

GastroIntestinal Biologic Agents Entyvio (vedolizumab) J3380 Prior Authorization Request Medicare Part B Form

Instructions: * Indicates required information – Form may be returned if required information is not provided. Please fax this request to the appropriate fax number listed at the bottom of the page.

□ Standard Request– (72 Hours)				Urgent Request (standard time frame could place the member's life, health or ability in serious jeopardy)			
	Date Requested						
		r Clinic name:				/ Fax	
MEMBER INFORMATION							
*Name: *ID#:						B:	
PRESCRIBER INFORMATION							
*Name:							
*Ado	dress:			*Fax:			
DISPENSING PROVIDER / ADMINISTRATION INFORMATION							
*Name: Phone:							
*Address:Fax:							
PROCEDURE / PRODUCT INFORMATION							
нс	PC Code	Name of Drug	Dos	e (Wt: kg Ht:)	Frequency	End Date if known
□ Self-administered □ Provider-administered □ Home Infusion							
Chart notes attached. Other important information:							
Diagnosis: ICD10: Description:							
\square Provider attests the diagnosis provided is an FDA-Approved indication for this drug							
CLINICAL INFORMATION							
 New Start or Initial Request: (Clinical documentation required for all requests) Provider has reviewed the attached "Criteria for Approval" and attests the member meets ALL required PA criteria. If not, please provide clinical rationale for formulary exception: 							
□ Continuation Requests: (Clinical documentation required for all requests)							
Patient had an <u>adequate response</u> or <u>significant improvement</u> while on this medication. If not, please provide clinical rationale for continuing this medication:							
ACKNOWLEDGEMENT							
Request By (Signature Required):							



Prior Authorization Group – Gastrointestinal-Biologic Agents PA

Drug Name(s): ENTYVIO VEDOLIZUMAB

Criteria for approval of Prior Authorization Drug:

- 1. Prescribed for an approved FDA diagnosis (as listed below):
- 2. Drug is being used appropriately per MCG GUIDELINES, CMS recognized compendia, authoritative medical literature, evidence-based guidelines and/or accepted standards of medical practice.
- 3. Member does not have any clinically relevant contraindications, or CMS/Plan exclusions, to the requested drug.
- If the member meets all these criteria, they may be approved by the Plan for the requested drug.
- Quantity limits and Tiering will be determined by the Plan.

Exclusion Criteria:

N/A

Prescriber Restrictions: N/A

Coverage Duration: Approval will be for 12 months

FDA Indications:

Entyvio

- Crohn's disease (Moderate to Severe), Active
- Ulcerative colitis (Moderate to Severe), Active

Off-Label Uses:

N/A

Age Restrictions:

Safety and effectiveness not established in pediatric patients

Other Clinical Considerations: N/A

Resources:

https://www.micromedexsolutions.com/micromedex2/librarian/CS/25D2A7/ND_PR/evidencexpert/ND_P/evidencexpert/DUPLICATIONSHIELDSYN C/DCE066/ND_PG/evidencexpert/ND_B/evidencexpert/ND_AppProduct/evidencexpert/ND_T/evidencexpert/PFActionId/evidencexpert.GoToDash board?docId=930961&contentSetId=100&title=Vedolizumab&servicesTitle=Vedolizumab&brandName=Entyvio&UserMdxSearchTerm=Entyvio&=n ull#

https://careweb.careguidelines.com/ed24/ac/ac_03177.htm