

Prior Authorization Criteria
Givlaari (givosiran)

All requests for Givlaari (givosiran) require a prior authorization and will be screened for medical necessity and appropriateness using the criteria listed below.

For all requests for Givlaari (givosiran) all of the following criteria must be met:

Coverage may be provided with a diagnosis of acute hepatic porphyria (AHP) and the following criteria is met:

- Member must be 18 years of age or older
- Must be prescribed by or in consultation with a provider who specializes in porphyria (i.e. hematologist, hepatologist, gastroenterologist)
- Member must have active disease defined as having at least 2 documented porphyria attacks requiring hospitalization, urgent care visits, or IV hemin administration within the last 6 months.
- Documentation the members has had elevated urinary or plasma porphobilinogen (PBG) or aminolevulinic acid (ALA) levels with the past year (reference range must be provided)
- The requested dose and frequency is in accordance with FDA-approved labeling, nationally recognized compendia, and/or evidence-based practice guidelines
- **Initial Duration of Approval:** 6 months
- **Reauthorization criteria**
 - Documentation from the prescriber indicating stabilization or improvement in the member's condition since starting the medication.
- **Reauthorization Duration of Approval:** 12 months

Coverage may be provided for any non-FDA labeled indication if it is determined that the use is a medically accepted indication supported by nationally recognized pharmacy compendia or peer-reviewed medical literature for treatment of the diagnosis(es) for which it is prescribed. These requests will be reviewed on a case by case basis to determine medical necessity.

When criteria are not met, the request will be forwarded to a Medical Director for review. The physician reviewer must override criteria when, in their professional judgment, the requested medication is medically necessary.

**GIVLAARI (GIVOSIRAN)
PRIOR AUTHORIZATION FORM**

Please complete and fax all requested information below including any progress notes, laboratory test results, or chart documentation as applicable to Highmark Wholecare Pharmacy Services. **FAX:** (888) 245-2049

If needed, you may call to speak to a Pharmacy Services Representative. **PHONE:** (800) 392-1147 Mon – Fri 8:30am to 5:00pm

PROVIDER INFORMATION

| | |
|----------------------|-----------------|
| Requesting Provider: | Provider NPI: |
| Provider Specialty: | Office Contact: |
| State license #: | Office NPI: |
| Office Address: | Office Phone: |
| | Office Fax: |

MEMBER INFORMATION

| | |
|--------------|------------------------|
| Member Name: | DOB: |
| Member ID: | Member weight: Height: |

REQUESTED DRUG INFORMATION

| | |
|---|--------------------|
| Medication: | Strength: |
| Directions: | Quantity: Refills: |
| Is the member currently receiving requested medication? <input type="checkbox"/> Yes <input type="checkbox"/> No Date Medication Initiated: | |

Billing Information

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| This medication will be billed: <input type="checkbox"/> at a pharmacy OR <input type="checkbox"/> medically, JCODE: |
| Place of Service: <input type="checkbox"/> Hospital <input type="checkbox"/> Provider's office <input type="checkbox"/> Member's home <input type="checkbox"/> Other |

Place of Service Information

| | |
|----------|--------|
| Name: | NPI: |
| Address: | Phone: |

MEDICAL HISTORY (Complete for ALL requests)

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| Diagnosis: | ICD Code: |
| Has the member had 2 or more porphyria attacks in the last 6 months that required at least one of the following: a hospitalization, an urgent care visit, or IV hemin administration? <input type="checkbox"/> Yes <input type="checkbox"/> No | |
| Please provide one of the following labs and reference range: | |
| Urinary or plasma porphobilinogen (PBG): _____ reference range: _____ date taken: _____ | |
| Aminolevulinic acid level (ALA): _____ reference range: _____ date taken: _____ | |

CURRENT or PREVIOUS THERAPY

| Medication Name | Strength/ Frequency | Dates of Therapy | Status (Discontinued & Why/Current) |
|-----------------|---------------------|------------------|-------------------------------------|
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REAUTHORIZATION

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| Has the member experienced a stabilization or improvement with treatment? <input type="checkbox"/> Yes <input type="checkbox"/> No |
| Please describe: |

SUPPORTING INFORMATION or CLINICAL RATIONALE

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| |

Prescribing Provider Signature

Date

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Updated: 05/2023
PARP Approved:05/2023