

# Rifaximin (Xifaxan®) and Rifamycin (Aemcolo®)

**Goal(s):**

- Promote use that is consistent with medical evidence and product labeling.

**Length of Authorization:**

- 3 days for traveler’s diarrhea caused by non-invasive strains of *E.Coli* for rifaximin or rifamycin.
- Up to 12 months for hepatic encephalopathy for rifaximin.

**Requires PA:**

- Rifaximin and Rifamycin

**Covered Alternatives:**

- Current PMPDP preferred drug list per OAR 410-121-0030 at [www.orpdl.org](http://www.orpdl.org)
- Searchable site for Oregon FFS Drug Class listed at [www.orpdl.org/drugs/](http://www.orpdl.org/drugs/)

Approval Criteria		
1. What diagnosis is being treated?	Record ICD10 code.	
2. Is this an FDA approved indication and is the indication funded by OHP?	<b>Yes:</b> Go to #3	<b>No:</b> Pass to RPh. Deny; medical appropriateness
3. Is the diagnosis traveler’s diarrhea caused by non-invasive strains of E.Coli?	<b>Yes:</b> Go to #4	<b>No:</b> Go to # 6
4. Will the prescriber consider a change to a preferred product?  Message: <ul style="list-style-type: none"> <li>• Preferred products do not require a PA.</li> <li>• Preferred products are evidence-based reviewed for comparative effectiveness and safety by the Oregon Pharmacy &amp; Therapeutics Committee.</li> <li>• Preferred products for traveler’s diarrhea are dependent on traveler’s destination and resistance patterns in that area. Refer to <b>Table 1</b> for adult treatment recommendations.</li> </ul>	<b>Yes:</b> Inform prescriber of covered alternatives in class.	<b>No:</b> Go to # 5
5. Does the patient have a contraindication or allergy to azithromycin or ciprofloxacin?	<b>Yes:</b> Approve for 3 days	<b>No:</b> Pass to RPh Deny; medical appropriateness

Approval Criteria		
6. Is the request for rifaximin to prevent or treat hepatic encephalopathy?	<b>Yes:</b> Go to #7	<b>No:</b> Pass to RPh. Deny; not funded by OHP or for medical appropriateness
7. Is the patient currently managed with a regularly scheduled daily regimen of lactulose?	<b>Yes:</b> Go to #9	<b>No:</b> Go to #8
8. Does the patient have a contraindication to lactulose?	<b>Yes:</b> Go to #9	<b>No:</b> Pass to RPh Deny; medical appropriateness  Note: studies demonstrate effectiveness of rifaximin as add-on therapy to lactulose.
9. Is the patient currently prescribed a benzodiazepine drug?	<b>Yes:</b> Go to #10	<b>No:</b> Approve for up to 12 months
10. Is the patient tapering off the benzodiazepine?  Note: tapering process may be several months	<b>Yes:</b> Approve for up to 12 months	<b>No:</b> Pass to RPh. Deny; medical appropriateness  Note: studies explicitly excluded use of benzodiazepines and benzodiazepine-like drugs because of their risk for precipitating an episode of hepatic encephalopathy.

**Table 1. Acute diarrhea treatment recommendations for adults<sup>1</sup>**

Antibiotic	Dose	Treatment Duration
Levofloxacin	500 mg orally	Single dose - If symptoms not resolved after 24 hours, complete a 3 day course
Ciprofloxacin	750 mg orally <b>OR</b> 500 mg orally once a day	Single dose - If symptoms not resolved after 24 hours, complete a 3 day course  3-day course
Ofloxacin	400 mg orally	Single dose - If symptoms not resolved after 24 hours, complete a 3 day course
Azithromycin <sup>a,b</sup>	1000 mg orally <b>OR</b> 500 mg once a day	Single dose - If symptoms not resolved after 24 hours, complete a 3 day course  3-day course <sup>b</sup>

Rifaximin <sup>c</sup>	200 mg orally three times a day	3-days (in patients > 12 years old)
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- a. Use empirically as first-line in Southeast Asia and India to cover fluoroquinolone resistant *Campylobacter* or in other geographic areas if *Campylobacter* or resistant enterotoxigenic *E. coli* are suspected.
- b. Preferred regimen for dysentery or febrile diarrhea.
- c. Do not use if clinical suspicion for *Campylobacter*, *Salmonella*, *Shigella*, or other causes of invasive diarrhea.

1. Riddle MS, DuPont HL, Connor BA. ACG Clinical Guideline: Diagnosis, Treatment, and Prevention of Acute Diarrheal Infections in Adults. Am J Gastroenterol. 2016;111(5):602-622

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*P&T/DUR Review: 11/19 (DM), 7/15; 5/15 (AG)*

*Implementation: 1/1/20; 10/15; 8/15*