

Epidiolex (Cannabidiol) Prior Authorization of Benefits Form

CONTAINS CONFIDENTIAL PATIENT INFORMATION

Complete form in its entirety and fax to: Prior Authorization of Benefits Center at 1-844-512-9004 or Provider Help Desk 1-800-454-3730

1. Patient information		2. Physician information	
Patient name: Patient ID #: Patient DOB: Date of Rx: Patient phone #: Patient email address:		Prescribing physician: Physician address: Physician phone #: Physician fax #: Physician specialty: Physician DEA: Physician NPI #: Physician email address:	
3. Medication	4. Strength	5. Directions	6. Quantity per 30 days
			Specify:

7. Diagnosis:

Epidiolex

8. Approval criteria: (Check all boxes that apply. Note: Any areas not filled out are considered not applicable to your patient and may affect the outcome of this request.)

Prior authorization (PA) is required for cannabidiol (Epidiolex). Payment will be considered under the following conditions:

1) Patient meets the FDA approved age; and

2) Baseline serum transaminases (ALT and AST) and total bilirubin levels have been obtained prior to initiating therapy (attach results); and

- 3) A diagnosis of Lennox-Gastaut syndrome with documentation of an adequate trial and inadequate response with at least two concomitant antiepileptic drugs (AEDs) from the following: valproic acid, lamotrigine, topiramate, felbamate, rufinamide, clobazam; or
- 4) A diagnosis of Dravet syndrome with documentation of an adequate trial and inadequate response with at least two concomitant AEDs from the following: clobazam, valproic acid, levetiracetam, topiramate; and
- 5) Is prescribed by or in consultation with a neurologist; and

6) The total daily dose does not exceed 20mg/kg/day.

If criteria for coverage are met, initial requests will be approved for three months. Additional PA requests will be considered when the following criteria are met:

•	ise to therapy (i.e. reduction in the frequency of seizures); and			
2) The total daily dose does not exc	eed 20mg/kg/day. when documented evidence is provided that use of these agents would be			
medically contraindicated.	when documented evidence is provided that use of these agents would be			
	Date obtained:			
Patient weight (kg): Date obtained: Is prescriber a neurologist?				
\Box Yes \Box No If no, note consultation wi	ith neurologist.			
	Physician name & phone:			
Have baseline serum transaminases (ALT and AST) and total bilirubin been obtained prior to initiating therapy?			
Yes (attach results)				
Lennox-Gastaut syndrome				
Document an adequate trial and inade	equate response with at least two concomitant AEDs from the following: valproic			
acid, lamotrigine, topiramate, felbama	ate, rufinamide, clobazam.			
Trial #1 drug name and dose:				
Trial dates:	Failure reason:			
Trial datas:	Failure reason:			
Trial dates:	Failure reason:			
Dravet syndrome				
Document an adequate trial and inade	equate response with at least two concomitant AEDs from the following:			
clobazam, valproic acid, levetiracetam	, topiramate.			
Trial #1 drug name and dose:				
Trial dates:	Failure reason:			
Trial #2 drug name and dose:				
Trial dates:	Failure reason:			
Renewals				
Document clinical response to therapy	/:			
 Patient weight (kg):	Date obtained:			
9. Physician signature				
Prescriber or authorized signature	Date			

Prior Authorization of Benefits is not the practice of medicine or the substitute for the independent medical judgment of a treating physician. Only a treating physician can determine what medications are appropriate for a patient. Please refer to the applicable plan for the detailed information regarding benefits, conditions, limitations and exclusions. The submitting provider certifies that the information provided is true, accurate and complete and the requested services are medically indicated and necessary to the health of the patient.

Note: Payment is subject to member eligibility. Authorization does not guarantee payment.