



**MEDICARE ADVANTAGE
PRIOR AUTHORIZATION REQUEST FORM**

Epidiolex - Medicare

Phone: 215-991-4300

Fax back to: 866-371-3239

Jefferson Health Plans manages the pharmacy drug benefit for your patient. Certain requests for coverage require review with the prescribing physician. Please answer the following questions and fax this form to the number listed above.

PLEASE NOTE: Any information (patient, prescriber, drug, labs) left blank, illegible, or not attached WILL delay the review process.

Member Name:	Prescriber Name:	
Member Number:	Fax:	Phone:
Date of Birth:	Office Contact:	
Line of Business: <input type="checkbox"/> Medicare Advantage	NPI:	State Lic ID:
Address:	Address:	
City, State ZIP:	City, State ZIP:	
Primary Phone:	Specialty/facility name (if applicable):	

REQUEST FOR EXPEDITED REVIEW: By checking this box and signing below, I certify that applying the 72 hour standard review timeframe may seriously jeopardize the life or health of the enrollee or the enrollee's ability to regain maximum function.

Drug Name:	
Strength:	
Directions / SIG:	

**Please attach any pertinent medical history including labs and information for this member that may support approval.
Please answer the following questions and sign.**

Q1. Does the patient have a hypersensitivity to cannabidiol or any of the ingredients in the product?

Yes

No

Q2. Does the patient have a documented diagnosis of Dravet syndrome (DS) or Lennox-Gastaut syndrome (LGS) or Tuberous Sclerosis Complex (TSC)?

Yes

No

Q3. Is Epidiolex being prescribed by a neurologist?

Yes

No

Q4. Is the patient 1 year of age or older?

Yes

No

Q5. Prior to initiation of therapy, are baseline serum transaminases (ALT and AST) and total bilirubin attached, and will these labs be monitored periodically during therapy?

Yes

No

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Member Name:	Prescriber Name:
Q6. Has the patient failed to become seizure-free with at least 2 antiepileptic drugs (specify drugs tried)? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Q7. Will Epidiolex be used as adjunctive therapy with other antiepileptic drug(s) (provide name of drug or drugs)? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Q8. Is the requested Epidiolex dose in accordance with FDA-approved labeled dose not exceeding 20 mg/kg/day for treatment of seizures associated with Lennox-Gastaut Syndrome and Dravet Syndrome or dose not exceeding 25 mg/kg/day for treatment of seizures associated with Tuberous Sclerosis Complex? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Q9. Requested Duration: <input type="checkbox"/> 12 Months <input type="checkbox"/> Other:	
Q10. Additional Information:	

Prescriber Signature

Date

v2025