

PHARMACY COVERAGE GUIDELINE

SABRIL® (vigabatrin) oral

Vigabatrin oral

VIGADRONE™ (vigabatrin) oral

VIGAFYDE™ (vigabatrin) oral

VIGPODER™ (vigabatrin) oral

This Pharmacy Coverage Guideline (PCG):

- Provides information about the reasons, basis, and information sources we use for coverage decisions
- Is not an opinion that a drug (collectively “Service”) is clinically appropriate or inappropriate for a patient
- Is not a substitute for a provider’s judgment (Provider and patient are responsible for all decisions about appropriateness of care)
- Is subject to all provisions e.g. (benefit coverage, limits, and exclusions) in the member’s benefit plan; and
- Is subject to change as new information becomes available.

Scope

- This PCG applies to Commercial and/or Marketplace plans
- This PCG does not apply to the Federal Employee Program, Medicare Advantage, Medicaid or members of out-of-state Blue Cross and/or Blue Shield Plans

Instructions & Guidance

- To determine whether a member is eligible for the Service, read the entire PCG.
- This PCG is used for FDA approved indications including, but not limited to, a diagnosis and/or treatment with dosing, frequency, and duration.
- Use of a drug outside the FDA approved guidelines, refer to the appropriate Off-Label Use policy.
- The “Criteria” section outlines the factors and information we use to decide if the Service is medically necessary as defined in the Member’s benefit plan.
- The “Description” section describes the Service.
- The “Definition” section defines certain words, terms or items within the policy and may include tables and charts.
- The “Resources” section lists the information and materials we considered in developing this PCG
- **We do not accept patient use of samples as evidence of an initial course of treatment, justification for continuation of therapy, or evidence of adequate trial and failure.**
- Information about medications that require prior authorization is available at www.azblue.com/pharmacy. You must fully complete the [request form](#) and provide chart notes, lab workup and any other supporting documentation. The prescribing provider must sign the form. Fax the form to BCBSAZ Pharmacy Management at (602) 864-3126 or email it to Pharmacyprecert@azblue.com.

Medical Necessity Requirements for **SABRIL** (vigabatrin), **Vigabatrin** generic, **VIGADRONE** (vigabatrin), **VIGAFYDE** (vigabatrin), **VIGPODER** (vigabatrin)

Criteria for Initial Therapy:

Indication

- Refractory Complex Partial Seizures and will be used as adjunctive therapy in individual who has inadequately responded to several alternative treatments

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- Infantile Spasms and will be used as monotherapy

Age Requirement

- 1 month to 2 years of age for infantile spasms
- 2 years or older for refractory partial complex seizures

Baseline Clinical Evaluation

- Baseline vision assessment before starting treatment. [**Note:** This requirement is waived if provider, member, and pharmacy are enrolled in the Risk Evaluation and Mitigation Strategies (REMS) program]

Alternative Therapies

- **Refractory Complex Partial Seizures:** Failure, contraindication, intolerance, or not a candidate for **BOTH** of the following:
 - TWO broad spectrum agents
 - Generic vigabatrin
- **Infantile Spasms:** Failure, contraindication, intolerance, or not a candidate for **BOTH** of the following:
 - ONE glucocorticoid (prednisone, prednisolone, methylprednisolone, or dexamethasone)
 - Generic vigabatrin

Safety

- Creatinine clearance is greater than 10 mL/min

Documentation Requirements

- A completed request form must be submitted, including:
 - Chart notes
 - Lab results (including creatinine clearance)
 - Supporting clinical documentation

Initial Therapy Criteria Approval Duration:

- 3 months OR end of plan year. Continuation requires documentation of significant clinical benefit
- **Note:** Because of the risk of visual loss, Sabril, Vigadrone, Vigafyde, Vigpoder, and generic vigabatrin should be withdrawn from individuals with refractory complex partial seizures who fail to show substantial clinical benefit within 3 months of initiation and within 2 to 4 weeks of initiation for individuals with infantile spasms, or sooner if treatment failure becomes obvious with either condition

Criteria for Continuation of Therapy (renewal therapy)

Note: Manufacturer assistance (e.g., coupons, samples, etc.) are not considered for continuation of therapy

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Clinical Response

- **Refractory Complex Partial Seizures:** Achieves and maintains **ONE** of the following:
 - At least a 50 percent reduction in frequency of complex partial seizures
 - At least a 50 percent reduction in secondary generalized seizures
- **Infantile Spasms:** Achieves and maintains **ONE** of the following:
 - Spasm freedom as assessed by caregiver
 - No spasms or hypersarrhythmia during closed circuit television electroencephalography (CCTV EEG)

Adherence

- Adherence to the prescribed therapy regimen has been documented

Safety

- Verification that Prescriber, Patient, and Pharmacy are enrolled in the REMS program
- Creatinine clearance is greater than 10 mL/min
- No development of adverse effects such as:
 - Vision loss
 - Neurotoxicity
 - Suicidal thoughts or behaviors
 - Emerging or worsening depression
 - Unusual changes in mood or behavior
 - Peripheral neuropathy

