

**ULCERATIVE COLITIS AGENTS PRIOR AUTHORIZATION FORM** (form effective 1/8/2024)

Prior authorization guidelines for **Ulcerative Colitis Agents** and **Quantity Limits/Daily Dose Limits** are available on the DHS Pharmacy Services website at <https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/default.aspx>.

<input type="checkbox"/> New request	<input type="checkbox"/> Renewal request	total pages: _____	Prescriber name:	
Name of office contact:			Specialty:	
Contact's phone number:		NPI:	State license #:	
LTC facility contact/phone:		Street address:		
Beneficiary name:		City/state/zip:		
Beneficiary ID#:	DOB:	Phone:	Fax:	

**CLINICAL INFORMATION**

Drug requested:	Dosage form:	Strength:	
Directions:		Quantity:	Refills:
Diagnosis ( <i>submit documentation</i> ):		Dx code ( <i>required</i> ):	
Is the beneficiary currently being treated with the requested medication?	<input type="checkbox"/> Yes – date of last dose: _____ <i>Submit documentation.</i> <input type="checkbox"/> No		

**Complete all sections that apply to the beneficiary and this request.**  
***Check all that apply and submit documentation for each item.***

**INITIAL requests**

- For a SPHINGOSINE 1-PHOSPHATE RECEPTOR (S1PR) MODULATOR (eg, VELSIPITY [etrasimod], ZEPOSIA [ozanimod]) for treatment of ulcerative colitis (UC):
  - Is prescribed the medication by or in consultation with an appropriate specialist (eg, a gastroenterologist)
  - Has moderate-to-severe UC
  - Has UC associated with multiple poor prognostic factors
  - Tried and failed to achieve remission with or has a contraindication or an intolerance to an induction course of corticosteroids
  - Tried and failed to maintain remission with or has a contraindication or an intolerance to conventional immunomodulators (eg, AZA, cyclosporine, 6-MP, MTX)
  - Has achieved remission with the requested medication AND:
    - Will be using the requested medication as maintenance therapy to maintain remission

Tried and failed or has a contraindication or an intolerance to the preferred Cytokine and CAM Antagonists that are FDA-approved or medically accepted for the treatment of UC. (Refer to <https://papdl.com/preferred-drug-list> for a list of preferred Cytokine and CAM Antagonists.)

**Request is for VELSIPITY (etrasimod) AND:**

Has a comorbid heart condition – describe: \_\_\_\_\_

Experienced any of the following in the past 6 months:

Myocardial infarction

Transient ischemic attack

Unstable angina

Decompensated heart failure requiring hospitalization

Stroke

Class III or IV heart failure

**Request is for ZEPOSIA (ozanimod) AND:**

Has severe untreated sleep apnea

Will be taking a monoamine oxidase (MAO) inhibitor while taking Zeposia (e.g., selegiline, phenelzine)

Has a comorbid heart condition – describe: \_\_\_\_\_

Experienced any of the following in the past 6 months:

Myocardial infarction

Transient ischemic attack

Unstable angina

Decompensated heart failure requiring hospitalization

Stroke

Class III or IV heart failure

**2. For all other NON-PREFERRED Ulcerative Colitis Agents:**

Tried and failed or has a contraindication or an intolerance to the preferred Ulcerative Colitis Agents approved or medically accepted for the beneficiary's condition (Refer to <https://papdl.com/preferred-drug-list> for a list of preferred and non-preferred drugs in this class.)

**RENEWAL requests**

**1. For a SPHINGOSINE 1-PHOSPHATE RECEPTOR (S1PR) MODULATOR (eg, VELSIPITY [etrasimod], ZEPOSIA [ozanimod]):**

Is prescribed the medication by or in consultation with an appropriate specialist (eg, a gastroenterologist)

Experienced improvement in disease activity or level of functioning since starting the requested medication

**Request is for VELSIPITY (etrasimod) AND:**

Has a comorbid heart condition – describe: \_\_\_\_\_

Experienced any of the following in the past 6 months:

Myocardial infarction

Transient ischemic attack

Unstable angina

Decompensated heart failure requiring hospitalization

Stroke

Class III or IV heart failure

**Request is for ZEPOSIA (ozanimod) AND:**

Has severe untreated sleep apnea

Will be taking a monoamine oxidase inhibitor while taking Zeposia (e.g., selegiline, phenelzine)

Has a comorbid heart condition – describe: \_\_\_\_\_

Experienced any of the following in the past 6 months:

Myocardial infarction

Transient ischemic attack

Unstable angina

Decompensated heart failure requiring hospitalization

Stroke

Class III or IV heart failure

**PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO 866-240-3712**

**Prescriber Signature:**

**Date:**

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