>propranolol oral capsule,extended release 24 hr</i>	T2	
<i>propranolol oral solution</i>	T1	
<i>propranolol oral tablet</i>	T1	
<i>quinapril</i>	T1	
<i>quinapril-hydrochlorothiazide</i>	T1	
<i>quinidine gluconate oral</i>	T4	
<i>quinidine sulfate oral tablet</i>	T2	
<i>ramipril</i>	T1	
<i>ranolazine</i>	T3	QL (62 EA per 31 days)
REPATHA PUSHTRONEX	T3	PA; QL (7 ML per 28 days)
REPATHA SURECLICK	T3	PA; QL (3 ML per 28 days)
REPATHA SYRINGE	T3	PA; QL (3 ML per 28 days)
<i>rosuvastatin</i>	T1	
<i>simvastatin oral tablet</i>	T1	
SORINE	T1	
SOTALOL AF	T1	
<i>sotalol oral</i>	T1	
<i>spironolactone</i>	T1	
<i>spironolacton-hydrochlorothiaz</i>	T1	
TAVALISSE	T5	PA; QL (62 EA per 31 days)
TAZTIA XT	T2	
<i>telmisartan</i>	T1	
<i>telmisartan-amlodipine</i>	T1	
<i>telmisartan-hydrochlorothiazid</i>	T1	
<i>terazosin</i>	T1	
TIADYLT ER	T2	
<i>timolol maleate oral</i>	T1	
<i>torseamide oral</i>	T1	
<i>trandolapril</i>	T1	
<i>triamterene</i>	T4	
<i>triamterene-hydrochlorothiazid</i>	T1	
UPTRAVI ORAL TABLET 1,000 MCG, 1,200 MCG, 1,400 MCG, 1,600 MCG, 400 MCG, 600 MCG, 800 MCG	T5	PA; QL (62 EA per 31 days)
UPTRAVI ORAL TABLET 200 MCG	T5	PA; QL (224 EA per 28 days)

Drug Name	Drug Tier	Requirements/Limits
UPTRAVI ORAL TABLETS,DOSE PACK	T5	PA; QL (400 EA per 365 days)
<i>valsartan oral tablet 160 mg, 40 mg, 80 mg</i>	T1	QL (62 EA per 31 days)
<i>valsartan oral tablet 320 mg</i>	T1	QL (31 EA per 31 days)
<i>valsartan-hydrochlorothiazide</i>	T2	QL (31 EA per 31 days)
VASCEPA ORAL CAPSULE 0.5 GRAM	T4	QL (248 EA per 31 days)
VASCEPA ORAL CAPSULE 1 GRAM	T4	QL (124 EA per 31 days)
VECAMYL	T4	
<i>verapamil oral</i>	T2	
VERQUVO	T4	PA; QL (31 EA per 31 days)
VYNDAMAX	T5	PA; QL (31 EA per 31 days)
VYNDAQEL	T5	PA; QL (124 EA per 31 days)
<i>warfarin</i>	T1	
XARELTO DVT-PE TREAT 30D START	T3	QL (51 EA per 30 days)
XARELTO ORAL SUSPENSION FOR RECONSTITUTION	T3	PA; QL (930 ML per 31 days)
XARELTO ORAL TABLET 10 MG, 20 MG	T3	QL (31 EA per 31 days)
XARELTO ORAL TABLET 15 MG	T3	QL (52 EA per 31 days)
XARELTO ORAL TABLET 2.5 MG	T3	QL (62 EA per 31 days)
ZONTIVITY	T4	
Dermatologicals/Topical Therapy		
ACANYA TOPICAL GEL WITH PUMP	T4	
ACCUTANE	T2	
<i>acitretin</i>	T4	PA
<i>acyclovir topical cream</i>	T3	
<i>acyclovir topical ointment</i>	T1	QL (30 GM per 30 days)
<i>adapalene topical gel 0.3 %</i>	T2	PA
<i>adapalene topical swab</i>	T2	PA
<i>adapalene-benzoyl peroxide</i>	T4	
ADBRY	T5	PA; QL (4 ML per 28 days)
ALA-CORT TOPICAL CREAM 1 %	T1	
ALA-CORT TOPICAL CREAM 2.5 %	T1	QL (30 GM per 28 days)
<i>alclometasone</i>	T1	
<i>amcinonide</i>	T2	
<i>ammonium lactate</i>	T2	
AMNESTEEM	T2	
AVITA	T4	PA; QL (45 GM per 28 days)
<i>azelaic acid</i>	T4	QL (50 GM per 28 days)
AZELEX	T4	QL (50 GM per 28 days)

Drug Name	Drug Tier	Requirements/Limits
<i>betamethasone dipropionate</i>	T1	
<i>betamethasone valerate topical cream</i>	T1	
<i>betamethasone valerate topical foam</i>	T4	
<i>betamethasone valerate topical lotion</i>	T1	
<i>betamethasone valerate topical ointment</i>	T1	
<i>betamethasone, augmented</i>	T2	
<i>calcipotriene scalp</i>	T2	QL (60 ML per 28 days)
<i>calcipotriene topical cream</i>	T2	QL (60 GM per 28 days)
<i>calcipotriene topical ointment</i>	T2	QL (60 GM per 28 days)
<i>calcipotriene-betamethasone topical ointment</i>	T4	ST; QL (400 GM per 28 days)
<i>calcipotriene-betamethasone topical suspension</i>	T5	ST; QL (400 GM per 28 days)
<i>calcitriol topical</i>	T4	ST
CAPEX	T4	QL (840 ML per 28 days)
CARAC	T5	PA
CIBINQO	T5	PA; QL (31 EA per 31 days)
<i>ciclopirox topical cream</i>	T2	QL (90 GM per 28 days)
<i>ciclopirox topical gel</i>	T2	QL (45 GM per 28 days)
<i>ciclopirox topical shampoo</i>	T2	QL (120 ML per 28 days)
<i>ciclopirox topical solution</i>	T2	
<i>ciclopirox topical suspension</i>	T3	QL (60 ML per 28 days)
CLARAVIS	T4	
<i>clindamycin phosphate topical foam</i>	T4	QL (100 GM per 28 days)
<i>clindamycin phosphate topical gel</i>	T2	QL (75 GM per 28 days)
<i>clindamycin phosphate topical lotion</i>	T2	QL (60 ML per 28 days)
<i>clindamycin phosphate topical solution</i>	T2	QL (60 ML per 28 days)
<i>clindamycin phosphate topical swab</i>	T2	
<i>clindamycin-benzoyl peroxide topical gel</i>	T2	
<i>clindamycin-benzoyl peroxide topical gel with pump 1.2-2.5 %</i>	T2	
<i>clindamycin-tretinoin</i>	T2	PA; QL (60 GM per 28 days)
<i>clobetasol scalp</i>	T2	QL (50 ML per 28 days)
<i>clobetasol topical cream</i>	T3	QL (60 GM per 28 days)
<i>clobetasol topical foam</i>	T2	QL (100 GM per 28 days)
<i>clobetasol topical gel</i>	T2	QL (60 GM per 28 days)
<i>clobetasol topical lotion</i>	T2	QL (118 ML per 28 days)
<i>clobetasol topical ointment</i>	T3	QL (60 GM per 28 days)
<i>clobetasol topical shampoo</i>	T4	QL (118 ML per 28 days)
<i>clobetasol topical spray,non-aerosol</i>	T2	QL (125 ML per 28 days)

Drug Name	Drug Tier	Requirements/Limits
<i>clobetasol-emollient topical cream</i>	T3	QL (60 GM per 28 days)
<i>clobetasol-emollient topical foam</i>	T3	QL (100 GM per 28 days)
<i>clocortolone pivalate</i>	T4	
CLODAN	T2	QL (118 ML per 28 days)
<i>clotrimazole topical cream</i>	T2	QL (45 GM per 28 days)
<i>clotrimazole topical solution</i>	T2	QL (30 ML per 28 days)
<i>clotrimazole-betamethasone topical cream</i>	T2	QL (45 GM per 28 days)
<i>clotrimazole-betamethasone topical lotion</i>	T2	QL (60 ML per 28 days)
COSENTYX (2 SYRINGES)	T5	PA; QL (2 ML per 28 days)
COSENTYX PEN (2 PENS)	T5	PA; QL (2 ML per 28 days)
COSENTYX SUBCUTANEOUS SYRINGE 75 MG/0.5 ML	T5	PA; QL (0.5 ML per 28 days)
CROTAN	T4	
<i>dapsone topical</i>	T4	QL (90 GM per 28 days)
DENAVIR	T3	
<i>desonide topical cream</i>	T3	QL (60 GM per 28 days)
<i>desonide topical gel</i>	T3	QL (60 GM per 28 days)
<i>desonide topical lotion</i>	T3	QL (118 ML per 28 days)
<i>desonide topical ointment</i>	T3	QL (60 GM per 28 days)
<i>desoximetasone topical cream</i>	T2	QL (100 GM per 28 days)
<i>desoximetasone topical gel</i>	T2	QL (60 GM per 28 days)
<i>desoximetasone topical ointment 0.25 %</i>	T2	QL (100 GM per 28 days)
<i>desoximetasone topical spray,non-aerosol</i>	T2	QL (100 ML per 28 days)
DESRX	T3	QL (60 GM per 28 days)
<i>diclofenac sodium topical gel 3 %</i>	T4	PA; QL (100 GM per 28 days)
DIFFERIN TOPICAL LOTION	T4	PA
<i>doxepin topical</i>	T4	PA; QL (45 GM per 28 days)
DUOBRII	T4	PA; QL (200 GM per 28 days)
DUPIXENT PEN SUBCUTANEOUS PEN INJECTOR 200 MG/1.14 ML	T5	PA; QL (2.28 ML per 28 days)
DUPIXENT PEN SUBCUTANEOUS PEN INJECTOR 300 MG/2 ML	T5	PA; QL (4 ML per 28 days)
DUPIXENT SYRINGE SUBCUTANEOUS SYRINGE 100 MG/0.67 ML	T5	PA; QL (1.34 ML per 28 days)
DUPIXENT SYRINGE SUBCUTANEOUS SYRINGE 200 MG/1.14 ML	T5	PA; QL (2.28 ML per 28 days)
DUPIXENT SYRINGE SUBCUTANEOUS SYRINGE 300 MG/2 ML	T5	PA; QL (8 ML per 28 days)
<i>econazole</i>	T2	QL (85 GM per 28 days)
ERY PADS	T2	

Drug Name	Drug Tier	Requirements/Limits
ERYGEL	T3	QL (60 GM per 28 days)
<i>erythromycin with ethanol topical gel</i>	T2	QL (60 GM per 28 days)
<i>erythromycin with ethanol topical solution</i>	T2	QL (60 ML per 28 days)
<i>erythromycin-benzoyl peroxide</i>	T2	
FINACEA TOPICAL FOAM	T4	QL (50 GM per 28 days)
<i>fluocinolone and shower cap</i>	T2	QL (118.28 ML per 28 days)
<i>fluocinolone topical cream 0.01 %</i>	T2	QL (60 GM per 28 days)
<i>fluocinolone topical cream 0.025 %</i>	T2	QL (120 GM per 28 days)
<i>fluocinolone topical ointment</i>	T2	QL (120 GM per 28 days)
<i>fluocinolone topical solution</i>	T2	QL (90 ML per 28 days)
<i>fluocinonide topical cream 0.05 %</i>	T2	QL (60 GM per 28 days)
<i>fluocinonide topical gel</i>	T2	QL (60 GM per 28 days)
<i>fluocinonide topical ointment</i>	T2	QL (60 GM per 28 days)
<i>fluocinonide topical solution</i>	T2	QL (60 ML per 28 days)
FLUOCINONIDE-E	T2	QL (60 GM per 28 days)
<i>fluorouracil topical cream 0.5 %</i>	T5	
<i>fluorouracil topical cream 5 %</i>	T2	
<i>fluorouracil topical solution</i>	T2	
<i>flurandrenolide topical cream</i>	T4	QL (120 GM per 28 days)
<i>flurandrenolide topical lotion</i>	T4	QL (120 ML per 28 days)
<i>fluticasone propionate topical cream</i>	T2	
<i>gentamicin topical</i>	T2	QL (60 GM per 28 days)
<i>halcinonide</i>	T4	
<i>halobetasol propionate topical cream</i>	T2	QL (50 GM per 28 days)
<i>halobetasol propionate topical ointment</i>	T2	QL (50 GM per 28 days)
HALOG TOPICAL OINTMENT	T4	
<i>hydrocortisone butyrate topical cream</i>	T2	QL (45 GM per 28 days)
<i>hydrocortisone butyrate topical lotion</i>	T4	QL (118 ML per 28 days)
<i>hydrocortisone butyrate topical ointment</i>	T2	QL (45 GM per 28 days)
<i>hydrocortisone butyrate topical solution</i>	T2	QL (60 ML per 28 days)
<i>hydrocortisone topical cream 1 %</i>	T1	
<i>hydrocortisone topical lotion 2.5 %</i>	T1	QL (118 ML per 28 days)
<i>hydrocortisone topical ointment 1 %, 2.5 %</i>	T1	
<i>hydrocortisone valerate</i>	T2	QL (60 GM per 28 days)
<i>imiquimod topical cream in packet 3.75 %</i>	T5	
<i>imiquimod topical cream in packet 5 %</i>	T2	
<i>isotretinoin</i>	T2	
<i>ketoconazole topical cream</i>	T2	QL (60 GM per 28 days)

Drug Name	Drug Tier	Requirements/Limits
<i>ketoconazole topical shampoo</i>	T2	
<i>lidocaine hcl mucous membrane solution 4 % (40 mg/ml)</i>	T2	PA; QL (50 ML per 28 days)
<i>lidocaine topical adhesive patch,medicated 5 %</i>	T3	PA; QL (93 EA per 31 days)
<i>lidocaine topical ointment</i>	T2	PA; QL (50 GM per 28 days)
LIDOCAINE VISCOUS	T2	
<i>lidocaine-prilocaine topical cream</i>	T2	PA; QL (30 GM per 28 days)
LIDODERM	T4	PA; QL (93 EA per 31 days)
<i>malathion</i>	T4	
MENTAX	T4	
<i>metronidazole topical cream</i>	T2	
<i>metronidazole topical gel 0.75 %</i>	T2	
<i>metronidazole topical gel 1 %</i>	T1	
<i>metronidazole topical lotion</i>	T2	
<i>mometasone topical</i>	T2	
<i>mupirocin</i>	T2	
MYORISAN	T2	
<i>naftifine topical cream 1 %</i>	T4	ST; QL (90 GM per 28 days)
<i>naftifine topical cream 2 %</i>	T4	ST; QL (60 GM per 28 days)
NAFTIN TOPICAL GEL 2 %	T4	ST; QL (60 GM per 28 days)
NEO-SYNALAR	T4	
NEUAC	T2	
NYAMYC	T2	QL (60 GM per 28 days)
<i>nystatin topical cream</i>	T2	QL (30 GM per 28 days)
<i>nystatin topical ointment</i>	T2	QL (30 GM per 28 days)
<i>nystatin topical powder</i>	T2	QL (60 GM per 28 days)
<i>nystatin-triamcinolone</i>	T3	
NYSTOP	T2	QL (60 GM per 28 days)
OPZELURA	T5	PA; QL (60 GM per 7 days)
OXISTAT TOPICAL LOTION	T4	ST; QL (60 ML per 28 days)
PANDEL	T4	
PANRETIN	T5	PA-NS
<i>permethrin</i>	T2	
<i>pimecrolimus</i>	T3	QL (100 GM per 28 days)
<i>podofilox</i>	T2	
<i>prednicarbate topical ointment</i>	T2	QL (60 GM per 28 days)
REGRANEX	T5	PA
RETIN-A MICRO PUMP TOPICAL GEL WITH PUMP 0.06 %	T5	PA; QL (50 GM per 28 days)

Drug Name	Drug Tier	Requirements/Limits
SANTYL	T3	QL (180 GM per 30 days)
<i>selenium sulfide topical lotion</i>	T1	
SILIQ	T5	PA; QL (6 ML per 28 days)
<i>silver sulfadiazine</i>	T1	
SKYRIZI SUBCUTANEOUS PEN INJECTOR	T5	PA; QL (1 ML per 84 days)
SKYRIZI SUBCUTANEOUS SYRINGE 150 MG/ML	T5	PA; QL (1 ML per 84 days)
SKYRIZI SUBCUTANEOUS SYRINGE KIT	T5	PA; QL (1 EA per 84 days)
SSD	T4	
STELARA SUBCUTANEOUS SOLUTION	T5	PA; QL (0.5 ML per 84 days)
STELARA SUBCUTANEOUS SYRINGE 45 MG/0.5 ML	T5	PA; QL (0.5 ML per 84 days)
STELARA SUBCUTANEOUS SYRINGE 90 MG/ML	T5	PA; QL (1 ML per 56 days)
<i>sulfacetamide sodium (acne)</i>	T1	
SULFAMYLON TOPICAL CREAM	T3	
<i>tacrolimus topical</i>	T2	QL (100 GM per 28 days)
TALTZ AUTOINJECTOR	T5	PA; QL (1 ML per 28 days)
TALTZ SYRINGE	T5	PA; QL (1 ML per 28 days)
<i>tazarotene topical cream</i>	T4	PA; QL (60 GM per 28 days)
<i>tazarotene topical gel</i>	T4	PA; QL (100 GM per 28 days)
TAZORAC TOPICAL CREAM 0.05 %	T4	PA; QL (60 GM per 28 days)
TOVET EMOLLIENT	T3	QL (100 GM per 28 days)
<i>tretinoin topical cream</i>	T2	PA; QL (45 GM per 28 days)
<i>tretinoin topical gel 0.01 %, 0.025 %</i>	T2	PA; QL (45 GM per 28 days)
<i>tretinoin topical gel 0.05 %</i>	T3	PA; QL (45 GM per 28 days)
<i>triamcinolone acetonide topical aerosol</i>	T4	
<i>triamcinolone acetonide topical cream</i>	T1	
<i>triamcinolone acetonide topical lotion</i>	T1	
<i>triamcinolone acetonide topical ointment 0.025 %, 0.1 %, 0.5 %</i>	T1	
<i>triamcinolone acetonide topical ointment 0.05 %</i>	T4	
TRIANEX	T4	
TRIDERM TOPICAL CREAM	T1	
TRITOCIN	T4	
VALCHLOR	T5	PA-NS
VEREGEN	T4	
ZENATANE ORAL CAPSULE 30 MG	T2	

Drug Name	Drug Tier	Requirements/Limits
ZYCLARA TOPICAL CREAM IN METERED-DOSE PUMP 2.5 %	T5	
ZYCLARA TOPICAL CREAM IN PACKET	T5	
Diagnostics / Miscellaneous Agents		
<i>acamprosate</i>	T2	
<i>anagrelide</i>	T3	
ARALAST NP INTRAVENOUS RECON SOLN 1,000 MG	T5	PA
AURYXIA	T5	PA; QL (372 EA per 31 days)
<i>bupropion hcl (smoking deter)</i>	T2	QL (62 EA per 31 days)
<i>carglumic acid</i>	T5	PA
<i>cevimeline</i>	T2	
CHEMET	T3	
CLINIMIX 4.25%/D5W SULFIT FREE	T4	PA-BvD
CLINIMIX E 2.75%/D5W SULF FREE	T4	PA-BvD
<i>d10 %-0.45 % sodium chloride</i>	T2	
<i>d2.5 %-0.45 % sodium chloride</i>	T2	
<i>d5 % and 0.9 % sodium chloride</i>	T2	
<i>d5 %-0.45 % sodium chloride</i>	T2	
<i>deferasirox oral tablet, dispersible 125 mg</i>	T4	PA
<i>deferasirox oral tablet, dispersible 250 mg, 500 mg</i>	T5	PA
<i>deferiprone</i>	T5	PA
<i>dextrose 10 % in water (d10w)</i>	T2	
<i>dextrose 5 % in water (d5w) intravenous parenteral solution</i>	T2	
<i>dextrose 5%-0.2 % sod chloride</i>	T2	
<i>disulfiram</i>	T2	
<i>droxidopa</i>	T5	PA
ENDARI	T4	PA; QL (180 EA per 30 days)
EXSERVAN	T5	PA; QL (62 EA per 31 days)
FERRIPROX ORAL SOLUTION	T5	PA
FOSRENOL ORAL POWDER IN PACKET	T5	
GLASSIA	T5	PA
INCRELEX	T5	PA
<i>lanthanum oral tablet, chewable 1,000 mg, 750 mg</i>	T4	
<i>lanthanum oral tablet, chewable 500 mg</i>	T5	
<i>levocarnitine (with sugar)</i>	T2	PA-BvD

Drug Name	Drug Tier	Requirements/Limits
<i>levocarnitine oral tablet</i>	T2	PA-BvD
LITHOSTAT	T4	
LOKELMA	T3	PA; QL (93 EA per 31 days)
<i>midodrine</i>	T2	
NICOTROL	T3	
NICOTROL NS	T4	
<i>nitisinone</i>	T5	
NITYR	T5	
ORFADIN ORAL CAPSULE 20 MG	T5	
ORFADIN ORAL SUSPENSION	T5	
OXBRYTA ORAL TABLET	T5	PA; QL (155 EA per 31 days)
OXBRYTA ORAL TABLET FOR SUSPENSION	T5	PA; QL (248 EA per 31 days)
PHEBURANE	T5	PA; QL (620 GM per 31 days)
<i>pilocarpine hcl oral</i>	T2	
PROLASTIN-C INTRAVENOUS RECON SOLN	T5	PA
PYRUKYND ORAL TABLET 20 MG, 50 MG	T5	PA; QL (62 EA per 31 days)
PYRUKYND ORAL TABLET 5 MG, 5 MG (4-WEEK PACK)	T5	PA
PYRUKYND ORAL TABLETS,DOSE PACK	T5	PA; QL (2 EA per 365 days)
RAVICTI	T5	PA
REVCOVI	T5	
<i>riluzole</i>	T4	
<i>risedronate oral tablet 30 mg</i>	T2	
<i>sevelamer carbonate</i>	T3	
<i>sevelamer hcl</i>	T4	
<i>sodium chloride 0.9 % intravenous parenteral solution</i>	T2	
<i>sodium chloride irrigation</i>	T2	
<i>sodium phenylbutyrate</i>	T5	
<i>sodium polystyrene sulfonate oral powder</i>	T2	
SPS (WITH SORBITOL) ORAL	T2	
TAVNEOS	T5	PA; QL (186 EA per 31 days)
THIOLA EC	T5	PA
TIGLUTIK	T5	PA
<i>tiopronin</i>	T5	PA
<i>trientine</i>	T3	QL (248 EA per 31 days)
<i>varenicline oral tablet</i>	T4	QL (60 EA per 30 days)

Drug Name	Drug Tier	Requirements/Limits
<i>varenicline oral tablets,dose pack</i>	T4	QL (106 EA per 365 days)
VELPHORO	T5	
VELTASSA	T3	PA; QL (30 EA per 30 days)
XURIDEN	T5	PA; QL (124 EA per 31 days)
ZEMAIRA	T5	PA
Ear, Nose / Throat Medications		
<i>acetic acid otic (ear)</i>	T2	
<i>azelastine nasal</i>	T2	QL (30 ML per 25 days)
<i>chlorhexidine gluconate mucous membrane</i>	T1	
CIPRO HC	T4	
<i>ciprofloxacin-dexamethasone</i>	T3	
<i>ciprofloxacin-fluocinolone</i>	T4	
<i>fluocinolone acetonide oil</i>	T2	
<i>hydrocortisone-acetic acid</i>	T2	
<i>ipratropium bromide nasal spray,non-aerosol 21 mcg (0.03 %)</i>	T1	QL (30 ML per 28 days)
<i>ipratropium bromide nasal spray,non-aerosol 42 mcg (0.06 %)</i>	T1	QL (15 ML per 28 days)
<i>neomycin-polymyxin-hc otic (ear)</i>	T2	
<i>ofloxacin otic (ear)</i>	T2	
<i>olopatadine nasal</i>	T2	QL (30.5 GM per 30 days)
OTOVEL	T4	
PERIOGARD	T1	
<i>triamcinolone acetonide dental</i>	T2	
Endocrine/Diabetes		
<i>acarbose</i>	T1	QL (93 EA per 31 days)
ACTHAR	T5	PA
ALCOHOL PADS	T2	
ALKINDI SPRINKLE ORAL CAPSULE, SPRINKLE 0.5 MG	T4	PA
ALKINDI SPRINKLE ORAL CAPSULE, SPRINKLE 1 MG, 2 MG, 5 MG	T5	PA
ANDRODERM	T3	PA
AVEED	T4	PA
BAQSIMI	T3	
BASAGLAR KWIKPEN U-100 INSULIN	T4	
<i>cabergoline</i>	T2	
<i>calcitonin (salmon) nasal</i>	T2	PA-BvD
<i>calcitriol oral capsule</i>	T2	PA-BvD

Drug Name	Drug Tier	Requirements/Limits
<i>calcitriol oral solution</i>	T3	PA-BvD
CERDELGA	T5	PA; QL (62 EA per 31 days)
<i>cinacalcet oral tablet 30 mg</i>	T3	PA-BvD; QL (62 EA per 31 days)
<i>cinacalcet oral tablet 60 mg</i>	T5	PA-BvD; QL (62 EA per 31 days)
<i>cinacalcet oral tablet 90 mg</i>	T5	PA-BvD; QL (124 EA per 31 days)
CORTROPHIN GEL	T5	PA
CYCLOSET	T4	
<i>danazol</i>	T2	
DEPO-TESTOSTERONE INTRAMUSCULAR OIL 100 MG/ML	T4	PA
<i>desmopressin nasal spray with pump</i>	T4	
<i>desmopressin oral</i>	T2	
<i>dexamethasone oral elixir</i>	T1	
<i>dexamethasone oral tablet</i>	T1	
<i>diazoxide</i>	T4	
<i>doxercalciferol oral capsule 0.5 mcg, 2.5 mcg</i>	T2	PA-BvD
<i>doxercalciferol oral capsule 1 mcg</i>	T4	PA-BvD
EMFLAZA	T5	PA
FIASP FLEXTOUCH U-100 INSULIN	T3	
FIASP PENFILL U-100 INSULIN	T3	
FIASP U-100 INSULIN	T3	
<i>fludrocortisone</i>	T2	
GALAFOLD	T5	PA; QL (14 EA per 28 days)
<i>glimepiride</i>	T1	
<i>glipizide</i>	T1	
<i>glipizide-metformin</i>	T1	
GLUCAGEN HYPOKIT	T3	
GLUCAGON EMERGENCY KIT (HUMAN)	T3	
<i>glyburide</i>	T2	
<i>glyburide micronized</i>	T2	
<i>glyburide-metformin</i>	T2	
GLYXAMBI	T3	QL (31 EA per 31 days)
GVOKE	T3	
GVOKE HYPOPEN 2-PACK	T3	
GVOKE PFS 1-PACK SYRINGE	T3	
HUMALOG JUNIOR KWIKPEN U-100	T3	
HUMALOG KWIKPEN INSULIN	T3	
HUMALOG MIX 50-50 INSULN U-100	T3	

Drug Name	Drug Tier	Requirements/Limits
HUMALOG MIX 50-50 KWIKPEN	T3	
HUMALOG MIX 75-25 KWIKPEN	T3	
HUMALOG MIX 75-25(U-100)INSULN	T3	
HUMALOG U-100 INSULIN	T3	
HUMULIN 70/30 U-100 INSULIN	T3	
HUMULIN 70/30 U-100 KWIKPEN	T3	
HUMULIN N NPH INSULIN KWIKPEN	T3	
HUMULIN N NPH U-100 INSULIN	T3	
HUMULIN R REGULAR U-100 INSULN	T3	
HUMULIN R U-500 (CONC) INSULIN	T3	
HUMULIN R U-500 (CONC) KWIKPEN	T3	
<i>hydrocortisone oral</i>	T1	
<i>insulin asp prt-insulin aspart</i>	T3	
<i>insulin aspart u-100</i>	T3	
<i>insulin lispro</i>	T3	
<i>insulin lispro protamin-lispro</i>	T3	
INVOKAMET	T3	QL (62 EA per 31 days)
INVOKAMET XR	T3	QL (62 EA per 31 days)
INVOKANA ORAL TABLET 100 MG	T3	QL (62 EA per 31 days)
INVOKANA ORAL TABLET 300 MG	T3	QL (31 EA per 31 days)
ISTURISA	T5	PA
JANUMET	T3	QL (62 EA per 31 days)
JANUMET XR ORAL TABLET, ER MULTIPHASE 24 HR 100-1,000 MG, 50-500 MG	T3	QL (31 EA per 31 days)
JANUMET XR ORAL TABLET, ER MULTIPHASE 24 HR 50-1,000 MG	T3	QL (62 EA per 31 days)
JANUVIA ORAL TABLET 100 MG, 50 MG	T3	QL (31 EA per 31 days)
JANUVIA ORAL TABLET 25 MG	T3	QL (93 EA per 31 days)
JARDIANCE ORAL TABLET 10 MG	T3	QL (62 EA per 31 days)
JARDIANCE ORAL TABLET 25 MG	T3	QL (31 EA per 31 days)
JAVYGTOR ORAL POWDER IN PACKET 100 MG	T5	PA
JENTADUETO	T3	QL (62 EA per 31 days)
JENTADUETO XR ORAL TABLET, IR - ER, BIPHASIC 24HR 2.5-1,000 MG	T3	QL (62 EA per 31 days)
JENTADUETO XR ORAL TABLET, IR - ER, BIPHASIC 24HR 5-1,000 MG	T3	QL (31 EA per 31 days)
JYNARQUE ORAL TABLET	T5	PA; QL (112 EA per 28 days)

Drug Name	Drug Tier	Requirements/Limits
JYNARQUE ORAL TABLETS, SEQUENTIAL	T5	PA; QL (56 EA per 28 days)
KORLYM	T5	PA; QL (124 EA per 31 days)
LANTUS SOLOSTAR U-100 INSULIN	T3	
LANTUS U-100 INSULIN	T3	
LEVEMIR FLEXTOUCH U-100 INSULN	T3	
LEVEMIR U-100 INSULIN	T3	
<i>levothyroxine oral capsule</i>	T4	
<i>levothyroxine oral tablet</i>	T1	
LEVOXYL ORAL TABLET 100 MCG, 112 MCG, 125 MCG, 137 MCG, 150 MCG, 175 MCG, 200 MCG, 25 MCG, 50 MCG, 75 MCG, 88 MCG	T3	
<i>liothyronine oral</i>	T2	
MEDROL ORAL TABLET 2 MG	T4	
<i>metformin oral solution</i>	T4	ST; QL (791 ML per 31 days)
<i>metformin oral tablet 1,000 mg, 500 mg, 850 mg</i>	T1	
<i>metformin oral tablet extended release 24 hr</i>	T1	
<i>metformin oral tablet extended release 24hr</i>	NF	
<i>metformin oral tablet,er gast.retention 24 hr</i>	NF	
<i>methimazole oral tablet 10 mg, 5 mg</i>	T2	
<i>methylprednisolone</i>	T2	
<i>miglitol</i>	T2	
<i>miglustat</i>	T5	PA; QL (93 EA per 31 days)
MOUNJARO	T3	QL (2 ML per 28 days)
MYALEPT	T5	PA
<i>nateglinide</i>	T1	QL (93 EA per 31 days)
NATPARA	T5	PA; QL (31 EA per 31 days)
NOVOLIN 70/30 U-100 INSULIN	T3	
NOVOLIN 70-30 FLEXPEN U-100	T3	
NOVOLIN N FLEXPEN	T3	
NOVOLIN N NPH U-100 INSULIN	T3	
NOVOLIN R FLEXPEN	T3	
NOVOLIN R REGULAR U-100 INSULN	T3	
NOVOLOG FLEXPEN U-100 INSULIN	T3	
NOVOLOG MIX 70-30 U-100 INSULN	T3	
NOVOLOG MIX 70-30FLEXPEN U-100	T3	
NOVOLOG PENFILL U-100 INSULIN	T3	
NOVOLOG U-100 INSULIN ASPART	T3	

Drug Name	Drug Tier	Requirements/Limits
ORLISSA ORAL TABLET 150 MG	T5	PA; QL (31 EA per 31 days)
ORLISSA ORAL TABLET 200 MG	T5	PA; QL (62 EA per 31 days)
<i>oxandrolone oral tablet 10 mg</i>	T4	PA
<i>oxandrolone oral tablet 2.5 mg</i>	T2	PA
OZEMPIC SUBCUTANEOUS PEN INJECTOR 0.25 MG OR 0.5 MG(2 MG/1.5 ML), 1 MG/DOSE (4 MG/3 ML), 2 MG/DOSE (8 MG/3 ML)	T3	QL (3 ML per 28 days)
PALYNZIQ	T5	PA
<i>paricalcitol oral</i>	T2	PA-BvD
<i>pioglitazone</i>	T1	QL (31 EA per 31 days)
<i>pioglitazone-glimepiride</i>	T1	QL (31 EA per 31 days)
<i>pioglitazone-metformin</i>	T1	QL (93 EA per 31 days)
<i>prednisolone oral solution</i>	T2	
<i>prednisolone sodium phosphate oral solution 10 mg/5 ml, 20 mg/5 ml (4 mg/ml), 25 mg/5 ml (5 mg/ml), 5 mg base/5 ml (6.7 mg/5 ml)</i>	T2	
<i>prednisolone sodium phosphate oral tablet, disintegrating</i>	T4	
<i>prednisone oral solution</i>	T2	
<i>prednisone oral tablet</i>	T1	
<i>prednisone oral tablets, dose pack</i>	T2	
<i>propylthiouracil</i>	T2	
RECORLEV	T5	PA; QL (248 EA per 31 days)
<i>repaglinide oral tablet 0.5 mg</i>	T1	QL (124 EA per 31 days)
<i>repaglinide oral tablet 1 mg</i>	T2	QL (124 EA per 31 days)
<i>repaglinide oral tablet 2 mg</i>	T2	QL (248 EA per 31 days)
RYBELSUS	T3	QL (31 EA per 31 days)
<i>sapropterin</i>	T5	PA
SOMAVERT	T5	PA
SYMLINPEN 120	T3	QL (10.8 ML per 28 days)
SYMLINPEN 60	T3	QL (6 ML per 28 days)
SYNAREL	T5	
SYNJARDY	T3	QL (62 EA per 31 days)
SYNJARDY XR ORAL TABLET, IR - ER, BIPHASIC 24HR 10-1,000 MG, 12.5-1,000 MG, 5-1,000 MG	T3	QL (62 EA per 31 days)
SYNJARDY XR ORAL TABLET, IR - ER, BIPHASIC 24HR 25-1,000 MG	T3	QL (31 EA per 31 days)
SYNTHROID	T3	

Drug Name	Drug Tier	Requirements/Limits
<i>testosterone cypionate intramuscular oil 100 mg/ml, 200 mg/ml, 200 mg/ml (1 ml)</i>	T2	PA
<i>testosterone enanthate</i>	T2	PA
<i>testosterone transdermal gel in metered-dose pump 20.25 mg/1.25 gram (1.62 %)</i>	T3	PA
<i>testosterone transdermal gel in packet 1.62 % (20.25 mg/1.25 gram), 1.62 % (40.5 mg/2.5 gram)</i>	T3	PA
<i>testosterone transdermal solution in metered pump w/app</i>	T4	PA
TIROSINT	T4	
TIROSINT-SOL	T4	
<i>tolvaptan</i>	T5	PA
TOUJEO MAX U-300 SOLOSTAR	T3	
TOUJEO SOLOSTAR U-300 INSULIN	T3	
TRADJENTA	T3	QL (31 EA per 31 days)
TRESIBA FLEXTOUCH U-100	T3	
TRESIBA FLEXTOUCH U-200	T3	
TRESIBA U-100 INSULIN	T3	
TRIJARDY XR ORAL TABLET, IR - ER, BIPHASIC 24HR 10-5-1,000 MG, 25-5-1,000 MG	T3	QL (31 EA per 31 days)
TRIJARDY XR ORAL TABLET, IR - ER, BIPHASIC 24HR 12.5-2.5-1,000 MG, 5-2.5-1,000 MG	T3	QL (62 EA per 31 days)
TRULICITY	T3	QL (2 ML per 28 days)
UNITHROID	T3	
VICTOZA 3-PAK	T3	QL (9 ML per 30 days)
XULTOPHY 100/3.6	T3	
ZEGALOGUE AUTOINJECTOR	T3	
ZEGALOGUE SYRINGE	T3	
Gastroenterology		
<i>alosetron oral tablet 0.5 mg</i>	T5	PA; QL (93 EA per 31 days)
<i>alosetron oral tablet 1 mg</i>	T5	PA; QL (62 EA per 31 days)
<i>amoxicil-clarithromy-lansopraz</i>	T4	
<i>aprepitant</i>	T4	PA-BvD
<i>balsalazide</i>	T2	
<i>betaine</i>	T3	
<i>budesonide oral capsule,delayed,extend.release</i>	T4	
<i>budesonide oral tablet,delayed and ext.release</i>	T5	
CARAFATE ORAL SUSPENSION	T3	

Drug Name	Drug Tier	Requirements/Limits
CHENODAL	T5	PA
CHOLBAM	T5	PA
<i>cimetidine</i>	T2	
<i>cimetidine hcl oral</i>	T2	
CIMZIA	T5	PA; QL (2 EA per 28 days)
CIMZIA POWDER FOR RECONST	T5	PA; QL (2 EA per 28 days)
CLENPIQ	T4	
COMPRO	T2	
CONSTULOSE	T2	
CREON	T3	
<i>cromolyn oral</i>	T4	
<i>dicyclomine oral capsule</i>	T2	
<i>dicyclomine oral solution</i>	T2	
<i>dicyclomine oral tablet</i>	T2	
<i>diphenoxylate-atropine oral liquid</i>	T3	
<i>diphenoxylate-atropine oral tablet</i>	T2	
<i>dronabinol</i>	T4	PA-BvD
EMEND ORAL SUSPENSION FOR RECONSTITUTION	T4	PA-BvD
ENULOSE	T2	
<i>esomeprazole magnesium oral capsule, delayed release(dr/ec)</i>	T2	QL (31 EA per 31 days)
<i>famotidine oral suspension</i>	T1	
<i>famotidine oral tablet 20 mg, 40 mg</i>	T1	
GATTEX 30-VIAL	T5	PA
GAVILYTE-C	T2	
GAVILYTE-G	T2	
GENERLAC	T2	
<i>glycopyrrolate oral tablet 1 mg, 2 mg</i>	T2	
<i>granisetron hcl oral</i>	T4	PA-BvD
<i>hydrocortisone rectal</i>	T3	
<i>hydrocortisone topical cream with perineal applicator 2.5 %</i>	T1	
<i>hydrocortisone-pramoxine rectal cream 1-1 %</i>	T4	
IBSRELA	T5	PA; QL (62 EA per 31 days)
<i>lactulose oral solution 10 gram/15 ml</i>	T1	
<i>lansoprazole oral capsule, delayed release(dr/ec) 15 mg</i>	T3	QL (31 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
<i>lansoprazole oral capsule, delayed release(dr/ec) 30 mg</i>	T3	QL (62 EA per 31 days)
LINZESS	T3	QL (31 EA per 31 days)
<i>loperamide oral capsule</i>	T2	
<i>lubiprostone</i>	T3	QL (62 EA per 31 days)
<i>meclizine oral tablet 12.5 mg, 25 mg</i>	T2	
<i>mesalamine oral capsule (with del rel tablets)</i>	T3	
<i>mesalamine oral capsule, extended release</i>	T3	
<i>mesalamine oral capsule, extended release 24hr</i>	T4	
<i>mesalamine oral tablet, delayed release (dr/ec) 1.2 gram</i>	T3	
<i>mesalamine oral tablet, delayed release (dr/ec) 800 mg</i>	T4	
<i>mesalamine rectal enema</i>	T4	
<i>methscopolamine</i>	T2	
<i>metoclopramide hcl oral solution</i>	T2	
<i>metoclopramide hcl oral tablet</i>	T2	
<i>misoprostol</i>	T2	
MOVANTI	T3	QL (31 EA per 31 days)
MYTESI	T5	QL (62 EA per 31 days)
<i>nizatidine oral capsule</i>	T2	
OICALIVA	T5	PA; QL (31 EA per 31 days)
<i>omeprazole oral capsule, delayed release(dr/ec)</i>	T1	
<i>ondansetron</i>	T2	PA-BvD
<i>ondansetron hcl oral solution</i>	T2	PA-BvD
<i>ondansetron hcl oral tablet 4 mg, 8 mg</i>	T2	PA-BvD
OSMOPREP	T4	
<i>pantoprazole oral tablet, delayed release (dr/ec)</i>	T1	
<i>peg 3350-electrolytes oral recon soln 236-22.74- 6.74 -5.86 gram</i>	T2	
<i>peg3350-sod sul-nacl-kcl-asb-c</i>	T4	
<i>peg-electrolyte soln</i>	T2	
PENTASA	T3	
<i>prochlorperazine</i>	T3	
<i>prochlorperazine maleate</i>	T2	
PROCTO-PAK	T2	
PROCTOSOL HC TOPICAL	T2	
PROCTOZONE-HC	T2	
PYLERA	T5	

Drug Name	Drug Tier	Requirements/Limits
<i>rabeprazole oral tablet, delayed release (dr/ec)</i>	T2	QL (62 EA per 31 days)
RECTIV	T4	
RELISTOR ORAL	T5	PA; QL (93 EA per 31 days)
RELISTOR SUBCUTANEOUS SOLUTION	T4	PA; QL (18.6 ML per 31 days)
RELISTOR SUBCUTANEOUS SYRINGE 12 MG/0.6 ML	T5	PA; QL (18.6 ML per 31 days)
RELISTOR SUBCUTANEOUS SYRINGE 8 MG/0.4 ML	T5	PA; QL (12.4 ML per 31 days)
SANCUSO	T4	
<i>scopolamine base</i>	T3	QL (10 EA per 30 days)
SKYRIZI SUBCUTANEOUS WEARABLE INJECTOR	T5	PA; QL (2.4 ML per 56 days)
SUCRAID	T5	
<i>sucralfate</i>	T2	
<i>sulfasalazine</i>	T2	
SUPREP BOWEL PREP KIT	T3	
SYMPROIC	T4	PA; QL (31 EA per 31 days)
<i>ursodiol oral capsule 300 mg</i>	T3	
<i>ursodiol oral tablet</i>	T3	
VARUBI	T4	PA-BvD
VIBERZI	T5	PA; QL (62 EA per 31 days)
ZENPEP ORAL CAPSULE, DELAYED RELEASE(DR/EC) 10,000-32,000 -42,000 UNIT, 15,000-47,000 -63,000 UNIT, 20,000-63,000- 84,000 UNIT, 3,000-10,000 -14,000-UNIT, 5,000-17,000- 24,000 UNIT	T3	
ZENPEP ORAL CAPSULE, DELAYED RELEASE(DR/EC) 25,000-79,000- 105,000 UNIT, 40,000-126,000- 168,000 UNIT	T5	
Immunology, Vaccines / Biotechnology		
ACTHIB (PF)	T3	
ACTIMMUNE	T5	PA
ADACEL(TDAP ADOLESN/ADULT)(PF)	T3	
ARANESP (IN POLYSORBATE) INJECTION SOLUTION 100 MCG/ML, 25 MCG/ML, 40 MCG/ML	T4	PA-BvD
ARANESP (IN POLYSORBATE) INJECTION SOLUTION 200 MCG/ML, 60 MCG/ML	T5	PA-BvD

