

HEALTHY U CHIP

PRIOR AUTHORIZATION REQUEST FORM

XIFAXAN®

For authorization, please answer each question and fax this form PLUS chart notes back to the Healthy U CHIP Prior Authorization Department at 385-425-4052.

Failure to submit clinical documentation to support this request will result in a dismissal of the request.

If you have prior authorization questions, please call for assistance: 385-425-5094

Disclaimer: Prior Authorization request forms are subject to change in accordance with Federal and State notice requirements.

Date:	Member Name:	ID#:
DOB:	Gender:	Physician:
Office Phone:	Office Fax:	Office Contact:

Height/Weight:

Member must try formulary preferred drugs before a request for a non-preferred drug may be considered. If treatment with preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria.

Product being requested: Xifaxan® (rifaximin)

Dosing/Frequency: _____

If the request is for reauthorization, proceed to reauthorization section

Questions	Yes	No	Comments/Notes
HEPATIC ENCEPHALOPATHY			
1. Is the request for Hepatic Encephalopathy?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
2. Is the member 18 years of age or older?	<input type="checkbox"/>	<input type="checkbox"/>	
3. Is the member currently using or severely intolerant to lactulose?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D)			
1. Does the member have IBS-D with recurrent abdominal pain for at least 1 day/week in the last 3 months?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
2. Is the abdominal pain associated with at least two of the following: related to defecation, associated with a change in frequency of stool, associated with a change in form/appearance of stool?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
3. Is the prescriber a gastroenterologist?	<input type="checkbox"/>	<input type="checkbox"/>	
4. Has the member shown trial and failure to nutritional and/or behavioral modifications (lactose restricted diet, gluten-free diet, low carb diet, elimination of fermentable oligo-di-monosaccharides and polyols (FODMAPS), increased physical activity)?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
5. Has the member shown trial and failure or contraindication to an antidiarrheal (loperamide, diphenoxylate)?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation

6. Has the member shown trial and failure or contraindication/intolerance to a tricyclic antidepressant (imipramine, despiramine)?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
7. Has serologic testing been performed to rule out celiac disease?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
8. Does documentation show that fecal calprotectin and C-reactive protein have been checked to rule out inflammatory bowel disease?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
TRAVELER'S DIARRHEA			
1. Is the request for Traveler's Diarrhea?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
2. Is the member 12 years of age or older?	<input type="checkbox"/>	<input type="checkbox"/>	
3. Is E. coli the suspected pathogen?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
4. Has the member shown trial and failure or contraindication to a quinolone (e.g., ciprofloxacin, levofloxacin, ofloxacin)?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
SMALL INTESTINAL BACTERIAL OVERGROWTH (SIBO)			
1. Is the medication prescribed by, or in consultation with, a gastroenterologist?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
2. Does the member have a documented clinical diagnosis of symptomatic (bloating, flatulence, abdominal discomfort, chronic diarrhea) SIBO by one of the following: <ul style="list-style-type: none"> • Glucose or lactulose breath testing • Duodenal culture resulting in colony count $\geq 10^3$ CFU/mL 	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
3. Has the member show an inadequate clinical response to at least TWO of the following antibiotic treatment regimens or contraindication to all: <ul style="list-style-type: none"> • Ciprofloxacin • Metronidazole • Amoxicillin-clavulanic acid • Trimethoprim-sulfamethoxazole • Doxycycline or tetracycline 	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
4. Has the member shown an Inadequate clinical response to diet modifications (low carbohydrate diet, low fermentable oligosaccharides/disaccharides/monosaccharides/and polyols (FODMAP) diet)?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
REAUTHORIZATION			
1. If the request is for reauthorization of therapy for treatment of hepatic encephalopathy, does updated documentation show a positive clinical response from therapy, such as a decrease in fasting serum ammonia levels and mental status?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
2. If the request is for reauthorization of therapy for IBS-D, is the member responding to treatment?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
3. If the request is for reauthorization of therapy for traveler's diarrhea, did the member have improved symptoms after 24-48 hours of therapy? <i>Please note that there is a limit of three 14-day treatment courses.</i>	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation

What medications and/or treatment modalities have been tried in the past for this condition? Please document name of treatment, reason for failure, treatment dates, etc.

Additional information:

Physician's Signature:

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Policy: PHARM-CHIP-078
Origination Date: 07/01/2024
Reviewed/Revised Date:
Next Review Date:
Current Effective Date: 07/01/2024

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