

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

Member Information (required)			Provider Information (required)		
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 (required)					
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 (required)					
<p>Select the diagnosis below:</p> <input type="checkbox"/> Active ankylosing spondylitis <input type="checkbox"/> Active psoriatic arthritis <input type="checkbox"/> Chronic severe plaque psoriasis <input type="checkbox"/> Crohn's disease <input type="checkbox"/> Moderately to severely active rheumatoid arthritis <input type="checkbox"/> Moderately to severely active ulcerative colitis <input type="checkbox"/> Sarcoidosis <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
<p>Clinical Information:</p> <p>Select if Inflectra is prescribed by or in consultation with one of the following specialists, if applicable for the patient's diagnosis: <input type="checkbox"/> Dermatologist <input type="checkbox"/> Gastroenterologist <input type="checkbox"/> Rheumatologist <input type="checkbox"/> Pulmonologist</p> <p>Select if infliximab be used in combination with the following: <input type="checkbox"/> Biologic DMARD [e.g., Enbrel (etanercept), Rituxan (rituximab), Orencia (abatacept), Kineret (anakinra), Cimzia (certolizumab)] <input type="checkbox"/> Janus kinase inhibitor [e.g., Xeljanz (tofacitinib)] <input type="checkbox"/> Not in combination with a biologic DMARD or janus kinase inhibitor</p> <p>For Inflectra and Renflexis requests only: Does the patient have history of failure, contraindication, or intolerance to Remicade? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>					
<p>For active ankylosing spondylitis, also answer the following: Does the patient have history of failure, contraindication, or intolerance to two non-steroidal anti-inflammatory drugs (NSAIDs)? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>					
<p>For Crohn's disease, also answer the following: Select if the patient has one of the following diagnoses: <input type="checkbox"/> Fistulizing Crohn's disease <input type="checkbox"/> Moderately to severely active Crohn's disease</p> <p>Select if the patient has history of failure, contraindication, or intolerance to the following: <input type="checkbox"/> 6-mercaptopurine (Purinethol) <input type="checkbox"/> Azathioprine (Imuran) <input type="checkbox"/> Corticosteroids (e.g., prednisone, methylprednisolone) <input type="checkbox"/> Methotrexate (Rheumatrex, Trexall)</p>					

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

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

Does the patient have history of failure, contraindication, or intolerance to methotrexate (Rheumatrex, Trexall)? Yes No

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

Select if the patient has history of failure, contraindication, or intolerance to the following:

- 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:

Does the patient have history of failure, contraindication, or intolerance to corticosteroids (e.g., prednisone)? Yes No

Does the patient have history of 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 questions:

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

Select if infliximab be used in combination with the following:

- Biologic DMARD [e.g., Enbrel (etanercept), Rituxan (rituximab), Orencia (abatacept), Kineret (anakinra), Cimzia (certolizumab)]
- Janus kinase inhibitor [e.g., Xeljanz (tofacitinib)]
- Not in combination with a biologic DMARD or janus kinase inhibitor

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.

Please fax this form to 1-844-712-8129 to initiate a prior authorization review for the member and medication above.

Please note: plan benefits may limit or exclude coverage of specific medications including those requested on this form.

I certify, to the best of my knowledge, the statements and information provided on this form are factual and correct.

Provider/Representative (and Title): _____ Date: _____

PROACT INTERNAL USE ONLY:

Clinical Review Decision

Approved, through

Denied (documentation attached, if necessary)

Tracking:

1 st Attempt		2 nd Attempt		Letter Mailed:	
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