Drug Name	Drug Tier	Requirements/Limits
ARANESP (IN POLYSORBATE) INJECTION SYRINGE 10 MCG/0.4 ML, 100 MCG/0.5 ML, 25 MCG/0.42 ML, 40 MCG/0.4 ML, 60 MCG/0.3 ML	T4	PA-BvD
ARANESP (IN POLYSORBATE) INJECTION SYRINGE 150 MCG/0.3 ML, 200 MCG/0.4 ML, 300 MCG/0.6 ML, 500 MCG/ML	T5	PA-BvD
ARCALYST	T5	PA
AVONEX INTRAMUSCULAR PEN INJECTOR KIT	T5	QL (4 EA per 28 days)
AVONEX INTRAMUSCULAR SYRINGE KIT	T5	QL (4 EA per 28 days)
<i>bcg vaccine, live (pf)</i>	T4	
BESREMI	T5	PA-NS; QL (2 ML per 28 days)
BETASERON SUBCUTANEOUS KIT	T5	QL (14 EA per 28 days)
BEXSERO	T3	
BIVIGAM	T5	PA
BOOSTRIX TDAP	T3	
DAPTACEL (DTAP PEDIATRIC) (PF)	T3	
EGRIFTA SV	T5	PA
ENGERIX-B (PF)	T3	PA-BvD
ENGERIX-B PEDIATRIC (PF)	T3	PA-BvD
EPOGEN INJECTION SOLUTION 2,000 UNIT/ML, 20,000 UNIT/2 ML, 20,000 UNIT/ML, 3,000 UNIT/ML, 4,000 UNIT/ML	T4	PA-BvD
EXTAVIA SUBCUTANEOUS KIT	T5	QL (15 EA per 30 days)
FLEBOGAMMA DIF INTRAVENOUS SOLUTION 10 %	T5	PA
FULPHILA	T5	
GAMMAGARD LIQUID	T5	PA
GAMMAGARD S-D (IGA < 1 MCG/ML)	T5	PA
GAMMAKED INJECTION SOLUTION 1 GRAM/10 ML (10 %)	T4	PA
GAMMAPLEX	T5	PA
GAMMAPLEX (WITH SORBITOL)	T5	PA
GAMUNEX-C INJECTION SOLUTION 1 GRAM/10 ML (10 %)	T3	PA
GARDASIL 9 (PF)	T3	
GENOTROPIN MINIQUICK SUBCUTANEOUS SYRINGE 0.2 MG/0.25 ML	T4	PA

Drug Name	Drug Tier	Requirements/Limits
GENOTROPIN MINIQUICK SUBCUTANEOUS SYRINGE 0.4 MG/0.25 ML, 0.6 MG/0.25 ML, 0.8 MG/0.25 ML, 1 MG/0.25 ML, 1.2 MG/0.25 ML, 1.4 MG/0.25 ML, 1.6 MG/0.25 ML, 1.8 MG/0.25 ML, 2 MG/0.25 ML	T5	PA
GENOTROPIN SUBCUTANEOUS CARTRIDGE 12 MG/ML (36 UNIT/ML)	T5	PA
GENOTROPIN SUBCUTANEOUS CARTRIDGE 5 MG/ML (15 UNIT/ML)	T4	PA
GRANIX	T5	
GRASTEK	T4	PA
HAVRIX (PF)	T3	
HIBERIX (PF)	T3	
HUMATROPE INJECTION CARTRIDGE 12 MG (36 UNIT), 24 MG (72 UNIT)	T5	PA
HUMATROPE INJECTION CARTRIDGE 6 MG (18 UNIT)	T4	PA
IMOVAX RABIES VACCINE (PF)	T3	PA-BvD
INFANRIX (DTAP) (PF) INTRAMUSCULAR SYRINGE	T3	
INTRON A INJECTION RECON SOLN 10 MILLION UNIT (1 ML)	T3	PA-NS
INTRON A INJECTION RECON SOLN 50 MILLION UNIT (1 ML)	T5	PA-NS
IPOL	T3	
IXIARO (PF)	T3	
KINRIX (PF) INTRAMUSCULAR SYRINGE	T3	
LEUKINE INJECTION RECON SOLN	T5	PA
MENACTRA (PF) INTRAMUSCULAR SOLUTION	T3	
MENQUADFI (PF)	T4	
MENVEO A-C-Y-W-135-DIP (PF) INTRAMUSCULAR KIT	T3	
M-M-R II (PF)	T3	
NEULASTA	T5	
NEUPOGEN INJECTION SOLUTION 300 MCG/ML	T4	
NEUPOGEN INJECTION SOLUTION 480 MCG/1.6 ML	T5	
NEUPOGEN INJECTION SYRINGE	T5	
NIVESTYM	T5	

Drug Name	Drug Tier	Requirements/Limits
NORDITROPIN FLEXP RO SUBCUTANEOUS PEN INJECTOR 10 MG/1.5 ML (6.7 MG/ML), 15 MG/1.5 ML (10 MG/ML), 30 MG/3 ML (10 MG/ML)	T5	PA
NORDITROPIN FLEXP RO SUBCUTANEOUS PEN INJECTOR 5 MG/1.5 ML (3.3 MG/ML)	T4	PA
NUTROPIN AQ NUSPIN	T5	PA
NYVEPRIA	T5	
OCTAGAM	T5	PA
ODACTRA	T4	PA
OMNITROPE SUBCUTANEOUS CARTRIDGE 10 MG/1.5 ML (6.7 MG/ML)	T5	PA
OMNITROPE SUBCUTANEOUS CARTRIDGE 5 MG/1.5 ML (3.3 MG/ML)	T4	PA
OMNITROPE SUBCUTANEOUS RECON SOLN	T5	PA
ORALAIR SUBLINGUAL TABLET 300 INDX REACTIVITY	T4	PA
PANZYGA	T5	PA
PEDIARIX (PF)	T3	PA-BvD
PEDVAX HIB (PF)	T3	
PEGASYS	T5	PA
PENTACEL (PF)	T3	
PLEGRIDY SUBCUTANEOUS PEN INJECTOR 125 MCG/0.5 ML	T5	QL (1 ML per 28 days)
PLEGRIDY SUBCUTANEOUS SYRINGE 125 MCG/0.5 ML	T5	QL (1 ML per 28 days)
PREHEVBRIO (PF)	T3	PA-BvD
PRIORIX (PF)	T3	
PRIVIGEN	T5	PA
PROCRIT INJECTION SOLUTION 10,000 UNIT/ML, 2,000 UNIT/ML, 3,000 UNIT/ML, 4,000 UNIT/ML	T3	PA-BvD
PROCRIT INJECTION SOLUTION 20,000 UNIT/ML, 40,000 UNIT/ML	T5	PA-BvD
PROQUAD (PF)	T3	
QUADRACEL (PF)	T3	
RABAVERT (PF)	T3	PA-BvD
REBIF (WITH ALBUMIN)	T5	QL (6 ML per 28 days)
REBIF REBIDOSE SUBCUTANEOUS PEN INJECTOR 22 MCG/0.5 ML, 44 MCG/0.5 ML	T5	QL (6 ML per 28 days)

Drug Name	Drug Tier	Requirements/Limits
REBIF REBIDOSE SUBCUTANEOUS PEN INJECTOR 8.8MCG/0.2ML-22 MCG/0.5ML (6)	T5	QL (4.2 ML per 365 days)
REBIF TITRATION PACK	T5	QL (8.4 ML per 365 days)
RECOMBIVAX HB (PF)	T3	PA-BvD
RETACRIT INJECTION SOLUTION 10,000 UNIT/ML, 2,000 UNIT/ML, 20,000 UNIT/2 ML, 3,000 UNIT/ML, 4,000 UNIT/ML	T3	PA-BvD
RETACRIT INJECTION SOLUTION 20,000 UNIT/ML, 40,000 UNIT/ML	T5	PA-BvD
ROTARIX	T3	
ROTATEQ VACCINE	T3	
SAIZEN	T5	PA
SAIZEN SAIZENPREP	T5	PA
SEROSTIM SUBCUTANEOUS RECON SOLN 4 MG, 5 MG, 6 MG	T5	PA
SHINGRIX (PF)	T3	QL (2 EA per 999 days)
TDVAX	T3	
TENIVAC (PF)	T3	
<i>tetanus,diphtheria tox ped(pf)</i>	T4	
TICOVAC	T3	
TRUMENBA	T3	
TWINRIX (PF)	T3	
TYPHIM VI	T3	
UDENYCA	T5	
VAQTA (PF)	T3	
VARIVAX (PF)	T3	
YF-VAX (PF)	T3	
ZARXIO	T5	
ZIEXTENZO	T5	
ZOMACTON	T4	PA
ZORBTIVE	T5	PA
Miscellaneous Supplies		
ASSURE ID INSULIN SAFETY SYRINGE 1 ML 29 GAUGE X 1/2"	T4	
GAUZE PAD TOPICAL BANDAGE 2 X 2 "	T2	
<i>insulin syringe-needle u-100 syringe 0.3 ml 29 gauge, 1 ml 29 gauge x 1/2", 1/2 ml 28 gauge</i>	T3	
<i>pen needle, diabetic needle 29 gauge x 1/2"</i>	T4	

Drug Name	Drug Tier	Requirements/Limits
Musculoskeletal / Rheumatology		
ACTEMRA ACTPEN	T5	PA; QL (3.6 ML per 28 days)
ACTEMRA SUBCUTANEOUS	T5	PA; QL (3.6 ML per 28 days)
<i>alendronate oral solution</i>	T1	
<i>alendronate oral tablet 10 mg, 35 mg, 70 mg</i>	T1	
<i>allopurinol oral tablet 100 mg, 300 mg</i>	T1	
BENLYSTA SUBCUTANEOUS	T5	PA; QL (4 ML per 28 days)
CUPRIMINE	T5	
ENBREL MINI	T5	PA; QL (7.84 ML per 28 days)
ENBREL SUBCUTANEOUS SOLUTION	T5	PA; QL (4 ML per 28 days)
ENBREL SUBCUTANEOUS SYRINGE 25 MG/0.5 ML (0.5)	T5	PA; QL (4 ML per 28 days)
ENBREL SUBCUTANEOUS SYRINGE 50 MG/ML (1 ML)	T5	PA; QL (7.84 ML per 28 days)
ENBREL SURECLICK	T5	PA; QL (7.84 ML per 28 days)
EVENITY SUBCUTANEOUS SYRINGE 210MG/2.34ML (105MG/1.17MLX2)	T5	PA; QL (2.34 ML per 28 days)
FORTEO SUBCUTANEOUS PEN INJECTOR 20 MCG/DOSE (600MCG/2.4ML)	T5	PA; QL (2.4 ML per 28 days)
HUMIRA PEN	T5	PA; QL (2 EA per 28 days)
HUMIRA PEN CROHNS-UC-HS START	T5	PA; QL (6 EA per 28 days)
HUMIRA PEN PSOR-UVEITS-ADOL HS	T5	PA; QL (4 EA per 28 days)
HUMIRA SUBCUTANEOUS SYRINGE KIT 40 MG/0.8 ML	T5	PA; QL (2 EA per 28 days)
HUMIRA(CF)	T5	PA; QL (2 EA per 28 days)
HUMIRA(CF) PEDI CROHNS STARTER SUBCUTANEOUS SYRINGE KIT 80 MG/0.8 ML	T5	PA; QL (3 EA per 28 days)
HUMIRA(CF) PEDI CROHNS STARTER SUBCUTANEOUS SYRINGE KIT 80 MG/0.8 ML-40 MG/0.4 ML	T5	PA; QL (2 EA per 28 days)
HUMIRA(CF) PEN	T5	PA; QL (2 EA per 28 days)
HUMIRA(CF) PEN CROHNS-UC-HS	T5	PA; QL (3 EA per 28 days)
HUMIRA(CF) PEN PEDIATRIC UC	T5	PA; QL (4 EA per 28 days)
HUMIRA(CF) PEN PSOR-UV-ADOL HS	T5	PA; QL (3 EA per 28 days)
<i>ibandronate oral</i>	T2	
KEVZARA	T5	PA; QL (2.28 ML per 28 days)
KINERET	T5	PA; QL (18.76 ML per 28 days)
<i>leflunomide</i>	T2	
MITIGARE	T3	QL (62 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
OLUMIANT ORAL TABLET 1 MG, 2 MG	T5	PA; QL (31 EA per 31 days)
ORENCIA CLICKJECT	T5	PA; QL (4 ML per 28 days)
ORENCIA SUBCUTANEOUS SYRINGE 125 MG/ML	T5	PA; QL (4 ML per 28 days)
ORENCIA SUBCUTANEOUS SYRINGE 50 MG/0.4 ML	T5	PA; QL (1.6 ML per 28 days)
ORENCIA SUBCUTANEOUS SYRINGE 87.5 MG/0.7 ML	T5	PA; QL (2.8 ML per 28 days)
OTEZLA	T5	PA; QL (62 EA per 31 days)
OTEZLA STARTER ORAL TABLETS,DOSE PACK 10 MG (4)-20 MG (4)-30 MG (47)	T5	PA; QL (55 EA per 28 days)
OTREXUP (PF)	T4	PA
<i>penicillamine</i>	T5	
<i>probenecid</i>	T2	
<i>probenecid-colchicine</i>	T2	
PROLIA	T4	PA; QL (1 ML per 180 days)
<i>raloxifene</i>	T3	
RASUVO (PF)	T4	PA
REDITREX (PF)	T4	PA
RIDAURA	T3	
RINVOQ ORAL TABLET EXTENDED RELEASE 24 HR 15 MG, 30 MG	T5	PA; QL (31 EA per 31 days)
RINVOQ ORAL TABLET EXTENDED RELEASE 24 HR 45 MG	T5	PA; QL (112 EA per 365 days)
<i>risedronate oral tablet 150 mg, 35 mg, 35 mg (12 pack), 35 mg (4 pack), 5 mg</i>	T2	
<i>risedronate oral tablet,delayed release (dr/ec)</i>	T2	
SAVELLA	T4	PA
SIMPONI SUBCUTANEOUS PEN INJECTOR 100 MG/ML	T5	PA; QL (1 ML per 28 days)
SIMPONI SUBCUTANEOUS PEN INJECTOR 50 MG/0.5 ML	T5	PA; QL (0.5 ML per 28 days)
SIMPONI SUBCUTANEOUS SYRINGE 100 MG/ML	T5	PA; QL (1 ML per 28 days)
SIMPONI SUBCUTANEOUS SYRINGE 50 MG/0.5 ML	T5	PA; QL (0.5 ML per 28 days)
<i>teriparatide</i>	T5	PA; QL (2.48 ML per 28 days)
TYMLOS	T5	PA; QL (1.56 ML per 30 days)
XELJANZ ORAL SOLUTION	T5	PA; QL (310 ML per 31 days)
XELJANZ ORAL TABLET	T5	PA; QL (62 EA per 31 days)
XELJANZ XR	T5	PA; QL (31 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
Obstetrics / Gynecology		
ALTAVERA (28)	T2	
ALYACEN 1/35 (28)	T2	
ANGELIQ ORAL TABLET 0.5-1 MG	T4	
APRI	T2	
ARANELLE (28)	T2	
AVIANE	T2	
BALZIVA (28)	T2	
BLISOVI 24 FE	T2	
BLISOVI FE 1.5/30 (28)	T2	
BRIELLYN	T2	
CAMILA	T2	
CLEOCIN VAGINAL SUPPOSITORY	T4	
CLIMARA PRO	T4	
<i>clindamycin phosphate vaginal</i>	T2	
CLINDESSE	T4	
CRINONE	T4	PA
CRYSELLE (28)	T2	
CYRED EQ	T2	
DEPO-ESTRADIOL	T4	
<i>desog-e.estradiol/e.estradiol</i>	T2	
<i>desogestrel-ethinyl estradiol</i>	T2	
DIVIGEL TRANSDERMAL GEL IN PACKET 0.5 MG/0.5 GRAM (0.1 %)	T4	
DOLISHALE	T2	
DOTTI	T2	
<i>drospirenone-ethinyl estradiol</i>	T2	
ELURYNG	T3	
EMOQUETTE	T2	
ENPRESSE	T2	
ENSKYCE	T2	
ERRIN	T2	
ESTARYLLA	T2	
<i>estradiol oral</i>	T1	
<i>estradiol transdermal patch semiweekly</i>	T2	
<i>estradiol transdermal patch weekly</i>	T2	
<i>estradiol vaginal</i>	T4	
<i>estradiol-norethindrone acet</i>	T2	

Drug Name	Drug Tier	Requirements/Limits
ESTRING	T4	
<i>ethynodiol diac-eth estradiol</i>	T2	
<i>etonogestrel-ethinyl estradiol</i>	T3	
EVAMIST	T4	
FEMRING	T4	
FEMYNOR	T2	
FYAVOLV	T4	
HAILEY 24 FE	T2	
ICLEVIA	T2	
INCASSIA	T2	
INTROVALE	T2	
ISIBLOOM	T2	
JASMIEL (28)	T2	
JINTELI	T2	
JULEBER	T2	
JUNEL 1.5/30 (21)	T2	
JUNEL 1/20 (21)	T2	
JUNEL FE 1.5/30 (28)	T2	
JUNEL FE 1/20 (28)	T2	
JUNEL FE 24	T2	
KAITLIB FE	T2	
KARIVA (28)	T2	
KELNOR 1/35 (28)	T2	
KELNOR 1-50 (28)	T2	
KURVELO (28)	T2	
LESSINA	T2	
LEVONEST (28)	T2	
<i>levonorgestrel-ethinyl estrad</i>	T2	
<i>levonorg-eth estrad triphasic</i>	T2	
LEVORA-28	T2	
LORYNA (28)	T2	
LOW-OGESTREL (28)	T2	
LUTERA (28)	T2	
LYLEQ	T2	
LYLLANA	T2	
LYZA	T2	
MARLISSA (28)	T2	
<i>medroxyprogesterone</i>	T2	

Drug Name	Drug Tier	Requirements/Limits
MENEST ORAL TABLET 0.3 MG, 0.625 MG, 1.25 MG	T4	
<i>metronidazole vaginal</i>	T3	
MICONAZOLE-3 VAGINAL SUPPOSITORY	T2	
MICROGESTIN 1.5/30 (21)	T2	
MICROGESTIN 1/20 (21)	T2	
MICROGESTIN FE 1.5/30 (28)	T2	
MICROGESTIN FE 1/20 (28)	T2	
MILI	T2	
MYFEMBREE	T5	PA; QL (31 EA per 31 days)
NECON 0.5/35 (28)	T2	
<i>noreth-ethinyl estradiol-iron</i>	T2	
<i>norethindrone (contraceptive)</i>	T2	
<i>norethindrone acetate</i>	T2	
<i>norethindrone ac-eth estradiol oral tablet 0.5-2.5 mg-mcg, 1-20 mg-mcg, 1-5 mg-mcg</i>	T2	
<i>norethindrone-e.estradiol-iron oral tablet 1 mg-20 mcg (21)/75 mg (7)</i>	T2	
<i>norgestimate-ethinyl estradiol oral tablet 0.18/0.215/0.25 mg-35 mcg (28), 0.25-35 mg-mcg</i>	T2	
NORTREL 0.5/35 (28)	T2	
NORTREL 1/35 (21)	T2	
NORTREL 1/35 (28)	T2	
NORTREL 7/7/7 (28)	T2	
NYLIA 1/35 (28)	T2	
NYLIA 7/7/7 (28)	T2	
NYMYO	T2	
ORIAHNN	T5	PA; QL (56 EA per 28 days)
PIMTREA (28)	T2	
PIRMELLA ORAL TABLET 1-35 MG-MCG	T2	
PORTIA 28	T2	
PREFEST	T4	
PREMARIN VAGINAL	T3	
<i>progesterone micronized</i>	T2	
RECLIPSEN (28)	T2	
SETLAKIN	T2	
SPRINTEC (28)	T2	
SRONYX	T2	

Drug Name	Drug Tier	Requirements/Limits
SYEDA	T2	
<i>terconazole</i>	T2	
<i>tranexamic acid oral</i>	T2	
TRI-ESTARYLLA	T2	
TRI-MILI	T2	
TRI-NYMYO	T2	
TRI-SPRINTEC (28)	T2	
TRIVORA (28)	T2	
TRI-VYLIBRA	T2	
VANDAZOLE	T3	
VELIVET TRIPHASIC REGIMEN (28)	T2	
VESTURA (28)	T2	
VIENVA	T2	
VYFEMLA (28)	T2	
VYLIBRA	T2	
YUVAFEM	T4	
ZOVIA 1-35 (28)	T2	
Ophthalmology		
<i>acetazolamide</i>	T2	
ACUVAIL (PF)	T4	
ALOCRIAL	T4	
ALOMIDE	T3	
ALPHAGAN P OPHTHALMIC (EYE) DROPS 0.1 %	T3	
<i>apraclonidine</i>	T2	
<i>atropine ophthalmic (eye) drops</i>	T2	
AZASITE	T4	
<i>azelastine ophthalmic (eye)</i>	T2	
<i>bacitracin ophthalmic (eye)</i>	T2	
<i>bacitracin-polymyxin b</i>	T2	
BESIVANCE	T4	
<i>betaxolol ophthalmic (eye)</i>	T2	
BETIMOL	T4	
BETOPTIC S	T4	
<i>bimatoprost ophthalmic (eye)</i>	T2	
BLEPHAMIDE S.O.P.	T3	
<i>brimonidine</i>	T2	
<i>brimonidine-timolol</i>	T3	

Drug Name	Drug Tier	Requirements/Limits
<i>brinzolamide</i>	T4	
<i>bromfenac</i>	T2	
<i>carteolol</i>	T2	
CILOXAN OPHTHALMIC (EYE) OINTMENT	T3	
<i>ciprofloxacin hcl ophthalmic (eye)</i>	T1	
COMBIGAN	T3	
<i>cromolyn ophthalmic (eye)</i>	T2	
<i>cyclosporine ophthalmic (eye)</i>	T3	QL (60 EA per 30 days)
CYSTADROPS	T5	PA; QL (20 ML per 28 days)
CYSTARAN	T5	PA; QL (60 ML per 28 days)
<i>dexamethasone sodium phosphate ophthalmic (eye)</i>	T2	
<i>diclofenac sodium ophthalmic (eye)</i>	T1	
<i>difluprednate</i>	T3	
<i>dorzolamide</i>	T2	
<i>dorzolamide-timolol</i>	T2	
<i>epinastine</i>	T2	
<i>erythromycin ophthalmic (eye)</i>	T2	
<i>fluorometholone</i>	T2	
<i>flurbiprofen sodium</i>	T2	
<i>gatifloxacin</i>	T3	
GENTAK OPHTHALMIC (EYE) OINTMENT	T2	
<i>gentamicin ophthalmic (eye) drops</i>	T1	
ILEVRO	T3	
IOPIDINE OPHTHALMIC (EYE) DROPPERETTE	T3	
<i>ketorolac ophthalmic (eye)</i>	T2	
LACRISERT	T4	
<i>latanoprost</i>	T1	
<i>levobunolol ophthalmic (eye) drops 0.5 %</i>	T1	
<i>levofloxacin ophthalmic (eye) drops 0.5 %</i>	T2	
LUMIGAN OPHTHALMIC (EYE) DROPS 0.01 %	T3	QL (5 ML per 31 days)
<i>methazolamide</i>	T2	
<i>moxifloxacin ophthalmic (eye) drops</i>	T3	
NATACYN	T3	
<i>neomycin-bacitracin-poly-hc</i>	T2	

Drug Name	Drug Tier	Requirements/Limits
<i>neomycin-bacitracin-polymyxin</i>	T2	
<i>neomycin-polymyxin b-dexameth</i>	T2	
<i>neomycin-polymyxin-gramicidin</i>	T2	
<i>neomycin-polymyxin-hc ophthalmic (eye)</i>	T2	
NEVANAC	T4	
<i>ofloxacin ophthalmic (eye)</i>	T2	
<i>olopatadine ophthalmic (eye)</i>	T3	
OXERVATE	T5	PA; QL (112 ML per 56 days)
<i>pilocarpine hcl ophthalmic (eye) drops 1 %, 2 %, 4 %</i>	T2	
<i>polymyxin b sulf-trimethoprim</i>	T1	
PRED-G S.O.P.	T4	
<i>prednisolone acetate</i>	T2	
<i>prednisolone sodium phosphate ophthalmic (eye)</i>	T2	
RESTASIS	T3	QL (60 EA per 30 days)
RESTASIS MULTIDOSE	T3	QL (5.5 ML per 27 days)
SIMBRINZA	T4	
<i>sulfacetamide sodium ophthalmic (eye) drops</i>	T2	
<i>sulfacetamide sodium ophthalmic (eye) ointment</i>	T1	
<i>sulfacetamide-prednisolone</i>	T2	
<i>timolol maleate ophthalmic (eye) drops</i>	T1	
<i>timolol maleate ophthalmic (eye) drops, once daily</i>	T4	
<i>timolol maleate ophthalmic (eye) gel forming solution</i>	T2	
TOBRADEX OPHTHALMIC (EYE) OINTMENT	T3	
<i>tobramycin ophthalmic (eye)</i>	T1	
<i>tobramycin-dexamethasone</i>	T2	
TOBREX OPHTHALMIC (EYE) OINTMENT	T3	
<i>travoprost</i>	T3	
<i>trifluridine</i>	T2	
XIIDRA	T3	QL (60 EA per 30 days)
ZIOPTAN (PF)	T4	
ZIRGAN	T4	ST
ZYLET	T4	
Respiratory And Allergy		
<i>acetylcysteine</i>	T2	PA-BvD

Drug Name	Drug Tier	Requirements/Limits
ADEMPAS	T5	PA; QL (93 EA per 31 days)
<i>albuterol sulfate inhalation hfa aerosol inhaler 90 mcg/actuation</i>	T3	QL (17 GM per 30 days)
<i>albuterol sulfate inhalation hfa aerosol inhaler 90 mcg/actuation (nda020503)</i>	T3	QL (13.4 GM per 30 days)
<i>albuterol sulfate inhalation hfa aerosol inhaler 90 mcg/actuation (nda020983)</i>	NF	
<i>albuterol sulfate inhalation solution for nebulization 0.63 mg/3 ml, 1.25 mg/3 ml, 2.5 mg/3 ml (0.083 %), 2.5 mg/0.5 ml</i>	T2	PA-BvD
<i>albuterol sulfate oral syrup</i>	T1	
<i>albuterol sulfate oral tablet</i>	T1	
ALYQ	T5	PA; QL (62 EA per 31 days)
<i>ambrisentan</i>	T5	PA; QL (31 EA per 31 days)
ANORO ELLIPTA	T3	QL (60 EA per 30 days)
<i>arformoterol</i>	T4	PA-BvD
ASMANEX HFA	T3	QL (13 GM per 30 days)
ASMANEX TWISTHALER INHALATION AEROSOL POWDR BREATH ACTIVATED 110 MCG/ ACTUATION (30), 220 MCG/ ACTUATION (120), 220 MCG/ ACTUATION (30), 220 MCG/ ACTUATION (60)	T3	QL (1 EA per 30 days)
ATROVENT HFA	T3	QL (25.8 GM per 30 days)
<i>azelastine-fluticasone</i>	T4	
BECONASE AQ	T4	
BERINERT INTRAVENOUS KIT	T5	PA
<i>bosentan</i>	T5	PA; QL (62 EA per 31 days)
BREO ELLIPTA	T3	QL (60 EA per 30 days)
BRONCHITOL	T5	PA; QL (560 EA per 28 days)
<i>budesonide inhalation suspension for nebulization 0.25 mg/2 ml, 0.5 mg/2 ml</i>	T3	PA-BvD
<i>budesonide inhalation suspension for nebulization 1 mg/2 ml</i>	T4	PA-BvD
<i>cetirizine oral solution 1 mg/ml</i>	T1	QL (310 ML per 31 days)
CINRYZE	T5	PA; QL (20 EA per 28 days)
CLARINEX-D 12 HOUR	T4	
COMBIVENT RESPIMAT	T4	QL (4 GM per 30 days)
<i>cromolyn inhalation</i>	T2	PA-BvD
<i>cyproheptadine</i>	T2	PA
DALIRESP ORAL TABLET 250 MCG	T4	QL (31 EA per 31 days)
DALIRESP ORAL TABLET 500 MCG	T3	QL (31 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
<i>desloratadine oral tablet</i>	T2	QL (31 EA per 31 days)
<i>desloratadine oral tablet, disintegrating</i>	T3	QL (31 EA per 31 days)
<i>epinephrine injection auto-injector</i>	T3	
ESBRIET ORAL CAPSULE	T5	PA; QL (279 EA per 31 days)
ESBRIET ORAL TABLET 267 MG	T5	PA; QL (279 EA per 31 days)
ESBRIET ORAL TABLET 801 MG	T5	PA; QL (93 EA per 31 days)
FASENRA	T5	PA; QL (1 ML per 56 days)
FASENRA PEN	T5	PA; QL (1 ML per 56 days)
<i>flunisolide</i>	T2	QL (50 ML per 25 days)
<i>fluticasone propionate nasal</i>	T1	QL (16 GM per 30 days)
<i>fluticasone propion-salmeterol inhalation aerosol powdr breath activated</i>	T3	QL (1 EA per 30 days)
<i>fluticasone propion-salmeterol inhalation blister with device</i>	T3	QL (60 EA per 30 days)
HAEGARDA	T5	PA
<i>hydroxyzine hcl oral solution 10 mg/5 ml</i>	T2	PA
<i>hydroxyzine hcl oral tablet</i>	T2	PA
<i>icatibant</i>	T5	PA; QL (18 ML per 30 days)
INCRUSE ELLIPTA	T3	QL (30 EA per 30 days)
<i>ipratropium bromide inhalation</i>	T1	PA-BvD
<i>ipratropium-albuterol</i>	T2	PA-BvD
KALYDECO ORAL GRANULES IN PACKET 25 MG	T5	PA; QL (62 EA per 31 days)
KALYDECO ORAL GRANULES IN PACKET 50 MG, 75 MG	T5	PA; QL (56 EA per 28 days)
KALYDECO ORAL TABLET	T5	PA; QL (62 EA per 31 days)
<i>levalbuterol hcl inhalation solution for nebulization 0.31 mg/3 ml, 0.63 mg/3 ml, 1.25 mg/3 ml</i>	T2	PA-BvD
<i>levalbuterol tartrate</i>	T3	QL (30 GM per 30 days)
<i>levocetirizine oral solution</i>	T2	QL (310 ML per 31 days)
<i>levocetirizine oral tablet</i>	T1	QL (31 EA per 31 days)
<i>mometasone nasal</i>	T3	QL (34 GM per 30 days)
<i>montelukast</i>	T2	QL (31 EA per 31 days)
NUCALA SUBCUTANEOUS AUTO-INJECTOR	T5	PA; QL (3 ML per 28 days)
NUCALA SUBCUTANEOUS RECON SOLN	T5	PA; QL (3 EA per 28 days)
NUCALA SUBCUTANEOUS SYRINGE 100 MG/ML	T5	PA; QL (3 ML per 28 days)

Drug Name	Drug Tier	Requirements/Limits
NUCALA SUBCUTANEOUS SYRINGE 40 MG/0.4 ML	T5	PA; QL (0.4 ML per 28 days)
OFEV	T5	PA; QL (62 EA per 31 days)
OMNARIS	T4	
OPSUMIT	T5	PA; QL (31 EA per 31 days)
ORKAMBI ORAL GRANULES IN PACKET 100-125 MG, 150-188 MG	T5	PA; QL (62 EA per 31 days)
ORKAMBI ORAL TABLET	T5	PA; QL (124 EA per 31 days)
ORLADEYO	T5	PA; QL (31 EA per 31 days)
<i>pirfenidone oral tablet 267 mg</i>	T5	PA; QL (279 EA per 31 days)
<i>pirfenidone oral tablet 801 mg</i>	T5	PA; QL (93 EA per 31 days)
<i>promethazine oral syrup</i>	T2	PA
<i>promethazine rectal suppository 12.5 mg, 25 mg</i>	T2	
PULMOZYME	T5	PA
QVAR REDHALER INHALATION HFA AEROSOL BREATH ACTIVATED 40 MCG/ACTUATION	T3	QL (10.6 GM per 30 days)
QVAR REDHALER INHALATION HFA AEROSOL BREATH ACTIVATED 80 MCG/ACTUATION	T3	QL (21.2 GM per 30 days)
RUCONEST	T5	PA
SAJAZIR	T5	PA; QL (18 ML per 30 days)
SEREVENT DISKUS	T3	QL (60 EA per 30 days)
<i>sildenafil (pulm.hypertension) oral suspension for reconstitution</i>	T4	PA; QL (224 ML per 31 days)
<i>sildenafil (pulm.hypertension) oral tablet</i>	T3	PA; QL (93 EA per 31 days)
SPIRIVA RESPIMAT	T3	QL (4 GM per 30 days)
SPIRIVA WITH HANDHALER	T3	QL (30 EA per 30 days)
STIOLTO RESPIMAT	T3	QL (4 GM per 30 days)
STRIVERDI RESPIMAT	T4	QL (4 GM per 30 days)
SYMBICORT	T3	QL (10.2 GM per 30 days)
SYMDEKO	T5	PA; QL (56 EA per 28 days)
<i>tadalafil (pulm. hypertension)</i>	T5	PA; QL (62 EA per 31 days)
TAKHZYRO	T5	PA; QL (4 ML per 28 days)
<i>terbutaline oral</i>	T4	
THEO-24	T4	
<i>theophylline oral solution</i>	T2	
<i>theophylline oral tablet extended release 12 hr 300 mg, 450 mg</i>	T2	
<i>theophylline oral tablet extended release 24 hr</i>	T2	

Drug Name	Drug Tier	Requirements/Limits
TRACLEER ORAL TABLET FOR SUSPENSION	T5	PA; QL (124 EA per 31 days)
TRELEGY ELLIPTA	T3	QL (60 EA per 30 days)
TRIKAFTA	T5	PA; QL (84 EA per 28 days)
TYVASO DPI INHALATION CARTRIDGE WITH INHALER 16 MCG (112)- 32 MCG (84)	T5	PA; QL (392 EA per 365 days)
TYVASO DPI INHALATION CARTRIDGE WITH INHALER 16 MCG, 32 MCG, 32-48 MCG, 48 MCG, 64 MCG	T5	PA
TYVASO DPI INHALATION CARTRIDGE WITH INHALER 16(112)-32(112) -48(28) MCG	T5	PA; QL (504 EA per 365 days)
VENTAVIS	T5	PA
VENTOLIN HFA	T3	QL (36 GM per 30 days)
WIXELA INHUB	T3	QL (60 EA per 30 days)
XOLAIR	T5	PA
YUPELRI	T5	PA-BvD
<i>zafirlukast oral tablet 10 mg</i>	T2	QL (93 EA per 31 days)
<i>zafirlukast oral tablet 20 mg</i>	T2	QL (62 EA per 31 days)
<i>zileuton</i>	T5	PA; QL (124 EA per 31 days)
ZYFLO	T4	PA; QL (124 EA per 31 days)
Urologicals		
<i>alfuzosin</i>	T2	QL (31 EA per 31 days)
<i>bethanechol chloride</i>	T2	
CIALIS ORAL TABLET 2.5 MG	T4	PA; QL (62 EA per 31 days)
CIALIS ORAL TABLET 5 MG	T4	PA; QL (31 EA per 31 days)
CYSTAGON	T3	
<i>darifenacin</i>	T3	QL (31 EA per 31 days)
<i>dutasteride</i>	T2	QL (31 EA per 31 days)
<i>dutasteride-tamsulosin</i>	T3	QL (31 EA per 31 days)
ELMIRON	T4	
<i>finasteride oral tablet 5 mg</i>	T2	
<i>flavoxate</i>	T2	
GELNIQUE TRANSDERMAL GEL IN PACKET	T4	QL (30 GM per 30 days)
MYRBETRIQ ORAL SUSPENSION,EXTENDED REL RECON	T3	QL (300 ML per 30 days)
MYRBETRIQ ORAL TABLET EXTENDED RELEASE 24 HR	T3	QL (31 EA per 31 days)

Drug Name	Drug Tier	Requirements/Limits
<i>oxybutynin chloride oral syrup</i>	T2	
<i>oxybutynin chloride oral tablet</i>	T2	
<i>oxybutynin chloride oral tablet extended release 24hr 10 mg, 5 mg</i>	T2	QL (31 EA per 31 days)
<i>oxybutynin chloride oral tablet extended release 24hr 15 mg</i>	T3	QL (62 EA per 31 days)
OXYTROL	T4	QL (8 EA per 28 days)
<i>potassium citrate oral tablet extended release</i>	T2	
PROCYSBI ORAL GRANULES DEL RELEASE IN PACKET	T5	PA
<i>silodosin</i>	T4	
<i>solifenacin</i>	T4	QL (31 EA per 31 days)
<i>tadalafil oral tablet 2.5 mg</i>	T4	PA; QL (62 EA per 31 days)
<i>tadalafil oral tablet 5 mg</i>	T4	PA; QL (31 EA per 31 days)
<i>tamsulosin</i>	T1	
<i>tolterodine oral capsule,extended release 24hr</i>	T3	QL (31 EA per 31 days)
<i>tolterodine oral tablet</i>	T3	QL (62 EA per 31 days)
<i>trospium oral capsule,extended release 24hr</i>	T2	QL (31 EA per 31 days)
<i>trospium oral tablet</i>	T2	QL (93 EA per 31 days)
Vitamins, Hematinics / Electrolytes		
<i>calcium acetate(phosphat bind)</i>	T2	
CLINIMIX 5%/D15W SULFITE FREE	T4	PA-BvD
CLINIMIX 4.25%/D10W SULF FREE	T4	PA-BvD
CLINIMIX 5%-D20W(SULFITE-FREE)	T4	PA-BvD
CLINIMIX E 4.25%/D10W SUL FREE	T4	PA-BvD
CLINIMIX E 4.25%/D5W SULF FREE	T4	PA-BvD
CLINIMIX E 5%/D15W SULFIT FREE	T4	PA-BvD
CLINIMIX E 5%/D20W SULFIT FREE	T4	PA-BvD
DOJOLVI	T5	PA
<i>fluoride (sodium) oral tablet</i>	T2	
INTRALIPID INTRAVENOUS EMULSION 20 %, 30 %	T4	PA-BvD
ISOLYTE S PH 7.4	T3	PA-BvD
ISOLYTE-P IN 5 % DEXTROSE	T3	PA-BvD
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Drug Name	Drug Tier	Requirements/Limits
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K-TAB ORAL TABLET EXTENDED RELEASE 10 MEQ, 20 MEQ	T4	
<i>magnesium sulfate injection</i>	T2	
PHOSLYRA	T4	
PLENAMINE	T3	PA-BvD
<i>potassium chlorid-d5-0.45%nacl</i>	T2	
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<i>potassium chloride in 5 % dex intravenous parenteral solution 20 meq/l</i>	T2	
<i>potassium chloride in lr-d5 intravenous parenteral solution 20 meq/l</i>	T2	
<i>potassium chloride in water intravenous piggyback 10 meq/100 ml, 20 meq/100 ml, 40 meq/100 ml</i>	T2	
<i>potassium chloride intravenous</i>	T2	
<i>potassium chloride oral capsule, extended release</i>	T1	
<i>potassium chloride oral liquid</i>	T2	
<i>potassium chloride oral tablet extended release</i>	T1	
<i>potassium chloride oral tablet,er particles/crystals</i>	T1	
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<i>potassium chloride-d5-0.9%nacl</i>	T2	
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PROSOL 20 %	T4	PA-BvD
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<i>clobetasol</i>	<i>d5 %-0.45 % sodium chloride</i>	50	DIFICID	7
<i>clobetasol-emollient</i>	<i>dalfampridine</i>	23	<i>diflunisal</i>	24
<i>clocortolone pivalate</i>	DALIRESP	73	<i>difluprednate</i>	71
CLODAN	DALVANCE	7	DIGITEK	39
<i>clomipramine</i>	<i>danazol</i>	53	<i>digoxin</i>	39
<i>clonazepam</i>	<i>dantrolene</i>	23	<i>dihydroergotamine</i>	24
<i>clonidine</i>	<i>dapsone</i>	7, 46	DILANTIN	24
<i>clonidine hcl</i>	DAPTACEL (DTAP		DILANTIN EXTENDED	24
<i>clopidogrel</i>	PEDIATRIC) (PF)	61	DILANTIN INFATABS	24
<i>clorazepate dipotassium</i>	<i>daptomycin</i>	7	DILANTIN-125	24
<i>clotrimazole</i>	<i>darifenacin</i>	76	<i>diltiazem hcl</i>	39
<i>clotrimazole-betamethasone</i>	DAURISMO	14	DILT-XR	39
<i>clozapine</i>	DAYTRANA	23	<i>dimethyl fumarate</i>	24
COARTEM	<i>deferasirox</i>	50	<i>diphenoxylate-atropine</i>	58
<i>colesevelam</i>	<i>deferiprone</i>	50	<i>disulfiram</i>	50
<i>colestipol</i>	DELSTRIGO	7	<i>divalproex</i>	24
<i>colistin (colistimethate na)</i>	<i>demeclocycline</i>	7	DIVIGEL	67
COMBIGAN	DENAVIR	46	<i>dofetilide</i>	39
COMBIVENT RESPIMAT	DEPO-ESTRADIOL	67	DOJOLVI	77
COMETRIQ	DEPO-TESTOSTERONE	53	DOLISHALE	67
COMPLERA	DESCOVY	7	<i>donepezil</i>	24
COMPRO	<i>desipramine</i>	23	DOPTELET (10 TAB PACK) ..	39
CONSTULOSE	<i>desloratadine</i>	74	DOPTELET (15 TAB PACK) ..	39
COPAXONE	<i>desmopressin</i>	53	DOPTELET (30 TAB PACK) ..	39
COPIKTRA	<i>desog-e.estradiol/e.estradiol</i>	67	<i>dorzolamide</i>	71
CORLANOR	<i>desogestrel-ethinyl estradiol</i>	67	<i>dorzolamide-timolol</i>	71
CORTROPHIN GEL	<i>desonide</i>	46	DOTTI	67
COSENTYX	<i>desoximetasone</i>	46	DOVATO	7
COSENTYX (2 SYRINGES) ..	DESRX	46	<i>doxazosin</i>	39
COSENTYX PEN (2 PENS)	<i>desvenlafaxine</i>	23	<i>doxepin</i>	24, 46
COTELLIC	<i>desvenlafaxine succinate</i>	23	<i>doxercalciferol</i>	53
CREON	<i>dexamethasone</i>	53	DOXY-100	7
CRESEMBA	<i>dexamethasone sodium</i>		<i>doxycycline hyclate</i>	7
CRINONE	<i>phosphate</i>	71	<i>doxycycline monohydrate</i>	7, 8
<i>cromolyn</i>	<i>dexmethylphenidate</i>	23	DRIZALMA SPRINKLE ...24, 25	

