



MEDICARE FORM

Entyvio® (vedolizumab) Injectable Medication Precertification Request

Page 1 of 3

(All fields must be completed and legible for precertification review.)

For Illinois MMP:

FAX: 1-855-320-8445

PHONE: 1-866-600-2139

For other lines of business:

Please use other form.

Note: Entyvio is preferred on MA and MAPD plans.

Please indicate: Start of treatment: Start date / / Continuation of therapy: Date of last treatment / /

Precertification Requested By: Phone: Fax:

A. PATIENT INFORMATION

Form section A: Patient Information. Fields include First Name, Last Name, Address, City, State, ZIP, Home Phone, Work Phone, Cell Phone, DOB, Allergies, Email, Current Weight, Height.

B. INSURANCE INFORMATION

Form section B: Insurance Information. Fields include Aetna Member ID #, Group #, Insured, Does patient have other coverage?, Carrier Name.

C. PRESCRIBER INFORMATION

Form section C: Prescriber Information. Fields include First Name, Last Name, Address, City, State, ZIP, Phone, Fax, St Lic #, NPI #, DEA #, UPIN, Office Contact Name, Phone.

D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION

Form section D: Dispensing Provider/Administration Information. Divided into Place of Administration and Dispensing Provider/Pharmacy. Includes checkboxes for Self-administered, Physician's Office, Outpatient Infusion Center, Home Infusion Center, Administration code(s) (CPT).

E. PRODUCT INFORMATION

Form section E: Product Information. Fields include Request is for Entyvio (vedolizumab): Dose, Frequency, HCPCS Code.

F. DIAGNOSIS INFORMATION – Please indicate primary ICD Code and specify any other where applicable.

Form section F: Diagnosis Information. Fields include Primary ICD Code, Secondary ICD Code, Other ICD Code.

G. CLINICAL INFORMATION – Required clinical information must be completed in its entirety for all precertification requests.

For Initiation Requests (clinical documentation required):

Form section G: Clinical Information. Note: Entyvio is preferred on MA and MAPD plans. Includes checkboxes for prior therapy and concomitant use.

Continued on next page



MEDICARE FORM

Entyvio® (vedolizumab) Injectable Medication Precertification Request

Page 2 of 3

(All fields must be completed and legible for precertification review.)

For Illinois MMP:

FAX: 1-855-320-8445

PHONE: 1-866-600-2139

For other lines of business:

Please use other form.

Note: Entyvio is preferred on MA and MAPD plans.

Patient First Name	Patient Last Name	Patient Phone	Patient DOB
--------------------	-------------------	---------------	-------------

G. CLINICAL INFORMATION (continued) – Required clinical information must be completed in its entirety for all precertification requests.

Crohn's Disease

Yes  No Does the patient have a diagnosis of fistulizing Crohn's disease? **If yes**, please indicate the date of the diagnosis: \_\_\_\_/\_\_\_\_/\_\_\_\_

    Please indicate the severity of the patient's Crohn's disease:  Mild  Moderate  Severe

Yes  No Is there clinical evidence that the disease is active?

Yes  No Is the Crohn's disease manifested by at least one of the following?

            Check all that apply:  abdominal pain  arthritis  bleeding  diarrhea  internal fistulae

intestinal obstruction  megacolon  perianal disease  spondylitis  weight loss

Yes  No Was treatment with corticosteroids ineffective?

Yes  No Was treatment with corticosteroids not tolerated or contraindicated?

not tolerated  contraindicated

                Which of the following corticosteroids was tried?  hydrocortisone  methylprednisolone

prednisone  Other: Please explain: \_\_\_\_\_

        Which of the following corticosteroids was tried?  hydrocortisone  methylprednisolone

prednisone  Other: Please explain: \_\_\_\_\_

Yes  No Was treatment with 6-mercaptopurine (6-MP) ineffective?

Yes  No Was treatment with 6-mercaptopurine (6-MP) not tolerated or contraindicated?

not tolerated  contraindicated

Yes  No Was treatment with azathioprine ineffective?

Yes  No Was treatment with azathioprine not tolerated or contraindicated?

not tolerated  contraindicated

Ulcerative Colitis

Yes  No Is the patient hospitalized fulminant ulcerative colitis?

    Please indicate the severity of the patient's ulcerative colitis:  Mild  Moderate  Severe

Yes  No Is there evidence that the disease is active?

Yes  No Is the patient refractory to immunosuppression with corticosteroids (e.g., hydrocortisone, methylprednisolone, prednisone)?

Yes  No Does the patient require continuous immunosuppression with corticosteroids (e.g., hydrocortisone, methylprednisolone, prednisone)?

            Name and dose: Name: \_\_\_\_\_ Dose: \_\_\_\_\_

            Please indicate the route:  Oral  IV

        Name and dose: Name: \_\_\_\_\_ Dose: \_\_\_\_\_

        Please indicate the route:  Oral  IV

Yes  No Was treatment with immunosuppressant agent (e.g., azathioprine, m6-mercaptopurine) ineffective?

Yes  No Was treatment with immunosuppressant agent (e.g., azathioprine, m6-mercaptopurine) not tolerated or contraindicated?

not tolerated  contraindicated

                Provide the name of the drug(s): \_\_\_\_\_

        Provide the name of the drug(s): \_\_\_\_\_

Yes  No Was treatment with 5-aminosalicylic acid agents (e.g., balsalazide, mesalamine, sulfasalazine) ineffective?

Yes  No Was treatment with 5-aminosalicylic acid agents (e.g., balsalazide, mesalamine, sulfasalazine) not tolerated or contraindicated?

not tolerated  contraindicated

                Provide the name of the drug(s): \_\_\_\_\_

        Provide the name of the drug(s): \_\_\_\_\_

    Please select the symptoms the patient exhibit:  more than 10 stools per day  continuous bleeding  abdominal pain  distension

acute, severe toxic symptoms, including fever and anorexia

For Continuation requests (clinical documentation required):

Yes  No Will Entyvio (vedolizumab) be used concomitantly with aprelimast, tofacitinib, or other biologic DMARDs (e.g., adalimumab, infliximab)?

Yes  No Is this continuation request a result of the patient receiving samples of Entyvio (vedolizumab)?

Yes  No Is there clinical documentation supporting disease stability?

Yes  No Is there clinical documentation supporting disease improvement?

Yes  No Has the patient received Entyvio (vedolizumab) within the past 6 months?

Yes  No Does the patient have a documented severe and/or potentially life-threatening adverse event that occurred during or following the previous infusion?

Yes  No Could the adverse reaction be managed through pre-medication in the home or office setting?

Continued on next page



**MEDICARE FORM**

**Entyvio® (vedolizumab) Injectable  
Medication Precertification Request**

Page 3 of 3

(All fields must be completed and legible for precertification review.)

**For Illinois MMP:**

**FAX:** 1-855-320-8445

**PHONE:** 1-866-600-2139

**For other lines of business:**

Please use other form.

**Note: Entyvio is preferred on MA and MAPD plans.**

Patient First Name	Patient Last Name	Patient Phone	Patient DOB
--------------------	-------------------	---------------	-------------

**H. ACKNOWLEDGEMENT**

**Request Completed By (Signature Required):** \_\_\_\_\_ **Date:** \_\_\_\_ / \_\_\_\_ / \_\_\_\_

Any person who knowingly files a request for authorization of coverage of a medical procedure or service with the intent to injure, defraud or deceive any insurance company by providing materially false information or conceals material information for the purpose of misleading, commits a fraudulent insurance act, which is a crime and subjects such person to criminal and civil penalties.

The plan may request additional information or clarification, if needed, to evaluate requests.