

PRIOR AUTHORIZATION PROGRAM REIMBURSEMENT REQUEST FORM
For biologic response modifier therapy: *Simponi (golimumab)*

Please fax form to:
1-866-840-1509

Please note that the patient **AND** physician must complete this form. **All fields are mandatory and must be completed.** Incomplete forms may result in your application being declined. Please retain a copy of this form for your records.

Instructions:

- 1. PLEASE PRINT CLEARLY AND COMPLETE ALL SECTIONS.**
2. The patient/plan member must complete section A.
3. Your physician must complete section B. The cost, if any, of completing this form is at the expense of the patient/plan member.
4. Please return the form to your insurance company via Pharmacy Services at TELUS Health (a service provider of your insurance company) by fax to **1-866-840-1509**.
5. If you have any questions on the application of this program or the decision on reimbursement, or to inquire on the status of your Reimbursement Request Form, please contact your insurer.

A. Information to be Completed by Patient		
Employee or Insured's Name	Drug Card Number ____ - ____ - ____ - ____	
Patient's Name	Patient's Date of Birth (DD/MMM/YYYY) __ / __ / ____	Relationship to Employee/Insured <input type="checkbox"/> Employee <input type="checkbox"/> Spouse <input type="checkbox"/> Dependent
<p>Please allow two business days for a response once all information is received and complete. Notification of the results of this request will occur Monday to Friday between 9 am and 4 pm Eastern Time.</p>		

Please provide contact information and indicate **ONE** method of preferred contact for notification of the results:

<input type="checkbox"/> E-mail	<input type="checkbox"/> Call me (and leave a message if I'm not there)	<input type="checkbox"/> Fax me at:
<input type="checkbox"/> Contact my pharmacy: Pharmacy Name	Pharmacy Phone Number	

I certify that the information provided by me is true, correct and complete to the best of my knowledge. I authorize my insurance company, TELUS Health (a service provider of my insurance company), their authorized representatives, agents and service providers to use and exchange this information needed for underwriting, administration and paying claims with any person or organization who has relevant information pertaining to this claim including health professionals, institutions and investigative agencies in the event of an audit. I authorize my insurance company and/or TELUS Health (a service provider of my insurance company) to contact any licensed physician, institution, pharmacy or person who has any records or knowledge of me or my health with respect to this submitted claim.

SIGNATURE OF PATIENT/PARENT/LEGAL GUARDIAN _____

Date: (DD/MMM/YYYY): __ / __ / ____

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B. Information to be Completed by Prescribing Physician

Drug Name	Strength	Dose
<i>Simponi (golimumab)</i>		

***Simponi (golimumab)* will be eligible for reimbursement only if the patient satisfies the conditions listed below and if the patient does not qualify for coverage under any other drug plan or government mandated program. If the patient is covered under another drug plan or government mandated program, the prior authorization program, as part of your drug benefits, may cover the portion not paid for by the primary plan. If “None of the above criteria” is indicated, the patient will not be eligible for reimbursement. For Quebec plan members, please refer to the RAMQ exception drug criteria, if applicable.**

Eligibility Criteria

Please indicate if the patient satisfies one of the following criteria:

- Ankylosing Spondylitis (9631008) (Simponi SC/IV): Patient has tried and failed conventional treatment (e.g. non-steroidal anti-inflammatory drugs [NSAIDs]), OR
- Psoriatic Arthritis (156370009) (Simponi SC/IV): Patient has tried and failed conventional treatment (e.g. disease-modifying antirheumatic drugs [DMARDs]), OR
- Rheumatoid Arthritis (69896004) (Simponi SC/IV): Patient has tried and failed a minimum 12 week trial of Methotrexate plus one other disease modifying anti-rheumatic drug (DMARD). Where combinations of non-biologic DMARDs are impossible (a rare situation), 3 consecutive non-biologic DMARDs would be acceptable; OR
- Ulcerative Colitis (64766004) (Simponi SC): Patient has tried and failed conventional treatment (including corticosteroids, aminosaliclates, azathioprine, or 6-mercaptopurine), OR
- Non-radiographic Axial Spondyloarthritis (713777005) (Simponi SC): Patient has objective signs of inflammation as indicated by elevated C-reactive protein (CRP) and/or magnetic resonance imaging (MRI) evidence; and has had an inadequate response to, or is intolerant to two (2) or more nonsteroidal anti-inflammatory drugs (NSAIDs).

OR

- None of the above criteria applies.

*Simponi IV is only approved for the treatment of Rheumatoid Arthritis, Psoriatic Arthritis and Ankylosing Sponylitis by Health Canada.

Relevant additional information _____

Physician Information

Physician's Name	License Number	Telephone Number	Fax Number
Address		City	Province
			Postal Code
Physician's Signature		Date: (DD/MMM/YYYY)	
		___ / ___ / ____	