<i>dronabinol</i>	58	ENSKYCE	67	EVAMIST	68
<i>drospirenone-ethinyl estradiol</i>	67	ENSPRYNG	14	EVENTITY	65
DROXIA	14	<i>entacapone</i>	25	<i>everolimus (antineoplastic)</i>	15
<i>droxidopa</i>	50	<i>entecavir</i>	8	<i>everolimus</i>	
<i>duloxetine</i>	25	ENTRESTO	39	<i>(immunosuppressive)</i>	15
DUOBRII	46	ENULOSE	58	EVOTAZ	8
DUOPA	25	ENVARUS XR	15	EVRYSDI	25
DUPIXENT PEN	46	EPCLUSA	8	<i>exemestane</i>	15
DUPIXENT SYRINGE	46	EPIDIOLEX	25	EXKIVITY	15
<i>dutasteride</i>	76	<i>epinastine</i>	71	EXSERVAN	50
<i>dutasteride-tamsulosin</i>	76	<i>epinephrine</i>	74	EXTAVIA	61
E.E.S. 400	8	EPITOL	25	<i>ezetimibe</i>	39
<i>econazole</i>	46	EPIVIR HBV	8	<i>ezetimibe-simvastatin</i>	39
EDARBYCLOR	39	<i>eplerenone</i>	39	<i>famciclovir</i>	8
EDURANT	8	EPOGEN	61	<i>famotidine</i>	58
<i>efavirenz</i>	8	EPRONTIA	25	FANAPT	26
<i>efavirenz-emtricitabin-tenofov</i>	8	EQUETRO	25	FASENRA	74
<i>efavirenz-lamivu-tenofov disop</i>	8	ERAXIS(WATER DILUENT) ..	8	FASENRA PEN	74
EGRIFTA SV	61	<i>ergoloid</i>	25	<i>felbamate</i>	26
<i>eletriptan</i>	25	<i>ergotamine-caffeine</i>	25	<i>felodipine</i>	40
ELIGARD	14	ERIVEDGE	15	FEMRING	68
ELIGARD (3 MONTH)	14	ERLEADA	15	FEMYNOR	68
ELIGARD (4 MONTH)	14	<i>erlotinib</i>	15	<i>fenofibrate</i>	40
ELIGARD (6 MONTH)	14	ERRIN	67	<i>fenofibrate micronized</i>	40
ELIQUIS	39	<i>ertapenem</i>	8	<i>fenofibrate nanocrystallized</i>	40
ELIQUIS DVT-PE TREAT		ERY PADS	46	<i>fenofibric acid (choline)</i>	40
30D START	39	ERYGEL	47	<i>fenopropfen</i>	26
ELMIRON	76	ERYPED 200	8	<i>fentanyl</i>	26
ELURYNG	67	ERY-TAB	8	<i>fentanyl citrate</i>	26
EMCYT	14	ERYTHROCIN	8	FENTORA	26
EMEND	58	ERYTHROCIN (AS		FERRIPROX	50
EMFLAZA	53	STEARATE)	8	FETZIMA	26, 27
EMGALITY PEN	25	<i>erythromycin</i>	8, 71	FIASP FLEXTOUCH U-100	
EMGALITY SYRINGE	25	<i>erythromycin ethylsuccinate</i>	8	INSULIN	53
EMOQUETTE	67	<i>erythromycin with ethanol</i>	47	FIASP PENFILL U-100	
EMSAM	25	<i>erythromycin-benzoyl peroxide</i> ..	47	INSULIN	53
<i>emtricitabine</i>	8	ESBRIET	74	FIASP U-100 INSULIN	53
<i>emtricitabine-tenofovir (tdf)</i>	8	<i>escitalopram oxalate</i>	25	FINACEA	47
EMTRIVA	8	<i>esomeprazole magnesium</i>	58	<i>finasteride</i>	76
EMVERM	8	ESTARYLLA	67	<i> fingolimod</i>	27
<i>enalapril maleate</i>	39	<i>estazolam</i>	25	FINTEPLA	27
<i>enalapril-hydrochlorothiazide</i>	39	<i>estradiol</i>	67	FIRDAPSE	27
ENBREL	65	<i>estradiol-norethindrone acet</i>	67	FIRMAGON KIT W	
ENBREL MINI	65	ESTRING	68	DILUENT SYRINGE	15
ENBREL SURECLICK	65	<i>eszopiclone</i>	25	FIRVANQ	8
ENDARI	50	<i>ethacrynic acid</i>	39	<i>flavoxate</i>	76
ENDOCET	25	<i>ethambutol</i>	8	FLEBOGAMMA DIF	61
ENGERIX-B (PF)	61	<i>ethosuximide</i>	25	<i>flecainide</i>	40
ENGERIX-B PEDIATRIC		<i>ethynodiol diac-eth estradiol</i>	68	FLECTOR	27
(PF)	61	<i>etodolac</i>	25	<i>fluconazole</i>	8
<i>enoxaparin</i>	39	<i>etonogestrel-ethinyl estradiol</i>	68	<i>fluconazole in nacl (iso-osm)</i>	8
ENPRESSE	67	<i>etravirine</i>	8	<i>flucytosine</i>	8

<i>fludrocortisone</i>	53	GAVILYTE-C	58	HIBERIX (PF)	62
<i>flunisolide</i>	74	GAVILYTE-G	58	HORIZANT	28
<i>fluocinolone</i>	47	GAVRETO	15	HUMALOG JUNIOR	
<i>fluocinolone acetonide oil</i>	52	GELNIQUE	76	KWIKPEN U-100	53
<i>fluocinolone and shower cap</i>	47	<i>gemfibrozil</i>	40	HUMALOG KWIKPEN	
<i>fluocinonide</i>	47	GENERLAC	58	INSULIN	53
FLUOCINONIDE-E	47	GENGRAF	15	HUMALOG MIX 50-50	
<i>fluoride (sodium)</i>	77	GENOTROPIN	62	INSULN U-100	53
<i>fluorometholone</i>	71	GENOTROPIN MINIQUICK			
<i>fluorouracil</i>	47	61, 62	HUMALOG MIX 50-50	
<i>fluoxetine</i>	27	GENTAK	71	KWIKPEN	54
<i>fluoxetine (pmd)</i>	27	<i>gentamicin</i>	9, 47, 71	HUMALOG MIX 75-25	
<i>fluphenazine decanoate</i>	27	<i>gentamicin in nacl (iso-osm)</i>	9	KWIKPEN	54
<i>fluphenazine hcl</i>	27	GENVOYA	9	HUMALOG MIX 75-25(U-	
<i>flurandrenolide</i>	47	GILOTRIF	15	100)INSULN	54
<i>flurazepam</i>	27	GLASSIA	50	HUMALOG U-100 INSULIN ..	54
<i>flurbiprofen</i>	27	<i>glatiramer</i>	27	HUMATROPE	62
<i>flurbiprofen sodium</i>	71	GLATOPA	27	HUMIRA	65
<i>fluticasone propionate</i>	47, 74	<i>glimepiride</i>	53	HUMIRA PEN	65
<i>fluticasone propion-salmeterol</i> ..	74	<i>glipizide</i>	53	HUMIRA PEN CROHNS-	
<i>fluvastatin</i>	40	<i>glipizide-metformin</i>	53	UC-HS START	65
<i>fluvoxamine</i>	27	GLUCAGEN HYPOKIT	53	HUMIRA PEN PSOR-	
<i>fondaparinux</i>	40	GLUCAGON EMERGENCY			
FORTEO	65	KIT (HUMAN)	53	UVEITS-ADOL HS	65
<i>fosamprenavir</i>	9	<i>glyburide</i>	53	HUMIRA(CF)	65
<i>fosfomycin tromethamine</i>	9	<i>glyburide micronized</i>	53	HUMIRA(CF) PEDI	
<i>fosinopril</i>	40	<i>glyburide-metformin</i>	53	CROHNS STARTER	65
<i>fosinopril-hydrochlorothiazide</i> ..	40	<i>glycopyrrolate</i>	58	HUMIRA(CF) PEN	65
FOSRENOL	50	GLYXAMBI	53	HUMIRA(CF) PEN	
FOTIVDA	15	<i>granisetron hcl</i>	58	CROHNS-UC-HS	65
FRAGMIN	40	GRANIX	62	HUMIRA(CF) PEN	
<i>frovatriptan</i>	27	GRASTEK	62	PEDIATRIC UC	65
FULPHILA	61	<i>griseofulvin microsize</i>	9	HUMIRA(CF) PEN PSOR-	
<i>furosemide</i>	40	<i>griseofulvin ultramicrosize</i>	9	UV-ADOL HS	65
FUZEON	9	<i>guanfacine</i>	27	HUMULIN 70/30 U-100	
FYAVOLV	68	GVOKE	53	INSULIN	54
FYCOMPA	27	GVOKE HYPOPEN 2-PACK ..	53	HUMULIN 70/30 U-100	
<i>gabapentin</i>	27	GVOKE PFS 1-PACK		KWIKPEN	54
GALAFOLD	53	SYRINGE	53	HUMULIN N NPH INSULIN	
<i>galantamine</i>	27	HAEGARDA	74	KWIKPEN	54
GAMMAGARD LIQUID	61	HAILEY 24 FE	68	HUMULIN N NPH U-100	
GAMMAGARD S-D (IGA < 1					
MCG/ML)	61	<i>halcinonide</i>	47	INSULIN	54
GAMMAKED	61	<i>halobetasol propionate</i>	47	HUMULIN R REGULAR U-	
GAMMAPLEX	61	HALOG	47	100 INSULN	54
GAMMAPLEX (WITH					
SORBITOL)	61	<i>haloperidol</i>	28	HUMULIN R U-500 (CONC)	
GAMUNEX-C	61	<i>haloperidol decanoate</i>	27	INSULIN	54
GARDASIL 9 (PF)	61	<i>haloperidol lactate</i>	28	HUMULIN R U-500 (CONC)	
<i>gatifloxacin</i>	71	HARVONI	9	KWIKPEN	54
GATTEX 30-VIAL	58	HAVRIX (PF)	62	<i>hydralazine</i>	40
GAUZE PAD	64	<i>heparin (porcine)</i>	40	<i>hydrochlorothiazide</i>	40
		HETLIOZ	28	<i>hydrocodone-acetaminophen</i>	28
		HETLIOZ LQ	28	<i>hydrocodone-ibuprofen</i>	28
				<i>hydrocortisone</i>	47, 54, 58
				<i>hydrocortisone butyrate</i>	47

<i>hydrocortisone valerate</i>	47	INVEGA TRINZA	29	KELNOR 1-50 (28)	68
<i>hydrocortisone-acetic acid</i>	52	INVOKAMET	54	KESIMPTA PEN	29
<i>hydrocortisone-pramoxine</i>	58	INVOKAMET XR	54	<i>ketoconazole</i>	9, 47, 48
<i>hydromorphone</i>	28	INVOKANA	54	<i>ketoprofen</i>	29
<i>hydroxychloroquine</i>	9	IOPIDINE	71	<i>ketorolac</i>	29, 71
<i>hydroxyurea</i>	15	IPOL	62	KEVZARA	65
<i>hydroxyzine hcl</i>	74	<i>ipratropium bromide</i>	52, 74	KINERET	65
<i>ibandronate</i>	65	<i>ipratropium-albuterol</i>	74	KINRIX (PF)	62
IBRANCE	15	<i>irbesartan</i>	40	KISQALI	16
IBSRELA	58	<i>irbesartan-hydrochlorothiazide</i> ..	40	KISQALI FEMARA CO-	
IBU	28	IRESSA	15	PACK	15, 16
<i>ibuprofen</i>	28	ISENTRESS	9	KLISYRI	16
<i>icatibant</i>	74	ISENTRESS HD	9	KLOR-CON	77
ICLEVIA	68	ISIBLOOM	68	KLOR-CON 10	77
ICLUSIG	15	ISOLYTE S PH 7.4	77	KLOR-CON 8	77
<i>icosapent ethyl</i>	40	ISOLYTE-P IN 5 %		KLOR-CON M10	77
IDHIFA	15	DEXTROSE	77	KLOR-CON M15	77
ILEVRO	71	<i>isoniazid</i>	9	KLOR-CON M20	78
<i>imatinib</i>	15	ISORDIL	40	KLOXXADO	29
IMBRUVICA	15	<i>isosorbide dinitrate</i>	40	KORLYM	55
<i>imipenem-cilastatin</i>	9	<i>isosorbide mononitrate</i>	40	KOSELUGO	16
<i>imipramine hcl</i>	28	<i>isosorbide-hydralazine</i>	41	K-TAB	78
<i>imipramine pamoate</i>	28	<i>isotretinoin</i>	47	KURVELO (28)	68
<i>imiquimod</i>	47	<i>isradipine</i>	41	KYNMOBI	29
IMOVAX RABIES		ISTURISA	54	<i>labetalol</i>	41
VACCINE (PF)	62	<i>itraconazole</i>	9	<i>lacosamide</i>	29
IMPAVIDO	9	<i>ivermectin</i>	9	LACRISERT	71
INBRIJA	28	IXIARO (PF)	62	<i>lactulose</i>	58
INCASSIA	68	JAKAFI	15	<i>lamivudine</i>	9
INCRELEX	50	JANTOVEN	41	<i>lamivudine-zidovudine</i>	9
INCRUSE ELLIPTA	74	JANUMET	54	<i>lamotrigine</i>	29
<i>indapamide</i>	40	JANUMET XR	54	<i>lansoprazole</i>	58, 59
INDOCIN	28	JANUVIA	54	<i>lanthanum</i>	50
<i>indomethacin</i>	28	JARDIANCE	54	LANTUS SOLOSTAR U-100	
INFANRIX (DTAP) (PF)	62	JASMIEL (28)	68	INSULIN	55
INGREZZA	28	JAVYGTOR	54	LANTUS U-100 INSULIN	55
INGREZZA INITIATION		JENTADUETO	54	<i>lapatinib</i>	16
PACK	28	JENTADUETO XR	54	<i>latanoprost</i>	71
INLYTA	15	JINTELI	68	LATUDA	29
INQOVI	15	JULEBER	68	<i>ledipasvir-sofosbuvir</i>	9
INREBIC	15	JULUCA	9	<i>leflunomide</i>	65
<i>insulin asp prt-insulin aspart</i>	54	JUNEL 1.5/30 (21)	68	<i>lenalidomide</i>	16
<i>insulin aspart u-100</i>	54	JUNEL 1/20 (21)	68	LENVIMA	16
<i>insulin lispro</i>	54	JUNEL FE 1.5/30 (28)	68	LESSINA	68
<i>insulin lispro protamin-lispro</i>	54	JUNEL FE 1/20 (28)	68	<i>letrozole</i>	16
<i>insulin syringe-needle u-100</i>	64	JUNEL FE 24	68	<i>leucovorin calcium</i>	16
INTELENCE	9	JUXTAPID	41	LEUKERAN	16
INTRALIPID	77	JYNARQUE	54, 55	LEUKINE	62
INTRON A	62	KAITLIB FE	68	<i>leuprolide</i>	16
INTROVALE	68	KALYDECO	74	<i>levalbuterol hcl</i>	74
INVEGA HAFYERA	28	KARIVA (28)	68	<i>levalbuterol tartrate</i>	74
INVEGA SUSTENNA	28, 29	KELNOR 1/35 (28)	68		

LEVEMIR FLEXTOUCH U-100 INSULN	55	LUPRON DEPOT (3 MONTH)	16	MENQUADFI (PF)	62
LEVEMIR U-100 INSULIN	55	LUPRON DEPOT (4 MONTH)	16	MENTAX	48
<i>levetiracetam</i>	29	LUPRON DEPOT (6 MONTH)	16	MENVEO A-C-Y-W-135-DIP (PF)	62
<i>levobunolol</i>	71	LUTERA (28)	68	<i>mercaptopurine</i>	16
<i>levocarnitine</i>	51	LYBALVI	29	<i>meropenem</i>	10
<i>levocarnitine (with sugar)</i>	50	LYLEQ	68	<i>mesalamine</i>	59
<i>levocetirizine</i>	74	LYLLANA	68	MESNEX	17
<i>levofloxacin</i>	9, 71	LYNPARZA	16	<i>metaxalone</i>	30
<i>levofloxacin in d5w</i>	9	LYSODREN	16	<i>metformin</i>	55
LEVONEST (28)	68	LYZA	68	<i>methadone</i>	30
<i>levonorgestrel-ethinyl estrad</i>	68	<i>magnesium sulfate</i>	78	<i>methamphetamine</i>	30
<i>levonorg-eth estrad triphasic</i>	68	<i>malathion</i>	48	<i>methazolamide</i>	71
LEVORA-28	68	<i>maraviroc</i>	10	<i>methenamine hippurate</i>	10
<i>levothyroxine</i>	55	MARLISSA (28)	68	<i>methimazole</i>	55
LEVOXYL	55	MARPLAN	29	<i>methotrexate sodium</i>	17
LEXIVA	9	MATULANE	16	<i>methotrexate sodium (pf)</i>	17
<i>lidocaine</i>	48	MATZIM LA	41	<i>methscopolamine</i>	59
<i>lidocaine hcl</i>	48	MAVENCLAD (10 TABLET PACK)	29	<i>methylphenidate hcl</i>	30, 31
LIDOCAINE VISCOUS	48	MAVENCLAD (4 TABLET PACK)	29	<i>methylprednisolone</i>	55
<i>lidocaine-prilocaine</i>	48	MAVENCLAD (5 TABLET PACK)	29	<i>metoclopramide hcl</i>	59
LIDODERM	48	MAVENCLAD (6 TABLET PACK)	30	<i>metolazone</i>	41
<i>linezolid</i>	9	MAVENCLAD (7 TABLET PACK)	30	<i>metoprolol succinate</i>	41
<i>linezolid in dextrose 5%</i>	10	MAVENCLAD (8 TABLET PACK)	30	<i>metoprolol ta-hydrochlorothiaz</i> ..	41
LINZESS	59	MAVENCLAD (9 TABLET PACK)	30	<i>metoprolol tartrate</i>	41
<i>liothyronine</i>	55	MAVENCLAD (10 TABLET PACK)	30	<i>metronidazole</i>	10, 48, 69
LIPOFEN	41	MAVENCLAD (11 TABLET PACK)	30	<i>metronidazole in nacl (iso-os)</i>	10
<i>lisinopril</i>	41	MAVENCLAD (12 TABLET PACK)	30	<i>metyrosine</i>	41
<i>lisinopril-hydrochlorothiazide</i>	41	MAVENCLAD (13 TABLET PACK)	30	<i>mexiletine</i>	41
<i>lithium carbonate</i>	29	MAVENCLAD (14 TABLET PACK)	30	<i>micafungin</i>	10
LITHOSTAT	51	MAVENCLAD (15 TABLET PACK)	30	MICONAZOLE-3	69
LIVALO	41	MAVENCLAD (16 TABLET PACK)	30	MICROGESTIN 1.5/30 (21)	69
LIVTENCITY	10	MAVENCLAD (17 TABLET PACK)	30	MICROGESTIN 1/20 (21)	69
LOKELMA	51	MAVENCLAD (18 TABLET PACK)	30	MICROGESTIN FE 1.5/30 (28)	69
LONSURF	16	MAVENCLAD (19 TABLET PACK)	30	MICROGESTIN FE 1/20 (28) ..	69
<i>loperamide</i>	59	MAVENCLAD (20 TABLET PACK)	30	<i>midodrine</i>	51
<i>lopinavir-ritonavir</i>	10	MAVENCLAD (21 TABLET PACK)	30	<i>miglitol</i>	55
<i>lorazepam</i>	29	MAVENCLAD (22 TABLET PACK)	30	<i>miglustat</i>	55
LORAZEPAM INTENSOL	29	MAVENCLAD (23 TABLET PACK)	30	MILI	69
LORBRENA	16	MAVENCLAD (24 TABLET PACK)	30	<i>minocycline</i>	10
LORYNA (28)	68	MAVENCLAD (25 TABLET PACK)	30	<i>minoxidil</i>	41
<i>losartan</i>	41	MAVENCLAD (26 TABLET PACK)	30	<i>mirtazapine</i>	31
<i>losartan-hydrochlorothiazide</i>	41	MAVENCLAD (27 TABLET PACK)	30	<i>misoprostol</i>	59
<i>lovastatin</i>	41	MAVENCLAD (28 TABLET PACK)	30	MITIGARE	65
LOW-OGESTREL (28)	68	MAVENCLAD (29 TABLET PACK)	30	M-M-R II (PF)	62
<i>loxapine succinate</i>	29	MAVENCLAD (30 TABLET PACK)	30	<i>modafinil</i>	31
<i>lubiprostone</i>	59	MAVENCLAD (31 TABLET PACK)	30	<i>moexipril</i>	41
LUCEMYRA	29	MAVENCLAD (32 TABLET PACK)	30	<i>molindone</i>	31
LUMAKRAS	16	MAVENCLAD (33 TABLET PACK)	30	<i>mometasone</i>	48, 74
LUMIGAN	71	MAVENCLAD (34 TABLET PACK)	30	<i>montelukast</i>	74
LUPKYNIS	16	MAVENCLAD (35 TABLET PACK)	30	<i>morphine</i>	31
LUPRON DEPOT	16	MAVENCLAD (36 TABLET PACK)	30		

<i>morphine concentrate</i>	31	NEVANAC	72	NOVOLOG MIX 70-	
MOUNJARO	55	<i>nevirapine</i>	10	30FLEXPEN U-100	55
MOVANTIK	59	NEXLETOL	41	NOVOLOG PENFILL U-100	
<i>moxifloxacin</i>	10, 71	NEXLIZET	41	INSULIN	55
<i>moxifloxacin-sod.chloride(iso)</i> ...	10	<i>niacin</i>	41	NOVOLOG U-100 INSULIN	
MULPLETA	41	NIACOR	41	ASPART	55
MULTAQ	41	<i>nicardipine</i>	41	NUBEQA	17
<i>mupirocin</i>	48	NICOTROL	51	NUCALA	74, 75
MYALEPT	55	NICOTROL NS	51	NUCYNTA	32
MYCAPSSA	17	<i>nifedipine</i>	41, 42	NUDEXTA	32
<i>mycophenolate mofetil</i>	17	<i>nilutamide</i>	17	NUPLAZID	32
<i>mycophenolate sodium</i>	17	<i>nimodipine</i>	42	NURTEC ODT	32
MYFEMBREE	69	NINLARO	17	NUTROPIN AQ NUSPIN	63
MYORISAN	48	<i>nisoldipine</i>	42	NUZYRA	10
MYRBETRIQ	76	<i>nitazoxanide</i>	10	NYAMYC	48
MYTESI	59	<i>nitisinone</i>	51	NYLIA 1/35 (28)	69
<i>nabumetone</i>	31	NITRO-BID	42	NYLIA 7/7/7 (28)	69
<i>nadolol</i>	41	<i>nitrofurantoin macrocrystal</i>	10	NYMALIZE	42
<i>nafcillin</i>	10	<i>nitrofurantoin monohyd/m-cryst</i> .	10	NYMYO	69
<i>naftifine</i>	48	<i>nitroglycerin</i>	42	<i>nystatin</i>	10, 48
NAFTIN	48	NITROSTAT	42	<i>nystatin-triamcinolone</i>	48
<i>naloxone</i>	31	NITYR	51	NYSTOP	48
<i>naltrexone</i>	31	NIVESTYM	62	NYVEPRIA	63
NAMENDA TITRATION		<i>nizatidine</i>	59	OCALIVA	59
PAK	31	NORDITROPIN FLEXPRO ...	63	OCTAGAM	63
NAMZARIC	31	<i>noreth-ethinyl estradiol-iron</i>	69	<i>octreotide acetate</i>	17
NAPRELAN CR	31	<i>norethindrone (contraceptive)</i>	69	ODACTRA	63
<i>naproxen</i>	31	<i>norethindrone acetate</i>	69	ODEFSEY	10
<i>naproxen sodium</i>	31	<i>norethindrone ac-eth estradiol</i> ...	69	ODOMZO	17
<i>naratriptan</i>	31, 32	<i>norethindrone-e.estradiol-iron</i> ...	69	OFEV	75
NARCAN	32	<i>norgestimate-ethinyl estradiol</i> ...	69	<i>ofloxacin</i>	10, 52, 72
NATACYN	71	NORTREL 0.5/35 (28)	69	<i>olanzapine</i>	32
<i>nateglinide</i>	55	NORTREL 1/35 (21)	69	<i>olanzapine-fluoxetine</i>	32
NATPARA	55	NORTREL 1/35 (28)	69	<i>olmesartan</i>	42
NAYZILAM	32	NORTREL 7/7/7 (28)	69	<i>olmesartan-amlodipin-hcthiazyd</i> .	42
<i>nebivolol</i>	41	<i>nortriptyline</i>	32	<i>olmesartan-hydrochlorothiazide</i> .	42
NECON 0.5/35 (28)	69	NORVIR	10	<i>olopatadine</i>	52, 72
<i>nefazodone</i>	32	NOURIANZ	32	OLUMIANT	66
<i>neomycin</i>	10	NOVOLIN 70/30 U-100		<i>omega-3 acid ethyl esters</i>	42
<i>neomycin-bacitracin-poly-hc</i>	71	INSULIN	55	<i>omeprazole</i>	59
<i>neomycin-bacitracin-polymyxin</i> ..	72	NOVOLIN 70-30 FLEXPEN		OMNARIS	75
<i>neomycin-polymyxin b-</i>		U-100	55	OMNITROPE	63
<i>dexameth</i>	72	NOVOLIN N FLEXPEN	55	<i>ondansetron</i>	59
<i>neomycin-polymyxin-gramicidin</i>	72	NOVOLIN N NPH U-100		<i>ondansetron hcl</i>	59
<i>neomycin-polymyxin-hc</i>	52, 72	INSULIN	55	ONGENTYS	32
NEORAL	17	NOVOLIN R FLEXPEN	55	ONUREG	17
NEO-SYNALAR	48	NOVOLIN R REGULAR U-		ONZETRA XSAIL	32
NERLYNX	17	100 INSULN	55	OPSUMIT	75
NEUAC	48	NOVOLOG FLEXPEN U-100		OPZELURA	48
NEULASTA	62	INSULIN	55	ORALAIR	63
NEUPOGEN	62	NOVOLOG MIX 70-30 U-100		ORENCIA	66
NEUPRO	32	INSULN	55	ORENCIA CLICKJECT	66

ORENITRAM	42	<i>penicillin v potassium</i>	11	<i>potassium chloride-d5-</i>	
ORFADIN	51	PENTACEL (PF)	63	<i>0.2%nacl</i>	78
ORGOVYX	17	<i>pentamidine</i>	11	<i>potassium chloride-d5-</i>	
ORIAHNN	69	PENTASA	59	<i>0.9%nacl</i>	78
ORLISSA	56	<i>pentoxifylline</i>	42	<i>potassium citrate</i>	77
ORKAMBI	75	<i>perindopril erbumine</i>	42	PRALUENT PEN	42
ORLADEYO	75	PERIOGARD	52	<i>pramipexole</i>	33
<i>oseltamivir</i>	10	<i>permethrin</i>	48	<i>prasugrel</i>	42
OSMOPREP	59	<i>perphenazine</i>	32	<i>pravastatin</i>	42
OTEZLA	66	<i>perphenazine-amitriptyline</i>	32	<i>praziquantel</i>	11
OTEZLA STARTER	66	PERSERIS	32	<i>prazosin</i>	42
OTOVEL	52	PHEBURANE	51	PRED-G S.O.P.	72
OTREXUP (PF)	66	<i>phenelzine</i>	33	<i>prednicarbate</i>	48
<i>oxacillin</i>	11	<i>phenobarbital</i>	33	<i>prednisolone</i>	56
<i>oxacillin in dextrose(iso-osm)</i>	11	<i>phenoxybenzamine</i>	42	<i>prednisolone acetate</i>	72
<i>oxandrolone</i>	56	PHENYTEK	33	<i>prednisolone sodium phosphate</i>	
<i>oxaprozin</i>	32	<i>phenytoin</i>	33	56, 72
<i>oxazepam</i>	32	<i>phenytoin sodium extended</i>	33	<i>prednisone</i>	56
OXBRYTA	51	PHOSLYRA	78	PREFEST	69
<i>oxcarbazepine</i>	32	PIFELTRO	11	<i>pregabalin</i>	33
OXERVATE	72	<i>pilocarpine hcl</i>	51, 72	PREHEVBRIO (PF)	63
OXISTAT	48	<i>pimecrolimus</i>	48	PREMARIN	69
OXTELLAR XR	32	<i>pimozide</i>	33	PRENATAL VITAMIN	
<i>oxybutynin chloride</i>	77	PIMTREA (28)	69	PLUS LOW IRON	78
<i>oxycodone</i>	32	<i>pindolol</i>	42	PREVALITE	42
<i>oxycodone-acetaminophen</i>	32	<i>pioglitazone</i>	56	PREVYMIS	11
OXYTROL	77	<i>pioglitazone-glimepiride</i>	56	PREZCOBIX	11
OZEMPIC	56	<i>pioglitazone-metformin</i>	56	PREZISTA	11
PACERONE	42	<i>piperacillin-tazobactam</i>	11	PRIFTIN	11
<i>paliperidone</i>	32	PIQRAY	17	<i>primaquine</i>	11
PALYNZIQ	56	<i>pirfenidone</i>	75	<i>primidone</i>	33
PANDEL	48	PIRMELLA	69	PRIORIX (PF)	63
PANRETIN	48	<i>piroxicam</i>	33	PRIVIGEN	63
<i>pantoprazole</i>	59	PLEGRIDY	63	<i>probenecid</i>	66
PANZYGA	63	PLENAMINE	78	<i>probenecid-colchicine</i>	66
<i>paricalcitol</i>	56	<i>podofilox</i>	48	PROCENTRA	33
<i>paromomycin</i>	11	<i>polymyxin b sulfate</i>	11	<i>prochlorperazine</i>	59
<i>paroxetine hcl</i>	32	<i>polymyxin b sulf-trimethoprim</i>	72	<i>prochlorperazine maleate</i>	59
PASER	11	POMALYST	17	PROCRIT	63
PEDIARIX (PF)	63	PONVORY	33	PROCTO-PAK	59
PEDVAX HIB (PF)	63	PONVORY 14-DAY		PROCTOSOL HC	59
<i>peg 3350-electrolytes</i>	59	STARTER PACK	33	PROCTOZONE-HC	59
<i>peg3350-sod sul-nacl-kcl-asb-c</i>	59	PORTIA 28	69	PROCYSBI	77
PEGASYS	63	<i>potassium chlorid-d5-</i>		<i>progesterone micronized</i>	69
<i>peg-electrolyte soln</i>	59	<i>0.45%nacl</i>	78	PROGRAF	17
PEMAZYRE	17	<i>potassium chloride</i>	78	PROLASTIN-C	51
<i>pen needle, diabetic</i>	64	<i>potassium chloride in 0.9%nacl</i>	78	PROLIA	66
<i>penicillamine</i>	66	<i>potassium chloride in 5 % dex</i>	78	PROMACTA	42, 43
<i>penicillin g pot in dextrose</i>	11	<i>potassium chloride in lr-d5</i>	78	<i>promethazine</i>	75
<i>penicillin g potassium</i>	11	<i>potassium chloride in water</i>	78	<i>propafenone</i>	43
<i>penicillin g procaine</i>	11	<i>potassium chloride-0.45 % nacl</i>	78	<i>propranolol</i>	43
<i>penicillin g sodium</i>	11			<i>propylthiouracil</i>	56

PROQUAD (PF)	63	REXULTI	33	<i>sildenafil (pulm.hypertension)</i>	75
PROSOL 20 %	78	REYATAZ	11	SILIQ	49
<i>protriptyline</i>	33	REYVOW	33	<i>silodosin</i>	77
PULMOZYME	75	<i>ribavirin</i>	11	<i>silver sulfadiazine</i>	49
PURIXAN	17	RIDAURA	66	SIMBRINZA	72
PYLERA	59	<i>rifabutin</i>	11	SIMPONI	66
<i>pyrazinamide</i>	11	<i>rifampin</i>	11	<i>simvastatin</i>	43
<i>pyridostigmine bromide</i>	33	<i>riluzole</i>	51	<i>sirolimus</i>	18
<i>pyrimethamine</i>	11	<i>rimantadine</i>	11	SIRTURO	12
PYRUKYND	51	RINVOQ	66	SIVEXTRO	12
QINLOCK	17	<i>risedronate</i>	51, 66	SKYRIZI	49, 60
QUADRACEL (PF)	63	RISPERDAL CONSTA	33, 34	<i>sodium chloride</i>	51
<i>quetiapine</i>	33	<i>risperidone</i>	34	<i>sodium chloride 0.45 %</i>	78
QUILLIVANT XR	33	<i>ritonavir</i>	12	<i>sodium chloride 0.9 %</i>	51
<i>quinapril</i>	43	<i>rivastigmine</i>	34	<i>sodium chloride 3 % hypertonic</i>	78
<i>quinapril-hydrochlorothiazide</i>	43	<i>rivastigmine tartrate</i>	34	<i>sodium chloride 5 % hypertonic</i>	78
<i>quinidine gluconate</i>	43	<i>rizatriptan</i>	34	<i>sodium phenylbutyrate</i>	51
<i>quinidine sulfate</i>	43	<i>ropinirole</i>	34	<i>sodium polystyrene sulfonate</i>	51
<i>quinine sulfate</i>	11	<i>rosuvastatin</i>	43	<i>sofosbuvir-velpatasvir</i>	12
QVAR REDHALER	75	ROTARIX	64	<i>solifenacin</i>	77
RABAVERT (PF)	63	ROTATEQ VACCINE	64	SOLTAMOX	18
<i>rabeprazole</i>	60	ROWEEPRA	34	SOMAVERT	56
RADICAVA ORS STARTER		ROZLYTREK	17	<i>sorafenib</i>	18
KIT SUSP	33	RUBRACA	17	SORINE	43
<i>raloxifene</i>	66	RUCONEST	75	<i>sotalol</i>	43
<i>ramelteon</i>	33	<i>rufinamide</i>	34	SOTALOL AF	43
<i>ramipril</i>	43	RUKOBIA	12	SOVALDI	12
<i>ranolazine</i>	43	RYBELSUS	56	SPIRIVA RESPIMAT	75
<i>rasagiline</i>	33	RYDAPT	17	SPIRIVA WITH	
RASUVO (PF)	66	RYTARY	34	HANDHALER	75
RAVICTI	51	SAIZEN	64	<i>spironolactone</i>	43
REBIF (WITH ALBUMIN)	63	SAIZEN SAIZENPREP	64	<i>spironolacton-hydrochlorothiaz</i>	43
REBIF REBIDOSE	63, 64	SAJAZIR	75	SPRINTEC (28)	69
REBIF TITRATION PACK	64	SANCUSO	60	SPRITAM	34
RECLIPSEN (28)	69	SANDIMMUNE	17	SPRYCEL	18
RECOMBIVAX HB (PF)	64	SANTYL	49	SPS (WITH SORBITOL)	51
RECORLEV	56	<i>sapropterin</i>	56	SRONYX	69
RECTIV	60	SAVELLA	66	SSD	49
REDITREX (PF)	66	SCSEMBLIX	17	STELARA	49
REGRANEX	48	<i>scopolamine base</i>	60	STIOLTO RESPIMAT	75
RELENZA DISKHALER	11	SECUADO	34	STIVARGA	18
RELISTOR	60	<i>selegiline hcl</i>	34	<i>streptomycin</i>	12
<i>repaglinide</i>	56	<i>selenium sulfide</i>	49	STRIBILD	12
REPATHA PUSHTRONEX	43	SELZENTRY	12	STRIVERDI RESPIMAT	75
REPATHA SURECLICK	43	SEREVENT DISKUS	75	SUCRAID	60
REPATHA SYRINGE	43	SEROSTIM	64	<i>sucralfate</i>	60
RESTASIS	72	<i>sertraline</i>	34	<i>sulfacetamide sodium</i>	72
RESTASIS MULTIDOSE	72	<i>sevelamer carbonate</i>	51	<i>sulfacetamide sodium (acne)</i>	49
RETACRIT	64	<i>sevelamer hcl</i>	51	<i>sulfacetamide-prednisolone</i>	72
RETEVMO	17	SHINGRIX (PF)	64	<i>sulfadiazine</i>	12
RETIN-A MICRO PUMP	48	SIGNIFOR	18	<i>sulfamethoxazole-trimethoprim</i> ..	12
REVCIVI	51			SULFAMYLON	49

<i>sulfasalazine</i>	60	<i>temazepam</i>	35	TOUJEO SOLOSTAR U-300
<i>sulindac</i>	34	TENCON	35	INSULIN
<i>sumatriptan</i>	34	TENIVAC (PF)	64	TOVET EMOLLIENT
<i>sumatriptan succinate</i>	34, 35	<i>tenofovir disoproxil fumarate</i>	12	TRACLEER
<i>sumatriptan-naproxen</i>	35	TEPMETKO	18	TRADJENTA
<i>sunitinib</i>	18	<i>terazosin</i>	43	<i>tramadol</i>
SUNOSI	35	<i>terbinafine hcl</i>	12	<i>tramadol-acetaminophen</i>
SUPRAX	12	<i>terbutaline</i>	75	<i>trandolapril</i>
SUPREP BOWEL PREP KIT	60	<i>terconazole</i>	70	<i>tranexamic acid</i>
SYEDA	70	<i>teriparatide</i>	66	<i>tranylcypramine</i>
SYMBICORT	75	<i>testosterone</i>	57	TRAVASOL 10 %
SYMDEKO	75	<i>testosterone cypionate</i>	57	<i>travoprost</i>
SYMFI	12	<i>testosterone enanthate</i>	57	<i>trazodone</i>
SYMFI LO	12	<i>tetanus, diphtheria tox ped(pf)</i>	64	TRECATOR
SYMLINPEN 120	56	<i>tetrabenazine</i>	35	TRELEGY ELLIPTA
SYMLINPEN 60	56	<i>tetracycline</i>	12	TRELSTAR
SYMPAZAN	35	THALOMID	18	TRESIBA FLEXTOUCH U-
SYMPROIC	60	THEO-24	75	100
SYMTUZA	12	<i>theophylline</i>	75	TRESIBA FLEXTOUCH U-
SYNAREL	56	THIOLA EC	51	200
SYNJARDY	56	<i>thioridazine</i>	35	TRESIBA U-100 INSULIN
SYNJARDY XR	56	<i>thiothixene</i>	35	<i>tretinoin</i>
SYNRIBO	18	TIADYLT ER	43	<i>tretinoin (antineoplastic)</i>
SYNTHROID	56	<i>tiagabine</i>	35	TREXALL
TABLOID	18	TIBSOVO	18	<i>triamcinolone acetonide</i>
TABRECTA	18	TICOVAC	64	<i>triamterene</i>
<i>tacrolimus</i>	18, 49	<i>tigecycline</i>	12	<i>triamterene-hydrochlorothiazid</i>
<i>tadalafil</i>	77	TIGLUTIK	51	TRIANEX
<i>tadalafil (pulm. hypertension)</i>	75	<i>timolol maleate</i>	43, 72	<i>triazolam</i>
TAFINLAR	18	<i>tiopronin</i>	51	TRIDERM
TAGRISSE	18	TIROSINT	57	<i>trientine</i>
TAKHZYRO	75	TIROSINT-SOL	57	TRI-ESTARYLLA
TALTZ AUTOINJECTOR	49	TIVICAY	12	<i>trifluoperazine</i>
TALTZ SYRINGE	49	TIVICAY PD	12	<i>trifluridine</i>
TALZENNA	18	<i>tizanidine</i>	35	<i>trihexyphenidyl</i>
<i>tamoxifen</i>	18	TOBI PODHALER	12	TRIJARDY XR
<i>tamsulosin</i>	77	TOBRADEX	72	TRIKAFTA
TASIGNA	18	<i>tobramycin</i>	12, 72	<i>trimethoprim</i>
TAVALISSE	43	<i>tobramycin in 0.225 % nacl</i>	12	TRI-MILI
TAVNEOS	51	<i>tobramycin sulfate</i>	12	<i>trimipramine</i>
<i>tazarotene</i>	49	<i>tobramycin-dexamethasone</i>	72	TRINTELLIX
TAZORAC	49	TOBREX	72	TRI-NYMYO
TAZTIA XT	43	<i>tolcapone</i>	35	TRI-SPRINTEC (28)
TAZVERIK	18	TOLSURA	12	TRITOCIN
TDVAX	64	<i>tolterodine</i>	77	TRIUMEQ
TEFLARO	12	<i>tolvaptan</i>	57	TRIUMEQ PD
TEGRETOL	35	<i>topiramate</i>	35	TRIVORA (28)
TEGRETOL XR	35	<i>toremifene</i>	18	TRI-VYLIBRA
TEGSEDI	35	<i>torseamide</i>	43	TRIZIVIR
<i>telmisartan</i>	43	TOUJEO MAX U-300		TROKENDI XR
<i>telmisartan-amlodipine</i>	43	SOLOSTAR	57	TROPHAMINE 10 %
<i>telmisartan-hydrochlorothiazid</i>	43			<i>trospium</i>

TRULICITY	57	<i>vigabatrin</i>	36	YF-VAX (PF)	64
TRUMENBA	64	VIGADRONE	36	YONSA	19
TRUSELTIQ	18, 19	VIIBRYD	36	YUPELRI	76
TUKYSA	19	VIJOICE	19	YUVAFEM	70
TURALIO	19	<i>vilazodone</i>	36	<i>zafirlukast</i>	76
TWINRIX (PF)	64	VIRACEPT	13	<i>zaleplon</i>	36
TYBOST	13	VIREAD	13	ZARXIO	64
TYMLOS	66	VITRAKVI	19	ZEBUTAL	36
TYPHIM VI	64	VIVITROL	36	ZEGALOGUE	
TYVASO DPI	76	VIVJOA	13	AUTOINJECTOR	57
UBRELVY	35	VIZIMPRO	19	ZEGALOGUE SYRINGE	57
UDENYCA	64	VONJO	19	ZEJULA	19
UNITHROID	57	<i>voriconazole</i>	13	ZELAPAR	36
UPTRAVI	43, 44	VOSEVI	13	ZELBORAF	19
<i>ursodiol</i>	60	VOTRIENT	19	ZEMAIRA	52
VABOMERE	13	VRAYLAR	36	ZEMDRI	13
<i>valacyclovir</i>	13	VTOL LQ	36	ZENATANE	49
VALCHLOR	49	VUMERITY	36	ZENPEP	60
<i>valganciclovir</i>	13	VYFEMLA (28)	70	ZENZEDI	36, 37
<i>valproic acid</i>	36	VYLIBRA	70	ZEPATIER	13
<i>valproic acid (as sodium salt)</i>	36	VYNDAMAX	44	ZEPOSIA	37
<i>valsartan</i>	44	VYNDAQEL	44	ZEPOSIA STARTER KIT	37
<i>valsartan-hydrochlorothiazide</i> ...	44	VYVANSE	36	ZEPOSIA STARTER PACK ...	37
VALTOCO	36	WAKIX	36	<i>zidovudine</i>	13
<i>vancomycin</i>	13	<i>warfarin</i>	44	ZIEXTENZO	64
VANDAZOLE	70	WELIREG	19	<i>zileuton</i>	76
VAQTA (PF)	64	WIXELA INHUB	76	ZIOPTAN (PF)	72
<i>varenicline</i>	51, 52	XALKORI	19	<i>ziprasidone hcl</i>	37
VARIVAX (PF)	64	XARELTO	44	<i>ziprasidone mesylate</i>	37
VARUBI	60	XARELTO DVT-PE TREAT		ZIRGAN	72
VASCEPA	44	30D START	44	ZOLINZA	20
VECAMYL	44	XATMEP	19	<i>zolmitriptan</i>	37
VELIVET TRIPHASIC		XCOPRI	36	<i>zolpidem</i>	37
REGIMEN (28)	70	XCOPRI MAINTENANCE		ZOMACTON	64
VELPHORO	52	PACK	36	ZOMIG	37
VELTASSA	52	XCOPRI TITRATION PACK	36	<i>zonisamide</i>	37
VEMLIDY	13	XELJANZ	66	ZONTIVITY	44
VENCLEXTA	19	XELJANZ XR	66	ZORBTIVE	64
VENCLEXTA STARTING		XENLETA	13	ZORTRESS	20
PACK	19	XERMELO	19	ZOSYN IN DEXTROSE	
<i>venlafaxine</i>	36	XGEVA	19	(ISO-OSM)	13
VENTAVIS	76	XIFAXAN	13	ZOVIA 1-35 (28)	70
VENTOLIN HFA	76	XIIDRA	72	ZUBSOLV	37
<i>verapamil</i>	44	XOFLUZA	13	ZYCLARA	50
VEREGEN	49	XOLAIR	76	ZYDELIG	20
VERQUVO	44	XOSPATA	19	ZYFLO	76
VERSACLOZ	36	XPOVIO	19	ZYKADIA	20
VERZENIO	19	XTANDI	19	ZYLET	72
VESTURA (28)	70	XULTOPHY 100/3.6	57	ZYPREXA RELPREVV	37
VIBERZI	60	XURIDEN	52	ZYVOX	13
VICTOZA 3-PAK	57	XYREM	36		
VIENVA	70	XYWAV	36		

acitretin

Products Affected

- *acitretin*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

actemra

Products Affected

- **ACTEMRA ACTPEN**
- **ACTEMRA SUBCUTANEOUS**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For rheumatoid arthritis, patients must have an inadequate response or intolerance to at least one DMARD (e.g., methotrexate, leflunomide). For giant cell arteritis, patients must have an adequate trial or intolerance to one systemic corticosteroid (e.g., prednisone). For polyarticular juvenile idiopathic arthritis, patients must have an inadequate response or intolerance to at least one DMARD (e.g., methotrexate, leflunomide) -OR- requires initial biologic therapy due to involvement of high-risk joints, high disease activity or at high risk of disabling joint damage. For systemic sclerosis-associated interstitial lung disease (SSc-ILD), patients must have an adequate trial or intolerance to one immunosuppressant (e.g., mycophenolate mofetil, corticosteroids, azathioprine, cyclophosphamide). Documentation of systemic juvenile idiopathic arthritis.
Age Restrictions	Deny if less than 18 years of age for systemic sclerosis-associated interstitial lung disease (SSc-ILD), Rheumatoid Arthritis, and Giant Cell Arteritis or less than 2 years of age for Polyarticular Juvenile Idiopathic Arthritis and Systemic Juvenile Idiopathic Arthritis
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

acthar h.p.

