	ETTER HEALTH® Policy/Guideline	♥aetna [™]		
Name: Proton Pump Inhibitors		tors Post Limit	Page:	1 of 3
Effective Date: 8/30/2024			Last Review Date: 7/2024	
Applies to:	⊠Illinois ⊠New Jersey ⊠Pennsylvania Kids	□Florida ⊠Maryland ⊠Virginia	⊠Florida Kids □Michigan □Kentucky PRMD	

Intent:

The intent of this policy/guideline is to provide information to the prescribing practitioner outlining the coverage criteria for Proton Pump Inhibitors Post Limit under the patient's prescription drug benefit.

Indication	AcipHex (rabeprazole)	AcipHex Sprinkles	Dexilant (dexlansoprazole)	Konvomep (omeprazole/	Nexium (esomeprazole)	Prevacid (lansoprazole)	Prilosec (omeprazole)	Protonix (pantoprazole)	Zegerid (omeprazole/
	((rabeprazole)	()	sodium bicarbonate)	()	(,	((sodium bicarbonate)
Short-term	✓					✓	✓		✓
treatment									
active									
duodenal ulcer									
H. pylori	✓				✓	✓	✓		
eradication									
reduce risk									
ulcer relapse									
Maintenance						✓			
healing									
duodenal									
ulcers									
Short-term				✓		✓	✓		✓
treatment									
gastric ulcer									
Short-term	✓	✓	✓		✓	✓	✓	✓	✓
treatment									
symptoms									
GERD									
Short-term	✓		✓		✓	✓	✓	✓	✓
treatment									
erosive									
esophagitis /									
GERD									
Maintenance	✓		✓		✓	✓	✓	✓	✓
healing									
erosive									
esophagitis									
Pathological	✓				✓	✓	✓	✓	
hypersecretory									
conditions									
Short-term						✓			
treatment									
NSAID-gastric									
ulcer									
Risk reduction					✓	✓			
of NSAID-									
gastric ulcer									
Risk reduction				✓					✓
upper GI bleed									Suspension
critically ill									

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Description:

Applicable Drug List:

Rabeprazole

Dexlansoprazole

Esomeprazole

Lansoprazole

Omeprazole

Pantoprazole

Omeprazole-Sodium Bicarbonate

Policy/Guideline:

The requested drug will be covered with prior authorization when the following criteria are met:

The requested drug is being prescribed for any of the following: A) Barrett's
esophagus as confirmed by biopsy, B) Hypersecretory syndrome, such as ZollingerEllison, confirmed with a diagnostic test

OR

 The requested drug is being prescribed for any of the following: A) Endoscopically verified peptic ulcer disease, B) Frequent and severe symptoms of chronic gastroesophageal reflux disease (GERD), C) Atypical symptoms or complications of GERD

OR

The patient is at high risk for gastrointestinal (GI) adverse events
 [Note: Risk factors for serious GI adverse events include, but are not limited to, the
 following: chronic nonsteroidal anti-inflammatory drug (NSAID) therapy, history of
 peptic ulcer disease and/or GI bleeding, treatment with oral corticosteroids,
 treatment with anticoagulants, poor general health status, or advanced age.]

Approval Duration and Quantity Restrictions:

Approval:

12 months

Quantity Level Limit:

Reference Formulary for drug specific quantity level limits

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