

Clinical Criteria Worksheet: Infliximab Products

Enrollee Information

Enrollee Last Name:

Enrollee First Name:

Date of Birth (MM/DD/YYYY):

--	--	--	--	--	--	--	--	--	--

Enrollee Medicaid ID (2 letters, 5 numbers, 1 letter):

--	--	--	--	--	--	--	--	--	--

Address:

City, Town or Post Office:

State:

ZIP Code:

Prescriber Information

Prescriber Last Name:

Prescriber First Name:

National Provider Identifier (NPI) Number: _____

Preferred Contact (Telephone Number): _____

Clinical Criteria – Drug Information

Drug Administration:

Provide the date of drug administration (MM/DD/YYYY):

--	--	--	--	--	--	--	--	--	--

Drug Name and Strength:

- infliximab (REMICADE®) 100 mg vial
- infliximab 100 mg vial
- infliximab-abda (RENFLEXIS®) 100 mg vial
- infliximab-axxq (AVSOLA®) 100 mg vial
- infliximab-dyyb (INFLECTRA®) 100 mg vial

Enrollee Last Name:

Enrollee First Name:

Clinical Criteria – Diagnosis

1. Diagnosis related to use (please select one diagnosis):

Food and Drug Administration Approved Indications:

- Ankylosing spondylitis
- Psoriatic arthritis
- Plaque psoriasis
- Rheumatoid arthritis, in combination with methotrexate (MTX)

Compendia-Supported Uses

- Adult-onset Still's disease
- Behcet's syndrome
- Graft versus host disease
- Polyarteritis Nodosa
- Refractory granulomatosis with polyangiitis, in combination with corticosteroids
- Severe, refractory hidradenitis suppurativa
- Refractory Kawasaki disease
- Severe, refractory synovitis, acne, pustulosis, hyperostosis, and osteitis syndrome (SAPHO syndrome)
- Refractory sarcoidosis (Adjunctive therapy)
- Synovitis
- Refractory Takayasu's disease
- Uveitis and refractory uveitis (Adjunctive therapy)
- Other: _____

2. Was the patient's medication record reviewed to confirm that the patient is not utilizing infliximab with other biological products to treat the same condition?

- Yes No

Enrollee Last Name:

Enrollee First Name:

Clinical Criteria

3. Please indicate if this request is for the initiation or continuation of infliximab therapy?

Initiation Continuation

4. Prior to initiation of infliximab therapy*, has the patient had a trial of a conventional agent, disease-modifying antirheumatic drug (DMARD) OR a tumor necrosis factor inhibitor (TNFi)?

Yes No

*Step Therapy does not apply to Ulcerative Colitis and Crohn's disease.

Attestation

I attest that this drug is medically necessary for this patient and that all of the information on this form is accurate to the best of my knowledge. I attest that documentation of the above diagnosis and medical necessity is available for review if requested by the New York State Medicaid Program.

Prescriber Signature (Required)

Date (MM/DD/YYYY)