Products Affected

- ACTHAR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	<p>Covered for the following indications: 1. Infantile spasms (West syndrome) in children less than 2 years of age. 2. Acute exacerbations of multiple sclerosis (MS) for patients receiving concurrent immunomodulator therapy (e.g., interferon beta, glatiramer acetate, dimethyl fumarate, fingolimod, teriflunomide) 3. Rheumatic disorders for patients receiving maintenance therapy with at least one NSAID, DMARD (e.g. leflunomide) or biologic (e.g. adalimumab) 4. Collagen diseases for members receiving maintenance therapy with at least one antimalarial (e.g. hydroxychloroquine) or immunosuppressant (e.g. azathioprine) 5. Dermatologic diseases 6. Allergic states (i.e. serum sickness and transfusion reaction due to serum protein reaction 7. Ophthalmic diseases 8. Respiratory diseases 9. Gout and unable to take first-line therapies. 10. Pediatric acquired epileptic aphasia. 11. Proteinuria in nephrotic syndrome and trial/failure or contraindication to two therapies from any of the following different classes: corticosteroids (e.g., cortisone or dexamethasone), calcineurin inhibitors (e.g., cyclosporine or tacrolimus, per DRUGDEX). 12. Diagnosis for adrenal insufficiency with trial/failure or contraindication to cosyntropin. For covered indications 2 through 10, limited/unsatisfactory response or intolerance (i.e. severe anaphylaxis) to two corticosteroids (i.e. IV methylprednisolone, IV dexamethasone, or high dose oral steroids) must be documented.</p>
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 month

PA Criteria	Criteria Details
Other Criteria	For reauthorization, the following (1. 2. and 3.) must be met. 1) Prescriber attestation that the member cannot use corticosteroids (e.g. IV methylprednisolone, high dose oral corticosteroids) due to unsatisfactory response, intolerance (e.g. severe anaphylaxis) or experienced a severe adverse event to corticosteroids (e.g. psychosis). 2) If the reauthorization is for the treatment of multiple sclerosis, a rheumatic disorder, dermatologic disease, or nephrotic syndrome, the prescriber attests that H.P. Acthar is being used for a new acute exacerbation and not on a routine basis to prevent an exacerbation as supported by Compendia. 3) If the reauthorization is for treatment of multiple sclerosis, a rheumatic disorder, or collagen disease, the member continues to receive maintenance therapy.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off Label Uses	Gout

actimmune

Products Affected

- ACTIMMUNE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Adbry

Products Affected

- ADBRY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For moderate to severe atopic dermatitis, documentation of all of the following (1-2): 1) trial & failure, intolerance, or contraindication to at least one topical corticosteroid -OR- atopic dermatitis of the face or anogenital involvement -OR- the member has severe atopic dermatitis evidenced by the extent of the member's body surface area affected which makes topical therapy impractical to apply - OR- severe atopic dermatitis evidenced by severely damaged skin 2) trial & failure, intolerance, or contraindication to topical tacrolimus or topical pimecrolimus -OR- the member has severe atopic dermatitis evidenced by the extent of the member's body surface area affected which makes topical therapy impractical to apply -OR- severe atopic dermatitis evidenced by severely damaged skin
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization
Other Criteria	For Atopic Dermatitis, patients must have an adequate trial/failure, intolerance, or contraindication to both preferred products: Dupixent and Rinvoq. For induction therapy, doses above plan quantity limit will be approved when aligned with recommended induction therapy dosing regimen. For reauthorization, attestation of positive clinical response to therapy.
Indications	All FDA-approved Indications.
Off Label Uses	

ADHD Drugs

Products Affected

- *clonidine hcl oral tablet extended release 12 hr*
- *guanfacine oral tablet extended release 24 hr*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of ADHD -AND- trial/failure, intolerance or contraindication to a stimulant
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

afinitor

Products Affected

- *everolimus (antineoplastic) oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg*
- *everolimus (antineoplastic) oral tablet for suspension 2 mg, 3 mg, 5 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) HR mutation status and HER2 mutation status 2) Alternatives tried/failed 3) Concomitant therapy 4) Candidacy for surgical resection
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

aimovig

Products Affected

- **AIMOVIG AUTOINJECTOR
SUBCUTANEOUS AUTO-INJECTOR
140 MG/ML, 70 MG/ML**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For Episodic Migraine, defined as 4-14 migraine days per month OR Chronic Migraine, defined as 15 or more headaches per month, the following criteria will apply (1-3). 1) Documentation of average monthly migraine days. 2) Attestation that headaches are not caused by medication rebound or overutilization (e.g. not taking triptans exceeding more than 18 doses per month) or lifestyle factors (e.g. sleep patterns, caffeine use). 3) Trial and failure or intolerance to one agent from 2 unique prophylactic migraine medication classes: e.g. Anti-epileptic drugs (e.g. topiramate), beta-blockers (e.g. propranolol), calcium-channel blockers (e.g. verapamil), tricyclic antidepressants (e.g. amitriptyline) -OR- contraindication to all prophylactic medication classes.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, attestation of reduction in average monthly migraine days or number of migraine episodes is required.
Indications	All FDA-approved Indications.
Off Label Uses	

ajovy

Products Affected

- AJOVY AUTOINJECTOR
- AJOVY SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For Episodic Migraine, defined as 4-14 migraine days per month OR Chronic Migraine, defined as 15 or more headaches per month, the following criteria will apply (1-3). 1) Documentation of average monthly migraine days. 2) Attestation that headaches are not caused by medication rebound or overutilization (e.g. not taking triptans exceeding more than 18 doses per month) or lifestyle factors (e.g. sleep patterns, caffeine use). 3) Trial and failure or intolerance to one agent from 2 unique prophylactic migraine medication classes: e.g. Anti-epileptic drugs (e.g. topiramate), beta-blockers (e.g. propranolol), calcium-channel blockers (e.g. verapamil), tricyclic antidepressants (e.g. amitriptyline) -OR- contraindication to all prophylactic medication classes.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, attestation of reduction in average monthly migraine days or number of migraine episodes is required.
Indications	All FDA-approved Indications.
Off Label Uses	

alecensa

Products Affected

- ALECENSA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of metastatic non-small cell lung cancer (NSCLC) that is anaplastic lymphoma kinase (ALK) positive
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

alkindi

Products Affected

- **ALKINDI SPRINKLE**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of adrenocortical insufficiency -AND- Therapeutic failure or intolerance to oral generic hydrocortisone tablets.
Age Restrictions	Deny if greater than 17 years old
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

ALPHA1-PROTEINASE INHIBITORS

Products Affected

- ARALAST NP INTRAVENOUS RECON SOLN 1,000 MG
- GLASSIA
- PROLASTIN-C INTRAVENOUS
- RECON SOLN
- ZEMAIRA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of panacinar emphysema AND documentation of a decline in forced expiratory volume in 1 second (fev1) despite optimal medical therapy (bronchodilators, corticosteroids, oxygen if indicated) AND documentation of phenotype (pi*zz, pi*znull or pi*nullnull) associated with causing serum alpha 1-antitrypsin of less than 80 mg/dl AND documentation of an alpha 1-antitrypsin serum level below the value of 35% of normal (less than 80 mg/dl).
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Covered under Part B when furnished incident to a physician service and is not self-administered.
Indications	All FDA-approved Indications.
Off Label Uses	

alunbrig

Products Affected

- ALUNBRIG ORAL TABLET 180 MG, 30 MG, 90 MG
- ALUNBRIG ORAL TABLETS,DOSE PACK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of non-small cell lung cancer (NSCLC) that is anaplastic lymphoma kinase (ALK) positive
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

ampyra

Products Affected

- *dalfampridine*

PA Criteria	Criteria Details
Exclusion Criteria	History of seizure disorder, Cr Cl less than 50ml/min
Required Medical Information	Documentation of diagnosis -AND- documentation that the patient is ambulatory and has walking impairment as evidenced by one of the following. 1. Functional status score (EDSS score). 2. Timed 25-foot Walk Test (T25W).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	3 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, documentation supporting improvement in walking impairment from baseline is required.
Indications	All FDA-approved Indications.
Off Label Uses	

anabolic steroids

Products Affected

- *oxandrolone*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- For the diagnosis of osteoporosis the trial/failure, intolerance or contraindication to at least 2 federal legend drugs indicated for use in osteoporosis.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

apokyn

Products Affected

- *apomorphine*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of Parkinson's disease -AND- for use in acute, intermittent treatment of hypomobility off episodes -AND- experiencing off episodes despite the use of oral carbidopa/levodopa -AND- Therapeutic failure, intolerance, or contraindication to the following (1. and 2.): 1) a generic pramipexole containing product or a generic ropinirole containing product. 2) generic entacapone, selegiline or rasagiline.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

arikayce

Products Affected

- ARIKAYCE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of Mycobacterium avium complex lung disease -AND- attestation of not achieving negative sputum cultures despite at least 6 months with a multidrug background regimen containing a macrolide - AND- Arikayce will be used in conjunction with a background multidrug regimen.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	18 months
Other Criteria	For reauthorization, attestation of positive sputum cultures -OR- negative sputum cultures for insufficient period of time (e.g. less than 12 months).
Indications	All FDA-approved Indications.
Off Label Uses	

attr-cm drugs

Products Affected

- VYNDAMAX
- VYNDAQEL

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of cardiomyopathy of wild type or hereditary transthyretin-mediated amyloidosis (ATTR-CM) with amyloid deposits on cardiac biopsy or scintigraphy with heart contralateral greater than 1.5 (Grade III) -AND- Cardiac involvement supported by cardiac magnetic resonance, echocardiography or serum cardiac biomarker (e.g. B-type natriuretic peptide, cardiac troponin) - AND- Primary (light chain) amyloidosis has been ruled out by immunohistochemistry, mass spectrometry or scintigraphy.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of improvement or delayed disease progression from baseline demonstrated by 6-minute walk test, cardiac function (e.g. LVEF, NYHA class), Kansas City Cardiomyopathy Questionnaire-Overall Summary, number of cardiovascular-related hospitalizations or serum cardiac biomarkers (e.g. B-type natriuretic peptide, cardiac troponin)
Indications	All FDA-approved Indications.
Off Label Uses	

atypical antipsychotics

Products Affected

- *aripiprazole*
- **REXULTI**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. If medication is being used for major depressive disorder, documentation of adjunctive therapy and an adequate trial of 1 alternative antidepressant is required (e.g. SSRI, SNRI, NDRIs, TCA, MAOI). For Rexulti, trial, intolerance, or contraindication to one other formulary generic atypical antipsychotic (e.g. quetiapine).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

aubagio

Products Affected

- AUBAGIO

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use of Aubagio and other disease modifying agents such as fingolimod, interferons, Copaxone, Tysabri
Required Medical Information	Documentation of relapsing form of multiple sclerosis (e.g. relapsing-remitting, clinically isolated syndrome, or active secondary progressive disease)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	5 years
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

auryxia

Products Affected

- AURYXIA

PA Criteria	Criteria Details
Exclusion Criteria	Treatment of iron deficiency anemia
Required Medical Information	Documentation of diagnosis
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

ayvakit

Products Affected

- **AYVAKIT**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For unresectable or metastatic Gastrointestinal Stromal Tumor (GIST), PDGFRA exon 18 mutation status. For Advanced Systemic Mastocytosis (AdvSM), platelet count greater than or equal to $50 \times 10^9/L$ AND aggressive systemic mastocytosis, systemic mastocytosis with an associated hematological neoplasm (SM-AHN), or Mast cell leukemia (MCL).
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

bafiertam

Products Affected

- **BAFIERTAM**

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use with other disease modifying agents such as interferons, Copaxone, Tysabri, Aubagio, Gilenya
Required Medical Information	Documentation of relapsing form of multiple sclerosis (e.g. relapsing-remitting, clinically isolated syndrome, or active secondary progressive disease) -AND- Therapeutic failure or intolerance to generic dimethyl fumarate
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	5 years
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

balversa

Products Affected

- **BALVERSA**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) FGFR3 or FGFR2 mutation status as detected by an FDA approved test 2) alternatives tried/failed within labelled time frame
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

banzel

Products Affected

- **BANZEL ORAL SUSPENSION**
- *rufinamide*

PA Criteria	Criteria Details
Exclusion Criteria	Patients with familial short QT syndrome
Required Medical Information	Documentation of seizures due to Lennox-Gastaut Syndrome -AND- documentation of adjunctive therapy -AND- an adequate trial or intolerance of a previous antiepileptic therapy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

benlysta

Products Affected

- BENLYSTA SUBCUTANEOUS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	<p>Documentation of active systemic lupus erythematosus (SLE) -AND- documentation of positive anti-nuclear antibody (ANA) titer (greater than or equal to 1:80) or anti-double-stranded DNA antibody (anti-dsDNA) greater than or equal to 30IU/mL -AND- trial, intolerance, or inadequate response to at least 2 of the following standard of care drug classes: 1.) corticosteroids (e.g. prednisone) 2.) antimalarials (e.g. hydroxychloroquine) 3.) immunosuppressants (e.g. azathioprine, mycophenolate mofetil, or methotrexate) -AND- member will continue to receive concomitant standard of care treatment with use of at least one of the following (alone or in combination): 1.) corticosteroids (e.g. prednisone) 2.) antimalarials (e.g. hydroxychloroquine) 3.) immunosuppressants (e.g. azathioprine, mycophenolate mofetil, or methotrexate) -OR- Documentation of active lupus nephritis -AND- documentation of positive ANA titer (greater than or equal to 1:80) or anti-dsDNA greater than or equal to 30 IU/mL -AND- trial, intolerance, or inadequate response to at least 2 of the following standard of care drug classes: 1.) corticosteroids (e.g. prednisone) 2.) antimalarials (e.g. hydroxychloroquine) 3.) immunosuppressants (e.g. azathioprine, mycophenolate mofetil, or methotrexate) -AND- member will continue to receive concomitant standard of care treatment which includes corticosteroids (e.g. prednisone) with at least one of the following: 1.) mycophenolate for induction followed by mycophenolate for maintenance 2.) cyclophosphamide for induction followed by azathioprine for maintenance</p>
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months

PA Criteria	Criteria Details
Other Criteria	For reauthorization, attestation of positive clinical response is required. For induction therapy dosing, doses above plan quantity limit will be approved aligned with recommended induction therapy dosing regimens per indication.
Indications	All FDA-approved Indications.
Off Label Uses	

berinert

Products Affected

- **BERINERT INTRAVENOUS KIT**

PA Criteria	Criteria Details
Exclusion Criteria	Member should not be on two acute therapies simultaneously and acute therapy should not be used as prophylactic therapy
Required Medical Information	For the treatment of acute abdominal, facial, or laryngeal attacks of hereditary angioedema (HAE) type I & II with the following (1-4): 1) Low C4 level of less than or equal to 14mg/dL or C4 below lower limit of laboratory reference range and 1 of the following (A or B). A) C1 inhibitor (C1INH) antigen level less than or equal to 19mg/dL or below lower limit of laboratory reference range. B) Normal C1INH antigen level and a low C1INH functional level below laboratory reference range. 2) Past medical history of at least 1 symptom of moderate or severe angioedema attack (e.g. airway swelling, painful facial distortion) in absence of concomitant hives. 3) Medications known to cause angioedema have been evaluated and discontinued. 4) Documentation of member's weight. For the treatment of acute abdominal, facial, or laryngeal attacks of hereditary angioedema (HAE) type III with the following (5-9): 5) Documentation of clinical laboratory performance C4, C1INH antigen, or C1INH functional level are within normal limits of laboratory reference ranges. 6) Documentation of family history of HAE or FXII mutation 7) Past medical history of at least 1 symptom of moderate or severe angioedema attack (e.g. airway swelling, painful facial distortion) in absence of concomitant hives. 8) Medications known to cause angioedema have been evaluated and discontinued. 9) Documentation of member's weight.
Age Restrictions	Deny if less than 5 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For 18 years of age or older, therapeutic failure, intolerance or contraindication to icatibant.

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off Label Uses	

Besremi

Products Affected

- **BESREMI**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of polycythemia vera
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

bosulif

Products Affected

- **BOSULIF**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis AND all of the following, if applicable to diagnosis: 1) phase of CML 2) Philadelphia chromosome status 3) alternatives tried/failed
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

braftovi

Products Affected

- BRAFTOVI ORAL CAPSULE 75 MG

PA Criteria	Criteria Details
Exclusion Criteria	Use in wild-type BRAF melanoma or wild-type BRAF CRC
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) BRAF V600E or V600K mutation status 2) alternatives tried/failed 3) concomitant therapy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

bronchitol

Products Affected

- BRONCHITOL

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of cystic fibrosis -AND- Passed a Bronchitol Tolerance Test -AND- Used in conjunction with standard therapies for the management of cystic fibrosis to improve pulmonary function (e.g. bronchodilators, antibiotics, anti-inflammatory therapy).
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of increase in FEV1
Indications	All FDA-approved Indications.
Off Label Uses	

brukinsa

Products Affected

- **BRUKINSA**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- if applicable to diagnosis, alternatives tried/failed
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

butrans

Products Affected

- *buprenorphine*
- **BUTRANS**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	<p>Documentation of moderate to severe chronic pain -AND- trial and failure of at least two previous federal legend medications for pain, including NSAIDs, tramadol, or opioid analgesics. For concomitant use of an opiate agonist and substance abuse therapy, documentation that the member has an acute pain condition (e.g. acute traumatic injury) in which treatment with other agents would cause insufficient pain control or if the member requires treatment for pain related to a terminal illness. For concomitant use of an opiate agonist, benzodiazepine and a centrally acting skeletal muscle relaxant, documentation that the member has tried/failed at least 2 other skeletal muscle relaxant (e.g. methocarbamol, metaxalone), understanding these skeletal muscle relaxants are high-risk medications in geriatric patients AND attestation of an intent to monitor and address concomitant drug-drug interaction adverse events. For concomitant use of an opiate agonist and other opiate potentiators (e.g. gabapentinoids, benzodiazepines) attestation of an intent to monitor and address concomitant drug-drug interaction adverse events. For long acting (e.g. extended release) opioid medications, the following apply (1-5). 1)Pain is severe enough to require daily, around-the-clock, long-term opioid treatment. 2)Patient is not long acting opioid naive. 3)Attestation that non-opiate alternative therapies have been explored (e.g. NSAIDs). 4)Attestation that controlled substance Rx history has been reviewed in the state Prescription Drug Monitoring Program. 5)Attestation of counseling on the potential adverse effects of opioid analgesics, including the risk of misuse, abuse, and addiction.</p>
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months

PA Criteria	Criteria Details
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

cablivi

Products Affected

- **CABLIVI INJECTION KIT**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of acquired thrombocytopenic purpura (aTTP) -AND- Used in conjunction with plasma exchange and immunosuppressive therapy (i.e. systemic corticosteroids or rituximab)
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	75 days initial authorization, 28 days reauthorization
Other Criteria	For reauthorization, attestation of remaining signs and symptoms of persistent disease (e.g. suppressed ADAMTS 13 activity level remain present)
Indications	All FDA-approved Indications.
Off Label Uses	

cabometyx

Products Affected

- **CABOMETYX**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) alternatives tried/failed 2) attestation of first line use 3) concomitant therapy 4) radioactive iodine refractory status
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

calquence

Products Affected

- CALQUENCE
- CALQUENCE (ACALABRUTINIB MAL)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- if applicable to diagnosis, alternatives tried/failed
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Camzyos

Products Affected

- CAMZYOS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of symptomatic New York Heart Association (NYHA) class II-III obstructive hypertrophic cardiomyopathy -AND- Left ventricular ejection fraction of greater than or equal to 55% -AND- Valsalva left ventricular outflow tract peak gradient of greater than or equal to 50 mmHg at rest or after provocation -AND- Therapeutic failure or intolerance to one of the following (1 or 2) or contraindication to all: 1) Non-vasodilating beta blocker (e.g. metoprolol) 2) non-dihydropyridine calcium channel blocker (e.g. diltiazem) -AND- Not currently treated with and attestation Camzyos will not be used concomitantly with disopyramide, ranolazine, or combination therapy of beta-blocker and calcium channel blocker
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of improvement in pV02 by greater than or equal to 1.5 mL/kg/min plus at least 1 NYHA class reduction or improvement in pV02 by greater than or equal to 3.0 mL/kg/min without NYHA class worsening -AND- Not currently treated with and attestation Camzyos will not be used concomitantly with disopyramide, ranolazine, or combination therapy of beta-blocker and calcium channel blocker
Indications	All FDA-approved Indications.
Off Label Uses	

caplyta

Products Affected

- CAPLYTA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- trial, intolerance, or contraindication to one other formulary generic atypical antipsychotic (e.g. quetiapine).
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

caprelsa

Products Affected

- CAPRELSA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of symptomatic or progressive medullary thyroid cancer in patients with unresectable locally advanced or metastatic disease
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

carac

Products Affected

- CARAC

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Trial and failure of 1 generic fluorouracil topical product (with shared indication) is required.
Indications	All FDA-approved Indications.
Off Label Uses	

carbaglu

Products Affected

- *carglumic acid*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of use as an adjunct therapy for acute hyperammonemia due to hepatic enzyme N-acetylglutamate synthase (NAGS) deficiency, propionic acidemia (PA), or methylmalonic acidemia (MMA) -OR- maintenance therapy for chronic hyperammonemia due to hepatic enzyme N-acetylglutamate synthase (NAGS) deficiency
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All Medically-accepted Indications.
Off Label Uses	

CERDELGA

Products Affected

- CERDELGA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of type 1 Gaucher disease confirmed by the following A. or B. A) With one of the following symptoms (1, 2, 3, 4, or, 5): 1)Hepatomegaly. 2)Splenomegaly. 3)Bone disease (i.e. osteonecrosis, osteopenia, secondary pathologic fractures, bone infarct). 4)Bone marrow complications as defined by anemia with hemoglobin less than or equal to 11.5 g/dL for females or 12.5 g/dL for males or thrombocytopenia with platelet count less than or equal to 120,000/mm ³ . 5)Symptomatic disease (e.g. bone pain, exertional limitation, cachexia). -OR- B) Attestation of deficiency in glucocerebrosidase activity in peripheral leukocytes or genetic testing confirms mutant alleles -AND- Documentation of CYP2D6 metabolizer status (e.g. intermediate metabolizer).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

CF drugs

Products Affected

- **BETHKIS**
 - **PULMOZYME**
 - **TOBI PODHALER INHALATION CAPSULE, W/INHALATION DEVICE**
- *tobramycin in 0.225 % nacl*
 - *tobramycin inhalation*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of cystic fibrosis. For Bethkis: failure on, intolerance to, or contraindication to generic tobramycin inhalation solution
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Inhalation solutions covered under Part B when administered in the home setting using a covered nebulizer (i.e. DME). For reauthorization of tobramycin products, attestation of decrease in sputum density of pseudomonas aeruginosa, increase in FEV1 or decrease in number of hospitalizations or pulmonary exacerbations. For reauthorization of Pulmozyme, attestation of increase in FEV1 or decrease in number of hospitalizations or pulmonary exacerbations.
Indications	All FDA-approved Indications.
Off Label Uses	

chenodal

Products Affected

- CHENODAL

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of radiolucent gallstones AND an inadequate response or intolerance to ursodiol tablets
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months for initial approval with an additional 12 months upon renewal
Other Criteria	Safety of use beyond 24 months is not established. For reauthorization, attestation of partial dissolution of gallstones
Indications	All FDA-approved Indications.
Off Label Uses	

chloroquine

Products Affected

- *chloroquine phosphate oral tablet 250 mg, 500 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. If using for diagnosis of malaria prophylaxis, documentation of duration of travel is required.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Malaria tx and amebiasis: 1 month. Malaria prophylaxis: Travel duration plus 10 wks
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

cholbam

Products Affected

- CHOLBAM

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of bile acid synthesis disorders due to single enzyme defects (e.g. AKR1D1, CTX, CYP7A1) -OR- documentation of use as adjunctive therapy for peroxisomal disorders (PDs), including Zellweger spectrum disorders, in patients who exhibit manifestations of liver disease, steatorrhea, or complications from decreased fat soluble vitamin absorption.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

cialis

Products Affected

- **CIALIS ORAL TABLET 2.5 MG, 5 MG**
- *tadalafil oral tablet 2.5 mg, 5 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of benign prostatic hyperplasia (BPH) and trial/failure of at least two alternative medications in the following classes (alpha-1 adrenergic blockers and/or 5-alpha reductase inhibitors)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For brand Cialis, trial and failure of generic tadalafil is required.
Indications	All FDA-approved Indications.
Off Label Uses	

Cibinqo

Products Affected

- CIBINQO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For moderate to severe refractory atopic dermatitis whose disease is not adequately controlled with other systemic drug products, documentation of all of the following (1-2): 1) trial & failure, intolerance, or contraindication to at least one topical corticosteroid -OR- atopic dermatitis of the face or anogenital involvement -OR- the member has severe atopic dermatitis evidenced by the extent of the member's body surface area affected which makes topical therapy impractical to apply -OR- severe atopic dermatitis evidenced by severely damaged skin, 2) trial & failure, intolerance, or contraindication to topical tacrolimus or topical pimecrolimus -OR- the member has severe atopic dermatitis evidenced by the extent of the member's body surface area affected which makes topical therapy impractical to apply -OR- severe atopic dermatitis evidenced by severely damaged skin.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For Atopic Dermatitis, patients must have an adequate trial/failure, intolerance, or contraindication to both preferred products: Dupixent and Rinvoq. For reauthorization, attestation of positive clinical response to therapy.
Indications	All FDA-approved Indications.
Off Label Uses	

cimzia

Products Affected

- **CIMZIA**
- **CIMZIA POWDER FOR RECONST**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For moderate to severe rheumatoid arthritis, inadequate response or intolerance to at least one DMARD (e.g. leflunomide). For moderate to severe Crohn's disease, inadequate response or intolerance to at least two immunosuppressants (e.g. corticosteroids, azathioprine). For ankylosing spondylitis, inadequate response or intolerance to one nonsteroidal anti-inflammatory drug (NSAID). For non-radiographic axial spondyloarthritis, inadequate response or intolerance to two nonsteroidal anti-inflammatory drugs (NSAIDs). For moderate to severe psoriasis, inadequate response or intolerance to one systemic therapy (e.g. methotrexate, cyclosporine) -OR- inadequate response to phototherapy. If not a candidate for phototherapy: treatment with systemic therapy has been ineffective, not tolerated, or is contraindicated.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months

PA Criteria	Criteria Details
Other Criteria	<p>For Crohn's disease, patients must have an adequate trial or intolerance to both preferred biologic products, Humira and Stelara. For Rheumatoid arthritis, patients must have an adequate trial or intolerance to 2 of the following preferred products Humira, Enbrel, Actemra, Xeljanz/Xeljanz XR and Rinvoq. For plaque psoriasis, patients must have an adequate trial or intolerance to 2 of the following preferred products Humira, Cosentyx, Otezla, Stelara, Skyrizi and Enbrel. For ankylosing spondylitis, patients must have an adequate trial or intolerance to 2 of the following preferred products Humira, Enbrel and Cosentyx. For Psoriatic arthritis, patients must have an adequate trial or intolerance to 2 of the following preferred products Cosentyx, Enbrel, Humira, Xeljanz/Xeljanz XR, Otezla, and Stelara. For initial and induction therapy dosing, doses above plan quantity limit will be approved aligned with recommended initial and induction therapy dosing regimens per indication.</p>
Indications	All FDA-approved Indications.
Off Label Uses	

cinryze

Products Affected

- CINRYZE

PA Criteria	Criteria Details
Exclusion Criteria	Member should not be on two prophylactic therapies simultaneously.
Required Medical Information	For the prophylactic treatment of abdominal, facial, or laryngeal attacks of hereditary angioedema (HAE) type I & II with the following (1-3): 1) Low C4 level of less than or equal to 14mg/dL or C4 below lower limit of laboratory reference range and 1 of the following (A or B). A) C1 inhibitor (C1INH) antigen level less than or equal to 19mg/dL or below lower limit of laboratory reference range. B) Normal C1INH antigen level and a low C1INH functional level below laboratory reference range. 2) Past medical history of at least 1 symptom of moderate or severe angioedema attack (e.g. airway swelling, painful facial distortion) in absence of concomitant hives. 3) Medications known to cause angioedema have been evaluated and discontinued. For the prophylactic treatment of acute abdominal, facial, or laryngeal attacks of hereditary angioedema (HAE) type III with the following (4-7): 4) Documentation of clinical laboratory performance C4, C1INH antigen, or C1INH functional level are within normal limits of laboratory reference ranges. 5) Documentation of family history of HAE or FXII mutation 6) Past medical history of at least 1 symptom of moderate or severe angioedema attack (e.g. airway swelling, painful facial distortion) in absence of concomitant hives. 7) Medications known to cause angioedema have been evaluated and discontinued.
Age Restrictions	Deny if less than 6 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	

cometriq

Products Affected

- COMETRIQ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of progressive, metastatic medullary thyroid cancer
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

copiktra

Products Affected

- COPIKTRA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- if applicable to diagnosis, alternatives tried/failed
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

corlanor

Products Affected

- CORLANOR ORAL SOLUTION
- CORLANOR ORAL TABLET 5 MG, 7.5 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis AND all of the following: 1) Normal sinus rhythm. 2) Resting heart rate greater than or equal to 70 beats per minute. 3) Left ventricular ejection fraction less than or equal to 35 percent, when applicable. 4) In adult patients (greater than or equal to 18 years), trial/failure of maximum tolerated dose of one beta-blocker used for treatment of heart failure (e.g., bisoprolol, carvedilol, metoprolol succinate) OR contraindication to beta-blocker use.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For oral solution, attestation of inability to swallow tablets is required.
Indications	All FDA-approved Indications.
Off Label Uses	

Cosentyx

Products Affected

- **COSENTYX (2 SYRINGES)**
- **COSENTYX PEN (2 PENS)**
- **COSENTYX SUBCUTANEOUS SYRINGE 75 MG/0.5 ML**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For moderate to severe psoriasis, inadequate response or intolerance to one systemic therapy (e.g. methotrexate) -OR- inadequate response to phototherapy -OR- contraindication to phototherapy and systemic therapy. For ankylosing spondylitis, inadequate response or intolerance to one nonsteroidal anti-inflammatory drug (NSAID). For non-radiographic axial spondyloarthritis, inadequate response or intolerance to 2 NSAIDs. For enthesitis-related arthritis, inadequate response or intolerance to at least 1 non-biologic disease modifying anti-rheumatic drugs (DMARD), or all are contraindicated.
Age Restrictions	Deny if less than 6 years of age for moderate to severe plaque psoriasis -OR- less than 2 years of age for psoriatic arthritis -OR- less than 4 years of age for enthesitis-related arthritis -OR- less than 18 years of age for all other indications
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For induction therapy dosing, doses above plan quantity limit will be approved aligned with recommended induction therapy dosing regimens per indication.
Indications	All FDA-approved Indications.
Off Label Uses	

cotellic

Products Affected

- COTELLIC

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) BRAF V600E or V600K mutation status 2) Concomitant therapy
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

crinone

Products Affected

- CRINONE

PA Criteria	Criteria Details
Exclusion Criteria	Use to promote fertility
Required Medical Information	Documentation of diagnosis
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Cysteamine Ophthalmic Drops

Products Affected

- CYSTADROPS
- CYSTARAN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of cystinosis -AND- Attestation of accumulation of corneal cystine crystals
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

daraprim

Products Affected

- *pyrimethamine*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For primary prophylaxis of toxoplasmosis gondii infection, CD4 count less than 100 cells/mm ³ -AND- Toxoplasma IgG positive -AND- failure, intolerance or contraindication to trimethoprim-sulfamethoxazole. For secondary prophylaxis of toxoplasmosis gondii infection, CD4 count less than 200 cells/mm ³ . For secondary prophylaxis of cystoisosporiasis with CD4 count less than 200 cells/mm ³ or acute cystoisosporiasis infection: failure, intolerance or contraindication to trimethoprim-sulfamethoxazole. For primary prophylaxis of Pneumocystis jirovecii pneumonia: diagnosis of HIV - AND- CD4 count less than 200 cells/mm ³ -AND- failure, intolerance or contraindication to trimethoprim-sulfamethoxazole.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All Medically-accepted Indications.
Off Label Uses	

daurismo

Products Affected

- **DAURISMO ORAL TABLET 100 MG, 25 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of newly diagnosed Acute Myeloid Leukemia -AND- Used in combination with cytarabine -AND- At least one comorbidity that preclude use of intensive induction chemotherapy defined as one of the following: 1) Age greater than or equal to 75 2) Severe cardiac or pulmonary comorbidity 3) Reduced renal function 4) Hepatic impairment 5.) Physician attests patient is not a candidate for intensive induction therapy
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

daytrana

Products Affected

- DAYTRANA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis
Age Restrictions	Deny if less than 6 years of age or greater than 17 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For ADHD, trial/failure or intolerance to 2 of the following generic medications: methylphenidate, atomoxetine, or dextroamphetamine/amphetamine is required.
Indications	All FDA-approved Indications.
Off Label Uses	

deferasirox

Products Affected

- *deferasirox oral tablet, dispersible*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For chronic iron overload due to blood transfusions, transfusion history of greater than or equal to 100 mL/kg of packed red blood cells (i.e. at least 20 units of packed red blood cells for a 40 kg person or more in individuals weighing more than 40 kg) -And- history of serum ferritin consistently greater than 1,000 mcg/L or liver iron concentration (LIC) greater than or equal to 7 iron per gram of liver dry weight (mg Fe/g dw). For Chronic Iron Overload in Non-Transfusion-Dependent Thalassemia (NTDT) Syndrome, LIC of at least 5 mg Fe/g dw -AND- serum ferritin greater than 300 mcg/L.
Age Restrictions	Deny if less than 2 years of age for chronic iron overload or less than 10 years of age chronic iron overload in NTDT
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization of chronic iron overload due to blood transfusion, continued requirement for regular blood transfusions -AND- serum ferritin level greater than or equal to 500mcg/L or LIC greater than or equal to 3 mg Fe/g dw. For reauthorization of chronic iron overload in NTDT syndrome, LIC greater than or equal to 3 mg Fe/g dw.
Indications	All FDA-approved Indications.
Off Label Uses	

diacomit

Products Affected

- **DIACOMIT ORAL CAPSULE 250 MG, 500 MG**
- **DIACOMIT ORAL POWDER IN PACKET 250 MG, 500 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of Dravets syndrome - AND- Used in combination with clobazam
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation supporting reduction in seizure frequency
Indications	All FDA-approved Indications.
Off Label Uses	

dihydroergotamine

Products Affected

- *dihydroergotamine nasal*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of acute migraine headaches with or without aura -AND- requires non-oral route of administration -AND- therapeutic failure or intolerance to generic sumatriptan nasal spray.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

dojolvi

Products Affected

- DOJOLVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of molecularly confirmed long-chain fatty acid oxidation disorders
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of positive clinical response to therapy
Indications	All FDA-approved Indications.
Off Label Uses	

doptelet

Products Affected

- **DOPTELET (10 TAB PACK)**
- **DOPTELET (15 TAB PACK)**
- **DOPTELET (30 TAB PACK)**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of thrombocytopenia and chronic liver disease - AND- beneficiary is scheduled to undergo a procedure -OR- Documentation of chronic immune thrombocytopenia -AND- Trial, intolerance, or inadequate response to corticosteroid therapy, immunoglobulin therapy or splenectomy -AND- One of the following (1 or 2): 1) Platelet count less than or equal to $50 \times 10^9/L$ and has significant mucous member bleeding or at least one risk factor for bleeding (e.g. hypertension, peptic ulcer disease). 2) Platelets count of less than or equal to $30 \times 10^9/L$
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	For thrombocytopenia with chronic liver disease- 1 mo. For chronic immune thrombocytopenia- 12 mo.
Other Criteria	Platelet count is provided for applicable dosing.
Indications	All FDA-approved Indications.
Off Label Uses	

doxepin cream

Products Affected

- *doxepin topical*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- course of therapy will not exceed 8 days -AND- experienced therapeutic failure, intolerance, or contraindication to one of the following (1 or 2): 1) at least 2 generic formulary topical corticosteroids. 2) generic topical tacrolimus or pimecrolimus for topic dermatitis with the facial or anogenital involvement.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	1 month
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

drizalma

Products Affected

- **DRIZALMA SPRINKLE ORAL
CAPSULE, DELAYED REL SPRINKLE
20 MG, 30 MG, 40 MG, 60 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- inability to swallow capsules. For fibromyalgia, members must also have widespread bilateral pain above and below the waist for greater than 3 months duration -AND- At least one (1) fibromyalgia-related symptom (e.g., cognitive impairment, fatigue, sleep disturbance, neurologic symptoms, exercise intolerance).
Age Restrictions	Deny if less than 18 years of age in the treatment of fibromyalgia, major depressive disorder, diabetic peripheral neuropathy and chronic musculoskeletal pain -OR- if less than 7 years of age in generalized anxiety disorder
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

duobrii

Products Affected

- DUOBRII

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of plaque psoriasis -AND- therapeutic failure or intolerance to generic tazarotene cream -AND- therapeutic failure or intolerance to 1 high-potency topical corticosteroid (e.g. betamethasone dipropionate 0.05%, halobetasol propionate 0.05%)
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	3 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

dupixent

Products Affected

- **DUPIXENT PEN SUBCUTANEOUS PEN INJECTOR 200 MG/1.14 ML, 300 MG/2 ML**
- **DUPIXENT SYRINGE SUBCUTANEOUS SYRINGE 100 MG/0.67 ML, 200 MG/1.14 ML, 300 MG/2 ML**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	<p>Documentation of all of the following (1-3): 1) moderate to severe atopic dermatitis 2) trial & failure, intolerance, or contraindication to at least one topical corticosteroid -OR- atopic dermatitis of the face or anogenital involvement -OR- the member has severe atopic dermatitis evidenced by a large proportion of the member's body surface area is affected making topical therapy impractical to apply -OR- severe atopic dermatitis evidenced by severely damaged skin 3) if 2 years or older: trial & failure, intolerance, or contraindication to topical tacrolimus or topical pimecrolimus -OR- the member has severe atopic dermatitis evidenced by a large proportion of the member's body surface area is affected making topical therapy impractical to apply -OR- severe atopic dermatitis evidenced by severely damaged skin -OR- Documentation of the following (4-7): 4) moderate-to-severe asthma 5) documented pretreatment FEV1 less than 80 percent predicted in adults or FEV1 less than 90 percent predicted in adolescents or FEV1 reversibility of at least 12% after albuterol administration 6) Blood eosinophils greater than or equal to 150 cells/uL -OR- patient is currently taking daily or alternate-day oral corticosteroids 7) using a medium- or high-dose inhaled corticosteroid and a long acting beta agonist -OR- Documentation of the following (8-9): 8) chronic rhinosinusitis with nasal polyposis (CRSwNP) 9) trial & failure, intolerance or contraindication to intra-nasal corticosteroid and 14 day course of oral corticosteroids -OR- Documentation of the following (10-13): 10) eosinophilic esophagitis 11) weight at least 40 kg 12) esophageal eosinophil count greater than or equal to 15 eos/hpf on esophageal biopsy 13) clinical symptoms of esophageal dysfunction.</p>
Age Restrictions	
Prescriber Restrictions	

PA Criteria	Criteria Details
Coverage Duration	6 months initial authorization, 12 months reauthorization
Other Criteria	For induction therapy, doses above plan quantity limit will be approved aligned with recommended induction therapy dosing regimen. For reauthorization, attestation of positive clinical response to therapy, for atopic dermatitis -OR- attestation of decreased in rescue medication or oral corticosteroid use, decrease in frequency of severe asthma exacerbation, increase in pulmonary function from baseline (e.g. FEV1) or reduction in reported asthma related symptoms, for asthma -OR- attestation of decrease in nasal polyp score or reduction in nasal congestion/obstruction severity score, for CRSwNP -OR- attestation of histological remission (less than 15 eos/hpf) on esophageal biopsy or reduced severity or frequency of clinical symptoms of esophageal dysfunction, for EoE
Indications	All FDA-approved Indications.
Off Label Uses	

egfr tyrosine kinase inhibitors

Products Affected

- *erlotinib*
- **EXKIVITY**
- **GILOTRIF**

PA Criteria	Criteria Details
Exclusion Criteria	Afatinib products: tumors with resistant EGFR mutations. Erlotinib products: use in NSCLC tumors with mutations other than those in FDA-approved indications. Use in combination with platinum based chemotherapy.
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis 1) Epidermal growth factor receptor (EGFR) mutations, including exon 19 deletions or exon 21 (L858R) substitution mutations 2) Alternatives tried/failed 3) Concomitant therapy 4) Line of therapy in which medication will be used
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

egrifta

Products Affected

- EGRIFTA SV

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documented diagnosis of HIV and lipodystrophy, member must actively be receiving antiretroviral therapy including protease inhibitors, nucleoside reverse transcriptase inhibitors, or non-nucleoside reverse transcriptase inhibitors
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

emflaza

Products Affected

- EMFLAZA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of Duchenne muscular dystrophy (DMD) with mutation of dystrophin gene -AND- onset of weakness or history of DMD starting before age 5 -AND- One of the following (1, 2, or 3). 1) Documented trial/failure, intolerance or contraindication to prednisone. 2) Documented trial of prednisone has resulted in intolerable adverse events (e.g. diabetes, hypertension that is difficult to manage, Cushingoid features, truncal obesity, greater than or equal to 10 percent increase in body weight over a 6 month period). 3) Documented severe behavioral adverse event while on prednisone that warrants prednisone dose reduction impacting efficacy for management of DMD (i.e. abnormal behavior, aggression, irritability, disturbance in mood)
Age Restrictions	Deny if less than 2 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

emgality

Products Affected

- **EMGALITY PEN**
- **EMGALITY SYRINGE
SUBCUTANEOUS SYRINGE 120
MG/ML, 300 MG/3 ML (100 MG/ML X 3)**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For Episodic Migraine, defined as 4-14 migraine days per month OR Chronic Migraine, defined as 15 or more headaches per month, the following criteria will apply (1-3). 1) Documentation of average monthly migraine days. 2) Attestation that headaches are not caused by medication rebound or overutilization (e.g. not taking triptans exceeding more than 18 doses per month) or lifestyle factors (e.g. sleep patterns, caffeine use). 3) Trial and failure or intolerance to one agent from 2 unique prophylactic migraine medication classes: e.g. Anti-epileptic drugs (e.g. topiramate), beta-blockers (e.g. propranolol), calcium-channel blockers (e.g. verapamil), tricyclic antidepressants (e.g. amitriptyline) -OR- contraindication to all prophylactic medication classes. For episodic cluster headaches, characterized by severe or very severe unilateral orbital, supraorbital, and/or temporal pain lasting 15 to 180 minutes when left untreated -AND- Attack frequency of at least one attack every other day during the cluster period.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization
Other Criteria	For induction therapy, doses above plan quantity limit will be approved aligned with recommended induction therapy dosing regimen. For reauthorization, attestation of reduction in average monthly migraine days or number of migraine episodes is required -OR- attestation of reduction in the number of mean weekly cluster headaches from baseline is required.
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	

enbrel

Products Affected

- ENBREL MINI
- ENBREL SUBCUTANEOUS SOLUTION
- ENBREL SUBCUTANEOUS SYRINGE
25 MG/0.5 ML (0.5), 50 MG/ML (1 ML)
- ENBREL SURECLICK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For rheumatoid arthritis, inadequate response or intolerance to at least one DMARD (e.g. methotrexate, leflunomide). For ankylosing spondylitis, inadequate response or intolerance to one nonsteroidal anti-inflammatory drug (NSAID). For juvenile idiopathic rheumatoid arthritis, inadequate response or intolerance to at least one DMARD (e.g., methotrexate, leflunomide) -OR- requires initial biologic therapy due to involvement of high-risk joints, high disease activity or at high risk of disabling joint damage. For moderate to severe plaque psoriasis, inadequate response or intolerance to one systemic therapy (e.g. methotrexate, cyclosporine) -OR- inadequate response to phototherapy. If not a candidate for phototherapy: treatment with systemic therapy has been ineffective, not tolerated, or is contraindicated.
Age Restrictions	Deny if less than 18 years of age for Rheumatoid Arthritis, Psoriatic Arthritis and Ankylosing Spondylitis or less than 2 years of age for Juvenile Idiopathic Arthritis or Less than 4 years of age for Plaque Psoriasis
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For plaque psoriasis induction therapy, doses above plan quantity limit will be approved aligned with recommended induction therapy dosing regimen.
Indications	All FDA-approved Indications.
Off Label Uses	

endari

Products Affected

- **ENDARI**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of Sickle Cell Disease with 2 or more sickle cell acute complications (e.g. vaso-occlusive crisis, acute anemia, acute chest syndrome, etc.) -AND-documentation of previous trial of hydroxyurea or plans of continued therapy while taking Endari
Age Restrictions	Deny if less than 5 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of stability in sickle cell acute complications or decrease in number of sickle cell acute complications is required (e.g. vaso-occlusive crisis, acute anemia, acute chest syndrome, etc.)
Indications	All FDA-approved Indications.
Off Label Uses	

enspryng

Products Affected

- ENSPRYNG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of neuromyelitis optica spectrum disorder (NMOSD) - AND- Attestation of anti-aquaporin-4 (AQP4) antibody positive -AND- Not used in combination with another monoclonal antibody used for the treatment of NMOSD.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of decrease in number of NMOSD relapses. For induction therapy dosing, doses above plan quantity limit will be approved aligned with recommended induction therapy dosing regimens
Indications	All FDA-approved Indications.
Off Label Uses	

epclusa

Products Affected

- **EPCLUSA ORAL PELLETS IN PACKET 150-37.5 MG, 200-50 MG**
- **EPCLUSA ORAL TABLET**
- *sofosbuvir-velpatasvir*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Criteria will be applied consistent with current AASLD/IDSA guidance
Age Restrictions	Deny if less than 3 years of age
Prescriber Restrictions	
Coverage Duration	Criteria/duration applied consistent with current AASLD-IDSA guidance
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

epidiolex

Products Affected

- EPIDIOLEX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of Lennox-Gastaut, Dravet syndromes or Tuberous Sclerosis Complex. For Lennox-Gastaut, trial and failure or intolerance of at least two standard of care treatments (e.g. lamotrigine, clobazam). For Lennox-Gastaut and Dravet syndromes, treatment is in combination with other conventional agents.
Age Restrictions	Deny if less than 1 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation supporting reduction in seizure frequency
Indications	All FDA-approved Indications.
Off Label Uses	

