

OPTIMA HEALTH PLAN

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

Directions: The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; **fax to 1-800-750-9692.** No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. **If information provided is NOT complete, correct, or legible, the authorization can be delayed.**

Gastrointestinal (GI) Motility Drugs

| | | |
|---|--|---|
| Drug Requested (select drug below that applies): | | |
| PREFERRED | | |
| <input type="checkbox"/> Linzess [®] (linaclotide) | <input type="checkbox"/> Movantik [™] (naloxegol) | <input type="checkbox"/> Symproic [®] (naldemedine) |
| Non-Preferred | | |
| <input type="checkbox"/> Amitiza [®] (lubiprostone) | <input type="checkbox"/> Relistor [®] (methylnaltrexone bromide) | <input type="checkbox"/> Trulance [®] (plecanatide) |

DRUG INFORMATION: Complete **all** information below or authorization will be delayed.

Drug Name/Form/Strength: _____
Dosing Schedule: _____ **Length of Therapy:** _____
Diagnosis: _____ **ICD Code, if applicable:** _____

CLINICAL CRITERIA: Check below **ALL** that apply. **ALL** criteria **must** be met for approval. To support each line checked, **ALL** documentation, including lab results, diagnostics, chart notes, and/or OTC medication trials, **must** be provided or request will be denied.

| | |
|---|--|
| <input type="checkbox"/> Approval of Movantik [™] <u>OR</u> Symproic [®] | |
| <input type="checkbox"/> Member has a diagnosis of opioid-induced constipation and is being treated for chronic, non-cancer pain | |
| AND | |
| <input type="checkbox"/> Trial and failure, contraindication, or intolerance to <u>one</u> of the following generics: | |
| <input type="checkbox"/> lactulose | <input type="checkbox"/> polyethylene glycol (generic MiraLAX [®]) |

| | |
|---|--|
| <input type="checkbox"/> Approval of Relistor [®] | |
| <input type="checkbox"/> Member has a diagnosis of opioid-induced constipation is being treated for chronic, non-cancer pain | |
| AND | |
| <input type="checkbox"/> Trial and failure, contraindication, or intolerance to <u>one</u> of the following generics: | |
| <input type="checkbox"/> lactulose | <input type="checkbox"/> polyethylene glycol (generic MiraLAX [®]) |

AND

(Continued on next page)

- Member had a 30-day trial of Movantik™ **OR** Symproic®

Approval of Linzess®

- Diagnosis of **Chronic Idiopathic Constipation (CIC)** **OR** **Irritable Bowel Syndrome with Constipation (IBS-C)**

AND

- Trial and failure, contraindication, or intolerance to **one** of the following **generics**:

- | | |
|------------------------------------|---|
| <input type="checkbox"/> lactulose | <input type="checkbox"/> polyethylene glycol (generic MiraLAX®) |
|------------------------------------|---|

Approval of Trulance® - Oral, 3 mg once daily.

- Irritable Bowel Syndrome with Constipation (IBS-C)**

AND

- Trial and failure, contraindication, or intolerance to **one** of the following **generics**:

- | | |
|------------------------------------|---|
| <input type="checkbox"/> lactulose | <input type="checkbox"/> polyethylene glycol (generic MiraLAX®) |
|------------------------------------|---|

AND

- Trial and failure, contraindication, or intolerance to Linzess®

Diagnosis for Amitiza®.

- Chronic Idiopathic Constipation (CIC)** (Oral-24mcg twice daily)

OR

- Irritable Bowel Syndrome with Constipation (IBS-C)** (Females ≥ 18 years: Oral 8mcg twice daily)

OR

- Opioid-induced constipation** (Oral – 24mcg twice daily)

AND

- Trial and failure, contraindication, or intolerance to **one** of the following **generics**:

- | | |
|------------------------------------|--|
| <input type="checkbox"/> lactulose | <input type="checkbox"/> polyethylene glycol (generic Miralax) |
|------------------------------------|--|

AND

- Trial and failure, contraindication, or intolerance to **one** of the following **Preferred** brands (**requires prior authorization**):

- | | | |
|-----------------------------------|------------------------------------|------------------------------------|
| <input type="checkbox"/> Linzess® | <input type="checkbox"/> Movantik® | <input type="checkbox"/> Symproic® |
|-----------------------------------|------------------------------------|------------------------------------|

(Continued on next page; signature page **MUST** be attached to form)

(Signature page **MUST** be included with request)

****Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.****

Previous therapies will be verified through pharmacy paid claims or submitted chart notes.

Member Name: _____

Member Optima #: _____ Date of Birth: _____

Prescriber Name: _____

Prescriber Signature: _____ Date: _____

Office Contact Name: _____

Phone Number: _____ Fax Number: _____

DEA OR NPI #: _____

*Approved by Pharmacy and Therapeutics Committee: 10/15/2015
REVISED/UPDATED/REFORMATTED: 8/2/2019