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Please note: All information below is required to process this request.

Mon-Fri: 5am to 10pm Pacific / Sat: 6am to 3pm Pacific

Inflectra[®], Remicade[®] & Renflexis[®] Prior Authorization Request Form (Page 1 of 2)

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Member Information <small>(required)</small>	Provider Information <small>(required)</small>
----------------------------------------------	------------------------------------------------

Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:

Medication Information <small>(required)</small>

Medication Name:	Strength:	Dosage Form:
<input type="checkbox"/> Check if requesting brand	Directions for Use:	
<input type="checkbox"/> Check if request is for continuation of therapy		

Clinical Information <small>(required)</small>

Select the diagnosis below:

Active ankylosing spondylitis (AS)

Active psoriatic arthritis (PsA)

Chronic severe plaque psoriasis (i.e., extensive and/or disabling)

Fistulizing Crohn's disease

Moderately to severely active Crohn's disease

Moderately to severely active rheumatoid arthritis (RA)

Moderately to severely active ulcerative colitis

Sarcoidosis

Other diagnosis: _____ ICD-10 Code(s): _____

Clinical Information:

Select if the requested medication is prescribed by or in consultation with one of the following specialists, if applicable for the patient's diagnosis:

Dermatologist Gastroenterologist Rheumatologist Pulmonologist

For Inflectra and Renflexis requests only:

Has the patient had a trial and failure, contraindication, or intolerance to Remicade, unless already receiving therapy? Yes No

For active ankylosing spondylitis (AS), also answer the following:

Has the patient had trial and failure, contraindication, or intolerance to two non-steroidal anti-inflammatory drugs (NSAIDs)? Yes No

For fistulizing Crohn's disease and moderately to severely active Crohn's disease, also answer the following:

Select if the patient has had trial and failure, contraindication, or intolerance to the following:

6-mercaptopurine (Purinethol)

Azathioprine (Imuran)

Corticosteroids (e.g., prednisone, methylprednisolone)

Methotrexate (Rheumatrex, Trexall)

For moderately to severely active rheumatoid arthritis (RA), also answer the following:

Is the patient receiving concurrent therapy with methotrexate (Rheumatrex, Trexall)? Yes No

Has the patient had trial and failure, contraindication, or intolerance to methotrexate (Rheumatrex, Trexall)? Yes No

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Inflectra[®], Remicade[®] & Renflexis[®] Prior Authorization Request Form (Page 2 of 2)

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For moderately to severely active ulcerative colitis, also answer the following:

Select if the patient has had trial and failure, contraindication, or intolerance to the following conventional therapies:

- 6-mercaptopurine (Purinethol)
- Aminosalicylate (e.g., mesalamine [Asacol, Pentasa, Rowasa], olsalazine [Dipentum], sulfasalazine [Azulfidine, Sulfazine])
- Azathioprine (Imuran)
- Corticosteroids (e.g., prednisone, methylprednisolone)

For sarcoidosis, also answer the following:

Has the patient had trial and failure, contraindication, or intolerance to corticosteroids (e.g., prednisone)? Yes No

Has the patient had trial and failure, contraindication, or intolerance to one immunosuppressant (e.g., methotrexate [Rheumatrex, Trexall], Cytoxan [cyclophosphamide], or Imuran [azathioprine])? Yes No

Reauthorization:

If this is a reauthorization request, answer the following question:

Is there documentation the patient has had a positive clinical response to infliximab therapy? Yes No

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note:

This request may be denied unless all required information is received.
For urgent or expedited requests please call 1-800-711-4555.
This form may be used for non-urgent requests and faxed to 1-800-527-0531.

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