

Livdelzi (seladelpar)
Prior Authorization Request Form
 Caterpillar Prescription Drug Benefit
 Phone: 877-228-7909 Fax: 800-424-7640

MEMBER'S LAST NAME: _____ **MEMBER'S FIRST NAME:** _____

Instructions: Please fill out all applicable sections completely and legibly. Attach any additional documentation that is important for the review (e.g., chart notes or lab data, to support the authorization request). Information contained in this form is Protected Health Information under HIPAA.

URGENT

MEMBER INFORMATION		
LAST NAME:	FIRST NAME:	
PHONE NUMBER:	DATE OF BIRTH:	
STREET ADDRESS:		
CITY:	STATE:	ZIP CODE:
PATIENT INSURANCE ID NUMBER:		

MALE **FEMALE** **HEIGHT (IN/CM):** _____ **WEIGHT (LB/KG):** _____ **ALLERGIES:** _____

IF YOU ARE NOT THE PATIENT OR THE PRESCRIBER, YOU WILL NEED TO SUBMIT A PHI DISCLOSURE AUTHORIZATION FORM WITH THIS REQUEST WHICH CAN BE FOUND AT THE FOLLOWING LINK: [PRIMETHERAPEUTICS.COM/NOPP](https://www.primetherapeutics.com/nopp)

PATIENT'S AUTHORIZED REPRESENTATIVE (IF APPLICABLE): _____
AUTHORIZED REPRESENTATIVE'S PHONE NUMBER: _____

PRESCRIBER INFORMATION			
LAST NAME:	FIRST NAME:		
PRESCRIBER SPECIALTY:	EMAIL ADDRESS:		
NPI NUMBER:	DEA NUMBER:		
PHONE NUMBER:	FAX NUMBER:		
STREET ADDRESS:			
CITY:	STATE:	ZIP CODE:	
REQUESTER (if different than prescriber):	OFFICE CONTACT PERSON:		

MEDICATION OR MEDICAL DISPENSING INFORMATION			
MEDICATION NAME:			
DOSE/STRENGTH:	FREQUENCY:	LENGTH OF THERAPY/REFILLS:	QUANTITY:
<input type="checkbox"/> NEW THERAPY <input type="checkbox"/> RENEWAL IF RENEWAL: DATE THERAPY INITIATED:			
DURATION OF THERAPY (SPECIFIC DATES):			

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1. HAS THE PATIENT TRIED ANY OTHER MEDICATIONS FOR THIS CONDITION? <input type="checkbox"/> YES (if yes, complete below) <input type="checkbox"/> NO		
MEDICATION/THERAPY (SPECIFY DRUG NAME AND DOSAGE):	DURATION OF THERAPY (SPECIFY DATES):	RESPONSE/REASON FOR FAILURE/ALLERGY:
2. LIST DIAGNOSES:		ICD-10:
<input type="checkbox"/> Primary biliary cholangitis (PBC) (previously known as primary biliary cirrhosis) <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s):		
3. REQUIRED CLINICAL INFORMATION: PLEASE PROVIDE ALL RELEVANT CLINICAL INFORMATION TO SUPPORT A PRIOR AUTHORIZATION.		
<u>Initial:</u> Will the patient be using the drug as a part of the clinical trial? <input type="checkbox"/> Yes <input type="checkbox"/> No		
Is the medication being prescribed by, or in consultation with, a hepatologist or gastroenterologist? <input type="checkbox"/> Yes <input type="checkbox"/> No		
Was the diagnosis of PBC was confirmed by any of the following criteria? <input type="checkbox"/> Yes <input type="checkbox"/> No <i>Please select all that apply.</i>		
<ul style="list-style-type: none">▪ Biochemical evidence of cholestasis with elevation of alkaline phosphatase (ALP) level for at least 6 months duration.▪ Presence of antimitochondrial antibodies (AMA) (titer >1:40 by immunofluorescence or immunoenzymatic reactivity) or PBC-specific antinuclear antibodies (ANA) (e.g., anti-gp210, antisp100).▪ Histologic evidence of PBC on liver biopsy (e.g., non-suppurative inflammation and destruction of interlobular and septal bile ducts)		
Does the patient has elevated serum ALP level prior to starting therapy with the requested agent? <input type="checkbox"/> Yes <input type="checkbox"/> No		
Does the patient have an alkaline phosphatase (ALP) level greater than 1.67 times the upper limit of normal? <input type="checkbox"/> Yes <input type="checkbox"/> No		
Does the patient have an aspartate aminotransferase (AST) less than 3 times time the upper limit of normal? <input type="checkbox"/> Yes <input type="checkbox"/> No		
Does the patient have an aalanine aminotransferase (ALT) less than 3 times the upper limit of normal? <input type="checkbox"/> Yes <input type="checkbox"/> No		

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Does the patient have an total bilirubin less than 2 times times the upper limit of normal?

Yes No

Has the patient has been receiving ursodiol therapy for at least 1 year and has had an inadequate response (documentation required)? Yes No

If no, was the patient unable to tolerate ursodiol therapy (documentation required)? Yes No

Does the patient have hepatic decompensation? Yes No

Have all other causes of chronic liver disease have been ruled out? Yes No

Will Livdelzi (seladelpar) be used in combination with Iqirvo (elafibranror)? Yes No

Renewal:

Has the patient had at least a 15% reduction in serum ALP level (documentation required)?

Yes No

Did the patient's ALP level decrease to 1.67 times upper limit of normal (documentation required)?

Yes No

Did the patient's total bilirubin decrease to less than or equal to upper limit of normal (documentation required)? Yes No

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: Not all drugs/diagnosis are covered on all plans. This request may be denied unless all required information is received.

ATTESTATION: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan, insurer, Medical Group or its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.

Prescriber Signature or Electronic I.D. Verification: _____ **Date:** _____

CONFIDENTIALITY NOTICE: The documents accompanying this transmission contain confidential health information that is legally privileged. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution, or action taken in reliance on the contents of these documents is strictly prohibited. If you have received this information in error, please notify the sender immediately (via return FAX) and arrange for the return or destruction of these documents.

FAX THIS FORM TO: 800-424-7640

MAIL REQUESTS TO: Prime Therapeutics Management Prior Authorization Program

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