Pharmacy Prior Authorization

AETNA BETTER HEALTH ILLINOIS FAMILY HEALTH PLAN (MEDICAID)

Cimzia (Medicaid)

This fax machine is located in a secure location as required by HIPAA regulations.
Complete/review information, sign and date. Fax signed forms to Aetna Better Health Illinois at **1-844-242-0908**.
When conditions are met, we will authorize the coverage of Cimzia (Medicaid).
Please note that all authorization requests will be reviewed as the AB rated generic (when available) unless states otherwise.

Drug Name ***(please circle)***

Cimzia

Other, please specify____

Quantity ___________ Frequency ________________ Strength ____________

Route of Administration ___________ Expected Length of therapy _______________

Patient Information

Patient Name: ____________________________________

Patient ID: ____________________________________

Patient Group No.: ____________________________________

Patient DOB: ____________________________________

Patient Phone: ____________________________________

Prescribing Physician

Physician Name: ____

Specialty: ___________________________ NPI Number: _________________________

Physician Fax: ___________________________ Physician Phone: _________________________

Physician Address: ___________________________ City, State, Zip: _________________________

Diagnosis:______________________________ **ICD Code:**______________________________

Please circle the appropriate answer for each question.

1. Has this plan authorized Cimzia in the past for this patient (i.e., previous authorization is on file under this plan)? Y N

   [If no, skip to question 5.]

2. Does the patient have a diagnosis of Crohn's? Y N

   [If no, skip to question 4.]

3. Is the patient in remission without requiring more than 5mg of prednisone daily? Y N

   [No further questions.]

4. Has the patient had at least a 20% improvement in symptoms? Y N

   [No further questions.]

5. Does the patient have a diagnosis of rheumatoid arthritis (RA) with moderate to high disease activity? Y N

   [If no, skip to question 8.]

6. Has the patient had failure to an adequate trial (3 months) of two disease modifying anti-rheumatic drugs (DMARDs) regimens (one must be methotrexate)? Y N

   If yes, list medications tried: _____________________________

   Note: Monotherapy regimen: methotrexate (MTX), hydroxychloroquine (HCQ), leflunomide (LEF), sulfasalazine (SSZ).

   Combination regimen: MTX+SSZ+HCQ; MTX+HCQ, MTX+LEF, MTX+SSZ, SSZ+HCQ

   [If yes, skip to question 21.]

7. Does the patient have a contraindication to methotrexate? Y N

   Note: Contraindications such as Pregnancy, alcoholism, Chronic liver disease, Leukopenia, thrombocytopenia, or anemia.

   If yes, please document contraindication:___________________

   [If no, then no further questions]

   [If yes, skip to question.21]

8. Does the patient have a diagnosis of ankylosing spondylitis (AS)? Y N

   [If no, skip to question 12.]

9. Does the patient have unacceptable disease activity despite an adequate trial (3 months) with at least 2 different NSAIDs? Y N

   If yes, please list medications tried:_______________________

   [If no, skip to question 11.]

10. Is the patient currently on or will continue taking an NSAID with the requested medication? Y N

[If yes, skip to question 21.]

11. Does the patient have contraindications to NSAIDs? Y N

    Note: Contraindications such as true allergic reaction to NSAIDs, history of worsening asthma symptoms after taking aspirin or NSAIDs, current GI bleed, severe renal dysfunction.

    If yes, please document contraindication:___________________

    [If yes, then skip to question 21.]

    [If no, then no further questions]

12. Does the patient have a diagnosis of psoriatic arthritis (PsA)? Y N

    [If no, skip to question 22.]

13. Does the patient have primarily axial disease (involving the spine) or active enthesitis/dactylitis? Y N

    [If no, skip to question 15.]

14. Has the patient tried an adequate trial (3 months) with at least 2 different NSAIDs and had inadequate response? Y N

    If yes, please list medications tried:_______________________

    [If yes, skip to question 19.]

    [If no, skip to question 20.]

15. Does the patient have active psoriatic arthritis? Y N

    [If no, then no further questions.]

16. Has the patient had failure to an adequate trial (3 months) of methotrexate? Y N

    [If yes, skip to question 19.]

17. Does the patient have a contraindication to methotrexate? Y N

    Note: Contraindications such as Pregnancy, alcoholism, Chronic liver disease, Leukopenia, thrombocytopenia, or anemia.

    If yes, please document contraindication:___________________

    [If no, then no further questions]

18. Has the patient had failure to an adequate trial (3 months) of sulfasalazine or leflunomide? Y N

    [If no, then no further questions.]

19. Is the patient currently on or will continue taking an NSAID with requested medication? Y N

    [If yes, then skip to question 21.]

20. Does the patient have contraindications to NSAIDs? Y N

    Note: Contraindications such as true allergic reaction to NSAIDs, history of worsening asthma symptoms after taking aspirin or NSAIDs, current GI bleed, severe renal dysfunction.

    If yes, please document contraindication:___________________

    [If no, then no further questions.]

21. Has the patient tried and failed BOTH Enbrel and Humira? Y N

    [If no, then no further questions.]

    [If yes, skip to question 28.]

22. Does the patient have a diagnosis of Crohn's Disease? Y N

    [If no, then no further questions.]

23. Has the patient had inadequate response or intolerable side effects to IV corticosteroids after 7-10 days or oral prednisone (dosed at 40mg or more per day) for 30 days)? Y N

    [If yes, skip to question 27.]

24. Does the patient have steroid-dependent Crohn's disease as evidenced by one of the following: A) Patient had a relapse within three months of stopping corticosteroids; OR B) Patient is unable to taper steroids to an acceptable dose after 3 months without having symptom recurrence? Y N

    [If no, then no further questions.]

25. Has the patient had failure to an adequate trial (3 months) of azathioprine (AZA), mercaptopurine (6-mp) or injectable methotrexate? Y N

    [If yes, skip to question 27.]

26. Does the patient have a contraindication to all of the following: azathioprine (AZA), injectable methotrexate and mercaptopurine (6-mp)? Y N

    If yes, please document contraindication(s): ________________

    [If no, then no further questions]

27. Has the patient had a trial and failure of Humira? Y N

    [If no, then no further questions.]

28. Is the patient at least 18 years of age? Y N

    [If no, then no further questions.]

29. Is Cimzia being prescribed by, or in consultation with a specialist, based on indication (rheumatologist or gastroenterologist)? Y N

    [If no, then no further questions.]

30. Has the patient been screened for latent tuberculosis (TB) and hepatitis B? Y N

    [If no, then no further questions.]

31. Does the patient have an active infection (including Hepatitis B and/or tuberculosis (TB)? Y N

    [If no, skip to question 33.]

32. Is the patient currently receiving or has completed treatment for latent TB infection or Hepatitis B? Y N

    [If no, then no further questions.]

33. Will Cimzia be given in combination with another biologic DMARD? Y N

    [If yes, then no further questions.]

34. Does the patient have CHF (NYHA class III or IV)? Y N

Comments:

I affirm that the information given on this form is true and accurate as of this date.

Prescriber (Or Authorized) Signature **Date**