Eprontia

Products Affected

- **EPRONTIA**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- inability to swallow pills or food. For migraine, therapeutic failure, intolerance, or contraindication to two generic preventive migraine therapies. For partial onset seizures, primary generalized tonic-clonic seizures, or adjunctive treatment of Lennox-Gastaut Syndrome, therapeutic failure, contraindication, or intolerance to two generic anti-epileptic drugs.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

erivedge

Products Affected

- **ERIVEDGE**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of advanced basal cell carcinoma (BCC), which includes metastatic -OR- locally advanced basal cell carcinoma, for whom surgery is inappropriate or in whom recurrence after surgery is documented - AND- is not a candidate for radiation
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

erleada

Products Affected

- ERLEADA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- the member meets one of the following (1 or 2) 1. Documentation of use in combination with a GnRH analog -OR- 2. The member has had a bilateral orchiectomy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

evenity

Products Affected

- **EVENITY SUBCUTANEOUS SYRINGE
210MG/2.34ML (105MG/1.17MLX2)**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- Postmenopausal woman at high risk for fracture, meeting one of the following (1. thru 3.) 1) History of previous hip or vertebral fracture. 2) T-score less than or equal to -2.5. 3) T-score between -1.0 and -2.5 -AND- meets FRAX calculation (A. or B.) A) 10-year risk of major osteoporotic fracture is greater than or equal to 20 percent or B) 10-year risk of hip fracture is greater than or equal to 3 percent.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Documentation of trial/failure or intolerance to at least one oral bisphosphonate or all are contraindicated. Covered under Part B for patients eligible for home health services when provider certifies that patient sustained bone fracture related to post-menopausal osteoporosis and is unable to learn the skills needed to self-administer the drug or is otherwise physically or mentally incapable of administering the drug or family/caregivers are unable or unwilling to administer the drug. A cumulative lifetime approval of romosozumab will be limited to a coverage duration of 12 months.
Indications	All FDA-approved Indications.
Off Label Uses	

evrysdi

Products Affected

- EVRYSDI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of spinal muscular atrophy -AND- Baseline motor function test results (e.g. MFM, CHOP, HINE, RULM, HFMSE, 6MWT) -AND- Not using concomitantly with Spinraza -AND- Molecular genetic testing of 5q SMA showing Homozygous gene deletion, Homozygous conversion mutation or Compound heterozygote
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	For reauthorization, attestation of stable or clinically significant improvement in Spinal Muscular Atrophy associated symptoms (e.g. stabilization or decreased decline in motor function compared to the predicted natural history trajectory of disease) -OR- Stable or improved motor function results compared to baseline (e.g. MFM, CHOP, HINE, RULM, HFMSE, 6MWT).
Indications	All FDA-approved Indications.
Off Label Uses	

Exservan

Products Affected

- **EXSERVAN**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of amyotrophic lateral sclerosis (ALS) -AND- Inability to swallow tablets.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of stability or improvement in symptoms of ALS.
Indications	All FDA-approved Indications.
Off Label Uses	

fasenra

Products Affected

- FASENRA
- FASENRA PEN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis of severe asthma -AND- history of at least one asthma exacerbations requiring oral or systemic corticosteroid treatment in the past 12 months -AND- documented pretreatment FEV1 less than 80 percent predicted in adults or FEV1 less than 90 percent predicted in adolescents or FEV1 reversibility of at least 12% after albuterol administration despite regular treatment with (a. or b.): a) high dose inhaled corticosteroid and additional asthma controller medication or b.) a medium or high dose inhaled corticosteroid plus a long-acting beta agonist with or without oral corticosteroids and additional asthma controller medication
Age Restrictions	Deny if less than 12 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Blood eosinophil count (in the absence of other potential causes of eosinophilia, including hypereosinophilic syndromes, neoplastic disease, and known or suspected parasitic infection) greater than or equal to 150 cells/microliter within 6 weeks of initiation of therapy. For induction therapy, doses above plan quantity limit will be approved aligned with recommended induction therapy dosing regimen.
Indications	All FDA-approved Indications.
Off Label Uses	

ferriprox

Products Affected

- *deferiprone*
- **FERRIPROX ORAL SOLUTION**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of transfusional iron overload due to thalassemia syndromes, sickle cell disease and other anemias -AND- transfusion history of greater than or equal to 100 mL/kg of packed red blood cells (i.e. at least 20 units of packed red blood cells for a 40 kg person or more in individuals weighing more than 40 kg) -AND- history of serum ferritin consistently greater than 1,000 mcg/L or liver iron concentration (LIC) greater than or equal to 7 iron per gram of liver dry weight (mg Fe/g dw)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Trial and failure of deferasirox (generic Exjade) is required. For reauthorization, continued requirement for regular blood transfusions - AND- serum ferritin level greater than or equal to 500mcg/L or LIC greater than or equal to 3 mg Fe/g dw
Indications	All Medically-accepted Indications.
Off Label Uses	

fetzima

Products Affected

- **FETZIMA ORAL CAPSULE,EXT REL HR 120 MG, 20 MG, 40 MG, 80 MG 24HR DOSE PACK**
- **FETZIMA ORAL CAPSULE,EXTENDED RELEASE 24**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of major depressive disorder and trial and failure of two other antidepressants.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	5 years
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

fintepla

Products Affected

- FINTEPLA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of Dravet syndrome or Lennox-Gastaut syndrome. For Lennox-Gastaut syndrome, therapeutic failure, contraindication, or intolerance to at least 2 standard of care treatments (e.g. lamotrigine, clobazam).
Age Restrictions	Deny if less than 2 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

firazyr

Products Affected

- *icatibant*
- SAJAZIR

PA Criteria	Criteria Details
Exclusion Criteria	Member should not be on two acute therapies simultaneously and acute therapy should not be used as prophylactic therapy
Required Medical Information	For the treatment of acute abdominal, facial, or laryngeal attacks of hereditary angioedema (HAE) type I & II with the following (1-3): 1) Low C4 level of less than or equal to 14mg/dL or C4 below lower limit of laboratory reference range and 1 of the following (A or B). A) C1 inhibitor (C1INH) antigen level less than or equal to 19mg/dL or below lower limit of laboratory reference range. B) Normal C1INH antigen level and a low C1INH functional level below laboratory reference range. 2) Past medical history of at least 1 symptom of moderate or severe angioedema attack (e.g. airway swelling, painful facial distortion) in absence of concomitant hives. 3) Medications known to cause angioedema have been evaluated and discontinued. For the treatment of acute abdominal, facial, or laryngeal attacks of hereditary angioedema (HAE) type III with the following (4-7): 4) Documentation of clinical laboratory performance C4, C1INH antigen, or C1INH functional level are within normal limits of laboratory reference ranges. 5) Documentation of family history of HAE or FXII mutation 6) Past medical history of at least 1 symptom of moderate or severe angioedema attack (e.g. airway swelling, painful facial distortion) in absence of concomitant hives. 7) Medications known to cause angioedema have been evaluated and discontinued.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

firdapse

Products Affected

- **FIRDAPSE**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of positive clinical response to therapy.
Indications	All FDA-approved Indications.
Off Label Uses	

flector

Products Affected

- *diclofenac epolamine*
- **FLECTOR**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis AND one of the following (1,2 or 3): 1) trial/failure, intolerance, or contraindication to 2 oral generic NSAIDs one of which must be diclofenac 2) hypersensitivity to oral NSAIDs 3) history or high risk for adverse gastrointestinal effects associated with oral NSAID use.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 month
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

forteo

Products Affected

- **FORTEO SUBCUTANEOUS PEN INJECTOR 20 MCG/DOSE (600MCG/2.4ML)**
- *teriparatide*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- at high risk for fracture, meeting one of the following (1. thru 4.) 1) History of previous hip or vertebral fracture. 2) T-score less than or equal to -2.5. 3) T-score between -1.0 and -2.5 -AND- meets FRAX calculation (A. or B.) A) 10-year risk of major osteoporotic fracture is greater than or equal to 20 percent or B) 10-year risk of hip fracture is greater than or equal to 3 percent. 4) Age 40 years or older with T-score between -1.0 and -2.5 -AND- History of glucocorticoid use for at least 3 months at a dose of 5mg per day or more of prednisone (or equivalent).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	24 months
Other Criteria	Documentation of trial/failure or intolerance to at least one oral bisphosphonate or all are contraindicated. Additional documentation of trial/failure, intolerance or contraindication to preferred parathyroid hormone analog Tymlos is required for applicable indication. Coverage of human parathyroid hormone related peptide analogs beyond 24 months will not be approved. A cumulative lifetime approval of Forteo/teriparatide will be limited to a coverage duration of 24 months in the absence of provider attestation that the member remains at or has returned to having a high risk for fracture
Indications	All FDA-approved Indications.
Off Label Uses	

Fotivda

Products Affected

- FOTIVDA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- if applicable to diagnosis, previous therapies tried/failed
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

gabapentin

Products Affected

- *gabapentin oral capsule 100 mg, 300 mg, 400 mg*
- *gabapentin oral solution 250 mg/5 ml*
- *gabapentin oral tablet 600 mg, 800 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. When using concomitantly with an opiate agonist, attestation of an intent to monitor and address concomitant drug-drug interaction adverse events for opiate potentiators
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

galafold

Products Affected

- GALAFOLD

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of Fabry disease confirmed by biochemical or genetic test -AND- Presence of an amenable GLA variant causing Fabry disease in the clinical context of the patient -AND- Will not be used concomitantly with enzyme replacement therapy (ERT) e.g. Fabrazyme.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of positive clinical response to therapy - AND- Will not be used concomitantly with enzyme replacement therapy (ERT) e.g. Fabrazyme.
Indications	All FDA-approved Indications.
Off Label Uses	

gattex

Products Affected

- **GATTEX 30-VIAL**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of short bowel syndrome (SBS) having less than 200 cm of functional small bowel AND if age 18 and older, dependence on parenteral nutrition or intravenous nutritional support for at least 12 months AND requiring parenteral nutrition at least 3 times per week -OR- Documentation of SBS having less than 200 cm of functional small bowel AND age 1 to 17 years of age AND Dependence on parenteral nutrition AND weight of at least 10 kg.
Age Restrictions	Deny if less than 1 year of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, continued dependence on parenteral nutrition/intravenous nutritional support -AND- attestation of increase in weight from baseline or decrease in intravenous parenteral nutrition requirements from baseline.
Indications	All FDA-approved Indications.
Off Label Uses	

gavreto

Products Affected

- **GAVRETO**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis 1) RET mutant or fusion status 2) Radioactive iodine-refractory (if radioactive iodine is appropriate)
Age Restrictions	Deny if less than 18 years of age for NSCLC or less than 12 years of age for thyroid cancer
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

gilenya

Products Affected

- *fingolimod*

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use of Gilenya and other disease modifying agents such as interferons, Copaxone, Tysabri
Required Medical Information	Documentation of relapsing form of multiple sclerosis (e.g. relapsing-remitting, clinically isolated syndrome, or active secondary progressive disease)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	5 years
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

gleevec

Products Affected

- *imatinib oral tablet 100 mg, 400 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis and all of the following, if applicable to diagnosis 1) Alternatives tried 2) Concomitant therapy 3) mutation status, if applicable to diagnosis.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

grastek

Products Affected

- **GRASTEK**

PA Criteria	Criteria Details
Exclusion Criteria	Asthma (severe, unstable or uncontrolled), concomitant sublingual or subcutaneous immunotherapy
Required Medical Information	Documentation of allergic rhinitis and use for Timothy grass pollen or cross reactive grass pollens (Sweet Vernal, Orchard, Perennial Rye, Timothy, Kentucky Blue Grass pollen, Redtop, or meadow fescue) -AND- allergic rhinitis with or without conjunctivitis has been confirmed by a pollen specific positive skin test or in vitro testing for pollen-specific IgE antibodies -AND- trial and failure or intolerance to an intranasal steroid and an oral non-sedating antihistamine, intranasal antihistamine or intranasal anticholinergic agent
Age Restrictions	Deny if less than 5 years of age or greater than 65 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Member must also be prescribed an epinephrine auto injector. For reauthorization, attestation of improved allergy symptoms is required.
Indications	All FDA-approved Indications.
Off Label Uses	

growth hormone

Products Affected

- GENOTROPIN
- GENOTROPIN MINIQUICK
- HUMATROPE INJECTION CARTRIDGE
- NORDITROPIN FLEXPPO
- NUTROPIN AQ NUSPIN
- OMNITROPE
- SAIZEN
- SAIZEN SAIZENPREP
- SEROSTIM SUBCUTANEOUS RECON SOLN 4 MG, 5 MG, 6 MG
- ZOMACTON
- ZORBTIVE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis, growth chart, bone age, growth velocity, response to stimulation test, when applicable to meet standard diagnostic criteria. Additionally for growth failure due to chronic kidney disease, glomerular filtration rate is less than 89ml/min per 1.73m*2. For HIV wasting and cachexia, Concurrent use of antiretroviral therapy -AND- weight loss of at least 10 percent from baseline. For short bowel syndrome, receiving optimal management for short bowel syndrome, including specialized nutritional support -AND- less than 200 cm of functional small bowel.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of growth velocity and bone age, as applicable to meet standard continuation of therapy guidelines. For reauthorization of HIV wasting and cachexia, attestation of increase in weight from start of therapy. For reauthorization of short bowel syndrome, continued dependence on parenteral nutrition/intravenous nutritional support -AND- attestation of increase in weight from baseline or decrease in intravenous parenteral nutrition requirements from baseline.
Indications	All FDA-approved Indications.
Off Label Uses	

haegarda

Products Affected

- HAEGARDA

PA Criteria	Criteria Details
Exclusion Criteria	Member should not be on two prophylactic therapies simultaneously.
Required Medical Information	<p>For the prophylactic treatment of abdominal, facial, or laryngeal attacks of hereditary angioedema (HAE) type I & II with the following (1-4): 1) Low C4 level of less than or equal to 14mg/dL or C4 below lower limit of laboratory reference range and 1 of the following (A or B). A) C1 inhibitor (C1INH) antigen level less than or equal to 19mg/dL or below lower limit of laboratory reference range. B) Normal C1INH antigen level and a low C1INH functional level below laboratory reference range. 2) Past medical history of at least 1 symptom of moderate or severe angioedema attack (e.g. airway swelling, painful facial distortion) in absence of concomitant hives. 3) Medications known to cause angioedema have been evaluated and discontinued. 4) Documentation of member's weight. For the prophylactic treatment of acute abdominal, facial, or laryngeal attacks of hereditary angioedema (HAE) type III with the following (5-9): 5) Documentation of clinical laboratory performance C4, C1INH antigen, or C1INH functional level are within normal limits of laboratory reference ranges. 6) Documentation of family history of HAE or FXII mutation 7) Past medical history of at least 1 symptom of moderate or severe angioedema attack (e.g. airway swelling, painful facial distortion) in absence of concomitant hives. 8) Medications known to cause angioedema have been evaluated and discontinued. 9) Documentation of member's weight.</p>
Age Restrictions	Deny if less than 6 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	

HARVONI

Products Affected

- **HARVONI ORAL PELLETS IN PACKET**
- **HARVONI ORAL TABLET 90-400 MG**
- *ledipasvir-sofosbuvir*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Criteria will be applied consistent with current AASLD/IDSA guidance
Age Restrictions	Deny if less than 3 years of age
Prescriber Restrictions	
Coverage Duration	Criteria/duration applied consistent with current AASLD-IDSA guidance
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

HETLIOZ

Products Affected

- HETLIOZ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Non-24 Sleep-Wake disorder in patient that is totally blind -AND- evidenced by all the following (1 through 4): 1) history of insomnia, excessive daytime sleepiness, or both alternating with asymptomatic episodes 2) symptoms persistent for at least 3 months 3) daily sleep logs for at least 1 month demonstrating a sleep/wake pattern that delays each day 4) sleep disturbances are not better explained by another current disorder or medication/substance use -OR- Documented diagnosis of Smith-Magenis Syndrome as confirmed by chromosome analysis -AND- patient is experiencing nighttime sleep disturbances (e.g. difficulty falling asleep, shortened sleep cycles, inability to enter REM sleep, or frequent awaking during the night and early in the morning)
Age Restrictions	Deny if less than 16 years of age for nighttime sleep disturbances in Smith-Magenis Syndrome or deny if less than 18 years of age for Non-24 Hour Sleep-Wake Disorder
Prescriber Restrictions	
Coverage Duration	3 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, attestation of increased total nighttime sleep or decreased daytime nap duration for Non-24 Sleep-Wake disorder -OR- attestation of positive clinical response to therapy with minimal side effects for Smith-Magenis Syndrome
Indications	All FDA-approved Indications.
Off Label Uses	

Hetlioz LQ

Products Affected

- HETLIOZ LQ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Smith-Magenis Syndrome as confirmed by chromosome analysis -AND- patient is experiencing nighttime sleep disturbances (e.g. difficulty falling asleep, shortened sleep cycles, inability to enter REM sleep, or frequent awaking during the night and early in the morning)
Age Restrictions	Deny if less than 3 or greater than 15 years of age.
Prescriber Restrictions	
Coverage Duration	3 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, attestation of positive clinical response to therapy with minimal side effects for Smith-Magenis Syndrome -AND- member is between 3 and 15 years of age
Indications	All FDA-approved Indications.
Off Label Uses	

high-risk meds

Products Affected

- *amitriptyline*
- *amitriptyline-chlordiazepoxide*
- *benztropine oral*
- *clomipramine*
- *cyclobenzaprine oral tablet 10 mg, 5 mg, 7.5 mg*
- *cyproheptadine*
- *doxepin oral capsule*
- *doxepin oral concentrate*
- *doxepin oral tablet*
- *ergoloid*
- *ergotamine-caffeine*
- *hydroxyzine hcl oral solution 10 mg/5 ml*
- *hydroxyzine hcl oral tablet*
- *imipramine hcl*
- *imipramine pamoate*
- *metaxalone*
- *perphenazine-amitriptyline*
- *promethazine oral syrup*
- *trimipramine*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	For all medications subject to this PA group, the following information (1 through 3) is required: 1. Documentation of diagnosis 2. Explanation of risk-benefit profile favoring use of the high-risk medication 3. Attestation of an intent to monitor and address treatment-related adverse events. For target tricyclic antidepressants (TCAs), in addition to criteria 1 through 3 above, trial and failure or documentation of intolerance or contraindication to at least 2 non-high risk alternative drugs for the same indication, if available, is required (e.g. SSRIs and SNRIs). If using a TCA for a medically-accepted indication not shared by the safer alternatives listed, then no trial of alternatives is required.
Age Restrictions	Automatic approval if less than 65 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Doxepin doses less than or equal to 6 mg per day will receive automatic approval.
Indications	All Medically-accepted Indications.
Off Label Uses	

High-risk meds phenobarbital

Products Affected

- *phenobarbital*

PA Criteria	Criteria Details
Exclusion Criteria	Coverage is not provided for use in sedation/insomnia
Required Medical Information	For use in seizures the following are required: 1. Explanation of risk-benefit profile favoring use of the high-risk medication 2. Attestation of an intent to monitor and address treatment-related adverse events. 3. For new starts, the trial and failure or documentation of intolerance or contraindication to at least 2 non-high risk alternative drugs used for seizures (e.g. carbamazepine, lamotrigine) is required.
Age Restrictions	Automatic approval if less than 65 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off Label Uses	Seizure disorders

homozygous fh

Products Affected

- **JUXTAPID ORAL CAPSULE 10 MG, 20 MG, 30 MG, 5 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of homozygous familial hypercholesterolemia (HoFH) supported by genetic confirmation of two mutant alleles at LDLR, APOB, PCSK9, or LDLRAP1 gene -OR- untreated LDL-C greater than 400mg/dL or TC greater than 500mg/dl with cutaneous or tendon xanthoma before age 10 yrs or HeFH in both parents -AND- The member has a current LDL-C greater than 100mg/dL despite use of maximally tolerated statin OR statin intolerance -AND- The member will continue to receive concurrent lipid-lowering therapies for the treatment of HoFH
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	Patients must have an adequate trial/failure or contraindication to the preferred product Repatha. For reauthorization, documentation showing an LDL-C reduction on Juxtapid therapy from baseline must be provided. Statin intolerance defined as follows: statin related rhabdomyolysis or skeletal muscle symptoms while receiving at least 2 separate trials of different statins which resolved upon discontinuation of statin or attestation of one of the following during any course of statin therapy: 1. CK increase to 10x upper limit of normal 2. LFTs increase to 3x upper limit of normal 3. Hospitalization due to severe statin-related AEs such as rhabdomyolysis.
Indications	All FDA-approved Indications.
Off Label Uses	

horizant

Products Affected

- **HORIZANT ORAL TABLET
EXTENDED RELEASE 300 MG, 600 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of moderate to severe active primary restless leg syndrome -AND- trial and failure of pramipexole or ropinirole -AND- trial and failure of an additional dopaminergic agent, clonidine or pregabalin -OR- Documentation of post herpetic neuralgia and trial and failure of generic gabapentin. When using concomitantly with an opiate agonist, attestation of an intent to monitor and address concomitant drug-drug interaction adverse events for opiate potentiators.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

humira

Products Affected

- HUMIRA PEN
 - HUMIRA PEN CROHNS-UC-HS START
 - HUMIRA PEN PSOR-UEVITS-ADOL HS
 - HUMIRA SUBCUTANEOUS SYRINGE KIT 40 MG/0.8 ML
 - HUMIRA(CF)
 - HUMIRA(CF) PEDI CROHNS
- STARTER SUBCUTANEOUS SYRINGE KIT 80 MG/0.8 ML, 80 MG/0.8 ML-40 MG/0.4 ML
 - HUMIRA(CF) PEN
 - HUMIRA(CF) PEN CROHNS-UC-HS
 - HUMIRA(CF) PEN PEDIATRIC UC
 - HUMIRA(CF) PEN PSOR-UV-ADOL HS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For moderate to severe rheumatoid arthritis, inadequate response or intolerance to at least one DMARD (e.g. methotrexate, leflunomide). For ankylosing spondylitis, inadequate response or intolerance to at least one nonsteroidal anti-inflammatory drug (NSAID). For moderate to severe juvenile idiopathic rheumatoid arthritis, inadequate response or intolerance to at least one DMARD (e.g., methotrexate, leflunomide) -OR- requires initial biologic therapy due to involvement of high-risk joints, high disease activity or at high risk of disabling joint damage. For moderate to severe plaque psoriasis, inadequate response or intolerance to one systemic therapy (e.g. methotrexate, cyclosporine) -OR- inadequate response to phototherapy. If not a candidate for phototherapy: treatment with systemic therapy has been ineffective, not tolerated, or is contraindicated. For uveitis, inadequate response or intolerance to 2 immunosuppressants.
Age Restrictions	Deny if less than 18 years of age for Rheumatoid Arthritis, Psoriatic Arthritis, Plaque Psoriasis, and Ankylosing Spondylitis or less than 12 years of age for Hidradenitis Suppurative or Less than 6 years of age for Crohn's disease or Less than 5 years of age for Ulcerative Colitis or less than 2 years of age for Juvenile Idiopathic Arthritis and Uveitis
Prescriber Restrictions	
Coverage Duration	12 months

PA Criteria	Criteria Details
Other Criteria	For Crohn's disease in adults (18 years or older), trial of 2 immunosuppressants (e.g. corticosteroids, azathioprine) or monotherapy with infliximab is required. For Crohn's disease in pediatrics, trial of 1 immunosuppressant (e.g. corticosteroids, azathioprine) or monotherapy with infliximab is required. For plaque psoriasis induction therapy, doses above plan quantity limit will be approved aligned with recommended induction therapy dosing regimen. For rheumatoid arthritis therapy without concomitant methotrexate, doses above plan quantity limit will be approved aligned with recommended weekly dosing regimen. For pediatric ulcerative colitis and hidradenitis suppurativa, doses above plan quantity limit will be approved to align with recommended dosing regimen. Induction therapy or treatment regimens for other indications are aligned with plan quantity limit on Humira starter kit.
Indications	All FDA-approved Indications.
Off Label Uses	

Ibrance

Products Affected

- **IBRANCE**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of HR-positive, HER2-negative advanced or metastatic breast cancer -AND- meets one of the following (1 or 2): 1) documentation of use with an aromatase inhibitor as initial endocrine-based therapy in postmenopausal women or men -OR- 2) documentation of use with fulvestrant in patients with disease progression following endocrine therapy.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Ibsrela

Products Affected

- **IBSRELA**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of irritable bowel syndrome with constipation -AND- failure or intolerance to Linzess -AND- if member is female, failure or intolerance to lubiprostone.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

iclusig

Products Affected

- ICLUSIG

PA Criteria	Criteria Details
Exclusion Criteria	Treatment of newly-diagnosed chronic phase CML
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) Philadelphia chromosome status 2) T315I status 3) Alternatives tried/failed 4) Candidacy for other tyrosine kinase inhibitor therapy
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

idhifa

Products Affected

- **IDHIFA ORAL TABLET 100 MG, 50 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of relapsed or refractory acute myeloid leukemia (AML) with an isocitrate dehydrogenase-2 (IDH2) mutation as detected by an FDA approved test
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Products Affected

- **BIVIGAM**
- **FLEBOGAMMA DIF INTRAVENOUS SOLUTION 10 %**
- **GAMMAGARD LIQUID**
- **GAMMAGARD S-D (IGA)**
- **GAMMAKED INJECTION SOLUTION 1 GRAM/10 ML (10 %)**
- **GAMMAPLEX**
- **GAMMAPLEX (WITH SORBITOL)**
- **GAMUNEX-C INJECTION SOLUTION 1 GRAM/10 ML (10 %)**
- **OCTAGAM**
- **PANZYGA**
- **PRIVIGEN**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	<p>Documentation of diagnosis. For select diagnoses the following apply- 1) For Myasthenia Gravis Syndrome, documentation that the patient is refractory to other standard therapies (e.g., cholinesterase inhibitors, corticosteroids, azathioprine) given in therapeutic doses over at least 3 months OR is intolerant of/has a contraindication to those standard therapies. 2) For Multiple Sclerosis, patient is refractory to other standard therapies (e.g., interferons) given in therapeutic doses over at least 3 months OR is intolerant of/has a contraindication to those standard therapies. 3) For Inflammatory Myopathies, the patient is refractory to corticosteroids given in therapeutic doses over at least 4 months OR is intolerant of/has a contraindication to corticosteroids or immunosuppressants. 4) For CLL with Associated Hypogammaglobulinemia, IgG level less than 600mg/dL or evidence of a specific antibody deficiency or recurrent bacterial infections. 5) For Bone Marrow Transplantation, the member is 20 years of age or older and within the first 100 days after transplantation. 6) For Dermatomyositis/Polymyositis, trial and failure, intolerance, or contraindication to standard first line therapy (i.e. corticosteroids or immunosuppressants). 7) For Pediatric HIV, the patient is less than 13 y.o. who have entry CD4 lymphocyte count greater than or equal to 200/mm³ and IgG less than 400mg/dL OR recurrent bacterial infections. 8) For Guillain-Barre syndrome, impaired function by objective assessment and/or objective findings on physical exam at the time of initial therapy and IVIG therapy must be initiated within 2 weeks of symptom onset. 9) For Autoimmune Mucocutaneous Blistering Diseases (e.g. Stevens-Johnson Syndrome), trial and failure, intolerance, or contraindication to conventional therapy (e.g. corticosteroids) or the patient has rapidly progressive disease in which a clinical response could not be affected quickly enough using conventional agents.</p>

PA Criteria	Criteria Details
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Covered under Part B when administered in the home to a member with a diagnosis of primary immunodeficiency disease
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off Label Uses	Myasthenia Gravis syndrome, Multiple Sclerosis, Inflammatory Myopathies, Polymyositis, Dermatomyositis, Bone Marrow Transplant, Pediatric HIV, Guillain-Barre syndrome, Autoimmune Mucocutaneous Blistering Diseases

imbruvica

Products Affected

- **IMBRUVICA ORAL CAPSULE 140 MG, 70 MG**
- **IMBRUVICA ORAL SUSPENSION**
- **IMBRUVICA ORAL TABLET**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) 17p deletion status 2) Alternatives tried/failed 3) concomitant therapy. For suspension, members must also have an inability to swallow oral tablets or oral capsules.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

inbrija

Products Affected

- **INBRIJA INHALATION CAPSULE,
W/INHALATION DEVICE**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of use for the treatment of intermittent off episodes of Parkinson's disease while on carbidopa/levodopa
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of positive clinical response
Indications	All FDA-approved Indications.
Off Label Uses	

increlex

Products Affected

- INCRELEX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis of severe primary IGF-1 deficiency and all of the following: 1) Normal or elevated response (greater than 10 ng/ml) to two (2) of the following standard growth hormone stimulation tests: arginine, clonidine, glucagon, insulin, levodopa, propranolol. 2) Serum IGF-1 concentration that is less than or equal to three (3) standard deviations below the normal value based on laboratory reference range. 3) Height less than or equal to three (3) standard deviations below normal (at or below the third percentile for gender and age). 4) If female, bone age less than or equal to 14 years. If male, bone age less than or equal to 16. - OR- Documentation of diagnosis of growth hormone deficiency caused by gene deletion and all of the following: 1) Growth velocity at least 2 standard deviations below the age-appropriate mean or height at least 2.25 standard deviations below the age-appropriate mean. 2) Subnormal response (less than 10 ng/mL) to two (2) of the following standard growth hormone stimulation tests: arginine, clonidine, glucagon, insulin, levodopa, propranolol. 3) Development of neutralizing antibodies to growth hormone product(s). 4) If female, bone age of less than or equal to 14 years. If male, bone age less than or equal to 16.
Age Restrictions	Deny if greater than 17 years old
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of growth velocity and bone age, as applicable to meet standard continuation of therapy guidelines
Indications	All FDA-approved Indications.
Off Label Uses	

ingrezza

Products Affected

- **INGREZZA INITIATION PACK**
- **INGREZZA ORAL CAPSULE 40 MG, 60 MG, 80 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of tardive dyskinesia
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

inlyta

Products Affected

- INLYTA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) Alternatives tried/failed 2) Concomitant therapy 3) Line of therapy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

inqovi

Products Affected

- INQOVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation chronic myelomonocytic leukemia. Documentation of de novo or secondary myelodysplastic syndrome -AND- One of the following (1 or 2): 1) French American-British MDS subtypes of refractory anemia, refractory anemia with ringed sideroblasts or refractory anemia with excess blasts. 2) International Prognostic Scoring System group of intermediate-1, intermediate-2 or high-risk.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

inrebic

Products Affected

- **INREBIC**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of intermediate-2 or high-risk myelofibrosis, including primary myelofibrosis, post-polycythemia vera myelofibrosis and post-essential thrombocythemia myelofibrosis per an accepted risk stratification tool for myelofibrosis (e.g., International Prognostic Scoring System [IPSS]) -AND- If a new start, baseline platelet count of greater than $50 \times 10^9/L$
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Baseline platelet count to be provided.
Indications	All FDA-approved Indications.
Off Label Uses	

interferon alfa

Products Affected

- **INTRON A INJECTION RECON SOLN
10 MILLION UNIT (1 ML), 50 MILLION
UNIT (1 ML)**
- **PEGASYS**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

interleukin-1b blockers

Products Affected

- ARCALYST

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For Deficiency of Interleukin-1 Receptor Antagonist (DIRA), documentation of need for maintenance of remission. For Recurrent Pericarditis (RP), documentation of trial/failure or intolerance to one, or contraindication to all of the following: oral nonsteroidal anti-inflammatory drug (NSAID), systemic corticosteroid, or colchicine.
Age Restrictions	Deny if less than 12 years of age for Recurrent Pericarditis and Cryopyrin-Associated Periodic Syndromes
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For DIRA: patient must weigh 10kg or more
Indications	All FDA-approved Indications.
Off Label Uses	

IPF AGENTS

Products Affected

- **ESBRIET ORAL CAPSULE**
 - **ESBRIET ORAL TABLET 267 MG, 801 MG**
 - **OFEV**
- *pirfenidone oral tablet 267 mg, 801 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of idiopathic pulmonary fibrosis -AND- baseline forced vital capacity (FVC) of at least 50% and a percent predicted diffusing capacity of the lungs of carbon monoxide (DLCO) of at least 30%. For Ofev only, documentation of systemic sclerosis-associated interstitial lung disease -AND- baseline forced vital capacity (FVC) of at least 40% and a percent predicted diffusing capacity of the lungs of carbon monoxide (DLCO) of at least 30% -AND- documentation of a high-resolution chest computed tomography (CT) scan demonstrating greater than or equal to 10% pulmonary fibrosis. For Ofev only, documentation of chronic fibrosing interstitial lung disease with progressive phenotype -AND- high resolution chest computing tomography (HRCT) scan demonstrating greater than 10% fibrosing disease -AND- baseline forced vital capacity (FVC) of at least 45% and a percent predicted diffusing capacity of the lungs of carbon monoxide (DLCO) of at least 30% -AND- disease progression in previous 24 months shown by one of the following : 1. Relative decline in FVC greater than or equal to 10% predicted 2. Relative decline in FVC greater than or equal to 5% but less than 10% predicted and either worsening of respiratory symptoms or increased extent of fibrotic changes on HRCT 3. Worsening of respiratory symptoms and increasing extent of fibrotic changes on HRCT
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	

iressa

Products Affected

- **IRESSA**

PA Criteria	Criteria Details
Exclusion Criteria	Use in tumors with EGFR mutations other than exon 19 deletions or exon 21 (L858R) substitution mutations.
Required Medical Information	Documentation of diagnosis -AND- the following: 1) EGFR exon 19 deletion mutations or exon 21 (L858R) mutations as detected by an FDA-approved test
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

isturisa

Products Affected

- **ISTURISA**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of Cushing's disease AND patient is not a candidate for pituitary surgery or surgery has not been curative
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, attestation of mean urine free cortisol (mUFC) less than starting baseline value.
Indications	All FDA-approved Indications.
Off Label Uses	

itraconazole

Products Affected

- *itraconazole*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. If using for diagnosis of onychomycosis, confirmation through positive laboratory testing (e.g. KOH preparation, fungal culture, or nail biopsy) is required.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Onychomycosis: 3 months. All other indications: 3 months initial, 12 months reauth
Other Criteria	Documentation of trial/failure or intolerance of amphotericin b must be provided for approval in patients with aspergillosis.
Indications	All FDA-approved Indications.
Off Label Uses	

jakafi

Products Affected

- **JAKAFI**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of intermediate or high-risk myelofibrosis, including primary myelofibrosis, post-polycythemia vera myelofibrosis and post-essential thrombocythemia myelofibrosis per an accepted risk stratification tool for myelofibrosis (e.g., International Prognostic Scoring System [IPSS]) and if a new start, baseline platelet count of greater than $50 \times 10^9/L$ -OR- documentation of polycythemia vera and inadequate response or intolerance to hydroxyurea -OR- Documentation of steroid refractory acute graft-versus-host disease and prior therapy with at least one systemic corticosteroid -OR- Documentation of chronic graft-versus-host disease with prior failure of at least one systemic therapy.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Baseline platelet count to be provided.
Indications	All FDA-approved Indications.
Off Label Uses	

jynarque

Products Affected

- JYNARQUE ORAL TABLET
- JYNARQUE ORAL TABLETS, SEQUENTIAL

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of rapidly progressing autosomal dominant polycystic kidney disease defined by one of the following: 1.) Historical decline in eGFR greater than or equal to 5mL/min/1.73 m*2 within a 12 month period. 2.) Decline in eGFR of greater than or equal to 2.5mL/min/1.73m*2 over a period of 5 years. 3.) 5% increase in total kidney volume per year by 3 repeat CT or MRI. 4.) Average kidney length greater than 16.5cm. 5.) Family history of end-stage renal disease before age 58. 6.) Mayo imaging classification of 1C, 1D, or 1E. 7.) Kidney bleeds.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, prescriber attestation of slowed decline of kidney function
Indications	All FDA-approved Indications.
Off Label Uses	

kalydeco

Products Affected

- **KALYDECO ORAL GRANULES IN PACKET 25 MG, 50 MG, 75 MG**
- **KALYDECO ORAL TABLET**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of cystic fibrosis (CF) in patients who have one mutation in the cystic fibrosis transmembrane conductance regulator (CFTR gene) that is responsive to ivacaftor based on clinical and or in vitro assay (e.g. G551D, G1244E, G1349D)
Age Restrictions	Granules- Deny if less than 4 months or greater than 5 years of age. Tablets- Deny if less than 6 years of age.
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, documentation supporting improvement or stabilization of FEV1 compared to baseline FEV1 -or- increase in body mass index -or- decreased pulmonary exacerbations -or- improved quality of life as demonstrated by CF Questionnaire is required.
Indications	All FDA-approved Indications.
Off Label Uses	

kesimpta

Products Affected

- **KESIMPTA PEN**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of relapsing form of multiple sclerosis (e.g. relapsing-remitting, clinically isolated syndrome, or active secondary progressive disease)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For induction therapy dosing, doses above plan quantity limit will be approved aligned with recommended induction therapy dosing regimens
Indications	All FDA-approved Indications.
Off Label Uses	

kevzara

Products Affected

- **KEVZARA**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For moderate to severe rheumatoid arthritis, inadequate response or intolerance to at least one DMARD (e.g. leflunomide).
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For rheumatoid arthritis, patients must have an adequate trial or intolerance to two of the following preferred products Humira, Enbrel, Actemra, Xeljanz/Xeljanz XR and Rinvoq.
Indications	All FDA-approved Indications.
Off Label Uses	

kineret

Products Affected

- **KINERET**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For moderate to severe rheumatoid arthritis, inadequate response or intolerance to at least one DMARD (e.g. methotrexate, leflunomide). For Deficiency of Interleukin-1 Receptor Antagonist (DIRA), therapeutic failure or intolerance to at least one (1) corticosteroid, or all corticosteroids are contraindicated.
Age Restrictions	Deny if less than 18 years of age for Rheumatoid Arthritis
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For rheumatoid arthritis, patients must have an adequate trial or intolerance to 2 of the following preferred products Humira, Enbrel, Actemra, Rinvoq and Xeljanz/Xeljanz XR.
Indications	All FDA-approved Indications.
Off Label Uses	

kisqali

Products Affected

- **KISQALI FEMARA CO-PACK ORAL TABLET 200 MG/DAY(200 MG X 1)-2.5 MG, 400 MG/DAY(200 MG X 2)-2.5 MG, 600 MG/DAY(200 MG X 3)-2.5 MG**
- **KISQALI ORAL TABLET 200 MG/DAY (200 MG X 1), 400 MG/DAY (200 MG X 2), 600 MG/DAY (200 MG X 3)**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following. 1) HR mutation status and HER2 mutation status 2) Alternatives tried/failed 3) Concomitant therapy, if applicable to diagnosis
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

klisyri

Products Affected

- **KLISYRI**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of actinic keratosis of the face or scalp -AND- Therapeutic failure or intolerance to 2 of the following 1) generic imiquimod 5% cream 2) fluorouracil 5% topical cream 3) fluorouracil topical solution
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	3 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

korlym

Products Affected

- **KORLYM**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of hyperglycemia secondary to hypercortisolism in patients with endogenous Cushing's syndrome -AND- Patient is not a candidate for surgery or where surgery has failed -AND- one of the following (1 or 2): 1) Diagnosis of diabetes with trial and failure, intolerance, or contraindication to one previous therapy for Type 2 Diabetes (e.g. metformin, sulfonylureas, insulin) or using in addition to a therapy for Type 2 diabetes. 2) Glucose intolerance
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

koselugo

Products Affected

- **KOSELUGO ORAL CAPSULE 10 MG,
25 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- for neurofibromatosis type 1 (NF1), documentation of symptomatic, inoperable plexiform neurofibromas (PN)
Age Restrictions	Deny if less than 2 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

kuvan

Products Affected

- **JAVYGTOR ORAL POWDER IN PACKET 100 MG**
- *sapropterin*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documented diagnosis of PKU -AND- documented baseline Phe level greater than 6 mg/dL -AND- clinical documentation of current weight -AND- sapropterin dihydrochloride dose does not exceed 20 mg/kg/day
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	3 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, initial therapy has resulted in decrease in phenylalanine levels from baseline or current phenylalanine levels within the range of 120-360 micromol/L -AND- clinical documentation of current weight is required -AND- sapropterin dihydrochloride dose does not exceed 20 mg/kg/day.
Indications	All FDA-approved Indications.
Off Label Uses	

kynmobi

Products Affected

- **KYNMOBI SUBLINGUAL FILM 10 MG, 15 MG, 20 MG, 25 MG, 30 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of Parkinson's disease -AND- for use in acute, intermittent treatment of hypomobility off episodes -AND- experiencing off episodes despite the use of oral carbidopa/levodopa -AND- Therapeutic failure, intolerance, or contraindication to the following (1. and 2.): 1) a generic pramipexole containing product or a generic ropinirole containing product. 2) generic entacapone, selegiline or rasagiline.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

latuda

Products Affected

- **LATUDA ORAL TABLET 120 MG, 20 MG, 40 MG, 60 MG, 80 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. If medication is being used for bipolar 1 disorder, documentation of trial and failure or intolerance to one other formulary medication indicated in bipolar 1 disorder (e.g. quetiapine)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

lenvima

Products Affected

- LENVIMA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following if applicable to diagnosis: 1) Radioactive iodine refractory status 2) Microsatellite instability-high status or mismatch repair deficient status 3) Alternatives tried/failed or attestation of first line use 4) Concomitant therapy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

leukine

Products Affected

- LEUKINE INJECTION RECON SOLN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis: following induction chemotherapy in patients who are 55 years or older with acute myelogenous leukemia (AML) -OR- mobilization of hematopoietic progenitor cells into peripheral blood for collection by leukapheresis and autologous transplantation -OR- acceleration of myeloid reconstitution following autologous bone marrow or peripheral blood progenitor cell transplantation -OR- acceleration of myeloid reconstitution following allogeneic BMT -OR- treatment of delayed neutrophil recovery or graft failure after autologous or allogeneic BMT -OR- following acute exposure to myelosuppressive doses of radiation (Hematopoietic Syndrome of Acute Radiation Syndrome [H-ARS]).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

leukotriene modifiers

Products Affected

- *zileuton*
- **ZYFLO**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of asthma -AND- trial/failure of generic montelukast and generic zafirlukast
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

lidoderm

Products Affected

- *lidocaine topical adhesive patch, medicated 5 %*
- **LIDODERM**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of postherpetic neuralgia (PHN) -AND- trial and failure of 1 other agent used to treat PHN (e.g. gabapentin), inability to swallow oral medication or unable to take an oral medication due to potential adverse events (e.g. sedation) -OR- documentation of diabetic peripheral neuropathy (DPN) -AND- trial and failure of one other agent used to treat DPN (e.g. duloxetine), inability to swallow oral medication or unable to take an oral medication due to potential adverse events (e.g. sedation)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off Label Uses	Diabetic peripheral neuropathy

