

## Proton Pump Inhibitors (PPIs) - Washington Prior Authorization Request Form

Please complete this <u>entire</u> form and fax it to: 866-940-7328. If you have questions, please call 800-310-6826. This form may contain multiple pages. Please complete all pages to avoid a delay in our decision.

Allow at least 24 hours for review.

Section A – Member Inform	ation									
First Name:	Last Name:			Member ID:						
Address:										
City:	State:	State:			ZIP Code:					
Phone:		DOB:	DOB:			Allergies:				
Primary Insurance Information (if any):										
Is the requested medication: □ New or □ Continuation of Therapy? If continuation, list start date:										
Is this patient currently ho	spitalized? 🗆	Yes □ No	If recently d	ischarged, list disch	narge d	date:				
Section B - Provider Inform	nation									
First Name:			Last Name:				M.D./D.O.			
Address:			City:		State:		ZIP code:			
Phone:	Fax:		NPI #:		Specia	Specialty:				
Office Contact Name / Fax atter	ntion to:									
Section C - Medical Informa	ation									
Medication:						Strength:				
Directions for use:						Quantity:				
Diagnosis (Please be specific &	s provide as much	information	as possible):			ICD-10 CODE:				
Is this member pregnant? ☐ Yes ☐ No										
		If yes,	what is this m	ember's due date?						
Section D - Previous Medic	ation Trials					Reason	for failure /			
			what is this m	ember's due date? Dates of Therapy	/		for failure / ntinuation			
Section D - Previous Medic	ation Trials				/					
Section D - Previous Medic	ation Trials				/					
Section D - Previous Medic	ation Trials				,					
Section D - Previous Medic	ation Trials				/					
Section D – Previous Medication Name  Medication Name  Section E – Additional infor	Strength Strength	Dire	ctions	Dates of Therapy	uld not	discor	ntinuation  patient's needs:			
Section D – Previous Medic  Medication Name  Section E – Additional infor	Strength Strength	Dire	ctions	Dates of Therapy	uld not	discor	ntinuation  patient's needs:			
Section D – Previous Medic  Medication Name  Section E – Additional infor	Strength Strength	Dire	ctions	Dates of Therapy	uld not	discor	ntinuation  patient's needs:			
Section D – Previous Medic  Medication Name  Section E – Additional infor	Strength Strength	Dire	ctions	Dates of Therapy	uld not	discor	ntinuation  patient's needs:			
Section D – Previous Medic  Medication Name  Section E – Additional infor	Strength Strength	Dire	ctions	Dates of Therapy	uld not	discor	ntinuation  patient's needs:			
Section D – Previous Medic  Medication Name  Section E – Additional infor	Strength Strength	Dire	ctions	Dates of Therapy	uld not	discor	ntinuation  patient's needs:			
Section D – Previous Medic  Medication Name  Section E – Additional infor	Strength Strength	Dire	ctions	Dates of Therapy	uld not	discor	ntinuation  patient's needs:			
Section D – Previous Medic  Medication Name  Section E – Additional infor	Strength Strength	Dire	ctions	Dates of Therapy	uld not	discor	ntinuation  patient's needs:			
Section D – Previous Medication Name  Medication Name  Section E – Additional infor	Strength Strength	Dire	ctions	Dates of Therapy	uld not	discor	ntinuation  patient's needs:			
Section D – Previous Medic  Medication Name  Section E – Additional infor	Strength Strength	Dire	ctions	Dates of Therapy	uld not	discor	ntinuation  patient's needs:			



## Proton Pump Inhibitors (PPIs) - Washington Prior Authorization Request Form

Community Plan		Prior Authorization Request Form
Member First name:	Member Last name:	Member DOB:
	Clinical and Drug Specific	Information
Prescribers should re-evaluate patient on endoscopy and discontinue the PPI	2 months during any 12-month personers with a diagnosis of gastroesophat.	eriod may be candidates for stepdown therapy.  ageal reflux disease (GERD) with negative findings  ditions, PPIs may be covered for more than 2
months per year with a prior authoriza	_	
ALL ADD	DITIONAL DOCUMENTATION REQU	JESTED IS REQUIRED
Please indicate client's diagnosis (chec	k all that apply):	
	-	nly 2 months of PPI therapy during any 12-monthet for additional information and a sample taper
<ul> <li>Pathological gastric acid hypersect</li> <li>Attach GI consultation note do</li> </ul>		rome
☐ Barrett's esophagus		
Attach clinical EGD report fron	n within the last 5 years	
☐ Peptic ulcer disease ☐ Duodenal ulcer: • Attach EGD report from • H. pylori test results (bi ☐ Gastric ulcer: • Attach EGD report from	n within last 12 months document iopsy, breath, or stool test). n within last 60 days documenting iopsy, breath, or stool test).	
<ul><li>Eosinophilic esophagitis</li><li>Attach EGD report from withir</li></ul>	n the last 12 months documenting	diagnosis.
<ul><li>Esophageal stenosis/stricture or S</li><li>Attach EGD report documention</li></ul>	_	
<ul> <li>Erosive/ulcerative esophagitis</li> <li>Attach EGD report from withir</li> <li>H. pylori test results (biopsy, b</li> </ul>	n last 16 months documenting LA coreath, or stool test).	classification <b>AND</b>
☐ <i>H. pylori</i> positive		
Attach <i>H. pylori</i> test results (bi	iopsy, breath, or stool test).	
<ul><li>Other (Specify)</li><li>Attach all specialist notes and</li></ul>	 current labs supporting continued	d use of PPI.
Indicate any concurrent medications pa	ug (NSAID). Specify drug:	
☐ High-dose systemic corticosteroid.		
<ul><li>Antiplatelet or anticoagulant. Spec</li><li>List risk factors for GI bleed:</li></ul>		<del>_</del>
☐ Daily aspirin. Dose per day (mg):		
History of GI bleed in last 10 years.		g GI bleed.
Was it ingested with full glass	·	oright afterward?   Yes   No
☐ Pancreatic enzyme. Specify drug: _		
☐ Cancer therapy. Specify regimen: _		
i - expected per ditation deeded to toler	are cancer inerany.	



## Proton Pump Inhibitors (PPIs) - Washington Prior Authorization Request Form

For requests over once daily dosing only:							
<ul> <li>Is patient increasing from once daily dosing to tw</li> <li>Has the patient experienced uncontrolle</li> <li>What was the duration of once daily do</li> </ul>	ed symptoms on once daily dosing? $\square$ Yes $\square$	⊒ No					
<ul> <li>If patient is currently on twice daily dosing, has once daily dosing been tried? ☐ Yes ☐ No</li> <li>What was the duration the once daily dosing was tried?</li> <li>What was the outcome?</li> </ul>							
Prescriber signature	Prescriber specialty	Date					

**Confidentiality Notice:** This transmission contains confidential information belonging to the sender and UnitedHealthcare. This information is intended only for the use of UnitedHealthcare. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution or action involving the contents of this document is prohibited. If you have received this telecopy in error, please notify the sender immediately.