



MEDICARE FORM

Entyvio® (vedolizumab) Injectable Medication Precertification Request

Page 1 of 3

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

For New Jersey HMO D-SNP:

FAX: 1-833-322-0034

PHONE: 1-844-362-0934

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 fields for address, phone, TIN, NPI, etc.

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 questions about prior therapy and concomitant use.

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| | | | |
|--------------------|-------------------|---------------|-------------|
| Patient First Name | Patient Last Name | Patient Phone | Patient DOB |
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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 apremilast, 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?

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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.