



Fax Referral To: 800-323-2445
Phone: 866-278-6634

Enbrel® (etanercept) Enrollment Form For Blue Cross Blue Shield of Rhode Island Members

Date: _____ Needs by Date (Please Specify): _____

Ship to: Patient Office Other: _____

PATIENT INFORMATION

(Complete the following or send patient demographic sheet)

Patient Name: _____
Address: _____
City, State, Zip: _____
Home Phone: _____
Alternate Phone: _____
SS #: _____
Insurance ID: _____
Date of Birth: _____ Gender: _____

PRESCRIBER INFORMATION

Prescriber's Name: _____
State License #: _____ UPIN: _____
DEA #: _____ NPI #: _____
Group or Hospital: _____
Address: _____
City, State Zip: _____
Phone: _____ Fax: _____
Contact Person: _____
Contact Phone: _____

INSURANCE INFORMATION (If available, please copy and attach the front and back of insurance and prescription drug card)

Primary Insurance: Subscriber: _____ Subscriber ID#: _____ Name of Insurer: **Blue Cross Blue Shield of RI**
Secondary Insurance: Subscriber: _____ Subscriber ID#: _____ Name of Insurer: _____

STATEMENT OF MEDICAL NECESSITY for BCBS of Rhode Island Members

Diagnosis (ICD-9 Code): 696.1 Psoriasis 696.0 Psoriatic Arthritis 714.0 Rheumatoid Arthritis Other: _____ • Date of Diagnosis: _____

APPROVAL CRITERIA: CHECK ALL BOXES THAT APPLY TO THE DIAGNOSIS.

NOTE: Areas not filled out are considered not applicable to your patient & MAY AFFECT THE OUTCOME of this request.

REQUIRED FOR ALL (none of the following can be present):

- Patient has a latex allergy Yes No
- Patient has tuberculosis or a history of recurrent, chronic current, or clinically important infection Yes No
- Patient had a tuberculin skin test to rule out latent tuberculosis Yes No
- Patient has a history of systemic malignancy within the last 5 years Yes No
- Patient has moderate to severe (NYHA Class III/IV) Congestive Heart Failure (CHF) Yes No
- Patient has Multiple Sclerosis or any other demyelinating disease Yes No
- Will the patient receive live vaccines while receiving Enbrel? Yes No
- Patient will use Enbrel in combination with other TNF agents or Kineret Yes No
- Patient is currently receiving systemic psoriasis therapy (except for methotrexate, glucocorticoids, salicylates, non-steroidal anti-inflammatory drugs, or analgesics),
Immunosuppressive therapy, or Anakinra Yes No
- Comments: _____

Ankylosing Spondylitis:

- Patient is 18 years of age or older Yes No
- Patient has diagnosis of Ankylosing Spondylitis Yes No
- Patient has failed, had inadequate response to, or is contraindicated for treatment with sulfasalazine, methotrexate, or non-steroidal anti-inflammatory drugs Yes No

Chronic Plaque Psoriasis:

- Patient is 18 years of age or older Yes No
- Patient has a diagnosis of moderate to severe plaque psoriasis with either of the following: Yes No
 - Patient has greater than 10% of body surface area with plaque psoriasis
 - Less than or equal to 10% body surface area with plaque psoriasis involving sensitive areas or areas that would significantly impact daily function (such as palms, soles of Feet, head/neck, or genitalia)
 - Other: _____
- Is the psoriasis controlled with topical therapy? Yes No
- Patient has failed, has a contraindication, or is intolerant to, but is otherwise a candidate for other systemic therapies (e.g., methotrexate, acitretin, or cyclosporine) or phototherapy Yes No

Moderate to severely active Juvenile Rheumatoid Arthritis (under 18 years of age):

- Has the patient failed or had an inadequate response to 1 or more DMARDs? Yes No
- If Yes, what DMARD has the patient tried?

<input type="checkbox"/> Azulfidine (sulfasalazine)	<input type="checkbox"/> Methotrexate	<input type="checkbox"/> Cytoxan
<input type="checkbox"/> Cuprimine/Depen (penicillamine)	<input type="checkbox"/> Plaquenil (hydroxychloroquine)	<input type="checkbox"/> Cyclosporine (Neoral or Sandimmune)
<input type="checkbox"/> Imuran (azathioprine)	<input type="checkbox"/> Ridaura (auranofin)	<input type="checkbox"/> Minocycline (Minocin or Dynacin)
<input type="checkbox"/> Lefkybinude (Arava)	<input type="checkbox"/> Gold Sodium Thiomalate (Myochrysin)	<input type="checkbox"/> Other: _____

Active Psoriatic Arthritis:

- Patient is 18 years of age or older Yes No
- Patient has active arthritis with at least 3 swollen and 3 tender joints Yes No
- Patient has arthritis in **any** of the following distributions: Yes No
 - Ankylosing Spondylitis-like Arthritis Asymmetric Arthritis
 - Arthritis Mutilans Distal Interphalangeal Joint Involvement
 - Polyarticular Arthritis, without Rheumatoid Nodules
- Patient has failure or contraindicated for disease-modifying antirheumatic drugs (DMARD) therapy, specifically methotrexate or sulfasalazine Yes No

Moderate to severely active Rheumatoid Arthritis:

- Patient has failed or had an inadequate response to 1 or more DMARDs? Yes No

If Yes, what DMARD has the patient tried?

- Azulfidine (sulfasalazine) Methotrexate Cytoxan
- Cuprimine/Depen (penicillamine) Plaquenil (hydroxychloroquine) Cyclosporine (Neoral or Sandimmune)
- Imuran (azothioprine) Ridaura (auranofin) Minocycline (Minocin or Dynacin)
- Lefkybinude (Arava) Gold Sodium Thiomalate (Myochrysine) Other: _____

Note: Enbrel (etanercept) may be administered with methotrexate if patient is not responding adequately to methotrexate alone.

PRESCRIPTION INFORMATION

MEDICATION	STRENGTH	DIRECTIONS	QUANTITY	REFILLS
<input type="checkbox"/> Enbrel [®] (etanercept)	<input type="checkbox"/> 50mg/ml Sureclick [™] Autoinjector <input type="checkbox"/> 50mg/ml Prefilled Syringe <input type="checkbox"/> 25mg/0.5ml Prefilled Syringe <input type="checkbox"/> 25mg Vial	<input type="checkbox"/> Inject 50mg SC TWICE a week (72-96 hours apart) <input type="checkbox"/> Inject 50mg SC ONCE a week <input type="checkbox"/> Inject 25mg SC TWICE a week (72-96 hours apart) <input type="checkbox"/> Other: _____		

PRODUCT SUBSTITUTION PERMITTED (Date)

DISPENSE AS WRITTEN (Date)