

Entyvio vial (intravenous) (vedolizumab)

If this is an URGENT request, please call (800) 882-4462 (800.88.CIGNA)

PHYSICIAN	PHYSICIAN INFORMATION		PAI	PATIENT INFORMATION			
* Physician Name:	Physician Name: **Due to privacy regulations we will not be able to respond with the outcome of our review unless all asterisked (*) iter						
Specialty:	* DEA, NPI or	TIN:	this form are completed.**				
Office Contact Person:			* Patient Name:				
Office Phone:			* Cigna ID:	* Date of Birth:	* Date of Birth:		
Office Fax:			* Patient Street Address:				
Office Street Address:		City:	State:	Zip:			
City:	State:	Zip:	Patient Phone:				
Urgency: ☐ Standard ☐ Urgent (In checking this box, I attest to the fact that applying the standard review time frame may seriously jeopardize the customer's life, health, or ability to regain maximum function)							
Medication Requested:							
Dose and Quantity:	se and Quantity: Duration of therapy: J-Code:						
Frequency of administration:	Frequency of administration: ICD10:						
Is this a new start or continuation of therapy? If your patient has already begun treatment with drug samples of Entyvio , please choose "new start of therapy".							
If continuation of therapy: (if Crohn's, continuation of therapy) Is there documentation your patient has had a beneficial response with the requested medication? Examples of beneficial response for Crohn's Disease include: decreased pain, fatigue, stool frequency, and/or blood in stool; or improvement via fecal markers (for example, fecal lactoferrin, fecal calprotectin), serum markers (for example, C-reactive protein), imaging studies (magnetic resonance enterography [MRE], computed tomography enterography [CTE]), endoscopic assessment, and/or reduced dose of corticosteroids.							
Examples of beneficial response for Ulcerative Colitis include decreased pain, fatigue, stool frequency, and/or decreased rectal bleeding; or improvement via fecal markers (for example, fecal calprotectin), serum markers (for example, C-reactive protein), endoscopic assessment, and/or reduced dose of corticosteroids.							
Besides the drug being requested, other biologics or targeted synthetic disease-modifying antirheumatic drugs (DMARDs) include Actemra, adalimumab (Humira and all biosimilars), Adbry, Bimzelx, Cibinqo, Cimzia, Cosentyx, Enbrel, Ilumya, Infliximab (Remicade and all biosimilars), Kevzara, Kineret, Olumiant, Omvoh, Orencia, Otezla, Rinvoq, Rituximab (Rituxan and all biosimilars), Siliq, Simponi Aria, Simponi, Skyrizi, Sotyktu, Stelara, Taltz, Tremfya, Tysabri, Velsipity, Xeljanz, Zeposia. Which of the following best describes your patient's situation?							
 ☐ The patient is NOT taking any other biologic or targeted synthetic DMARD at this time, nor will they in the future. The requested drug is the only biologic or targeted synthetic DMARD the patient is/will be using. ☐ The patient is currently on another biologic or targeted synthetic DMARD, but this drug will be stopped and the requested drug will be started. ☐ The patient is currently on another biologic or targeted synthetic DMARD, and the requested drug will be added. The patient may continue to take both drugs together. ☐ The patient is currently on BOTH the requested drug AND another biologic or targeted synthetic DMARD. ☐ other/unknown 							
(if other/more than Entyvio) Please provide the rationale for concurrent use.							
(Please note: there are different preferred products depending on your patient's plan. Please refer to the applicable Cigna health care professional resource [e.g. cignaforhcp.com] to determine benefit availability and the terms and conditions of coverage)							