Livtencity

Products Affected

- LIVTENCITY

PA Criteria	Criteria Details
Exclusion Criteria	Diagnosis of CMV prophylaxis.
Required Medical Information	Documentation of refractory cytomegalovirus infection or disease as evidenced by antigenemia or polymerase chain reaction (PCR) test -AND- all of the following (1-3): 1) member weighs at least 35 kg, 2) member is a recipient of hematopoietic stem cell transplant -OR- solid organ transplant. 3) member has experienced therapeutic failure to one of the following: ganciclovir, valganciclovir, cidofovir, or foscarnet.
Age Restrictions	Deny if less than 12 years of age
Prescriber Restrictions	
Coverage Duration	3 months
Other Criteria	For reauthorization, attestation of a previous reduction in CMV DNA level -AND- documentation of one of the following (1-2): 1) new onset symptomatic CMV infection -OR- 2) virologic relapse with treatment-emergent maribavir resistance.
Indications	All FDA-approved Indications.
Off Label Uses	

Lokelma

Products Affected

- LOKELMA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of hyperkalemia as defined by serum potassium level between 5.1 and 7.4 mmol/L on at least two (2) screenings -AND- Modification of medications to reduce serum potassium levels were not successful, when applicable
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, attestation of reduction in serum potassium levels following Lokelma administration and continued treatment for hyperkalemia is required.
Indications	All FDA-approved Indications.
Off Label Uses	

Ionsurf

Products Affected

- LONSURF

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of metastatic colorectal cancer in patients who have previously been treated with fluoropyrimidine-, oxaliplatin-, and irinotecan-based chemotherapy, an anti-VEGF therapy, and if RAS wild-type, an anti-EGFR therapy -OR- documentation of metastatic gastric or gastroesophageal junction adenocarcinoma previously treated with at least two prior lines of chemotherapy that included a fluoropyrimidine, a platinum, either a taxane or irinotecan, and if appropriate, HER2/neu-targeted therapy
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

lorbrena

Products Affected

- **LORBRENA ORAL TABLET 100 MG,
25 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of ALK-positive metastatic non-small cell lung cancer (NSCLC)
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Iotronex

Products Affected

- *alosetron oral tablet 0.5 mg, 1 mg*

PA Criteria	Criteria Details
Exclusion Criteria	For irritable bowel syndrome (IBS): Exclude if male gender
Required Medical Information	Documentation of chronic severe diarrhea-predominant IBS -AND- trial & failure, intolerance, or contraindication to a generic anti-diarrheal agent (e.g. loperamide)
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	Initial: 12 weeks. Reauth: 6 months
Other Criteria	For reauthorization, attestation that symptoms of IBS continue to persist AND positive clinical response.
Indications	All FDA-approved Indications.
Off Label Uses	

Lumakras

Products Affected

- LUMAKRAS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis and all of the following, if applicable to diagnosis 1) KRAS G12C mutation status, as detected by an FDA-approved test 2) Alternatives tried
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

lupkynis

Products Affected

- LUPKYNIS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of active lupus nephritis -AND- Concurrent systemic lupus erythematosus documented by positive ANA titer (greater than or equal to 1:80) or anti-dsDNA greater than or equal to 30 IU/mL -AND- Trial, intolerance, or inadequate response to at least 2 of the following standard of care drug classes: 1.) corticosteroids (e.g. prednisone) 2.) antimalarials (e.g. hydroxychloroquine) 3.) immunosuppressants (e.g. azathioprine, mycophenolate mofetil, or methotrexate) -AND- Member will continue to receive concomitant standard of care treatment with corticosteroids (e.g. prednisone) and mycophenolate mofetil.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	24 weeks initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, attestation of disease stability or disease improvement
Indications	All FDA-approved Indications.
Off Label Uses	

Lybalvi

Products Affected

- LYBALVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of schizophrenia or bipolar I disorder -AND- One of the following (1 or 2): 1) Therapeutic failure, intolerance or contraindication to generic risperidone and generic quetiapine 2) Currently stable and responding to olanzapine but experiencing weight gain from the medication
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

lynparza

Products Affected

- LYNPARZA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- All of the following, if applicable to diagnosis: 1) BRCA mutations 2) Genomic instability status 3) Homologous recombinant repair gene mutations 4) HER2 status 5) Alternatives tried/failed 6) Concomitant therapy
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

lyrica

Products Affected

- *pregabalin oral capsule 100 mg, 150 mg, 200 mg, 225 mg, 25 mg, 300 mg, 50 mg, 75 mg*
- *pregabalin oral solution*
- *pregabalin oral tablet extended release 24 hr*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	For immediate release and controlled release tablets, documentation of DPN and trial/failure or intolerance to duloxetine -OR- PHN and trial/failure or intolerance to gabapentin. For immediate release capsules, documentation of seizures and trial/failure or intolerance to two AEDS - OR- neuropathic pain associated with spinal cord injury -OR- documentation to support a diagnosis of fibromyalgia and trial/failure or intolerance to duloxetine. When using pregabalin products concomitantly with an opiate agonist, attestation of an intent to monitor and address concomitant drug-drug interaction adverse events for opiate potentiators.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

mavenclad

Products Affected

- MAVENCLAD (10 TABLET PACK)
- MAVENCLAD (4 TABLET PACK)
- MAVENCLAD (5 TABLET PACK)
- MAVENCLAD (6 TABLET PACK)
- MAVENCLAD (7 TABLET PACK)
- MAVENCLAD (8 TABLET PACK)
- MAVENCLAD (9 TABLET PACK)

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use of Mavenclad and other disease modifying agents such as interferons, Copaxone, Tysabri. Treatment duration greater than 24 months.
Required Medical Information	Documentation of diagnosis of relapse-remitting multiples sclerosis or active secondary progressive disease -AND- therapeutic failure or intolerance to one other disease modifying therapy (e.g. Avonex, Gilenya, Copaxone)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	24 months
Other Criteria	Coverage beyond 24 months will not be approved.
Indications	All FDA-approved Indications.
Off Label Uses	

mavyret

Products Affected

- MAVYRET ORAL PELLETS IN PACKET
- MAVYRET ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Criteria will be applied consistent with current AASLD/IDSA guidance
Age Restrictions	Deny if less than 3 years of age
Prescriber Restrictions	
Coverage Duration	Criteria/duration applied consistent with current AASLD-IDSA guidance
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

mayzent

Products Affected

- MAYZENT ORAL TABLET 0.25 MG, 1 MG, 2 MG
- MAYZENT STARTER(FOR 1MG MAINT)
- MAYZENT STARTER(FOR 2MG MAINT)

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use of Mayzent and other disease modifying agents such as interferons, Copaxone, Tysabri.
Required Medical Information	Documentation of relapsing form of multiple sclerosis (e.g. relapsing-remitting, clinically isolated syndrome, or active secondary progressive disease) -AND- Testing for CYP2C9 variants has confirmed member does not have CYP2C9*3/*3 genotype
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	24 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

megace

Products Affected

- *megestrol oral suspension 400 mg/10 ml (40 mg/ml), 625 mg/5 ml (125 mg/ml)*
- *megestrol oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

mekinist

Products Affected

- MEKINIST

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following. 1) BRAF mutations, if applicable to diagnosis. 2) Alternatives tried/failed. 3) Concomitant therapy, if applicable to diagnosis.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

mektovi

Products Affected

- MEKTOVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) BRAF mutation status 2) Concomitant therapy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

methamphetamine

Products Affected

- *methamphetamine*

PA Criteria	Criteria Details
Exclusion Criteria	Obesity
Required Medical Information	Documentation of diagnosis
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For ADHD, trial/failure or intolerance to 2 of the following generic medications: methylphenidate, atomoxetine, or dextroamphetamine/amphetamine is required.
Indications	All FDA-approved Indications.
Off Label Uses	

mulpleta

Products Affected

- MULPLETA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of thrombocytopenia and chronic liver disease - AND- beneficiary is scheduled to undergo a procedure -AND- trial and failure or intolerance to Doptelet
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 month
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

myalept

Products Affected

- MYALEPT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of congenital or acquired generalized lipodystrophy - AND- Leptin levels less than 8 ng/mL for males or less than 12 ng/mL for females -AND- the member has a diagnosis of diabetes or fasting insulin levels greater than 30uU/mL or fasting hypertriglyceridemia greater than 200mg/dL -AND- Therapeutic failure to one previous diabetes therapy (e.g. metformin, insulin) or hypertriglyceridemia therapy (e.g. statin, fibrate)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, decrease in HbA1c from baseline, decrease in fasting plasma glucose from baseline or decrease in fasting triglycerides from baseline
Indications	All FDA-approved Indications.
Off Label Uses	

mycapssa

Products Affected

- MYCAPSSA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of acromegaly -AND- High pretreatment insulin-like growth factor-1 (IGF-1) based on laboratory reference range -AND- Previous response to and tolerated treatment with octreotide or lanreotide.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of decreased or normalized IGF-1 from baseline
Indications	All FDA-approved Indications.
Off Label Uses	

Myfembree

Products Affected

- MYFEMBREE

PA Criteria	Criteria Details
Exclusion Criteria	Diagnosis of severe hepatic impairment or osteoporosis.
Required Medical Information	Documentation of Diagnosis. For premenopausal woman with uterine leiomyomas, experiencing heavy menstrual bleeding -AND- Treatment with a gonadotropin-releasing hormone receptor antagonist in combination with estrogen and progestin does not exceed 24 months. For moderate to severe pain associated with endometriosis in premenopausal woman, if the patient is a female of childbearing age, the patient is not pregnant - AND - Therapeutic failure, contraindication, or intolerance to 2 standard of care treatments (i.e. generic NSAID, combined hormonal contraceptive, progestin, GnRH agonist) -AND- Treatment with a gonadotropin-releasing hormone receptor antagonist in combination with estrogen and progestin does not exceed 24 months
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 18 months reauthorization
Other Criteria	For reauthorization of uterine leiomyomas, attestation of continued experience of heavy menstrual bleeding -AND- Attestation of decrease in menstrual blood loss -AND- Treatment with a gonadotropin-releasing hormone receptor antagonist in combination with estrogen and progestin does not exceed 24 months. For reauthorization of endometriosis, continued pain associated with endometriosis -AND- attestation of reduction in pain -AND- For women of childbearing age, attestation of not pregnant -AND- Treatment with a gonadotropin-releasing hormone receptor antagonist in combination with estrogen and progestin does not exceed 24 months
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	

namenda

Products Affected

- **NAMENDA TITRATION PAK**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis and trial/failure of generic memantine
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

namzarin

Products Affected

- **NAMZARIC**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis and trial/failure of generic memantine and generic donepezil
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

natpara

Products Affected

- NATPARA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of use as an adjunct to control hypocalcemia in patients with hypoparathyroidism -AND- Serum calcium concentration is greater than 7.5mg/dL -AND- Attestation of sufficient 25-hydroxyvitamin D stores
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, total serum calcium level is greater than 7.5 mg/dL and less than or equal to 10.6 mg/dL
Indications	All FDA-approved Indications.
Off Label Uses	

nayzilam

Products Affected

- NAYZILAM

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of seizure clusters or acute repetitive seizures -AND- Therapeutic failure, contraindication or intolerance to generic diazepam rectal gel delivery system
Age Restrictions	Deny if less than 12 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

nerlynx

Products Affected

- **NERLYNX**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following. 1) HER2 mutations, if applicable to diagnosis. 2) Alternatives tried/failed. 3) Concomitant therapy, if applicable to diagnosis.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

nexavar

Products Affected

- *sorafenib*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For locally recurrent or metastatic, progressive, differentiated thyroid carcinoma, refractory to radioactive iodine treatment
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

nexletol

Products Affected

- NEXLETOL
- NEXLIZET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	<p>1.HeFH supported by presence of causal mutation of FH by genetic testing OR untreated LDL-C greater than or equal to 190 mg/dL or untreated LDL-C greater than or equal to 160 mg/dL before 20 years of age with physical signs of FH (e.g. xanthomas, xanthelasma) OR diagnosis based on WHO criteria/Dutch Lipid Clinical Network criteria with score greater than 8 points, or definite Simon Broome register criteria, or definite on the Make Early Diagnosis to Prevent Early Deaths tool AND LDL-C greater than 100 mg/dL despite use of maximally tolerated statin or statin intolerance AND therapeutic failure, intolerance or contraindication to ezetimibe AND must be used with maximally tolerated statin dose or documentation of statin intolerance.</p> <p>2.Hypercholesterolemia ASCVD AND LDL-C greater than 70 mg/dL despite use of maximally tolerated statin or statin intolerance AND therapeutic failure, intolerance or contraindication to ezetimibe AND must be used with maximally tolerated statin dose or documentation of statin intolerance.</p>
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months

PA Criteria	Criteria Details
Other Criteria	For reauthorization, documentation showing an LDL-C reduction from baseline AND attestation of continued use of Nexletol or Nexlizet with a maximally tolerated statin, unless statin intolerant. Statin intolerance defined as follows: statin related rhabdomyolysis or skeletal muscle symptoms while receiving at least 2 separate trials of different statin which resolved upon discontinuation of statin or documentation of one of the following during any course of statin therapy: 1. CK increase to 10x upper limit of normal 2. LFTs increase to 3x upper limit of normal 3. Hospitalization due to severe statin-related AEs such as rhabdomyolysis.
Indications	All FDA-approved Indications.
Off Label Uses	

ninlaro

Products Affected

- NINLARO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of multiple myeloma -AND- previous treatment with at least 1 prior therapy -AND- used in combination with lenalidomide and dexamethasone
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

NORTHERA

Products Affected

- *droxidopa*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of neurogenic orthostatic hypotension caused by primary autonomic failure (e.g., Parkinson's disease, multiple system atrophy, or pure autonomic failure), dopamine beta-hydroxylase deficiency or non-diabetic autonomic neuropathy -AND- documentation of inadequate response, intolerance or contraindication to preferred generic alternative midodrine.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

nourianz

Products Affected

- NOURIANZ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of Parkinson's disease experiencing off episodes -AND- trial/failure, contraindication or intolerance to two (2) of the following generic products: ropinirole, pramipexole, entacapone, selegiline, or rasagiline -AND- Used as adjunct to levodopa/carbidopa.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

nubeqa

Products Affected

- NUBEQA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following if applicable to diagnosis: 1. Concomitant therapy 2. History of a bilateral orchiectomy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

nucala

Products Affected

- NUCALA SUBCUTANEOUS AUTO-INJECTOR
- NUCALA SUBCUTANEOUS RECON SOLN
- NUCALA SUBCUTANEOUS SYRINGE 100 MG/ML, 40 MG/0.4 ML

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	<p>Documentation of diagnosis of severe asthma evidenced by pretreatment FEV1 less than 80 percent predicted in adults or FEV1 less than 90 percent predicted in adolescents or FEV1 reversibility of at least 12% after albuterol administration -AND- 1.) A history of 2 or more exacerbations in the previous year or inadequate symptom control with inhaled corticosteroid in combination with 3 months of controller medication (e.g. long-acting beta2-agonist [LABA], leukotriene receptor antagonist [LTRA], theophylline), unless intolerant of or contraindication to all agents -AND- 2 or 3. 2) Greater than or equal to 150 cells/uL screening within 6 weeks of dosing. 3) Greater than or equal to 300 cells/uL within 12 months of screening. -OR- Documentation of eosinophilic granulomatosis with polyangitis (EGPA) in patients who have a history of relapsing or refractory disease and will be receiving concomitant glucocorticoid treatment with or without immunosuppressive therapy -OR- Documentation of hypereosinophilic syndrome (HES) without an identifiable non-hematologic secondary cause for greater than or equal to 6 months -AND- At least 2 HES flares (HES-related worsening of clinical symptoms or blood eosinophil counts requiring an escalation in therapy) within the past 12 months -AND- Stable on HES therapy for at least 4 weeks (chronic or episodic oral corticosteroids, immunosuppressive or cytotoxic therapy) -OR- Documentation of chronic rhinosinusitis with nasal polyps (CRSwNP) and trial/failure, contraindication, or intolerance to an intranasal corticosteroid.</p>
Age Restrictions	Deny if less than 6 years old for asthma or less than 12 years old for hypereosiniphilic syndrome
Prescriber Restrictions	
Coverage Duration	12 months

PA Criteria	Criteria Details
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

nuedexta

Products Affected

- NUEDEXTA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation supporting improvement in symptoms is required.
Indications	All FDA-approved Indications.
Off Label Uses	

nuplazid

Products Affected

- NUPLAZID

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of hallucinations and delusions associated with Parkinson's disease psychosis
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

nuvigil

Products Affected

- *armodafinil*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	<p>Diagnosis of shift work sleep disorder (SWSD) as defined by a minimum of 5 night shifts per month and the shift is 6 to 12 hours in duration occurring between 10pm and 8am. Diagnosis of narcolepsy -AND- Documentation of baseline data of excessive daytime sleepiness (EDS) via the Epworth Sleepiness Scale (ESS) or Maintenance of Wakefulness Test (MWT) -AND- Documentation of the following (1, 2, or 3): 1) Hypocretin-1 deficiency defined by (A or B), A) Cerebrospinal fluid hypocretin-1 less than 110 pg/mL. B) Cerebrospinal fluid hypocretin-1 less than 1/3 of the normal value based on laboratory reference range - OR- 2) Multiple sleep latency test (MSLT) documenting MSL less than 8 minutes and 2 sleep-onset rapid eye movement periods (SOREMP) -OR- 3) MSLT documenting MSL less than 8 minutes and 1 SOREMP and Polysomnography substantiating 1 SOREMP. Diagnosis of obstructive sleep apnea/hypopnea syndrome (OSAHS) documented by objective polysomnography as established in accordance with ICSD or DSM V criteria acceptable for all indications</p>
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, provider attestation of improvement in daytime sleepiness is required.
Indications	All FDA-approved Indications.
Off Label Uses	

ocaliva

Products Affected

- OCALIVA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of primary biliary cholangitis -AND- trial and failure, contraindication, or intolerance to ursodiol monotherapy -AND- will use concomitantly with ursodiol unless contraindicated or intolerant.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

octreotide

Products Affected

- *octreotide acetate injection solution*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For acromegaly, high pretreatment insulin-like growth factor-1 (IGF-1) based on laboratory reference range -AND- therapeutic failure or cannot be treated with surgical resection, pituitary irradiation or bromocriptine mesylate at maximally tolerated doses.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization of acromegaly, decreased or normalized IGF-1 from baseline
Indications	All FDA-approved Indications.
Off Label Uses	

odactra

Products Affected

- ODACTRA

PA Criteria	Criteria Details
Exclusion Criteria	Asthma (severe, unstable or uncontrolled), concomitant sublingual or subcutaneous immunotherapy
Required Medical Information	Documentation of allergic rhinitis due to house dust mites -AND- allergic rhinitis with or without conjunctivitis has been confirmed by skin testing for licensed house dust mite allergen extracts or in vitro testing for IgE antibodies to D. pteronyssinus or D. Farina -AND- trial and failure or intolerance to an intranasal steroid and an oral non-sedating antihistamine, intranasal antihistamine or intranasal anticholinergic agent
Age Restrictions	Deny if less than 18 years of age or greater than 65 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Member must also be prescribed an epinephrine auto injector. For reauthorization, attestation of improved allergy symptoms is required.
Indications	All FDA-approved Indications.
Off Label Uses	

odomzo

Products Affected

- ODOMZO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of locally advanced basal cell carcinoma (laBCC) that has recurred following surgery or radiation therapy or for use in patients who are not candidates for surgery or radiation therapy
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

olumiant

Products Affected

- **OLUMIANT ORAL TABLET 1 MG, 2 MG**

PA Criteria	Criteria Details
Exclusion Criteria	Part A covered for Covid-19 in hospitalized patients
Required Medical Information	Documentation of diagnosis. For moderate to severe rheumatoid arthritis an inadequate response or intolerance to at least one non-biologic DMARD (e.g., methotrexate, leflunomide). For severe alopecia areata, inadequate response or intolerance to an intralesional corticosteroid or high potency topical corticosteroid, or contraindication to all.
Age Restrictions	Deny if less than 18 years of age for Rheumatoid Arthritis and Alopecia Areata
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For rheumatoid arthritis, patients must have an adequate trial or intolerance to 2 of the following preferred products Humira, Enbrel, Actemra, Rinvoq and Xeljanz/Xeljanz XR, with at least 1 being a tumor necrosis factor blocker.
Indications	All FDA-approved Indications.
Off Label Uses	

onfi

Products Affected

- *clobazam oral suspension*
- *clobazam oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of seizures due to Lennox-Gastaut Syndrome -AND- documentation of adjunctive therapy -AND- adequate trial or intolerance of a previous antiepileptic therapy
Age Restrictions	Deny if less than 2 years old
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

ongentys

Products Affected

- **ONGENTYS**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of Parkinson's disease experiencing off episodes -AND- Used as adjunct to levodopa/carbidopa -AND- trial/failure, contraindication or intolerance to the following (1. and 2.): 1) entacapone 2) one of the following generic products: rasagiline, pramipexole, ropinirole, selegiline
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

onureg

Products Affected

- ONUREG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of acute myeloid leukemia that has achieved first complete remission or complete remission with incomplete blood count recovery following intensive induction chemotherapy -AND- Inability to complete intensive curative therapy
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Opzelura

Products Affected

- OPZELURA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of all of the following (1-4): 1) mild to moderate atopic dermatitis (AD), 2) attestation of up to 20 percent of body surface area (excluding scalp) with AD involvement, 3) trial & failure, intolerance, or contraindication to at least one topical corticosteroid -OR- documentation of facial or anogenital involvement, 4) trial & failure, intolerance, or contraindication to topical generic tacrolimus or topical generic pimecrolimus -OR- Documentation of all of the following (5-7): 5) Nonsegmental vitiligo, 6) body surface area with vitiligo involvement does not exceed 10 percent, 7) trial & failure or intolerance to at least 1 generic, formulary high or ultrahigh potency topical corticosteroid -OR- vitiligo with facial or anogenital involvement
Age Restrictions	Deny if less than 12 years of age
Prescriber Restrictions	
Coverage Duration	Initial authorization: 8 weeks for AD, 24 weeks for vitiligo. Reauthorization: 12 months
Other Criteria	For reauthorization, attestation of positive clinical response to therapy for atopic dermatitis -OR- meaningful repigmentation of affected areas for vitiligo
Indications	All FDA-approved Indications.
Off Label Uses	

oralair

Products Affected

- **ORALAIR SUBLINGUAL TABLET 300
INDX REACTIVITY**

PA Criteria	Criteria Details
Exclusion Criteria	Asthma (severe, unstable or uncontrolled), concomitant sublingual or subcutaneous immunotherapy, therapy initiation during active allergy season
Required Medical Information	Documentation of allergic rhinitis and use for Sweet Vernal, Orchard, Perennial Rye, Timothy or Kentucky Blue Grass pollens -AND- allergic rhinitis with or without conjunctivitis has been confirmed by a pollen specific positive skin test or in vitro testing for pollen-specific IgE antibodies -AND- trial and failure or intolerance to an intranasal steroid and an oral non-sedating antihistamine, intranasal antihistamine or intranasal anticholinergic agent
Age Restrictions	Deny if less than 5 years of age or greater than 65 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Member must also be prescribed an epinephrine auto injector. For reauthorization, attestation of improved allergy symptoms is required.
Indications	All FDA-approved Indications.
Off Label Uses	

orencia

Products Affected

- **ORENCIA CLICKJECT**
- **ORENCIA SUBCUTANEOUS SYRINGE**
125 MG/ML, 50 MG/0.4 ML, 87.5 MG/0.7 ML

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For moderate to severe rheumatoid arthritis, inadequate response or intolerance to at least one DMARD (e.g. methotrexate, leflunomide). For juvenile idiopathic rheumatoid arthritis, inadequate response or intolerance to at least one DMARD (e.g., methotrexate, leflunomide) -OR- requires initial biologic therapy due to involvement of high-risk joints, high disease activity or at high risk of disabling joint damage.
Age Restrictions	Deny if less than 18 years of age for Rheumatoid Arthritis, and Psoriatic Arthritis or less than 2 years of age for Juvenile Idiopathic Arthritis
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For rheumatoid arthritis, patients must have an adequate trial or intolerance to 2 of the following preferred products Humira, Enbrel, Actemra, Rinvoq and Xeljanz/Xeljanz XR. For psoriatic arthritis, patients must have an adequate trial or intolerance to 2 of the following preferred products Humira, Enbrel, Cosentyx, Xeljanz/Xeljanz XR, Otezla and Stelara. For juvenile idiopathic arthritis, patients must have an adequate trial or intolerance to 2 of the following preferred products Humira, Enbrel, Xeljanz/Xeljanz solution and Actemra.
Indications	All FDA-approved Indications.
Off Label Uses	

orgovyx

Products Affected

- **ORGOVYX**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of advanced prostate cancer -AND- the member is appropriate to receive androgen deprivation therapy by meeting one of the following (1, 2, or 3) 1. Biochemical (prostate specific antigen) or clinical relapse following local primary intervention 2. Newly diagnosed castration-sensitive metastatic disease 3. Advanced local disease
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For induction therapy dosing, doses above plan quantity limit will be approved aligned with recommended induction therapy dosing regimens per indication.
Indications	All FDA-approved Indications.
Off Label Uses	

oriahnn

Products Affected

- **ORIAHNN**

PA Criteria	Criteria Details
Exclusion Criteria	Diagnosis of severe hepatic impairment or osteoporosis.
Required Medical Information	Documentation of premenopausal woman with uterine leiomyomas - AND- Experiencing heavy menstrual bleeding -AND- Treatment with a gonadotropin-releasing hormone receptor antagonist in combination with estrogen and progestin does not exceed 24 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 18 months reauthorization
Other Criteria	For reauthorization, attestation of continued experience of heavy menstrual bleeding -AND- Attestation of decrease in menstrual blood loss -AND- Treatment with a gonadotropin-releasing hormone receptor antagonist in combination with estrogen and progestin does not exceed 24 months.
Indications	All FDA-approved Indications.
Off Label Uses	

orilissa

Products Affected

- **ORILISSA ORAL TABLET 150 MG, 200 MG**

PA Criteria	Criteria Details
Exclusion Criteria	Diagnosis of severe hepatic impairment or osteoporosis.
Required Medical Information	Documentation of female with diagnosis of endometriosis with moderate to severe pain -AND- For women of childbearing age, attestation of not pregnant -AND- Inadequate response, failure or contraindication to 2 standard of care treatments (e.g. NSAIDS, combined hormonal contraceptives, progestin, GnRH agonist, Danazol) -AND- Treatment with Orilissa does not exceed 24 months
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 18 months reauthorization
Other Criteria	For reauthorization, Orilissa is continued to be used for pain associated with endometriosis -AND- attestation of reduction in pain -AND- Total cumulative duration of therapy does not exceed 24 months.
Indications	All FDA-approved Indications.
Off Label Uses	

orkambi

Products Affected

- **ORKAMBI ORAL TABLET**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of cystic fibrosis and homozygous F508del mutation
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, documentation supporting improvement or stabilization of FEV1 compared to baseline FEV1 -or- increase in body mass index -or- decreased pulmonary exacerbations -or- improved quality of life as demonstrated by CF Questionnaire is required.
Indications	All FDA-approved Indications.
Off Label Uses	

orkambi granules

Products Affected

- **ORKAMBI ORAL GRANULES IN PACKET 100-125 MG, 150-188 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of cystic fibrosis and homozygous F508del mutation
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, documentation supporting improvement or stabilization of FEV1 compared to baseline FEV1 -or- increase in body mass index -or- decreased pulmonary exacerbations -or- improved quality of life as demonstrated by CF Questionnaire is required.
Indications	All FDA-approved Indications.
Off Label Uses	

orladeyo

Products Affected

- ORLADEYO

PA Criteria	Criteria Details
Exclusion Criteria	Member should not be on two prophylactic therapies simultaneously.
Required Medical Information	For the prophylactic treatment of attacks of hereditary angioedema (HAE) type I & II with the following (1-3): 1) Low C4 level of less than or equal to 14mg/dL or C4 below lower limit of laboratory reference range and 1 of the following (A or B). A) C1 inhibitor (C1INH) antigen level less than or equal to 19mg/dL or below lower limit of laboratory reference range. B) Normal C1INH antigen level and a low C1INH functional level below laboratory reference range. 2) Past medical history of at least 1 symptom of moderate or severe angioedema attack (e.g. airway swelling, painful facial distortion) in absence of concomitant hives. 3) Medications known to cause angioedema have been evaluated and discontinued. For the prophylactic treatment of attacks of hereditary angioedema (HAE) type III with the following (4-7): 4) Documentation of clinical laboratory performance C4, C1INH antigen, or C1INH functional level are within normal limits of laboratory reference ranges. 5) Documentation of family history of HAE or FXII mutation 6) Past medical history of at least 1 symptom of moderate or severe angioedema attack (e.g. airway swelling, painful facial distortion) in absence of concomitant hives. 7) Medications known to cause angioedema have been evaluated and discontinued.
Age Restrictions	Deny if less than 12 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

OTEZLA

Products Affected

- **OTEZLA**
- **OTEZLA STARTER ORAL TABLETS,DOSE PACK 10 MG (4)-20 MG (4)-30 MG (47)**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For plaque psoriasis, inadequate response or intolerance to one systemic therapy (e.g. methotrexate) -OR- inadequate response to phototherapy -OR- contraindication to phototherapy and systemic therapy. For oral ulcers associated with Behcet's Disease, inadequate response or intolerance to topical triamcinolone for acute flare-up of oral ulcers -AND- inadequate response or intolerance to colchicine for prevention of recurrent oral ulcers
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

otrexup

Products Affected

- OTREXUP (PF)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis AND trial/failure, intolerance, or contraindication to oral generic methotrexate tablets
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

oxbryta

Products Affected

- OXBRYTA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of sickle cell disease -AND- Failure, intolerance or contraindication to hydroxyurea
Age Restrictions	Deny if less than 4 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of improvement of sickle cell disease signs, symptoms or complications
Indications	All FDA-approved Indications.
Off Label Uses	

Oxbryta for Suspension

Products Affected

- **OXBRYTA ORAL TABLET FOR SUSPENSION**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of sickle cell disease -AND- Failure, intolerance or contraindication to hydroxyurea -AND- one of the following (1 or 2): 1) is eleven years of age or younger -OR- 2) has an inability to swallow tablets
Age Restrictions	Deny if less than 4 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of improvement of sickle cell disease signs, symptoms or complications
Indications	All FDA-approved Indications.
Off Label Uses	

oxervate

Products Affected

- OXERVATE

PA Criteria	Criteria Details
Exclusion Criteria	Treatment duration greater than 8 weeks per eye
Required Medical Information	Documentation of diagnosis -AND- affected eye (e.g. right eye, both eyes).
Age Restrictions	Deny if less than 2 years of age
Prescriber Restrictions	
Coverage Duration	8 weeks
Other Criteria	Coverage beyond 8 weeks per eye will not be approved
Indications	All FDA-approved Indications.
Off Label Uses	

palynziq

Products Affected

- PALYNZIQ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of phenylketonuria. Member meets the following criteria 1.) Baseline Phe level greater than 600 micrometers/L -AND- 2.) Failure or intolerance to existing management (i.e. Kuvan therapy) -AND- 3.) Has a prescription for epinephrine agent unless contraindicated.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of reduction in baseline pretreatment Phe levels -OR- blood Phe levels are less than or equal to 600 micrometers/L -OR- attestation that additional therapy with Palynziq is needed to allow adequate trial of maximum dose of 60mg per day for 16 weeks or member is not currently on 60mg per day dose.
Indications	All FDA-approved Indications.
Off Label Uses	

Panretin

Products Affected

- PANRETIN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of cutaneous lesions in patients with AIDS-related Kaposi Sarcoma (KS) who are not receiving systemic therapy for KS.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

pemazyre

Products Affected

- PEMAZYRE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) FGFR2 fusion or other rearrangement as detected by an FDA-approved test 2) Previous therapies tried/failed
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Pheburane

Products Affected

- PHEBURANE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- therapeutic failure or intolerance to generic sodium phenylbutyrate
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

phenoxybenzamine

Products Affected

- *phenoxybenzamine*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of excessive sweating and hypertension associated with pheochromocytoma supported by one of the following (1. or 2.): 1. Elevated metanephrines in plasma or urine. 2. Tumor evidence from CT scan or MRI
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

piqray

Products Affected

- **PIQRAY ORAL TABLET 200 MG/DAY (200 MG X 1), 250 MG/DAY (200 MG X1-50 MG X1), 300 MG/DAY (150 MG X 2)**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of HR-positive, HER2-negative advanced or metastatic breast cancer in men and postmenopausal women with disease progression on or after endocrine-based therapy -AND- Used in combination with fulvestrant - AND- PI3K mutation positive as detected by an FDA approved test.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

pomalyst

Products Affected

- POMALYST

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of multiple myeloma, and combination use with dexamethasone, and previous trial of at least 2 therapies including lenalidomide and a proteasome inhibitor, and disease progression on or within 60 days of completion of the last therapy -OR- Documentation of AIDS-related Kaposi sarcoma (KS) after failure of highly active antiretroviral therapy (HAART) or in patients with KS who are HIV negative
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Ponvory

Products Affected

- **PONVORY**
- **PONVORY 14-DAY STARTER PACK**

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use with other disease modifying agents such as interferons, Copaxone, Tysabri, Aubagio, Gilenya
Required Medical Information	Documentation of relapsing form of multiple sclerosis (e.g. relapsing-remitting, clinically isolated syndrome, or active secondary progressive disease)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	24 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

praluent

Products Affected

- PRALUENT PEN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	<p>1.HoFH supported by genetic confirmation of two mutant alleles at LDLR, APOB, PCSK9, or LDLRAP1 gene OR untreated LDL-C greater than 400mg/dL or TC greater than 500mg/dl with cutaneous or tendon xanthoma before age 10 yrs or HeFH in both parents AND The member has a current LDL-C of greater than 135 mg/dL (if 17 years of age or younger) or greater than 100mg/dL (18 years of age or older) despite use of maximally tolerated statin or statin intolerance AND The member will continue to receive concurrent lipid-lowering therapies for the treatment of HoFH. 2.HeFH supported by presence of causal mutation of FH by genetic testing OR untreated LDL-C greater than or equal to 190 mg/dL or untreated LDL-C greater than or equal to 160 mg/dL before 20 years of age with physical signs of FH (e.g. xanthomas, xanthelasma) OR diagnosis based on WHO criteria/Dutch Lipid Clinical Network criteria with score greater than 8 points, or definite on Simon Broome register, or definite on the Make Early Diagnosis to Prevent Early Deaths tool AND LDL-C greater than 100 mg/dL despite use of maximally tolerated statin or statin intolerance. 3.Hypercholesterolemia ASCVD or Primary Hyperlipidemia AND LDL-C greater than 70 mg/dL despite use of maximally tolerated statin or statin intolerance</p>
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization

PA Criteria	Criteria Details
Other Criteria	<p>Patients must have an adequate trial/failure or contraindication to the preferred product Repatha. Do not used with another PCSK9 inhibitor. For reauthorization, documentation showing an LDL-C reduction on Praluent therapy from baseline must be provided. Statin intolerance defined as follows: statin related rhabdomyolysis or skeletal muscle symptoms while receiving at least 2 separate trials of different statins which resolved upon discontinuation of statin or attestation of one of the following during any course of statin therapy: 1. CK increase to 10x upper limit of normal 2. LFTs increase to 3x upper limit of normal 3. Hospitalization due to severe statin-related AEs such as rhabdomyolysis.</p>
Indications	All FDA-approved Indications.
Off Label Uses	

prenatal vitamins

Products Affected

- **PRENATAL VITAMIN PLUS LOW IRON**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of nutritional supplementation required in a female of child-bearing potential during pre-conception, pregnancy, or lactation
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

prescription drug combo

Products Affected

- acetaminophen-codeine oral solution 120-12 mg/5 ml
- acetaminophen-codeine oral tablet
- **ALPRAZOLAM INTENSOL**
- alprazolam oral tablet 0.25 mg, 0.5 mg, 1 mg, 2 mg
- alprazolam oral tablet extended release 24 hr 0.5 mg, 1 mg, 2 mg, 3 mg
- butalbital-acetaminop-caf-cod oral capsule 50-300-40-30 mg, 50-325-40-30 mg
- **ENDOCET ORAL TABLET 10-325 MG, 5-325 MG, 7.5-325 MG**
- eszopiclone
- fentanyl transdermal patch 72 hour 100 mcg/hr, 12 mcg/hr, 25 mcg/hr, 50 mcg/hr, 75 mcg/hr
- hydrocodone-acetaminophen oral tablet 10-325 mg, 5-325 mg, 7.5-325 mg
- hydrocodone-ibuprofen oral tablet 10-200 mg, 5-200 mg
- hydromorphone oral liquid
- hydromorphone oral tablet
- methadone oral solution 10 mg/5 ml, 5 mg/5 ml
- methadone oral tablet 10 mg, 5 mg
- morphine concentrate oral solution
- morphine oral capsule, er multiphase 24 hr 120 mg, 30 mg, 45 mg, 60 mg, 75 mg, 90 mg
- morphine oral solution 10 mg/5 ml, 20 mg/5 ml (4 mg/ml)
- morphine oral tablet
- morphine oral tablet extended release 100 mg, 15 mg, 200 mg, 30 mg, 60 mg
- **NUCYNTA**
- oxycodone oral capsule
- oxycodone oral concentrate
- oxycodone oral solution
- oxycodone oral tablet 10 mg, 15 mg, 20 mg, 30 mg, 5 mg
- oxycodone-acetaminophen oral tablet 10-325 mg, 2.5-325 mg, 5-325 mg, 7.5-325 mg
- tramadol oral tablet 100 mg, 50 mg
- tramadol-acetaminophen
- triazolam
- zaleplon oral capsule 10 mg, 5 mg
- zolpidem

PA Criteria	Criteria Details
Exclusion Criteria	

PA Criteria	Criteria Details
Required Medical Information	<p>For concomitant use of an opiate agonist and substance abuse therapy, documentation that the member has an acute pain condition (e.g. acute traumatic injury) in which treatment with other agents would cause insufficient pain control or if the member requires treatment for pain related to a terminal illness. For concomitant use of an opiate agonist, benzodiazepine, and a centrally acting skeletal muscle relaxant, documentation that the member has tried/failed at least 2 other skeletal muscle relaxants (e.g. methocarbamol, metaxalone), understanding these skeletal muscle relaxants are high-risk medications in geriatric patients AND attestation of an intent to monitor and address concomitant drug-drug interaction adverse events. For concomitant use of an opiate agonist and other opiate potentiators (e.g. gabapentinoids, benzodiazepines, sedative-hypnotics) attestation of an intent to monitor and address concomitant drug-drug interaction adverse events. For long acting (e.g. extended release) opioid medications, the following apply (1-5). 1)Pain is severe enough to require daily, around-the-clock, long-term opioid treatment. 2)Patient is not opioid naive. 3)Attestation that non-opiate alternative therapies have been explored (e.g. NSAIDs). 4)Attestation that controlled substance Rx history has been reviewed in the state Prescription Drug Monitoring Program. 5)Attestation of counseling on the potential adverse effects of opioid analgesics, including the risk of misuse, abuse, and addiction.</p>
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	<p>Opiate tx for pain+subs. abuse, app. opiate x 1mo. All other combos and dx approve x 12mo.</p>
Other Criteria	<p>Opiate agonists will receive automatic approval if no recent claims for a substance abuse therapy (e.g. buprenorphine-naloxone) OR a benzodiazepine (e.g. triazolam, alprazolam) OR a benzodiazepine with a centrally acting skeletal muscle relaxant (e.g., carisoprodol) OR a gabapentinoid OR a sedative-hypnotic. Benzodiazepines (e.g. triazolam, alprazolam) will receive automatic approval if no recent claims for an opiate agonist (e.g. oxycodone, hydrocodone, oxymorphone) or an opiate agonist with a centrally acting skeletal muscle relaxant (e.g. carisoprodol). Sedative-hypnotics (e.g. zolpidem) will receive automatic approval if no recent claims for an opiate agonist. Infusible opiate agonists will be covered under Part B when administered via infusion pump.</p>
Indications	<p>All FDA-approved Indications.</p>

PA Criteria	Criteria Details
Off Label Uses	

procysbi

Products Affected

- **PROCYSBI ORAL GRANULES DEL
RELEASE IN PACKET**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- previous trial and failure, intolerance, or contraindication to Cystagon (cysteamine bitartrate immediate-release)
Age Restrictions	Deny if less than 1 year of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For granules, attestation of inability to swallow capsules or gastrostomy tube (g-tube) placement is required.
Indications	All FDA-approved Indications.
Off Label Uses	

prolia

Products Affected

- PROLIA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -And- For osteoporosis at high risk for fracture, meeting one of the following (1. thru 4.) 1) History of previous hip or vertebral fracture. 2) T-score less than or equal to -2.5. 3) T-score between -1.0 and -2.5 (i.e. osteopenia) -AND- meets FRAX calculation (A. or B.) A) 10-year risk of major osteoporotic fracture is greater than or equal to 20 percent or B) 10-year risk of hip fracture is greater than or equal to 3 percent. 4) Age 40 years or older with T-score between -1.0 and -2.5 -AND- History of glucocorticoid use for at least 3 months at a dose of 5mg per day or more of prednisone (or equivalent).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For osteoporosis and osteopenia, documentation of trial/failure or intolerance to at least one oral bisphosphonate or all are contraindicated. Covered under Part B for patients eligible for home health services when provider certifies that patient sustained bone fracture related to post-menopausal osteoporosis and is unable to learn the skills needed to self-administer the drug or is otherwise physically or mentally incapable of administering the drug or family/caregivers are unable or unwilling to administer the drug.
Indications	All FDA-approved Indications.
Off Label Uses	

provigil

Products Affected

- *modafinil*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of shift work sleep disorder (SWSD) as defined by a minimum of 5 night shifts per month with at least 3 of those nights occurring consecutively and the shift is 6 to 12 hours in duration occurring between 10pm and 8am. Diagnosis of narcolepsy -AND- Documentation of baseline data of excessive daytime sleepiness (EDS) via the Epworth Sleepiness Scale (ESS) or Maintenance of Wakefulness Test (MWT) - AND- Documentation of the following (1, 2, or 3): 1) Hypocretin-1 deficiency defined by (A or B), A) Cerebrospinal fluid hypocretin-1 less than 110 pg/mL. B)Cerebrospinal fluid hypocretin-1 less than 1/3 of the normal value based on laboratory reference range -OR- 2) Multiple sleep latency test (MSLT) documenting MSL less than 8 minutes and 2 sleep-onset rapid eye movement periods (SOREMP) -OR- 3) MSLT documenting MSL less than 8 minutes and 1 SOREMP and Polysomnography substantiating 1 SOREMP. Diagnosis of obstructive sleep apnea/hypopnea syndrome (OSAHS) documented by objective polysomnography as established in accordance with ICSD or DSM V criteria acceptable for all indications. Diagnosis of fatigue associated with Multiple Sclerosis (MS)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, provider attestation of improvement in daytime sleepiness is required.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off Label Uses	Fatigue associated with Multiple Sclerosis (MS)

pulmonary arterial hypertension

Products Affected

- **ADEMPAS**
- **ALYQ**
- *ambrisentan*
- *bosentan*
- **OPSUMIT**
- **ORENITRAM ORAL TABLET EXTENDED RELEASE 0.125 MG, 0.25 MG, 1 MG, 2.5 MG, 5 MG**
- *sildenafil (pulm.hypertension) oral suspension for reconstitution*
- *sildenafil (pulm.hypertension) oral tablet*
- *tadalafil (pulm. hypertension)*
- **TRACLEER ORAL TABLET FOR SUSPENSION**
- **UPTRAVI ORAL TABLET 1,000 MCG, 1,200 MCG, 1,400 MCG, 1,600 MCG, 200 MCG, 400 MCG, 600 MCG, 800 MCG**
- **UPTRAVI ORAL TABLETS,DOSE PACK**
- **VENTAVIS**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of pulmonary arterial hypertension, substantiated by results from right heart catheterization (RHC), defined as a mean pulmonary arterial pressure (mPAP) of greater than 20 mmHg at rest, with a pulmonary capillary wedge pressure (PWP) of less than or equal to 15 mmHg, and a PVR greater than or equal to 3 Wood units -AND- WHO Group. For bosentan in pediatric individuals, an exception to RHC will be allowed when the risk of RHC outweighs the benefit -AND- prescriber attests alternative studies have been completed (i.e. CT, MRI or specified test ruling out other causes of pulmonary hypertension). For Adempas, additional diagnosis of CTEPH as documented by right heart catheterization and V/Q scan substantiating mPAP greater than 20 mmHg at rest and PWP less than or equal to 15 mmHg and documented presence of occlusive thrombi within the pulmonary arteries will be approved.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Ventavis covered under Part B when using via nebulizer in the home setting.
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	