Documentation Requirements

- Chart notes
- Supporting clinical documentation with evidence of improvement in given indication
- Lab values that confirm safe use

Continuation Therapy Criteria Approval Duration:

- 12 months OR end of plan year. Continuation requires documentation of significant clinical benefit

Criteria for Off-Label Use Requests:

Criteria for a request for non-FDA use or indication, treatment with dosing, frequency, or duration outside the FDA-approved dosing, frequency, and duration, refer to one of the following Pharmacy Coverage Guideline:

1. Off-Label Use of Non-Cancer Medications
2. Off-Label Use of Cancer Medications

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Description:

Sabril (vigabatrin), Vigadrone (vigabatrin), Vigpoder (vigabatrin), and generic vigabatrin are indicated as adjunctive therapy for adults and pediatric patients 2 years of age or older with refractory complex partial seizures who have inadequately responded to several alternative treatments, they are not indicated as a first line agent for complex partial seizures. They are also indicated as monotherapy for pediatric patients with infantile spasms 1 month to 2 years of age. Vigafyde (vigabatrin) is indicated as monotherapy for pediatric patients with infantile spasms 1 month to 2 years of age.

Epilepsy is a neurological disorder where brief disturbances in the electrical function of the brain result in seizures. These seizures may affect consciousness, bodily movements or sensations for a short time. There are several different types of seizure that occur in epilepsy including partial (affecting one area of the brain), generalized (affecting nerve cells throughout the brain), and unclassified.

Anti-epileptic drugs (AED) are effective in controlling seizures. There is insufficient evidence to conclude that one AED is superior to another in controlling partial and generalized seizures or in improving outcomes. The evidence is also insufficient to conclude that branded AED are more effective than generic AED in terms of reducing seizure frequency or improving outcomes. In addition, the evidence is insufficient to support any relevant negative outcome (such as increased seizure frequency, hospitalizations, and mortality) when switching from a branded to a generic medication. However, switching between different manufacturers could lead to variations in serum concentrations and it is suggested that prescription refills should be from the same manufacturer. The FDA maintains that there is no convincing evidence that people with epilepsy have less seizure control when taking generic medications.

All AED are associated with an increased risk of suicidal ideation and suicidal behavior when used in patients with epilepsy. While there is a high degree of variability in tolerability to AEDs, no specific AED is considered to be the safest or best tolerated. Adverse events are common to all AED and include confusion, dizziness, somnolence, ataxia, nausea, and vomiting. Individual AEDs are associated with serious, but rare adverse events. Sabril (vigabatrin) carries a boxed warning and has a REMS program for risk of irreversible vision loss. Practice guidelines suggest that choice of treatment should be individualized based on several factors such as drug effectiveness for the seizure type, patient age, concomitant medications, tolerability, safety, response to previous therapy, potential adverse effects of the drug, interactions with other medications, comorbid medical conditions, gender, lifestyle, patient preferences, and cost. Treatment should begin with a single agent with dose titration to achieve control of seizures or development of unacceptable side effects. If seizures persist, another agent is used as monotherapy; some recommend attempting a second alternative before using multiple drugs to control seizures. Achieving a seizure-free state is difficult and many patients may have to try multiple regimens and combination therapies to achieve control of seizures.

Partial seizures are divided into simple partial, complex partial, and partial seizures that evolve into secondary generalized seizures. The difference between simple and complex seizures is that during simple partial seizures, patients retain awareness; during complex partial seizures, they lose awareness.

Infantile spasms (also known as West's syndrome) is a rare epileptic disorder with main characteristics of infantile spasms, mental retardation, and hypsarrhythmia, a specific abnormal pattern detected by an electroencephalogram

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(EEG) that is described as slow waves of high voltage and random pattern of spikes that vary in duration and location. Infantile spasms are characterized by sudden jerking and bending forward of the body, followed by stiffening of the body. Spasms usually last around 1-5 seconds but can range from 2-500 spasms at any given time.

Use of Sabril (vigabatrin), Vigadrone (vigabatrin), Vigafyde (vigabatrin), Vigpoder (vigabatrin), and vigabatrin are subject to a Risk Evaluation and Mitigation Strategies (REMS) program that requires provider, patient, and dispensing pharmacy be enrolled into the program. Only providers and Pharmacies enrolled into the REMS may prescribe and dispense the drug, respectively, to individuals who are also in the program. A REMS program attempts to manage known or potentially serious risks associated with a drug product and is required by the Food and Drug Administration (FDA) for some drugs to ensure that the benefits of a drug outweigh its risks.

