

**Clinical Policy: Lacosamide (Vimpat)** 

Reference Number: HIM.PA.49

Effective Date: 12.01.14 Last Review Date: 08.17

Line of Business: Health Insurance Marketplace Revision Log

See <u>Important Reminder</u> at the end of this policy for important regulatory and legal information.

### **Description**

Lacosamide (Vimpat®) is an anticonvulsant.

### **FDA** Approved Indication(s)

Vimpat is indicated for the treatment of partial-onset seizures in patients 4 years of age and older.

As the safety of Vimpat injection has not been established in pediatric patients, Vimpat injection is indicated for the treatment of partial-onset seizures only in adult patients (17 years of age and older).

### Policy/Criteria

Provider <u>must</u> submit documentation (which may include office chart notes and lab results) supporting that member has met all approval criteria

### I. Initial Approval Criteria

### A. Partial Onset Seizures (must meet all):

- 1. Diagnosis of partial seizures;
- 2. Age  $\geq$  4 years;
- 3. Failure of 3 formulary alternatives (e.g., carbamazepine, diazepam, felbamate, gabapentin, lamotrigine, valproic acid, etc.) unless all are contraindicated or clinically significant adverse effects are experienced;
- 4. Dose does not exceed (a or b):
  - a. Age  $\geq$  17 years or weight  $\geq$  50 kg: 400 mg/day;
  - b. Age 4 to < 17 years (i or ii):
    - i. Weight 30 kg to < 50 kg: 8 mg/kg/day;
    - ii. Weight 11 kg to < 30 kg: 12 mg/kg/day.

### **Approval duration: 12 months**

### **B.** Other diagnoses/indications

1. Refer to HIM.PHAR.21 if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized).

#### **II.** Continued Therapy

### A. Partial Onset Seizures (must meet all):

1. Currently receiving medication via Centene benefit, or documentation supports that member is currently receiving Vimpat for partial-onset seizures;

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- 2. Documentation of positive response to therapy;
- 3. If request is for a dose increase, new dose does not exceed (a or b):
  - a. Age  $\geq$  17 years or weight  $\geq$  50 kg: 400 mg/day;
  - b. Age 4 to < 17 years (i or ii):
    - i. Weight 30 kg to < 50 kg: 8 mg/kg/day;
    - ii. Weight 11 kg to < 30 kg: 12 mg/kg/day.

## **Approval duration: 12 months**

### **B.** Other diagnoses/indications (must meet 1 or 2):

1. Currently receiving medication via Centene benefit and documentation supports positive response to therapy.

Approval duration: Duration of request or 12 months (whichever is less); or

2. Refer to HIM.PHAR.21if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized).

### III. Diagnoses/Indications for which coverage is NOT authorized:

**A.** Non-FDA approved indications, which are not addressed in this policy, unless there is sufficient documentation of efficacy and safety according to the off-label use policy – HIM.PHAR.21 or evidence of coverage documents.

### IV. Appendices/General Information

Appendix A: Abbreviation/Acronym Key FDA: Food and Drug Administration

Appendix B: Therapeutic Alternatives N/A

#### V. References

1. Vimpat Prescribing Information. Smyrna, GA: UCB, Inc.; November 2017. Available at: www.vimpat.com. Accessed November 15, 2017.

Reviews, Revisions, and Approvals	Date	P&T Approval Date
Changed guideline to new format.	08.16	08.16
Converted to new template	04.17	08.17
Added max dose.		
Updated references.		
Dosing updated per FDA expanded indication for	11.15.17	
pediatric patients. Age requirement added per safety		
guidance endorsed by Centene Medical Affairs.		

### **Important Reminder**

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program

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approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. The Health Plan makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved. "Health Plan" means a health plan that has adopted this clinical policy and that is operated or administered, in whole or in part, by Centene Management Company, LLC, or any of such health plan's affiliates, as applicable.

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This clinical policy is effective as of the date determined by the Health Plan. The date of posting may not be the effective date of this clinical policy. This clinical policy may be subject to applicable legal and regulatory requirements relating to provider notification. If there is a discrepancy between the effective date of this clinical policy and any applicable legal or regulatory requirement, the requirements of law and regulation shall govern. The Health Plan retains the right to change, amend or withdraw this clinical policy, and additional clinical policies may be developed and adopted as needed, at any time.

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