

Request for Prior Authorization



MARALIXIBAT (LIVMARLI)

Provider Help Desk 1 (844) 236-1464

FAX Completed Form To 1 (877) 733-3195

(PLEASE PRINT – ACCURACY IS IMPORTANT)

IA Medicaid Member ID #	Patient name	DOB	
Patient address			
Provider NPI	Prescriber name	Phone	
Prescriber address		Fax	
Pharmacy name	Address	Phone	
Prescriber must complete all information above. It must be legible, correct, and complete or form will be returned.			
Pharmacy NPI	Pharmacy fax	NDC	
	d for maralixibat (Livmarli). Requests for ence is provided that the use of the prefe		

contraindicated. Payment will be considered for an FDA approved or compendia indicated diagnosis for the requested drug when the following conditions are met:

- 1) Request adheres to all FDA approved labeling for requested drug and indication, including age, dosing, contraindications, warnings and precautions, drug interactions, and use in specific populations; and
- 2) Patient has a diagnosis of Alagille syndrome (ALGS) confirmed by genetic testing demonstrating a JAGI or NOTCH2 mutation or deletion; and
- 3) Patient has cholestasis with moderate to severe pruritis; and
- 4) Is prescribed by or consultation with a hepatologist, gastroenterologist, or a prescriber who specializes in ALGS; and
- 5) Documentation of previous trials and therapy failures, at a therapeutic dose, with at least two of the following agents:
 - a. Ursodeoxycholic acid (ursodiol)
 - b. Cholestyramine
 - c. Rifampin; and
- 6) Patient's current weight in kilograms (kg) is provided.

The required trials may be overridden when documented evidence is provided that the use of these agents would be medically contraindicated.

If criteria for coverage are met, initial authorization will be given for 6 months to assess the response to treatment. Request for continuation of therapy will require documentation of an improvement in pruritis symptoms and patient's current weight in kg.

Non-Preferred

Livmarli

Strength

Dosage Instructions

Quantity

Days Supply

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Diagnosis (Attach copy of genetic testing):	
Prescriber Specialty: Hepatologist Gastroe Other (specify):	
If other, note consultation with hepatologist, gastroenterologist, o	or prescriber specializing in ALGS:
Consultation date:	
Physician name, specialty & phone:	
Does patient have cholestasis with moderate to sev	ere pruritis? 🗌 No 📋 Yes
Patient's current weight in kg:	
Document trials, at a therapeutic dose, with two of	the following agents:
Ursodeoxycholic acid (ursodiol) Trial:	
Dose:	Trial dates:
Failure reason:	
Cholestyramine Trial:	
Dose:	Trial dates:
Failure reason:	
Rifampin Trial:	
Dose:	Trial dates:
Failure reason:	
RenewalRequests:	
Patient's current weight in kg:	
Document an improvement in pruritis symptoms: _	

Attach lab results and other documentation as necessary.

Prescriber signature (Must match prescriber listed above.)	Date of submission

IMPORTANT NOTE: In evaluating requests for prior authorization the consultant will consider the treatment from the standpoint of medical necessity only. If approval of this request is granted, this does not indicate that the member continues to be eligible for Medicaid. It is the responsibility of the provider who initiates the request for prior authorization to establish by inspection of the member's Medicaid eligibility card and, if necessary by contact with the county Department of Human Services, that the member continues to be eligible for Medicaid.