Purified Cortrophin Gel

Products Affected

- CORTROPHIN GEL

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	<p>Covered for the following indications: 1. Acute exacerbations of multiple sclerosis (MS) for patients receiving concurrent immunomodulator therapy (e.g., interferon beta, glatiramer acetate, dimethyl fumarate, fingolimod, teriflunomide) 2. Rheumatic disorders for patients receiving maintenance therapy with at least one NSAID, DMARD (e.g. leflunomide) or biologic (e.g. adalimumab) 3. Collagen diseases for members receiving maintenance therapy with at least one antimalarial (e.g. hydroxychloroquine) or immunosuppressant (e.g. azathioprine) 4. Dermatologic diseases, if using for severe psoriasis, then the member is concurrently receiving maintenance therapy with phototherapy, systemic therapy (e.g., methotrexate), or a biologic (e.g. adalimumab). 5. Allergic states (i.e. serum sickness and transfusion reaction due to serum protein reaction), if using for atopic dermatitis, then the member is concurrently receiving maintenance therapy with a topical corticosteroid, topical calcineurin inhibitor, topical PDE-4 inhibitor, or dupilumab. 6. Ophthalmic diseases 7. Respiratory diseases 8. Gout and unable to take first-line therapies. 9. Pediatric acquired epileptic aphasia. 10. Proteinuria in nephrotic syndrome and trial/failure or contraindication to two therapies from any of the following different classes: corticosteroids (e.g., cortisone or dexamethasone), calcineurin inhibitors (e.g., cyclosporine or tacrolimus, per DRUGDEX). For covered indications 1 through 9, limited/unsatisfactory response or intolerance (i.e. severe anaphylaxis) to two corticosteroids (i.e. IV methylprednisolone, IV dexamethasone, or high dose oral steroids) must be documented.</p>
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 month

PA Criteria	Criteria Details
Other Criteria	For reauthorization, the following (1. 2. and 3.) must be met. 1) Prescriber attestation that the member cannot use corticosteroids (e.g. IV methylprednisolone, high dose oral corticosteroids) due to unsatisfactory response, intolerance (e.g. severe anaphylaxis) or experienced a severe adverse event to corticosteroids (e.g. psychosis). 2) If the reauthorization is for the treatment of multiple sclerosis, a rheumatic disorder, dermatologic disease, or nephrotic syndrome, the prescriber attests that H.P. Acthar is being used for a new acute exacerbation and not on a routine basis to prevent an exacerbation as supported by Compendia. 3) If the reauthorization is for treatment of multiple sclerosis, a rheumatic disorder, or collagen disease, the member continues to receive maintenance therapy.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off Label Uses	Epileptic Aphasia

Pyrukynd

Products Affected

- **PYRUKYND ORAL TABLET 20 MG, 5 MG, 5 MG (4-WEEK PACK), 50 MG**
- **PYRUKYND ORAL TABLETS,DOSE PACK**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of hemolytic anemia with pyruvate kinase deficiency, as supported by one of the following (1-2): 1) mutations in the PKLR gene, 2) reduced activity of the pyruvate kinase enzyme.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	24 weeks initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, attestation of increase in hemoglobin level from baseline -OR- decrease in transfusion burden from baseline.
Indications	All FDA-approved Indications.
Off Label Uses	

qinlock

Products Affected

- QINLOCK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of advanced gastrointestinal stromal tumor -AND- Prior treatment with imatinib and 2 other kinase inhibitors.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

quinine

Products Affected

- *quinine sulfate*

PA Criteria	Criteria Details
Exclusion Criteria	Treatment or prevention of leg cramps
Required Medical Information	Documentation of diagnosis
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	10 days
Other Criteria	Doses for duration greater than 10 days will not be approved
Indications	All FDA-approved Indications.
Off Label Uses	

Radicava

Products Affected

- **RADICAVA ORS STARTER KIT SUSP**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of amyotrophic lateral sclerosis (ALS)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

rasuvo

Products Affected

- **RASUVO (PF)**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis AND trial/failure, intolerance, or contraindication to oral generic methotrexate tablets
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

ravicti

Products Affected

- RAVICTI

PA Criteria	Criteria Details
Exclusion Criteria	Urea cycle disorders due to N-acetylglutamate synthetase deficiency, Treatment of acute hyperammonemia in urea cycle disorders
Required Medical Information	Documentation of chronic management of a urea cycle disorders (UCDs)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Recorlev

Products Affected

- **RECORLEV**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of endogenous Cushing's syndrome AND patient is not a candidate for pituitary surgery or surgery has not been curative
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of mean urine free cortisol (mUFC) less than starting baseline value.
Indications	All FDA-approved Indications.
Off Label Uses	

reditrex

Products Affected

- REDITREX (PF)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis AND trial/failure, intolerance, or contraindication to oral generic methotrexate tablets
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

regranex

Products Affected

- **REGRANEX**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of lower-extremity diabetic neuropathic ulcer(s) that extends into the subcutaneous tissue or beyond and have an adequate blood supply -AND- being used as an adjunct to standard ulcer care practices (e.g. sharp debridement, non-weight bearing regimen, infection control) -AND- attestation of a wound care plan.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	20 weeks
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

relistor

Products Affected

- **RELISTOR ORAL**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of opioid induced constipation (OIC) due to any of the following (1-2): 1) chronic non-cancer pain, or 2) chronic pain related to prior cancer or its treatment who do not require frequent (e.g. weekly) opioid dosage escalation -AND- documentation of opioid medication use for at least one month -AND- trial and failure, contraindication, or intolerance to 2 of the following (3-5): 3) Laxatives, 4) lubiprostone, 5) Movantik
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

relistor SC

Products Affected

- **RELISTOR SUBCUTANEOUS SOLUTION**
- **RELISTOR SUBCUTANEOUS SYRINGE 12 MG/0.6 ML, 8 MG/0.4 ML**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of opioid induced constipation (OIC) due to any of the following (1-3): 1) chronic non-cancer pain, 2) advanced illness or active cancer in palliative care or 3) chronic pain related to prior cancer or its treatments who do not require frequent (e.g., weekly) opioid dosage escalation -AND- documentation of opioid medication use for at least one month -AND- trial and failure, contraindication, or intolerance to 2 of the following (4 to 6): 4) Laxatives 5) lubiprostone 6) Movantik.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

repatha

Products Affected

- **REPATHA PUSHTRONEX**
- **REPATHA SURECLICK**
- **REPATHA SYRINGE**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	<p>1.HoFH supported by genetic confirmation of two mutant alleles at LDLR, APOB, PCSK9, or LDLRAP1 gene OR untreated LDL-C greater than 400mg/dL or TC greater than 500mg/dl with cutaneous or tendon xanthoma before age 10 yrs or HeFH in both parents AND The member has a current LDL-C of greater than 135 mg/dL (if 17 years of age or younger) or greater than 100mg/dL (18 years of age or older) despite use of maximally tolerated statin or statin intolerance AND The member will continue to receive concurrent lipid-lowering therapies for the treatment of HoFH. 2.HeFH supported by presence of causal mutation of FH by genetic testing OR untreated LDL-C greater than or equal to 190 mg/dL or untreated LDL-C greater than or equal to 160 mg/dL before 20 years of age with physical signs of FH (e.g. xanthomas, xanthelasma) OR diagnosis based on WHO criteria/Dutch Lipid Clinical Network criteria with score greater than 8 points, or definite on Simon Broome register, or definite on the Make Early Diagnosis to Prevent Early Deaths tool, AND LDL-C greater than 100 mg/dL if 18 and older or LDL-C greater than 130 if 17 and younger despite use of maximally tolerated statin or statin intolerance. If 17 and younger will continue to receive concurrent lipid-lowering therapies. 3.Hypercholesterolemia ASCVD or Primary Hyperlipidemia AND LDL-C greater than 70 mg/dL despite use of maximally tolerated statin or statin intolerance</p>
Age Restrictions	Deny if less than 18 years of age for ASCVD and Primary Hyperlipidemia, or less than 10 years of age for HoFH and HeFH.
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization

PA Criteria	Criteria Details
Other Criteria	Do not used with another PCSK9 inhibitor. For reauthorization, documentation showing an LDL-C reduction on Repatha therapy from baseline must be provided. Statin intolerance defined as follows: statin related rhabdomyolysis or skeletal muscle symptoms while receiving at least 2 separate trials of different statins which resolved upon discontinuation of statin or attestation of one of the following during any course of statin therapy: 1. CK increase to 10x upper limit of normal 2. LFTs increase to 3x upper limit of normal 3. Hospitalization due to severe statin-related AEs such as rhabdomyolysis.
Indications	All FDA-approved Indications.
Off Label Uses	

retevmo

Products Affected

- **RETEVMO ORAL CAPSULE 40 MG, 80 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis 1) RET fusion status 2) Radioactive iodine-refractory (if radioactive iodine is appropriate)
Age Restrictions	Deny if less than 18 years of age for NSCLC or less than 12 years of age for thyroid cancer
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

revlimid

Products Affected

- *lenalidomide*

PA Criteria	Criteria Details
Exclusion Criteria	Documentation of chronic lymphocytic leukemia outside of a controlled clinical trial
Required Medical Information	Diagnosis of multiple myeloma in combination with dexamethasone -OR- diagnosis of multiple myeloma, as maintenance following autologous hematopoietic stem cell transplant (auto-HSCT) -OR- diagnosis of transfusion-dependent anemia due to low- or intermediate-1-risk myelodysplastic syndromes (MDS) associated with a deletion 5q abnormality with or without additional cytogenetic abnormalities -OR- diagnosis of mantle cell lymphoma (MCL) in which disease has relapsed or progressed after two prior therapies, one of which included bortezomib -OR- diagnosis of follicular lymphoma in combination with a rituximab product -OR- diagnosis of marginal zone lymphoma in combination with a rituximab product after previous treatment.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All Medically-accepted Indications.
Off Label Uses	

rinvoq

Products Affected

- **RINVOQ ORAL TABLET EXTENDED
RELEASE 24 HR 15 MG, 30 MG, 45 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of moderate to severe Rheumatoid Arthritis, moderate to severe Refractory Atopic Dermatitis, Ankylosing Spondylitis, Psoriatic Arthritis, or Ulcerative Colitis. For moderate to severe rheumatoid arthritis, an inadequate response or intolerance to at least one non-biologic DMARD (e.g., leflunomide, methotrexate). For moderate to severe refractory atopic dermatitis whose disease is not adequately controlled with other systemic drug products, documentation of all of the following (1-2): 1) trial & failure, intolerance, or contraindication to at least one topical corticosteroid -OR- atopic dermatitis of the face or anogenital involvement -OR- the member has severe atopic dermatitis evidenced by the extent of the member's body surface area affected which makes topical therapy impractical to apply -OR- severe atopic dermatitis evidenced by severely damaged skin, 2) trial & failure, intolerance, or contraindication to topical tacrolimus or topical pimecrolimus -OR- the member has severe atopic dermatitis evidenced by the extent of the member's body surface area affected which makes topical therapy impractical to apply -OR- severe atopic dermatitis evidenced by severely damaged skin. For ankylosing spondylitis, inadequate response or intolerance to one nonsteroidal anti-inflammatory drug (NSAID).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For diagnoses in which tumor necrosis factor (TNF) blockers are also indicated (e.g., Rheumatoid Arthritis, Psoriatic Arthritis), the member has experienced therapeutic failure or intolerance to at least 1 TNF blocker.
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	

rozlytrek

Products Affected

- **ROZLYTREK ORAL CAPSULE 100 MG, 200 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For metastatic non-small cell lung cancer, the tumor status is ROS1-positive. For solid tumors with NTRK gene fusion without a known acquired resistance mutation, the tumors are metastatic or surgical resection is likely to result in severe morbidity - AND- There are no satisfactory alternative treatments or the tumors have progressed following treatment.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

rubraca

Products Affected

- RUBRACA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following. 1) BRCA mutations, if applicable to diagnosis. 2) Alternatives tried/failed. 3) Concomitant therapy, if applicable to diagnosis.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

ruconest

Products Affected

- RUCONEST

PA Criteria	Criteria Details
Exclusion Criteria	Member should not be on two acute therapies simultaneously and acute therapy should not be used as prophylactic therapy
Required Medical Information	<p>For the treatment of attacks of hereditary angioedema (HAE) type I & II with the following (1-4): 1) Low C4 level of less than or equal to 14mg/dL or C4 below lower limit of laboratory reference range and 1 of the following (A or B). A) C1 inhibitor (C1INH) antigen level less than or equal to 19mg/dL or below lower limit of laboratory reference range. B) Normal C1INH antigen level and a low C1INH functional level below laboratory reference range. 2) Past medical history of at least 1 symptom of moderate or severe angioedema attack (e.g. airway swelling, painful facial distortion) in absence of concomitant hives. 3) Medications known to cause angioedema have been evaluated and discontinued. 4) Documentation of member's weight. For the treatment of attacks of hereditary angioedema (HAE) type III with the following (5-9): 5) Documentation of clinical laboratory performance C4, C1INH antigen, or C1INH functional level are within normal limits of laboratory reference ranges. 6) Documentation of family history of HAE or FXII mutation 7) Past medical history of at least 1 symptom of moderate or severe angioedema attack (e.g. airway swelling, painful facial distortion) in absence of concomitant hives. 8) Medications known to cause angioedema have been evaluated and discontinued. 9) Documentation of member's weight.</p>
Age Restrictions	Deny if less than 13 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For 18 years of age or older, therapeutic failure, intolerance or contraindication to icatibant.
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	

rydapt

Products Affected

- RYDAPT

PA Criteria	Criteria Details
Exclusion Criteria	Use as single agent induction therapy for AML
Required Medical Information	Documentation of diagnosis -AND- All of the following, if applicable to diagnosis: 1) FLT3 mutations 2) Concomitant therapy
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

sabril

Products Affected

- *vigabatrin*
- **VIGADRONE**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of refractory complex partial seizures -AND- documentation of adjunctive therapy -AND- an adequate trial or intolerance to at least two alternative treatments (e.g. carbamazepine, lamotrigine, levetiracetam, oxcarbazepine, tiagabine) -OR- documentation of use as monotherapy in treatment of infantile spasms
Age Restrictions	Deny if less than 2 years of age in treatment of refractory complex partial seizures -OR- if less than 1 month old and greater than 2 years of age in treatment of infantile spasms
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

samsca

Products Affected

- *tolvaptan*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of symptomatic hypervolemic or euvolemic hyponatremia evidenced by (1. or 2.): 1.) Serum Na less than 125 mEq/L -OR- 2.) Serum NA less than 135mEq/L with symptoms (e.g. nausea, malaise, lethargy, headache, seizures)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 month
Other Criteria	Doses must be initiated in the hospital setting to closely monitor serum sodium. Treatment should be limited to 30 days to minimize risk of liver injury. For reauthorization, treatment is for a new episode of a clinically significant euvolemic or hypervolemic hyponatremia -AND- on of the following (1. or 2.) 1.) Serum Na less than 125 mEq/L -OR- 2.) Serum NA less than 135mEq/L with symptoms (e.g. nausea, malaise, lethargy, headache, seizures)
Indications	All FDA-approved Indications.
Off Label Uses	

savella

Products Affected

- SAVELLA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation to support a diagnosis of fibromyalgia and trial/failure or intolerance to duloxetine
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Scemblix

Products Affected

- **SCSEMBLIX ORAL TABLET 20 MG, 40 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) T3151 mutation status 2) Philadelphia chromosome status 3) Alternatives tried/failed
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

signifor

Products Affected

- SIGNIFOR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of Cushing's disease AND patient is not a candidate for pituitary surgery or surgery has not been curative
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of decrease in urinary free cortisol levels from baseline
Indications	All FDA-approved Indications.
Off Label Uses	

siliq

Products Affected

- SILIQ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For moderate to severe plaque psoriasis, inadequate response or intolerance to one systemic therapy (e.g. methotrexate, cyclosporine) -OR- inadequate response to phototherapy. If not a candidate for phototherapy: treatment with systemic therapy has been ineffective, not tolerated, or is contraindicated.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	4 months initial authorization, 12 months reauthorization
Other Criteria	If on Siliq therapy greater than or equal to 4 months or reauthorization, provider must attest to improvement in the physician's global assessment score, psoriasis area severity index score, or a decrease in the affected body surface area of psoriatic plaque lesions. For psoriasis, patients must have an adequate trial or intolerance to 2 preferred products Humira, Cosentyx, Otezla, Stelara, Enbrel, and Skyrizi. For psoriasis induction therapy, doses above plan quantity limit will be approved aligned with recommended induction therapy dosing regimen.
Indications	All FDA-approved Indications.
Off Label Uses	

simponi

Products Affected

- **SIMPONI SUBCUTANEOUS PEN INJECTOR 100 MG/ML, 50 MG/0.5 ML**
- **SIMPONI SUBCUTANEOUS SYRINGE 100 MG/ML, 50 MG/0.5 ML**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. Simponi 50mg: For moderate to severe rheumatoid arthritis, inadequate response or intolerance to at least one DMARD (e.g. leflunomide) and Simponi will be used in combination with methotrexate. For ankylosing spondylitis, inadequate response or intolerance to one nonsteroidal anti-inflammatory drug (NSAID). Diagnosis of psoriatic arthritis Simponi 100mg: Diagnosis of moderate to severe ulcerative colitis.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For Rheumatoid arthritis, patients must have an adequate trial or intolerance to 2 of the following preferred products Humira, Enbrel, Actemra, Xeljanz/Xeljanz XR and Rinvoq. For psoriatic arthritis, patients must have an adequate trial or intolerance to 2 of the following preferred products Humira, Enbrel, Cosentyx, Xeljanz/Xeljanz XR, Otezla, and Stelara. For ankylosing spondylitis, patients must have an adequate trial or intolerance to 2 of the following preferred products Humira, Enbrel and Cosentyx. For ulcerative colitis, patients must have an adequate trial or intolerance to the preferred products Humira, Stelara and Xeljanz/Xeljanz XR. For ulcerative colitis indication therapy, doses above plan quantity limit will be approved aligned with recommended induction therapy dosing regimen.
Indications	All FDA-approved Indications.
Off Label Uses	

skyrizi

Products Affected

- SKYRIZI SUBCUTANEOUS PEN INJECTOR
- SKYRIZI SUBCUTANEOUS SYRINGE 150 MG/ML
- SKYRIZI SUBCUTANEOUS SYRINGE KIT
- SKYRIZI SUBCUTANEOUS WEARABLE INJECTOR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For moderate to severe plaque psoriasis, inadequate response or intolerance to one systemic therapy (e.g. methotrexate, cyclosporine) -OR- inadequate response to phototherapy. If not a candidate for phototherapy: treatment with systemic therapy has been ineffective, not tolerated, or is contraindicated. For Crohn's disease, attestation of clinical response or remission following IV administration of Skyrizi within 3 months of initiating therapy with Skyrizi SC.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For induction therapy, doses above plan quantity limit will be approved when aligned with recommended induction therapy dosing regimen.
Indications	All FDA-approved Indications.
Off Label Uses	

solaraze

Products Affected

- *diclofenac sodium topical gel 3 %*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- trial and failure, intolerance, or contraindication to topical fluorouracil
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	3 months
Other Criteria	For reauthorization, attestation of 30 day washout period since optimal therapeutic effect may not be evident until 30 days following cessation of therapy AND attestation of previous response to diclofenac sodium 3% topical gel therapy
Indications	All FDA-approved Indications.
Off Label Uses	

somavert

Products Affected

- SOMAVERT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For acromegaly, high pretreatment insulin-like growth factor-1 (IGF-1) based on laboratory reference range -AND- inadequate or partial response to surgery or radiotherapy or not a candidate for surgery or radiotherapy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization of acromegaly, decreased or normalized IGF-1 from baseline
Indications	All FDA-approved Indications.
Off Label Uses	

sovaldi

Products Affected

- SOVALDI ORAL PELLETS IN PACKET
- SOVALDI ORAL TABLET 400 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Criteria will be applied consistent with current AASLD/IDSA guidance - AND- the member has a contraindication to or is otherwise not a candidate for all regimens recommended by the AASLD/IDSA guidelines containing the following agents: ledipasvir/sofosbuvir, sofosbuvir/velpatasvir, glecaprevir/pibrentasvir.
Age Restrictions	Deny if less than 3 years of age
Prescriber Restrictions	
Coverage Duration	Criteria/duration applied consistent with current AASLD-IDSA guidance
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

sprycel

Products Affected

- SPRYCEL

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- All of the following, if applicable to diagnosis: 1) Philadelphia Chromosome status (e.g. positive) 2) Alternatives tried/failed 3) Concomitant therapy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

stelara

Products Affected

- **STELARA SUBCUTANEOUS SOLUTION**
- **STELARA SUBCUTANEOUS SYRINGE 45 MG/0.5 ML, 90 MG/ML**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- documentation of member weight and prescribed dose. For moderate to severe plaque psoriasis, inadequate response or intolerance to one systemic therapy (e.g. methotrexate, cyclosporine) -OR- inadequate response to phototherapy. If not a candidate for phototherapy: treatment with systemic therapy has been ineffective, not tolerated, or is contraindicated. For Crohn's Disease, inadequate response or intolerance to two immunosuppressants (e.g. corticosteroids, azathioprine) -OR- intolerance to a TNF inhibitor (e.g. Humira) -OR- Contraindication to a TNF inhibitor due to demyelinating disease or heart failure -OR- attestation of clinical response or remission following IV administration of Stelara within 2 months of initiating therapy with Stelara SC. For Ulcerative colitis, attestation of clinical response or remission following IV administration of Stelara within 2 months of initiating therapy with Stelara SC
Age Restrictions	Deny if less than 18 years of age for Crohn's Disease and Ulcerative Colitis or less than 6 years of age for Plaque Psoriasis and Psoriatic Arthritis
Prescriber Restrictions	
Coverage Duration	12 months

PA Criteria	Criteria Details
Other Criteria	<p>Must follow recommended dosing guidelines based upon weight.</p> <p>Psoriasis: For patients weighing less than 100 kilograms (220 pounds), 45 mg dosing will be approved. For patients weighing more than 100 kilograms (220 pounds), 90 mg dosing will be approved. Psoriatic Arthritis: 45 mg dosing will be approved. For patients with co-existent moderate to severe plaque psoriasis weighing greater than 100 kilograms (220 pounds), 90 mg dosing will be approved. Induction therapy, doses above plan quantity limit will be approved aligned with recommended induction therapy dosing regimen.</p>
Indications	All FDA-approved Indications.
Off Label Uses	

stivarga

Products Affected

- **STIVARGA**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of metastatic colorectal cancer and trial of a fluoropyrimidine-, oxaliplatin-, and irinotecan-containing chemotherapy, AND an anti-VEGF therapy AND if RAS wild-type, an anti-EGFR therapy -OR- documentation of locally advanced, unresectable or metastatic gastrointestinal stromal tumor (GIST) after treatment with both imatinib and sunitinib -OR- documentation of hepatocellular cancer AND previous treatment with sorafenib
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

sunosi

Products Affected

- **SUNOSI**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of narcolepsy -AND- Documentation of baseline data of excessive daytime sleepiness (EDS) via the Epworth Sleepiness Scale (ESS) or Maintenance of Wakefulness Test (MWT) -AND- Documentation of the following (1, 2, or 3): 1) Hypocretin-1 deficiency defined by (A or B), A) Cerebrospinal fluid hypocretin-1 less than 110 pg/mL. B) Cerebrospinal fluid hypocretin-1 less than 1/3 of the normal value based on laboratory reference range -OR- 2) Multiple sleep latency test (MSLT) documenting MSL less than 8 minutes and 2 sleep-onset rapid eye movement periods (SOREMP) -OR- 3) MSLT documenting MSL less than 8 minutes and 1 SOREMP and Polysomnography substantiating 1 SOREMP. Diagnosis of obstructive sleep apnea/hypopnea syndrome (OSAHS) documented by objective polysomnography as established in accordance with ICSD or DSM V criteria acceptable for all indications
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For narcolepsy and OSAHS, documentation of trial and failure, contraindication or intolerance to modafinil and armodafinil. For reauthorization, provider attestation of improvement in daytime sleepiness is required.
Indications	All FDA-approved Indications.
Off Label Uses	

sutent

Products Affected

- *sunitinib*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) disease progression on or intolerance to imatinib mesylate 2) high risk of recurrent RCC following nephrectomy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

symdeko

Products Affected

- **SYMDEKO**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of cystic fibrosis (CF) in patients who have either the homozygous F508del mutation or another mutation in the cystic fibrosis transmembrane conductance regulator (CFTR gene) that is responsive to tezacaftor/ivacaftor based on clinical and/or in vitro assay (e.g. E56K, R117C, A455E)
Age Restrictions	Deny if less than 6 years of age
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, documentation supporting improvement or stabilization of FEV1 compared to baseline FEV1 -or- increase in body mass index -or- decreased pulmonary exacerbations -or- improved quality of life as demonstrated by CF Questionnaire is required.
Indications	All FDA-approved Indications.
Off Label Uses	

sympazan

Products Affected

- SYMPAZAN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of seizures due to Lennox-Gastaut Syndrome -AND- documentation of adjunctive therapy -AND- adequate trial or intolerance of a previous antiepileptic therapy -AND- unable to tolerate generic clobazam
Age Restrictions	Deny if less than 2 years old
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

symproic

Products Affected

- SYMPROIC

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of opioid induced constipation (OIC) due to any of the following (1-2): 1) chronic non-cancer pain, or 2) chronic pain related to prior cancer or its treatments who do not require frequent (e.g., weekly) opioid dosage escalation -AND- documentation of opioid medication use for at least one month -AND- trial and failure, contraindication, or intolerance to at least 2 of the following (3-5): 3) Laxatives, 4) lubiprostone, 5) Movantik
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

tabrecta

Products Affected

- TABRECTA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of metastatic non-small cell lung cancer with MET exon 14 skipping mutation as detected by an FDA approved test.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

tagrisso

Products Affected

- TAGRISSO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following. 1) EGFR mutations, if applicable to diagnosis. 2) Alternatives tried/failed. 3) Concomitant therapy, if applicable to diagnosis.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

takhzyro

Products Affected

- TAKHZYRO

PA Criteria	Criteria Details
Exclusion Criteria	Member should not be on two prophylactic therapies simultaneously.
Required Medical Information	For the prophylactic treatment of attacks of hereditary angioedema (HAE) type I & II with the following (1-3): 1) Low C4 level of less than or equal to 14mg/dL or C4 below lower limit of laboratory reference range and 1 of the following (A or B). A) C1 inhibitor (C1INH) antigen level less than or equal to 19mg/dL or below lower limit of laboratory reference range. B) Normal C1INH antigen level and a low C1INH functional level below laboratory reference range. 2) Past medical history of at least 1 symptom of moderate or severe angioedema attack (e.g. airway swelling, painful facial distortion) in absence of concomitant hives. 3) Medications known to cause angioedema have been evaluated and discontinued. For the prophylactic treatment of attacks of hereditary angioedema (HAE) type III with the following (4-7): 4) Documentation of clinical laboratory performance C4, C1INH antigen, or C1INH functional level are within normal limits of laboratory reference ranges. 5) Documentation of family history of HAE or FXII mutation 6) Past medical history of at least 1 symptom of moderate or severe angioedema attack (e.g. airway swelling, painful facial distortion) in absence of concomitant hives. 7) Medications known to cause angioedema have been evaluated and discontinued.
Age Restrictions	Deny if less than 12 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

taltz

Products Affected

- TALTZ AUTOINJECTOR
- TALTZ SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For moderate to severe plaque psoriasis, inadequate response or intolerance to one systemic therapy (e.g. methotrexate, cyclosporine) -OR- inadequate response to phototherapy. If not a candidate for phototherapy: treatment with systemic therapy has been ineffective, not tolerated, or is contraindicated. For ankylosing spondylitis, inadequate response or intolerance to one nonsteroidal anti-inflammatory drug (NSAID). For non-radiographic axial spondyloarthritis, inadequate response or intolerance to 2 NSAIDs.
Age Restrictions	Deny if less than 18 years of age for Psoriatic Arthritis, Ankylosing Spondylitis and non-radiographic axial spondyloarthritis or less than 6 years of age for Plaque Psoriasis
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For psoriatic arthritis, patients must have an adequate trial or intolerance to 2 of the following preferred products Humira, Enbrel, Cosentyx, Xeljanz/Xeljanz XR, Otezla, and Stelara. For plaque psoriasis patients must have an adequate trial or intolerance to 2 of the following preferred products Humira, Cosentyx, Otezla, Stelara, Enbrel and Skyrizi. For ankylosing spondylitis, patients must have an adequate trial or intolerance to 2 of the following preferred products Humira, Enbrel and Cosentyx. For induction therapy, doses above plan quantity limit will be approved when aligned with recommended induction therapy dosing regimen.
Indications	All FDA-approved Indications.
Off Label Uses	

talzenna

Products Affected

- **TALZENNA ORAL CAPSULE 0.25 MG, 0.5 MG, 0.75 MG, 1 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of deleterious or suspected deleterious gBRCAm, HER2-negative locally advanced or metastatic breast cancer
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

targretin

Products Affected

- *bexarotene oral*
- *bexarotene topical*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of cutaneous manifestations of cutaneous T-cell lymphoma in patients who are refractory to at least one prior systemic therapy.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

tasigna

Products Affected

- TASIGNA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- All of the following, if applicable to diagnosis: 1) Philadelphia Chromosome status (e.g. positive) 2) Alternatives tried/failed
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

tavalisse

Products Affected

- TAVALISSE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For diagnosis of ITP, the following criteria apply (1 and 2): 1) trial, intolerance, or inadequate response to a corticosteroid, immunoglobulin, or splenectomy. 2) One of the following (A or B): A) Platelet count less than or equal to $50 \times 10^9/L$ and has significant mucous member bleeding or at least one risk factor for bleeding (e.g. hypertension, peptic ulcer disease). B) Platelet count of less than or equal to $30 \times 10^9/L$.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Tavneos

Products Affected

- TAVNEOS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of severe active granulomatosis with polyangiitis or severe active microscopic polyangiitis -AND- Prescriber attests to positive test for anti-PR3 or positive test for anti-MPO -AND- Member will continue to receive concomitant standard of care treatment with systemic glucocorticoids (e.g. prednisone) and immunosuppressives (e.g. mycophenolate mofetil, azathioprine).
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, attestation of therapeutic response defined by disease stability or disease improvement -AND- Member will continue to receive concomitant standard of care treatment with glucocorticoids (e.g. prednisone) and immunosuppressives (e.g. mycophenolate mofetil, azathioprine).
Indications	All FDA-approved Indications.
Off Label Uses	

tazorac

Products Affected

- *tazarotene topical cream*
- *tazarotene topical gel*
- **TAZORAC TOPICAL CREAM 0.05 %**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of 1 of the following (A or B). A) Documentation of plaque psoriasis -AND- trial and failure or intolerance to at least one topical corticosteroid (e.g. fluocinonide, mometasone, triamcinolone, betamethasone). B) Documentation of acne vulgaris -AND- trial and failure or intolerance of at least two topical acne medications (e.g. adapalene, clindamycin, sulfacetamide, erythromycin) one of which must be generic topical tretinoin
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

tazverik

Products Affected

- TAZVERIK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of locally advanced or metastatic epithelioid sarcoma - AND- Disease is not eligible for complete resection. Documentation of relapsed or refractory follicular lymphoma -AND-Tumors are EZH2 mutation positive, as detected by FDA approved test, in a member that has received at least 2 prior systemic therapies -OR- Prescriber attests there are no satisfactory alternative treatment options.
Age Restrictions	Deny if less than 16 years of age for epithelioid sarcoma or deny if less than 18 years of age for follicular lymphoma
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

tecfidera

Products Affected

- *dimethyl fumarate oral capsule, delayed release(dr/ec) 120 mg, 120 mg (14)- 240 mg (46), 240 mg*

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use with other disease modifying agents such as interferons, Copaxone, Tysabri, Aubagio, Gilenya
Required Medical Information	Documentation of relapsing form of multiple sclerosis (e.g. relapsing-remitting, clinically isolated syndrome, or active secondary progressive disease)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	5 years
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

tegsedi

Products Affected

- **TEGSEDI**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of polyneuropathy associated with hereditary TTR (hATTR) amyloidosis with mutation in TTR gene confirmed by genetic testing -AND- Neurologic examination shows clinical signs and symptoms of the disease (e.g. peripheral/autonomic neuropathy, motor disability, carpal tunnel, etc.) -AND- Not simultaneously utilizing other gene targeted therapy for polyneuropathy of hATTR
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of positive clinical response
Indications	All FDA-approved Indications.
Off Label Uses	

tepmetko

Products Affected

- **TEPMETKO**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of metastatic non-small cell lung cancer with a MET exon 14 skipping alteration
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

testosterone (androgens)

Products Affected

- **ANDRODERM**
- **AVEED**
- **DEPO-TESTOSTERONE INTRAMUSCULAR OIL 100 MG/ML**
- *testosterone cypionate intramuscular oil 100 mg/ml, 200 mg/ml, 200 mg/ml (1 ml)*
- *testosterone enanthate*
- *testosterone transdermal gel in metered-dose pump 20.25 mg/1.25 gram (1.62 %)*
- *testosterone transdermal gel in packet 1.62 % (20.25 mg/1.25 gram), 1.62 % (40.5 mg/2.5 gram)*
- *testosterone transdermal solution in metered pump w/app*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of primary or secondary hypogonadism in males with testicular failure due to cryptorchidism, bilateral torsions, orchitis, vanishing testis syndrome, orchidectomy, Klinefelter's syndrome, chemotherapy, radiation or toxic damage -OR- documentation of primary or secondary hypogonadism in males with multiple symptoms of hypogonadism including at least one of the following specific symptoms: height loss due to vertebral fractures, low trauma fractures, low bone density, incomplete or delayed sexual development, breast discomfort, loss of axillar and/or pubic body hair, hot flushes -OR- documentation of HIV infection in men with weight loss -OR- documentation of chronic steroid treatment in men. In all previously noted indications, members must also have documented low total testosterone level below the normal range for the laboratory -OR- a total testosterone level near the lower limit of the normal range with a low free testosterone level which is less than normal based upon the laboratory reference range -OR- the member is not producing any testosterone. Additional approvable indications include female patients with metastatic breast cancer (testosterone enanthate only), primary or secondary hypogonadism in males with testicular failure due to double orchidectomy, and delayed puberty in males (testosterone enanthate only).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off Label Uses	HIV Wasting

thalomid

Products Affected

- **THALOMID ORAL CAPSULE 100 MG, 150 MG, 200 MG, 50 MG**

PA Criteria	Criteria Details
Exclusion Criteria	Use as monotherapy for ENL treatment in the presence of moderate to severe neuritis
Required Medical Information	Documentation of multiple myeloma in combination with dexamethasone -OR- documentation for use in the treatment of cutaneous manifestations of moderate to severe erythema nodosum leprosum (ENL) -OR- documentation of therapy for prevention and suppression of the cutaneous manifestations of ENL recurrence
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

thiola

Products Affected

- **THIOLA EC**
- *tiopronin*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following criteria must be met (1-3) 1) member is at least 9 years of age or weighs at least 20kg, 2) Confirmation of cystinuria by at least one 24-hour urine collection with measurement of urinary cysteine levels greater than 400 mg/day, 3) Attestation of failure of urine alkalization with potassium citrate (to achieve pH of 7.0).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, attestation of urine cystine concentration less than 250 mg/L-OR- decrease in production of cystine stones is required.
Indications	All FDA-approved Indications.
Off Label Uses	

thrombopoiesis stimulating agents

Products Affected

- **PROMACTA ORAL POWDER IN PACKET 12.5 MG, 25 MG**
- **PROMACTA ORAL TABLET 12.5 MG, 25 MG, 50 MG, 75 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis of chronic immune idiopathic thrombocytopenia purpura and trial and failure of corticosteroid or immunoglobulin therapy or splenectomy -OR- documentation of thrombocytopenia in patients with chronic hepatitis C to allow the initiation and maintenance of interferon-based therapy (eltrombopag only)-OR- severe aplastic anemia who have had an insufficient response to immunosuppressive therapy -OR- documentation of first line treatment for severe aplastic anemia and used in combination with at least two immunosuppressive therapies.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Platelet count to be provided
Indications	All FDA-approved Indications.
Off Label Uses	

tibsovo

Products Affected

- TIBSOVO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) IDH1 mutation status, 2) alternatives tried/failed, 3) comorbidity that precludes use of intensive induction chemotherapy (i.e. age greater than or equal to 75 years, severe cardiac or pulmonary comorbidity, reduced renal function, hepatic impairment, or physician attestation)
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

tiglutik

Products Affected

- **TIGLUTIK**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of amyotrophic lateral sclerosis (ALS) -AND- Inability to swallow tablets.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of stability or improvement in symptoms of ALS.
Indications	All FDA-approved Indications.
Off Label Uses	

tolsura

Products Affected

- TOLSURA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- Unable to tolerate generic itraconazole capsules -AND- Prescriber provides rationale for clinical need of SUBA technology
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation the member is still unable to tolerate generic itraconazole capsules
Indications	All FDA-approved Indications.
Off Label Uses	

topical lidocaine

Products Affected

- *lidocaine hcl mucous membrane solution 4 % (40 mg/ml)*
- *lidocaine topical ointment*
- *lidocaine-prilocaine topical cream*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

transmucosal fentanyl citrate

Products Affected

- *fentanyl citrate buccal lozenge on a handle*
1,200 mcg, 1,600 mcg, 200 mcg, 400 mcg, 600 mcg, 800 mcg
- *fentanyl citrate buccal tablet, effervescent*
100 mcg, 200 mcg, 400 mcg, 600 mcg, 800 mcg
- *FENTORA BUCCAL TABLET, EFFERVESCENT 100 MCG, 200 MCG, 400 MCG, 600 MCG, 800 MCG*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of therapeutic use and the member is currently receiving and tolerant to long acting opioid therapy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

trelstar

Products Affected

- **TRELSTAR INTRAMUSCULAR
SUSPENSION FOR RECONSTITUTION**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

tretinoin

Products Affected

- *adapalene topical gel 0.3 %*
- *adapalene topical swab*
- **AVITA**
- *clindamycin-tretinoin*
- **DIFFERIN TOPICAL LOTION**
- **RETIN-A MICRO PUMP TOPICAL GEL WITH PUMP 0.06 %**
- *tretinoin*

PA Criteria	Criteria Details
Exclusion Criteria	Cosmetic use
Required Medical Information	Documentation of acne vulgaris -AND- trial and failure or intolerance of at least two topical acne medications (e.g. clindamycin, sulfacetamide, erythromycin)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

trikafta

Products Affected

- **TRIKAFTA**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of cystic fibrosis (CF) in patients who have at least one F508del mutation or another mutation in the cystic fibrosis transmembrane conductance regulator (CFTR gene) that is responsive to elexacaftor/tezacaftor/ivacaftor based on in vitro assay (e.g. E56K, R117C, A455E)
Age Restrictions	Deny if less than 6 years of age
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, documentation supporting improvement or stabilization of FEV1 compared to baseline FEV1 -or- increase in body mass index -or- decreased pulmonary exacerbations -or- improved quality of life as demonstrated by CF Questionnaire is required.
Indications	All FDA-approved Indications.
Off Label Uses	

Truseltiq

Products Affected

- **TRUSELTIQ ORAL CAPSULE 100 MG/DAY (100 MG X 1), 125 MG/DAY(100 MG X1-25MG X1), 50 MG/DAY (25 MG X 2), 75 MG/DAY (25 MG X 3)**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) FGFR2 fusion or other rearrangement as detected by an FDA-approved test 2) Previous therapies tried/failed
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

tukysa

Products Affected

- TUKYSA ORAL TABLET 150 MG, 50 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- All of the following, if applicable to diagnosis: 1) HER2 mutations 2) Alternatives tried/failed 3) Concomitant therapy
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

turalio

Products Affected

- TURALIO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of symptomatic tenosynovial giant cell tumor associated with severe morbidity and functional limitations -AND- patient is not amenable to improvement with surgery or not a candidate for surgery
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	24 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

tykerb

Products Affected

- *lapatinib*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- All of the following, if applicable to diagnosis: 1) HER2 mutations 2) Alternatives tried/failed 3) Concomitant therapy
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

tymlos

Products Affected

- TYMLOS

PA Criteria	Criteria Details
Exclusion Criteria	Treatment duration greater than 24 months.
Required Medical Information	Documentation of diagnosis -AND- at high risk for fracture, meeting one of the following (1. thru 3.) 1) History of previous hip or vertebral fracture. 2) T-score less than or equal to -2.5. 3) T-score between -1.0 and -2.5 -AND- meets FRAX calculation (A. or B.) A) 10-year risk of major osteoporotic fracture is greater than or equal to 20 percent or B) 10-year risk of hip fracture is greater than or equal to 3 percent.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	24 months
Other Criteria	Documentation of trial/failure or intolerance to at least one oral bisphosphonate or all are contraindicated. Coverage of human parathyroid hormone related peptide analogs beyond 24 months will not be approved. A cumulative lifetime approval of Tymlos will be limited to a coverage duration of 24 months.
Indications	All FDA-approved Indications.
Off Label Uses	

Tyvaso DPI

Products Affected

- **TYVASO DPI INHALATION CARTRIDGE WITH INHALER 16 MCG, 16 MCG (112)- 32 MCG (84), 16(112)-32(112) -48(28) MCG, 32 MCG, 32-48 MCG, 48 MCG, 64 MCG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- results of a right heart catheterization (RHC) meeting all of the following (1., 2., and 3.): 1) mean pulmonary arterial pressure (mPAP) of greater than 20 mmHg at rest, 2) pulmonary capillary wedge pressure (PWP) of less than or equal to 15 mmHg, 3) PVR greater than or equal to 3 Wood units -AND- WHO Group
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

