

### 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)				<b>Urgent Request</b> (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							
<ul> <li>New Start or Initial Request: (Clinical documentation required for all requests)</li> <li>Provider has reviewed the attached "Criteria for Approval" and attests the member meets         ALL required PA criteria.     </li> <li>If not, please provide clinical rationale for formulary exception:</li> </ul>							
□ 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