Where will this medication be ☐ Accredo Specialty Pharmacy** ☐ Hospital Outpatient ☐ Retail pharmacy ☐ Other (please specify):	e obtained?	☐ Home Health / Home Infusion☐ Physician's office stock (billin claim form) **Cigna's nationally preferred sp	g on a medical			
**Medication orders can be placed with Accredo via E-prescribe - Accredo (1620 Century Center Pkwy, Memphis, TN 38134-8822 NCPDP 4436920), Fax 888.302.1028, or Verbal 866.759.1557						
Facility and/or doctor dispen Facility Name: Address (City, State, Zip Code):	sing and administering medica State:	ation: Tax ID#:				
Where will this drug be admi ☐ Patient's Home ☐ Hospital Outpatient	nistered?	☐ Physician's Office ☐ Other (please specify):				
NOTE: Per some Cigna pi	ans, infusion of medication MUST o	ccur in the least intensive, medically appropri	ate setting.			
Is this patient a candidate for re-direction to an alternate setting (such as alternate infusion site, physician's office, home) with assistance of a Specialty Care Options Case Manager? Yes No (provide medical necessity rationale):						
Is the requested medication for a contract the patient?	chronic or long-term condition for wh	ich the prescription medication may be neces	sary for the life of ☐ Yes ☐ No			
Diagnosis related to use (ple	ase specify):					
☐ Crohn's disease (CD)	ulcerative colitis (UC)	Other:				
Clinical Information: Is the requested medication being	prescribed by (or in consultation wit	h) a gastroenterologist?	☐ Yes ☐ No			
If ulcerative colitis:						
	ogic or targeted synthetic DMARD (i emicade, biosimilars], Rinvoq, Simp	tsDMARD) for Ulcerative Colitis (for example, oni, Stelara, Xeljanz/XR)?	adalimumab SC ☐ Yes ☐ No			
Does your patient have Pouchitis?	☐ Yes ☐ No					
	ried therapy with an antibiotic (for ex ine (enema or suppository)?	cample, metronidazole, ciprofloxacin), corticos	steroid (enema or Yes No			
The covered alternative is ONE conventional systemic therapy (for example, 6-mercaptopurine, azathioprine, cyclosporine, tacrolimus, or a corticosteroid such as prednisone or methylprednisolone). If your patient has tried this drug, please provide drug strength, date(s) taken and for how long, and what the documented results were of taking this drug, including any intolerances or adverse reactions your patient experienced. If your patient has NOT tried this drug, please provide details why your patient can't try this alternative.						
(if UC, no biologic) Per the information provided above, which of the following is true for your patient in regard to the covered alternative? The patient tried the alternative, but it didn't work. The patient tried the alternative, but they did not tolerate it. The patient cannot try the alternative because of a contraindication to this drug. Other: Please specify						
If Crohn's:						
Prior to the start of Entyvio therapy Infliximab IV products, Stelara, Sk		ogic for Crohn's disease (Adalimumab SC pro	oducts, Cimzia, ☐ Yes ☐ No			
Will vedolizumab (Entyvio) be take	en concurrently with a corticosteroid?	?	☐ Yes ☐ No			
Will vedolizumab (Entyvio) be taken concurrently with a conventional systemic therapy?						

Per the information provided above, which of the following is true for your patient in regard to the covered alternatives? The patient tried one of the alternatives, but it didn't work The patient tried one of the alternatives, but they did not tolerate it The patient cannot try one of these alternatives because of a contraindication to this drug Other				
Does your patient have any of the following? Severe disease needing hospitalization Involvement of the upper GI tract Smoker Less than 40 years of age Stricturing disease Perianal disease Other enterocutaneous fistula Extraintestinal manifestations (ankylosing spondylitis, pyoderma gangrenosum, erythema nodosum) Previous Crohn's disease-related surgery (for example, ileocolonic resection to reduce the chance of Crohn's disease recurrence) Bowel obstruction History of abscess or perforation (after healing) None of the above				
Additional pertinent information: Please provide any additional pertinent clinical information, including: alternatives tried and any reason(s) alternatives cannot be tried; if the patient is currently on the requested drug (with dates of use) and how they have been receiving it (samples, out of pocket, etc).				
Attestation: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan or insurer its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form. Prescriber Signature: Date:				
Save Time! Submit Online at: www.covermymeds.com/main/prior-authorization-forms/cigna/ or via SureScripts in your EHR.				
Our standard response time for prescription drug coverage requests is 5 business days. If your request is urgent, it is important that you call us to expedite the request. View our Prescription Drug List and Coverage Policies online at cigna.com.				

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