VALCHLOR

Products Affected

- VALCHLOR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of Stage IA or IB mycosis fungoides-type cutaneous T-cell lymphoma in patients who have received prior skin-directed therapy (e.g. topical corticosteroids, topical chemotherapy, local radiation and topical retinoids).
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

valtoco

Products Affected

- VALTOCO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of seizure clusters or acute repetitive seizures -AND- Therapeutic failure, contraindication or intolerance to generic diazepam rectal gel delivery system
Age Restrictions	Deny if less than 6 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

veltassa

Products Affected

- VELTASSA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of hyperkalemia as defined by serum potassium level between 5.1 and 6.4 mmol/L on at least two (2) screenings -AND- Modification of medications to reduce serum potassium levels were not successful, when applicable.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	6 months initial authorization, 12 months reauthorization
Other Criteria	For reauthorization, attestation of reduction in serum potassium levels following Veltassa administration and continued treatment for hyperkalemia is required.
Indications	All FDA-approved Indications.
Off Label Uses	

venclexta

Products Affected

- VENCLEXTA
- VENCLEXTA STARTING PACK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis and concomitant therapy, if applicable to diagnosis. For newly-diagnosed AML, presence of at least one comorbidity that precludes use of intensive induction chemotherapy (i.e. age greater than or equal to 75 years, severe cardiac or pulmonary comorbidity, reduced renal function, hepatic impairment, or physician attestation) is required.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

verquvo

Products Affected

- VERQUVO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of heart failure (NYHA Class II to IV) -AND- Left ventricular ejection fraction less than 45% -AND- Hospitalization for heart failure or received outpatient IV diuretics for heart failure -AND- Used in combination with a angiotensin-converting enzyme inhibitor, angiotensin II receptor blocker or Entresto -AND- Used in combination with bisoprolol, carvedilol IR/ER or metoprolol succinate ER.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

verzenio

Products Affected

- VERZENIO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following: 1) HR mutation status and HER2 mutation status 2) Alternatives tried/failed 3) Concomitant therapy, if applicable to diagnosis 4) Ki-67 score greater than or equal to 20% per an FDA-approved test, if applicable to diagnosis.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

viberzi

Products Affected

- **VIBERZI**

PA Criteria	Criteria Details
Exclusion Criteria	Severe (Child-Pugh C) hepatic impairment
Required Medical Information	Documentation of diarrhea predominant, irritable bowel syndrome (IBS-D) -AND- trial/failure or intolerance to one of the following medications for IBS-D or documentation of contraindication to all: antidiarrheal (e.g., loperamide), antispasmodic (e.g., dicyclomine, hyoscyamine), tricyclic antidepressant (e.g., amitriptyline, nortriptyline).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

viibryd

Products Affected

- **TRINTELLIX**
- **VIIBRYD ORAL TABLETS,DOSE
PACK 10 MG (7)- 20 MG (23)**
- *vilazodone*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis major depressive disorder and trial and failure of one other antidepressant
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	5 years
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Vioice

Products Affected

- **VIJOICE ORAL TABLET 125 MG, 250 MG/DAY (200 MG X1-50 MG X1), 50 MG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of severe manifestations of PIK3CA Related Overgrowth Spectrum (PROS)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

vitrakvi

Products Affected

- VITRAKVI ORAL CAPSULE 100 MG, 25 MG
- VITRAKVI ORAL SOLUTION

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of a neurotrophic receptor tyrosine kinase (NTRK) gene fusion without a known acquired resistance mutation -AND- Tumors are metastatic or surgical resection is likely to result in severe morbidity - AND- There are no satisfactory alternative treatments or tumors have progressed following treatment.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Vivjoa

Products Affected

- VIVJOA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of recurrent vulvovaginal candidiasis defined as at least 3 episodes of vulvovaginal candidiasis in less than one year -AND Documentation the member is NOT of reproductive potential defined as postmenopausal or another reason for permanent infertility (e.g., tubal ligation, hysterectomy, salpingo-oophorectomy).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Pending CMS Review
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

vizimpro

Products Affected

- VIZIMPRO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of metastatic non-small cell lung cancer -AND- one of the following (1 or 2): 1. Epidermal growth factor (EGFR) exon 19 deletions - OR- 2. Epidermal growth factor receptor (EGFR) exon 21 L858R substitution mutations.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Vonjo

Products Affected

- VONJO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis of intermediate or high-risk myelofibrosis - AND- attestation of a platelet count of less than $50 \times 10^9/L$.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

voriconazole

Products Affected

- *voriconazole intravenous*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- attestation that the beneficiary cannot take oral voriconazole
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	3 months
Other Criteria	For reauthorization, attestation of continued indicators of active disease (e.g. histopathology, positive cultures) is required
Indications	All FDA-approved Indications.
Off Label Uses	

vosevi

Products Affected

- VOSEVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Criteria will be applied consistent with current AASLD/IDSA guidance - AND- the member has a contraindication to or is otherwise not a candidate for all regimens recommended by the AASLD/IDSA guidelines containing the following agents: ledipasvir/sofosbuvir, sofosbuvir/velpatasvir, glecaprevir/pibrentasvir.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	Criteria/duration applied consistent with current AASLD-IDSA guidance
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

votrient

Products Affected

- VOTRIENT

PA Criteria	Criteria Details
Exclusion Criteria	Documentation of adipocytic soft tissue sarcoma or gastrointestinal stromal tumor
Required Medical Information	Documentation of diagnosis -AND- if applicable to diagnosis, alternatives tried/failed
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

vraylar

Products Affected

- VRAYLAR ORAL CAPSULE
- VRAYLAR ORAL CAPSULE,DOSE PACK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- trial, intolerance, or contraindication to one other formulary generic atypical antipsychotic (e.g. quetiapine).
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

vumerity

Products Affected

- VUMERITY

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use with other disease modifying agents such as interferons, Copaxone, Tysabri, Aubagio, Gilenya
Required Medical Information	Documentation of relapsing form of multiple sclerosis (e.g. relapsing-remitting, clinically isolated syndrome, or active secondary progressive disease) -AND- Therapeutic failure or intolerance to generic dimethyl fumarate
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	5 years
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

wakix

Products Affected

- WAKIX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of narcolepsy -AND- Documentation of baseline data of excessive daytime sleepiness (EDS) via the Epworth Sleepiness Scale (ESS) or Maintenance of Wakefulness Test (MWT) - AND- Documentation of the following (1, 2, or 3): 1) Hypocretin-1 deficiency defined by (A or B), A) Cerebrospinal fluid hypocretin-1 less than 110 pg/mL. B) Cerebrospinal fluid hypocretin-1 less than 1/3 of the normal value based on laboratory reference range -OR- 2) Multiple sleep latency test (MSLT) documenting MSL less than 8 minutes and 2 sleep-onset rapid eye movement periods (SOREMP) -OR- 3) MSLT documenting MSL less than 8 minutes and 1 SOREMP and Polysomnography substantiating 1 SOREMP. If the member has a diagnosis of cataplexy provision of baseline number of cataplexy episodes is required.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	If no diagnosis of cataplexy, trial and failure, intolerance, or contraindication to generic modafinil and a generic CNS stimulant indicated for use in narcolepsy (e.g. methylphenidate, amphetamine salts) is required -OR- Prescriber attests a significant concern about the potential for illegal drug diversion. For reauthorization, provider attestation of improvement in symptoms of narcolepsy or improvement in symptoms of cataplexy (if applicable).
Indications	All FDA-approved Indications.
Off Label Uses	

Welireg

Products Affected

- **WELIREG**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of von Hippel Lindau (VHL) syndrome -AND- one of the following diagnoses not requiring immediate surgery (1, 2, or 3): 1) Renal cell carcinoma. 2) CNS hemangioblastoma. 3) Pancreatic neuroendocrine tumor.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

xalkori

Products Affected

- **XALKORI**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) documentation of ALK or ROS1 mutation status
Age Restrictions	Deny if less than 18 years of age for NSCLC. Deny if less than 1 year of age or greater than 21 years of age for ALCL.
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

Xarelto Suspension

Products Affected

- **XARELTO ORAL SUSPENSION FOR RECONSTITUTION**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- Inability to swallow Xarelto (Rivaroxaban) tablets.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, member continues to have an inability to swallow Xarelto (Rivaroxaban) tablets.
Indications	All FDA-approved Indications.
Off Label Uses	

xcopri

Products Affected

- **XCOPRI 150MG X1)**
- **XCOPRI MAINTENANCE PACK ORAL • XCOPRI TITRATION PACK
TABLET 250MG/DAY(150 MG X1-
100MG X1), 350 MG/DAY (200 MG X1-**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of partial-onset seizures -AND- Therapeutic failure, intolerance or contraindication to 1 other anti-epileptic drug (e.g. carbamazepine, levetiracetam)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

xeljanz

Products Affected

- XELJANZ ORAL TABLET
- XELJANZ XR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For moderate to severe rheumatoid arthritis and an inadequate response or intolerance to methotrexate. Xeljanz immediate release for juvenile idiopathic arthritis, inadequate response or intolerance to at least one DMARD (e.g., methotrexate, leflunomide) -OR- requires initial biologic therapy due to involvement of high-risk joints, high disease activity or at high risk of disabling joint damage. For ankylosing spondylitis, inadequate response or intolerance to at least one nonsteroidal anti-inflammatory drug (NSAID).
Age Restrictions	Deny if less than 18 years of age for rheumatoid arthritis, psoriatic arthritis, ulcerative colitis, ankylosing spondylitis, or less than 2 years of age for juvenile idiopathic arthritis
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	The member has experienced inadequate response or intolerance to at least 1 tumor necrosis factor (TNF) blocker. Doses greater than 10 mg per day for Xeljanz and 11 mg per day for Xeljanz XR will not be approved for rheumatoid arthritis and psoriatic arthritis. Doses greater than 20mg per day for Xeljanz and 22 mg per day for Xeljanz XR will not be approved for ulcerative colitis.
Indications	All FDA-approved Indications.
Off Label Uses	

xeljanz solution

Products Affected

- XELJANZ ORAL SOLUTION

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of juvenile idiopathic arthritis -AND- Inadequate response or intolerance to at least one DMARD (e.g., methotrexate, leflunomide) or requires initial biologic therapy due to involvement of high-risk joints, high disease activity or at high risk of disabling joint damage.
Age Restrictions	Deny if less than 2 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	The member has experienced therapeutic failure or intolerance to at least 1 tumor necrosis factor (TNF) blocker.
Indications	All FDA-approved Indications.
Off Label Uses	

xenazine

Products Affected

- *tetrabenazine oral tablet 12.5 mg, 25 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- attestation that the beneficiary is not actively suicidal
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	In patients with comorbid depression, attestation of adequate treatment for depression is required.
Indications	All FDA-approved Indications.
Off Label Uses	

xermelo

Products Affected

- XERMELO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of carcinoid syndrome diarrhea AND used in combination with a somatostatin analog AND trial and failure of somatostatin analog monotherapy
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

xgeva

Products Affected

- XGEVA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis. For hypercalcemia of malignancy, refractory to bisphosphonates. For giant cell tumor of bone, unresectable or surgical resection is likely to result in severe morbidity -AND- one of the following (1. or 2.)- 1.) the member is 18 years old or older -OR- 2.) the member is a skeletally mature adolescent (e.g. has at least one mature long bone)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

xifaxan

Products Affected

- XIFAXAN ORAL TABLET 550 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of 1 or 2. 1) Diagnosis of hepatic encephalopathy AND trial/failure, intolerance, or contraindication to lactulose. 2) Diagnosis of Irritable Bowel Syndrome with Diarrhea (IBS-D) AND trial/failure, intolerance to one of the following medications for IBS-D or documentation of contraindication to all: antidiarrheal (e.g., loperamide), antispasmodic (e.g., dicyclomine, hyoscyamine), tricyclic antidepressant (e.g., amitriptyline, nortriptyline).
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	Hepatic encephalopathy: 1 year. IBS-D: 14 days.
Other Criteria	No more than three courses of rifaximin for the treatment of IBS-D will be approved per lifetime.
Indications	All FDA-approved Indications.
Off Label Uses	

xolair

Products Affected

- XOLAIR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of chronic idiopathic urticaria with trial/failure or intolerance of a second-generation non-sedating H1 antihistamine at the maximum recommended doses (e.g., cetirizine, fexofenadine, loratadine, desloratadine, levocetirizine) -OR- Documentation of moderate to severe persistent asthma in patients with a positive skin test or in vitro reactivity to a perennial aeroallergen -AND- Baseline IgE titre greater than or equal to 30 IU/mL -AND- symptoms that are inadequately controlled despite a 3 month trial of both 1. and 2. 1) medium-dose inhaled corticosteroid or systemic steroid 2) a long-acting beta-agonist or leukotriene antagonist - AND- patient is currently on the optimal dose of a long-acting beta2-agonist, leukotriene modifier, or theophylline -OR- Documentation of add-on maintenance treatment for nasal polyps -AND- trial & failure, intolerance or contraindication to intra-nasal corticosteroids
Age Restrictions	Deny if less than 12 years of age in treatment for chronic idiopathic urticaria -OR- deny if less than 6 years of age for severe persistent asthma -OR- deny if less than 18 years of age for nasal polyps
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	For reauthorization, attestation of decreased rescue medication or oral corticosteroid use, decrease frequency of exacerbation, increased pulmonary function from baseline or reduction in asthma related symptoms in treatment of asthma -OR- improved symptoms in treatment of CIU -OR- reduction in nasal polyp score or nasal congestion/obstruction severity score in treatment of nasal polyps
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	

xospata

Products Affected

- XOSPATA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- FLT3 mutations, if applicable to diagnosis
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

xpovio

Products Affected

- **XPOVIO ORAL TABLET 100 MG/WEEK (50 MG X 2), 40 MG/WEEK (40 MG X 1), 40MG TWICE WEEK (40 MG X 2), 60 MG/WEEK (60 MG X 1), 60MG TWICE WEEK (120 MG/WEEK), 80 MG/WEEK (40 MG X 2), 80MG TWICE WEEK (160 MG/WEEK)**

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of use in combination with dexamethasone for relapse or refractory multiple myeloma with failure, intolerance or contraindication to 5 therapies (e.g. bortezomib, carfilzomib, lenalidomide, pomalidomide and daratumumab) -OR- Documentation of use in combination with both bortezomib and dexamethasone for relapse or refractory multiple myeloma after receiving 1 prior multiple myeloma therapy -OR- Documentation of relapsing or refractory diffuse large B-cell lymphoma with failure, intolerance or contraindication to at least 2 lines of systemic therapy
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

xtandi

Products Affected

- XTANDI ORAL CAPSULE
- XTANDI ORAL TABLET 40 MG, 80 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- one of the following (1. or 2.): 1. Concomitant GnRH analog 2. The member has had a bilateral orchiectomy
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

xuriden

Products Affected

- XURIDEN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of hereditary orotic aciduria
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

xyrem

Products Affected

- XYREM

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of narcolepsy -AND- Documentation of baseline data of excessive daytime sleepiness (EDS) via the Epworth Sleepiness Scale (ESS) or Maintenance of Wakefulness Test (MWT) - AND- Documentation of the following (1, 2, or 3): 1) Hypocretin-1 deficiency defined by (A or B), A) Cerebrospinal fluid hypocretin-1 less than 110 pg/mL. B) Cerebrospinal fluid hypocretin-1 less than 1/3 of the normal value based on laboratory reference range -OR- 2) Multiple sleep latency test (MSLT) documenting MSL less than 8 minutes and 2 sleep-onset rapid eye movement periods (SOREMP) -OR- 3) MSLT documenting MSL less than 8 minutes and 1 SOREMP and Polysomnography substantiating 1 SOREMP. If the member has a diagnosis of cataplexy provision of baseline number of cataplexy episodes is required.
Age Restrictions	Deny if less than 7 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	If no diagnosis of cataplexy- trial and failure, intolerance, or contraindication to generic modafinil AND a generic CNS stimulant indicated for use in narcolepsy (e.g. methylphenidate, amphetamine salts) is required. For reauthorization, attestation supporting improvement in symptoms of narcolepsy and cataplexy (if applicable) is required.
Indications	All FDA-approved Indications.
Off Label Uses	

xywav

Products Affected

- XYWAV

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of narcolepsy -AND- Documentation of baseline data of excessive daytime sleepiness (EDS) via the Epworth Sleepiness Scale (ESS) or Maintenance of Wakefulness Test (MWT) - AND- Documentation of the following (1, 2, or 3): 1) Hypocretin-1 deficiency defined by (A or B), A) Cerebrospinal fluid hypocretin-1 less than 110 pg/mL. B) Cerebrospinal fluid hypocretin-1 less than 1/3 of the normal value based on laboratory reference range -OR- 2) Multiple sleep latency test (MSLT) documenting MSL less than 8 minutes and 2 sleep-onset rapid eye movement periods (SOREMP) -OR- 3) MSLT documenting MSL less than 8 minutes and 1 SOREMP and Polysomnography substantiating 1 SOREMP -AND- Sensitivity to sodium intake due to heart failure, hypertension or impaired renal function. If the member has a diagnosis of cataplexy, provision of baseline number of cataplexy episodes is required. -OR- (II) Diagnosis of idiopathic hypersomnia - AND- Documentation of baseline data of excessive daytime sleepiness (EDS) via the Epworth Sleepiness Scale (ESS) or Maintenance of Wakefulness Test (MWT) -AND- documentation the member does not have cataplexy -AND- documentation of less than 2 SOREMPs -AND- documentation of the following (1, 2, or 3): 1) MSLT documenting MSL less than or equal to 8 minutes -OR- 2) polysomnography demonstrating total sleep time greater than or equal to 660 minutes per 24 hours -OR- 3) wrist actigraphy demonstrating total sleep time greater than or equal to 660 minutes per 24 hours.
Age Restrictions	For narcolepsy, deny if less than 7 years of age. For idiopathic hypersomnia, deny if less than 18 years of age.
Prescriber Restrictions	
Coverage Duration	12 months

PA Criteria	Criteria Details
Other Criteria	If diagnosis of narcolepsy without cataplexy- trial and failure, intolerance, or contraindication to generic modafinil AND a generic CNS stimulant indicated for use in narcolepsy (e.g. methylphenidate, amphetamine salts) is required. For reauthorization, attestation supporting improvement in symptoms of narcolepsy, idiopathic hypersomnia and cataplexy (if applicable) is required.
Indications	All FDA-approved Indications.
Off Label Uses	

yonsa

Products Affected

- YONSA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- all of the following: 1) Alternatives tried/failed 2) Concomitant therapy
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

zavesca

Products Affected

- *miglustat*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of mild to moderate type 1 Gaucher disease confirmed by the following (A. or B.) A. (1, 2, 3, 4, or, 5): 1)Hepatomegaly. 2)Splenomegaly. 3)Bone disease (i.e. osteonecrosis, osteopenia, secondary pathologic fractures, bone infarct). 4)Bone marrow complications as defined by anemia with hemoglobin less than or equal to 11.5 g/dL for females or 12.5 g/dL for males or thrombocytopenia with platelet count less than or equal to 120,000/mm ³ -OR- 5)Symptomatic disease (e.g. bone pain, exertional limitation, cachexia). -OR- B. Attestation of deficiency in glucocerebrosidase activity in peripheral leukocytes or genetic testing confirms mutant alleles.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	Documentation of trial/failure or intolerance to at least one enzyme replacement therapy product including Cerezyme, Elelyso, or VPRIV
Indications	All FDA-approved Indications.
Off Label Uses	

zejula

Products Affected

- ZEJULA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- Alternatives tried/failed
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

zelboraf

Products Affected

- TAFINLAR
- ZELBORAF

PA Criteria	Criteria Details
Exclusion Criteria	Wild-type BRAF melanoma
Required Medical Information	Documentation of diagnosis -AND- both of the following. 1) BRAF mutations, if applicable to diagnosis. 2) Alternatives tried/failed and concomitant therapy, if applicable to diagnosis (e.g. diagnosis of V600K metastatic melanoma and drug regimen of Zelboraf and Cotellic)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

zepatier

Products Affected

- ZEPATIER

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Criteria will be applied consistent with current AASLD/IDSA guidance - AND- the member has a contraindication to or is otherwise not a candidate for all regimens recommended by the AASLD/IDSA guidelines containing the following agents: ledipasvir/sofosbuvir, sofosbuvir/velpatasvir, glecaprevir/pibrentasvir.
Age Restrictions	Deny if less than 12 years of age
Prescriber Restrictions	
Coverage Duration	Criteria/duration applied consistent with current AASLD-IDSA guidance
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

zeposia

Products Affected

- **ZEPOSIA**
- **ZEPOSIA STARTER KIT**
- **ZEPOSIA STARTER PACK**

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use of Zeposia and other disease modifying agents such as interferons, Copaxone, Tysabri.
Required Medical Information	Documentation of diagnosis. For moderate to severe active ulcerative colitis, inadequate response or intolerance to one systemic therapy (e.g. methotrexate, cyclosporine).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Ulcerative Colitis: 12 months, Multiple Sclerosis: 24 months
Other Criteria	For moderate to severe active ulcerative colitis, patients must have an adequate trial or intolerance to 2 of the following preferred products Humira, Xeljanz/Xeljanz XR and Stelara SC.
Indications	All FDA-approved Indications.
Off Label Uses	

zolinza

Products Affected

- ZOLINZA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of cutaneous manifestations in patients with cutaneous T-cell lymphoma (CTCL) who have progressive, persistent, or recurrent disease on or following 2 systemic therapies. Systemic therapies include bexarotene, interferon alpha, extracorporeal photochemotherapy, PUVA, single agent or combination chemotherapies (e.g. cyclophosphamide, vinblastine, romidepsin)
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

ZYDELIG

Products Affected

- ZYDELIG

PA Criteria	Criteria Details
Exclusion Criteria	First line treatment. Combination use with benadmustine and/or rituximab for the treatment of FL.
Required Medical Information	Documentation of diagnosis -AND- all of the following, if applicable to diagnosis: 1) Alternatives tried/failed 2) Concomitant therapy 3) Candidacy for use of rituximab alone
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

zykadia

Products Affected

- ZYKADIA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- ALK mutations, as detected by an FDA approved test.
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

zytiga

Products Affected

- *abiraterone oral tablet 250 mg, 500 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Documentation of diagnosis -AND- concomitant therapy
Age Restrictions	Deny if less than 18 years of age
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

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<i>acetaminophen-codeine oral tablet</i>	230	BAFIERTAM	24
<i>acitretin</i>	1	BALVERSA	25
ACTEMRA ACTPEN	2	BANZEL ORAL SUSPENSION	26
ACTEMRA SUBCUTANEOUS	2	BENLYSTA SUBCUTANEOUS	27
ACTHAR	3	<i>benztropine oral</i>	115
ACTIMMUNE	5	BERINERT INTRAVENOUS KIT	29
<i>adapalene topical gel 0.3 %</i>	307	BESREMI	31
<i>adapalene topical swab</i>	307	BETHKIS	47
ADBRY	6	<i>bexarotene oral</i>	287
ADEMPAS	236	<i>bexarotene topical</i>	287
AIMOVIG AUTOINJECTOR		BIVIGAM	125
SUBCUTANEOUS AUTO-INJECTOR		<i>bosentan</i>	236
140 MG/ML, 70 MG/ML	9	BOSULIF	32
AJOVY AUTOINJECTOR	10	BRAFTOVI ORAL CAPSULE 75 MG	33
AJOVY SYRINGE	10	BRONCHITOL	34
ALECENSA	11	BRUKINSA	35
ALKINDI SPRINKLE	12	<i>buprenorphine</i>	36
<i>alosetron oral tablet 0.5 mg, 1 mg</i>	162	<i>butalbital-acetaminop-caf-cod oral capsule 50-300-40-30 mg, 50-325-40-30 mg</i>	230
ALPRAZOLAM INTENSOL	230	BUTRANS	36
<i>alprazolam oral tablet 0.25 mg, 0.5 mg, 1 mg, 2 mg</i>	230	CABLIVI INJECTION KIT	38
<i>alprazolam oral tablet extended release 24 hr 0.5 mg, 1 mg, 2 mg, 3 mg</i>	230	CABOMETYX	39
ALUNBRIG ORAL TABLET 180 MG, 30 MG, 90 MG	14	CALQUENCE	40
ALUNBRIG ORAL TABLETS,DOSE PACK	14	CALQUENCE (ACALABRUTINIB MAL)	40
ALYQ	236	CAMZYOS	41
<i>ambrisentan</i>	236	CAPLYTA	42
<i>amitriptyline</i>	115	CAPRELSA	43
<i>amitriptyline-chlordiazepoxide</i>	115	CARAC	44
ANDRODERM	296	<i>carglumic acid</i>	45
<i>apomorphine</i>	17	CERDELGA	46
ARALAST NP INTRAVENOUS RECON SOLN 1,000 MG	13	CHENODAL	48
ARCALYST	135	<i>chloroquine phosphate oral tablet 250 mg, 500 mg</i>	49
ARIKAYCE	18	CHOLBAM	50
<i>aripiprazole</i>	20	CIALIS ORAL TABLET 2.5 MG, 5 MG	51
<i>armodafinil</i>	196	CIBINQO	52
AUBAGIO	21	CIMZIA	53
AURYXIA	22	CIMZIA POWDER FOR RECONST	53
AVEED	296	CINRYZE	55
		<i>clindamycin-tretinoin</i>	307
		<i>clobazam oral suspension</i>	202
		<i>clobazam oral tablet</i>	202

<i>clomipramine</i>	115	<i>doxepin topical</i>	72
<i>clonidine hcl oral tablet extended release</i> <i>12 hr</i>	7	DRIZALMA SPRINKLE ORAL CAPSULE, DELAYED REL SPRINKLE 20 MG, 30 MG, 40 MG, 60 MG	73
COMETRIQ	57	<i>droxidopa</i>	189
COPIKTRA	58	DUOBRII	74
CORLANOR ORAL SOLUTION	59	DUPIXENT PEN SUBCUTANEOUS PEN INJECTOR 200 MG/1.14 ML, 300 MG/2 ML	75
CORLANOR ORAL TABLET 5 MG, 7.5 MG	59	DUPIXENT SYRINGE SUBCUTANEOUS SYRINGE 100 MG/0.67 ML, 200 MG/1.14 ML, 300 MG/2 ML	75
CORTROPHIN GEL	238	EGRIFTA SV	78
COSENTYX (2 SYRINGES)	60	EMFLAZA	79
COSENTYX PEN (2 PENS)	60	EMGALITY PEN	80
COSENTYX SUBCUTANEOUS SYRINGE 75 MG/0.5 ML	60	EMGALITY SYRINGE SUBCUTANEOUS SYRINGE 120 MG/ML, 300 MG/3 ML (100 MG/ML X 3)	80
COTELLIC	61	ENBREL MINI	82
CRINONE	62	ENBREL SUBCUTANEOUS SOLUTION	82
<i>cyclobenzaprine oral tablet 10 mg, 5 mg,</i> <i>7.5 mg</i>	115	ENBREL SUBCUTANEOUS SYRINGE 25 MG/0.5 ML (0.5), 50 MG/ML (1 ML) ..	82
<i>cyproheptadine</i>	115	ENBREL SURECLICK	82
CYSTADROPS	63	ENDARI	83
CYSTARAN	63	ENDOCET ORAL TABLET 10-325 MG, 5-325 MG, 7.5-325 MG	230
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DAURISMO ORAL TABLET 100 MG, 25 MG	65	EPCLUSA ORAL PELLETS IN PACKET 150-37.5 MG, 200-50 MG	85
DAYTRANA	66	EPCLUSA ORAL TABLET	85
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DIACOMIT ORAL CAPSULE 250 MG, 500 MG	68	<i>ergotamine-caffeine</i>	115
DIACOMIT ORAL POWDER IN PACKET 250 MG, 500 MG	68	ERIVEDGE	88
<i>diclofenac epolamine</i>	99	ERLEADA	89
<i>diclofenac sodium topical gel 3 %</i>	270	<i>erlotinib</i>	77
DIFFERIN TOPICAL LOTION	307	ESBRIET ORAL CAPSULE	136
<i>dihydroergotamine nasal</i>	69	ESBRIET ORAL TABLET 267 MG, 801 MG	136
<i>dimethyl fumarate oral capsule, delayed</i> <i>release(dr/ec) 120 mg, 120 mg (14)- 240</i> <i>mg (46), 240 mg</i>	293	<i>eszopiclone</i>	230
DOJOLVI	70		
DOPTELET (10 TAB PACK)	71		
DOPTELET (15 TAB PACK)	71		
DOPTELET (30 TAB PACK)	71		
<i>doxepin oral capsule</i>	115		
<i>doxepin oral concentrate</i>	115		
<i>doxepin oral tablet</i>	115		

EVENITY SUBCUTANEOUS	
SYRINGE 210MG/2.34ML (
105MG/1.17MLX2)	90
<i>everolimus (antineoplastic) oral tablet 10</i>	
<i>mg, 2.5 mg, 5 mg, 7.5 mg</i>	8
<i>everolimus (antineoplastic) oral tablet for</i>	
<i>suspension 2 mg, 3 mg, 5 mg</i>	8
EVRYSDI	91
EXKIVITY	77
EXSERVAN	92
FASENRA	93
FASENRA PEN	93
<i>fentanyl citrate buccal lozenge on a handle</i>	
<i>1,200 mcg, 1,600 mcg, 200 mcg, 400 mcg,</i>	
<i>600 mcg, 800 mcg</i>	305
<i>fentanyl citrate buccal tablet, effervescent</i>	
<i>100 mcg, 200 mcg, 400 mcg, 600 mcg, 800</i>	
<i>mcg</i>	305
<i>fentanyl transdermal patch 72 hour 100</i>	
<i>mcg/hr, 12 mcg/hr, 25 mcg/hr, 50 mcg/hr,</i>	
<i>75 mcg/hr</i>	230
FENTORA BUCCAL TABLET,	
EFFERVESCENT 100 MCG, 200 MCG,	
400 MCG, 600 MCG, 800 MCG	305
FERRIPROX ORAL SOLUTION	94
FETZIMA ORAL CAPSULE,EXT REL	
24HR DOSE PACK	95
FETZIMA ORAL	
CAPSULE,EXTENDED RELEASE 24	
HR 120 MG, 20 MG, 40 MG, 80 MG	95
<i> fingolimod</i>	106
FINTEPLA	96
FIRDAPSE	98
FLEBOGAMMA DIF INTRAVENOUS	
SOLUTION 10 %	125
FLECTOR	99
FORTEO SUBCUTANEOUS PEN	
INJECTOR 20 MCG/DOSE	
(600MCG/2.4ML)	100
FOTIVDA	101
<i>gabapentin oral capsule 100 mg, 300 mg,</i>	
<i>400 mg</i>	102
<i>gabapentin oral solution 250 mg/5 ml</i>	102
<i>gabapentin oral tablet 600 mg, 800 mg</i>	102
GALAFOLD	103
GAMMAGARD LIQUID	125
GAMMAGARD S-D (IGA < 1	
MCG/ML)	125
GAMMAKED INJECTION	
SOLUTION 1 GRAM/10 ML (10 %)	125
GAMMAPLEX	125
GAMMAPLEX (WITH SORBITOL)	125
GAMUNEX-C INJECTION	
SOLUTION 1 GRAM/10 ML (10 %)	125
GATTEX 30-VIAL	104
GAVRETO	105
GENOTROPIN	109
GENOTROPIN MINIQUICK	109
GILOTRIF	77
GLASSIA	13
GRASTEK	108
<i>guanfacine oral tablet extended release 24</i>	
<i>hr</i>	7
HAEGARDA	110
HARVONI ORAL PELLETS IN	
PACKET	112
HARVONI ORAL TABLET 90-400 MG	
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HETLIOZ	113
HETLIOZ LQ	114
HORIZANT ORAL TABLET	
EXTENDED RELEASE 300 MG, 600	
MG	118
HUMATROPE INJECTION	
CARTRIDGE	109
HUMIRA PEN	119
HUMIRA PEN CROHNS-UC-HS	
START	119
HUMIRA PEN PSOR-UVEITS-ADOL	
HS	119
HUMIRA SUBCUTANEOUS	
SYRINGE KIT 40 MG/0.8 ML	119
HUMIRA(CF)	119
HUMIRA(CF) PEDI CROHNS	
STARTER SUBCUTANEOUS	
SYRINGE KIT 80 MG/0.8 ML, 80	
MG/0.8 ML-40 MG/0.4 ML	119
HUMIRA(CF) PEN	119
HUMIRA(CF) PEN CROHNS-UC-HS ..	119
HUMIRA(CF) PEN PEDIATRIC UC	119
HUMIRA(CF) PEN PSOR-UV-ADOL	
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<i>hydrocodone-ibuprofen oral tablet 10-200</i>	KEVZARA	145
<i>mg, 5-200 mg</i>	KINERET	146
<i>hydromorphone oral liquid</i>	KISQALI FEMARA CO-PACK ORAL	
<i>hydromorphone oral tablet</i>	TABLET 200 MG/DAY(200 MG X 1)-	
<i>hydroxyzine hcl oral solution 10 mg/5 ml</i> ..	2.5 MG, 400 MG/DAY(200 MG X 2)-2.5	
<i>hydroxyzine hcl oral tablet</i>	MG, 600 MG/DAY(200 MG X 3)-2.5	
IBRANCE	MG	147
IBSRELA	KISQALI ORAL TABLET 200	
<i>icatibant</i>	MG/DAY (200 MG X 1), 400 MG/DAY	
ICLUSIG	(200 MG X 2), 600 MG/DAY (200 MG X	
IDHIFA ORAL TABLET 100 MG, 50	3)	147
MG	KLISYRI	148
<i>imatinib oral tablet 100 mg, 400 mg</i>	KORLYM	149
IMBRUVICA ORAL CAPSULE 140	KOSELUGO ORAL CAPSULE 10 MG,	
MG, 70 MG	25 MG	150
IMBRUVICA ORAL SUSPENSION	KYNMOBI SUBLINGUAL FILM 10	
IMBRUVICA ORAL TABLET	MG, 15 MG, 20 MG, 25 MG, 30 MG	152
<i>imipramine hcl</i>	<i>lapatinib</i>	312
<i>imipramine pamoate</i>	LATUDA ORAL TABLET 120 MG, 20	
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60 MG, 80 MG	155
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INQOVI	<i>% (40 mg/ml)</i>	304
INREBIC	<i>lidocaine topical adhesive patch,medicated</i>	
INTRON A INJECTION RECON	<i>5 %</i>	157
SOLN 10 MILLION UNIT (1 ML), 50	<i>lidocaine topical ointment</i>	304
MILLION UNIT (1 ML)	<i>lidocaine-prilocaine topical cream</i>	304
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MAYZENT ORAL TABLET 0.25 MG, 1 MG, 2 MG	170	NUCYNTA	230
MAYZENT STARTER(FOR 1MG MAINT)	170	NUDEXTA	194
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<i>megestrol oral tablet</i>	171	OCALIVA	197
MEKINIST	172	OCTAGAM	125
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<i>metaxalone</i>	115	ODACTRA	199
<i>methadone oral solution 10 mg/5 ml, 5 mg/5 ml</i>	230	ODOMZO	200
<i>methadone oral tablet 10 mg, 5 mg</i>	230	OFEV	136
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<i>modafinil</i>	235	ONGENTYS	203
<i>morphine concentrate oral solution</i>	230	ONUREG	204
<i>morphine oral capsule, er multiphase 24 hr 120 mg, 30 mg, 45 mg, 60 mg, 75 mg, 90 mg</i>	230	OPSUMIT	236
<i>morphine oral solution 10 mg/5 ml, 20 mg/5 ml (4 mg/ml)</i>	230	OPZELURA	205
<i>morphine oral tablet</i>	230	ORALAIR SUBLINGUAL TABLET 300 INDX REACTIVITY	206
<i>morphine oral tablet extended release 100 mg, 15 mg, 200 mg, 30 mg, 60 mg</i>	230	ORENCIA CLICKJECT	207
MULPLETA	175	ORENCIA SUBCUTANEOUS SYRINGE 125 MG/ML, 50 MG/0.4 ML, 87.5 MG/0.7 ML	207
MYALEPT	176	ORENITRAM ORAL TABLET EXTENDED RELEASE 0.125 MG, 0.25 MG, 1 MG, 2.5 MG, 5 MG	236
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<i>oxandrolone</i>	16	PROMACTA ORAL POWDER IN PACKET 12.5 MG, 25 MG	300
OXBRYTA ORAL TABLET	216	PROMACTA ORAL TABLET 12.5 MG, 25 MG, 50 MG, 75 MG	300
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<i>oxycodone oral capsule</i>	230	<i>pyrimethamine</i>	64
<i>oxycodone oral concentrate</i>	230	PYRUKYND ORAL TABLET 20 MG, 5 MG, 5 MG (4-WEEK PACK), 50 MG	240
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<i>oxycodone oral tablet 10 mg, 15 mg, 20 mg, 30 mg, 5 mg</i>	230	QINLOCK	241
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PHEBURANE	222	RELISTOR ORAL	249
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PIQRAY ORAL TABLET 200 MG/DAY (200 MG X 1), 250 MG/DAY (200 MG X1-50 MG X1), 300 MG/DAY (150 MG X 2)	224	REPATHA PUSHTRONEX	251
<i>pirfenidone oral tablet 267 mg, 801 mg</i>	136	REPATHA SURECLICK	251
POMALYST	225	REPATHA SYRINGE	251
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SEROSTIM SUBCUTANEOUS RECON SOLN 4 MG, 5 MG, 6 MG	109	TALZENNA ORAL CAPSULE 0.25 MG, 0.5 MG, 0.75 MG, 1 MG	286
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SIMPONI SUBCUTANEOUS SYRINGE 100 MG/ML, 50 MG/0.5 ML	268	TAZORAC TOPICAL CREAM 0.05 %	291
SKYRIZI SUBCUTANEOUS PEN INJECTOR	269	TAZVERIK	292
SKYRIZI SUBCUTANEOUS SYRINGE 150 MG/ML	269	TEGSEDI	294
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<i>sorafenib</i>	185	<i>testosterone transdermal gel in metered-dose pump 20.25 mg/1.25 gram (1.62 %)</i> ...296	
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SPRYCEL	273	<i>tetrabenazine oral tablet 12.5 mg, 25 mg</i> ...340	
STELARA SUBCUTANEOUS SOLUTION	274	THALOMID ORAL CAPSULE 100 MG, 150 MG, 200 MG, 50 MG	298
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		TRACLEER ORAL TABLET FOR SUSPENSION	236
		<i>tramadol oral tablet 100 mg, 50 mg</i>	230
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<i>trimipramine</i>	115	VONJO	327
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VERZENIO	320	XELJANZ ORAL SOLUTION	339
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<i>vilazodone</i>	322	XOLAIR	344
		XOSPATA	346
		XPOVIO ORAL TABLET 100 MG/WEEK (50 MG X 2), 40 MG/WEEK (40 MG X 1), 40MG TWICE WEEK (40 MG X 2), 60 MG/WEEK (60 MG X 1), 60MG TWICE WEEK (120 MG/WEEK), 80 MG/WEEK (40 MG X 2), 80MG TWICE WEEK (160 MG/WEEK)	347
		XTANDI ORAL CAPSULE	348
		XTANDI ORAL TABLET 40 MG, 80 MG	348
		XURIDEN	349

XYREM	350
XYWAV	351
YONSA	353
<i>zaleplon oral capsule 10 mg, 5 mg</i>	230
ZEJULA	355
ZELBORAF	356
ZEMAIRA	13
ZEPATIER	357
ZEPOSIA	358
ZEPOSIA STARTER KIT	358
ZEPOSIA STARTER PACK	358
<i>zileuton</i>	156
ZOLINZA	359
<i>zolpidem</i>	230
ZOMACTON	109
ZORBTIVE	109
ZYDELIG	360
ZYFLO	156
ZYKADIA	361

brand adhd

Products Affected

- **APTENSIO XR 10 MG CAPSULE,EXTENDED RELEASE SPRINKLE**
- **APTENSIO XR 15 MG CAPSULE,EXTENDED RELEASE SPRINKLE**
- **APTENSIO XR 20 MG CAPSULE,EXTENDED RELEASE SPRINKLE**
- **APTENSIO XR 30 MG CAPSULE,EXTENDED RELEASE SPRINKLE**
- **APTENSIO XR 40 MG CAPSULE,EXTENDED RELEASE SPRINKLE**
- **APTENSIO XR 50 MG CAPSULE,EXTENDED RELEASE SPRINKLE**
- **APTENSIO XR 60 MG CAPSULE,EXTENDED RELEASE SPRINKLE**
- **QUILLIVANT XR 5 MG/ML (25 MG/5 ML) ORAL SUSPENSION,EXTEND RELEASE 24HR**
- **VYVANSE 10 MG CAPSULE**
- **VYVANSE 20 MG CAPSULE**
- **VYVANSE 30 MG CAPSULE**
- **VYVANSE 40 MG CAPSULE**
- **VYVANSE 50 MG CAPSULE**
- **VYVANSE 60 MG CAPSULE**
- **VYVANSE 70 MG CAPSULE**

Details

Criteria	Require a 1 month trial of 2 of the following generic ADHD medications (Step 1 drug) when being utilized for the same medically accepted indication: methylphenidate, atomoxetine, dextroamphetamine/amphetamine or dexmethylphenidate in the last 180 days
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celecoxib

Products Affected

- *celecoxib 100 mg capsule*
- *celecoxib 200 mg capsule*
- *celecoxib 400 mg capsule*
- *celecoxib 50 mg capsule*

Details

Criteria	Require a 1 month trial of 2 formulary generic NSAIDs (Step 1 drug) in the last 180 days
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copaxone

Products Affected

- **COPAXONE 20 MG/ML
SUBCUTANEOUS SYRINGE**
- **COPAXONE 40 MG/ML
SUBCUTANEOUS SYRINGE**

Details

Criteria	Require a 1 month trial of generic glatiramer and Glatopa in the last 180 days
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herpetic keratitis

Products Affected

- **ZIRGAN 0.15 % EYE GEL**

Details

Criteria	Require a 1 month trial of generic trifluridine eye drops (Step 1 drug) in the last 90 days
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lupron

Products Affected

- **LUPRON DEPOT 11.25 MG (3 MONTH) INTRAMUSCULAR SYRINGE KIT**
- **LUPRON DEPOT 22.5 MG (3 MONTH) INTRAMUSCULAR SYRINGE KIT**
- **LUPRON DEPOT 3.75 MG INTRAMUSCULAR SYRINGE KIT**
- **LUPRON DEPOT 30 MG (4 MONTH) INTRAMUSCULAR SYRINGE KIT**
- **LUPRON DEPOT 45 MG (6 MONTH) INTRAMUSCULAR SYRINGE KIT**
- **LUPRON DEPOT 7.5 MG INTRAMUSCULAR SYRINGE KIT**

Details

Criteria	Require a trial of Eligard (Step 1 drug) in the last 180 days when being utilized for the same medically accepted indication
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Riomet

Products Affected

- *metformin 500 mg/5 ml oral solution*

Details

Criteria	Require a 1 month trial of generic metformin IR tablets in the last 90 days -OR- documentation supporting the inability to swallow or difficulty swallowing tablets containing metformin
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rytary

Products Affected

- **RYTARY 23.75 MG-95 MG
CAPSULE,EXTENDED RELEASE**
- **RYTARY 36.25 MG-145 MG
CAPSULE,EXTENDED RELEASE**
- **RYTARY 48.75 MG-195 MG
CAPSULE,EXTENDED RELEASE**
- **RYTARY 61.25 MG-245 MG
CAPSULE,EXTENDED RELEASE**

Details

Criteria	Require a trial of generic carbidopa/levodopa product (Step 1 drug) in the last 90 days
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suboxone

Products Affected

- *buprenorphine 2 mg-naloxone 0.5 mg sublingual tablet*
- *buprenorphine 8 mg-naloxone 2 mg sublingual tablet*

Details

Criteria	Require a 1 month trial of Zubsolv (Step 1 drug) in the last 90 days
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Topical Antifungals

Products Affected

- *naftifine 1 % topical cream*
- *naftifine 2 % topical cream*
- **NAFTIN 2 % TOPICAL GEL**
- **OXISTAT 1 % LOTION**

Details

Criteria	Require a 1 month trial of generic econazole cream and one of the following: generic ketoconazole cream or ketoconazole shampoo (Step 1 drugs), in the last 180 days
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Topical Psoriasis

Products Affected

- *calcipotriene-betamethasone 0.005 %-0.064 % topical suspension*
- *calcipotriene-betamethasone 0.005 %-0.064 % topical ointment*
- *calcitriol 3 mcg/gram topical ointment*
- *calcipotriene-betamethasone 0.005 %-0.064 % topical suspension*

Details

Criteria	Require a 1 month trial of generic calcipotriene cream, ointment or solution (Step 1 drug) in the last 90 days
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LUPRON DEPOT 11.25 MG (3 MONTH) INTRAMUSCULAR SYRINGE KIT.....	5	VYVANSE 60 MG CAPSULE.....	1
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