



PRIOR AUTHORIZATION PROGRAM FORM

Remicade® (Influximab)

Instructions:

1. Section 1 to be completed by Plan Member / Patient
2. Section 2 to be completed by Physician (expenses incurred by the completion of this form is at plan member's expense)
3. Section 3 to be completed by Pharmacist

Please fax completed form to NexgenRx Formulary Management to 1-877-639-4369 or mail to 145 The West Mall P.O. Box 110 U, Toronto, Ontario, M8Z-5M4

Section 1: To be completed by Patient

Member's Name: (Last, First)	Card ID Number:										
Patient's Name: (Last, First)	Patient's Date of Birth (dd/mm/yy)	Relationship to Member (please circle)			PATIENT CODE						
		Employee	Spouse	Dependent							

Results of this request to be communicated to:

<input type="checkbox"/> Pharmacy	<input type="checkbox"/> Patient / Legal Guardian Named Below	<input type="checkbox"/> email:
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I authorize NexgenRx Inc. (a) to use the personal information disclosed on this form, and any other personal information known to NexgenRx Inc. regarding the above-named patient, for the purpose of assessing this prior authorization request and any related claim and administering the benefit plan under which any such claim is made, and (b) to contact, and to obtain any such personal information from and to disclose any such personal information to, any physician, pharmacist or other health care professional having knowledge of such patient's health relevant to this request and any related claim.

Patient / Legal Guardian Name: _____ Telephone Number: _____

Signature of Patient/ Legal Guardian: _____ Date (dd/mm/yy): _____

Section 2: To be completed by Physician

Drug Name & Strength: REMICADE 100 MG VIAL	DIN: 02244016	Dosage Instructions:
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Eligibility for drug coverage is dependent upon the patient meeting at least one of the qualifying criteria listed below. If the patient has another drug plan, prior authorization may cover some or all of the excess not paid by the primary plan.

Please indicate if the patient meets at least one of the following qualifying criteria for drug coverage:

- To reduce the signs and symptoms and inhibit the progression of structural damage and improvement in physical function in moderate to severe active rheumatoid arthritis, in combination with methotrexate, in adults.
- To reduce the signs and symptoms and improvement in physical function in patients with active ankylosing spondylitis.
- To reduce the signs and symptoms, induce and maintain clinical remission and mucosal healing and reduction of corticosteroid use in adult patients with moderate to severe active Crohn's disease or ulcerative colitis who have had an inadequate response to standard therapy.
- To treat fistulising Crohn's disease in adults who have failed to respond to a full and adequate course of standard therapy.
- To reduce the signs and symptoms, induce and maintain clinical remission, mucosal healing and reduction of corticosteroid use in pediatric patients with moderate to severe active Crohn's disease or ulcerative colitis who have had an inadequate response to standard therapy.
- To reduce the signs and symptoms and inhibit the progression of structural damage and improvement in physical function in patients with psoriatic arthritis.
- To treat moderate to severe active polyarticular juvenile rheumatoid arthritis in patients aged 4-17 who have failed to have an adequate response to standard DMARDS.
- To treat chronic moderate to severe plaque psoriasis in adults who are candidates for systemic therapy or phototherapy. Remicade should be used after phototherapy has been shown to be ineffective or inappropriate.

Physician Name: (Last, First)	License Number:
Address: (Street, City, Province, Postal Code)	Telephone Number : () -
	Fax Number: () -
Signature of Physician:	Date (dd/mm/yy)

Section 3: To be completed by Pharmacist

Pharmacy Name:	Provider Number:
Pharmacy Address: (Street, City, Province, Postal Code)	Telephone Number : () -
	Fax Number: () -
Signature of Pharmacist: _____	
Pharmacist's Name: (Print Last, First)	

Internal Office Use Only:

Date Received:		Date & Approved by:
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