



HEALTH PARTNERS PLANS
PRIOR AUTHORIZATION REQUEST FORM

Health Partners Plans

Bile Salts

Phone: 215-991-4300

Fax back to: 866-240-3712

Health Partners 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.

Form with fields: Patient Name, Prescriber Name, HPP Member Number, Date of Birth, Patient Primary Phone, Address, City, State ZIP, Line of Business, Drug Name, Quantity, Directions, Diagnosis Code, Diagnosis, Strength, Refills, Specialty Pharmacy.

HPP's maximum approval time is 12 months but may be less depending on the drug.

Please attach any pertinent medical history including labs and information for this member that may support approval.

Please answer the following questions and sign.

Questions Q1-Q7 regarding bile salt request, Ocaliva, and liver function tests.

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Patient Name:	Prescriber Name:
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<p>Q8. Does the patient have recent HDL-C monitoring?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>Q9. Does the patient have complete biliary obstruction?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>Q10. Does the patient have a history of therapeutic failure, contraindication or intolerance to the preferred bile salts (i.e., Cholbam, ursodiol)?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>Q11. Is Ocaliva prescribed by or in consultation with a hepatologist or gastroenterologist?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>Q12. Is the patient being treated for a diagnosis that is indicated in the U. S. Food and Drug Administration (FDA)-approved package insert OR a medically accepted indication?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>Q13. Is the diagnosis documented by medical history and laboratory results?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>Q14. Does the patient have documented baseline liver function tests, including AST, ALT, GGT, alkaline phosphatase, bilirubin, and INR?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>Q15. Does the patient have a documented baseline HDL-C?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>Q16. Does the patient have a documented history of therapeutic failure of optimally-titrated doses of ursodeoxycholic acid (UDCA)?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>Q17. Will Ocaliva be prescribed in combination with ursodeoxycholic acid (UDCA)?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>Q18. Does the patient have a contraindication or intolerance to ursodeoxycholic acid (UDCA)?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>Q19. Is this request for continuation of therapy with Cholbam?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>

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Patient Name: Prescriber Name:

Q20. Does the patient have documented improvement in liver function within the first 3 months of treatment?
Q21. Does the patient have documented AST, ALT, GGT, alkaline phosphatase, bilirubin and INR monitoring as recommended per prescribing information?
Q22. Does the patient have complete biliary obstruction, persistent clinical or laboratory indicators of worsening liver function or cholestasis?
Q23. Is Cholbam prescribed by or in consultation with a hepatologist or gastroenterologist?
Q24. Is the patient being treated for a condition that is indicated in the U.S. Food and Drug Administration (FDA)-approved package insert or a medically accepted indication?
Q25. Is the condition documented by medical history and laboratory results?
Q26. Will the patient's AST, ALT, GGT, alkaline phosphatase, bilirubin and INR be monitored according to the prescribing information?

Prescriber Signature

Date

Updated for 2020