The precise mechanism of vigabatrin’s anti-seizure effect is unknown, but it is believed to be the result of its action as an irreversible inhibitor of γ -aminobutyric acid transaminase (GABA-T), the enzyme responsible for the metabolism of the inhibitory neurotransmitter GABA. Inhibition of GABA-T results in increased levels of GABA in the central nervous system.

Definitions:

Sabril (vigabatrin), Vigadrone (vigabatrin), Vigafyde (vigabatrin) Vigpoder (vigabatrin), vigabatrin REMS items:

- Enrollment and agreement information
- Ophthalmologic assessment requirements
- Treatment initiation information
- Treatment maintenance information
- Pharmacy requirements and responsibilities

Therapeutic spectrum of antiseizure drugs:

<p>Broad spectrum: Drugs used to treat a broad range of seizure types (both focal and generalized onset)</p>	<p>Narrow spectrum (focal): Drugs used primarily for focal-onset seizures (including focal evolving to bilateral convulsive seizures*)</p>	<p>Narrow spectrum (absence): Absence seizures only (a type of generalized seizure)</p>
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Brivaracetam Clobazam Felbamate Lamotrigine§ Levetiracetam Perampanel Rufinamide Topiramate Valproate Zonisamide	Carbamazepine¶ Cenobamate EslicarbazepineΔ GabapentinΔ Lacosamide OxcarbazepineΔ Phenobarbital¶ Phenytoin¶ Pregabalin Primidone¶ Stiripentol TiagabineΔ VigabatrinΔ	Ethosuximide
<p>* Previously referred to as secondary generalized seizures ¶ Some evidence of efficacy for generalized-onset tonic-clonic seizures, but may also worsen certain generalized seizure types § May worsen or precipitate myoclonic seizures Δ Potential to worsen certain generalized seizure types</p>		

Resources:

Sabril (vigabatrin) tablet and powder for oral solution product information, revised by Lundbeck Pharmaceuticals, LLC. 10-2021. Available at DailyMed <http://dailymed.nlm.nih.gov>. Accessed May 13, 2025.

Vigafyde oral solution product information, revised by Upsher-Smith Laboratories, LLC. 07-2024. Available at DailyMed <http://dailymed.nlm.nih.gov>. Accessed May 13, 2025.

Vigabatrin tablet product information, revised by Amneal Pharmaceuticals. 12-2022. Available at DailyMed <http://dailymed.nlm.nih.gov>. Accessed May 13, 2025.

Vigabatrin powder for oral solution product information, revised by Camber Pharmaceuticals, Inc. 04-2022. Available at DailyMed <http://dailymed.nlm.nih.gov>. Accessed May 13, 2025.

Vigadrone (vigabatrin) tablet product information, revised by Upsher-Smith Laboratories, LLC. 03-2023. Available at DailyMed <http://dailymed.nlm.nih.gov>. Accessed May 13, 2025.

Vigadrone (vigabatrin) powder for oral solution product information, revised by Upsher-Smith Laboratories, LLC. 01-2024. Available at DailyMed <http://dailymed.nlm.nih.gov>. Accessed May 13, 2025.

Vigpoder (vigabatrin) powder for oral solution product information, revised by Pyros Pharmaceuticals, Inc. 07-2023. Available at DailyMed <http://dailymed.nlm.nih.gov>. Accessed May 13, 2025.

Takacs DS, Katayayan A. Infantile epileptic spasms syndrome: Clinical features and diagnosis. In: UpToDate, Nordli DR, Dashe JF (Eds), UpToDate, Waltham MA.: UpToDate Inc. Available at <http://uptodate.com>. Literature current through May 2025. Topic last updated August 12, 2024. Accessed June 19, 2025.

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Takacs DS, Katayan A. Infantile epileptic spasms syndrome: Management and prognosis. In: UpToDate, Nordli DR, Dashe JF (Eds), UpToDate, Waltham MA.: UpToDate Inc. Available at <http://uptodate.com>. Literature current through May 2025. Topic last updated August 10, 2023. Accessed June 19, 2025.

Schachter SC. Overview of the management of epilepsy in adults. In: UpToDate, Garcia P, Dashe JF (Eds), UpToDate, Waltham MA.: UpToDate Inc. Available at <http://uptodate.com>. Literature current through May 2025. Topic last updated April 03, 2025. Accessed June 19, 2025.

Karceski S, Shih T Initial treatment of epilepsy in adults. In: UpToDate, Garcia P, Dashe JF (Eds), UpToDate, Waltham MA.: UpToDate Inc. Available at <http://uptodate.com>. Literature current through May 2025. Topic last updated October 22, 2024. Accessed June 19, 2025.

Sirven JI. Evaluation and management of drug resistant epilepsy. In: UpToDate, Garcia P, Dashe JF (Eds), UpToDate, Waltham MA.: UpToDate Inc. Available at <http://uptodate.com>. Literature current through May 2025. Topic last updated November 04, 2024. Accessed June 19, 2025.