

## SPECIAL AUTHORITY REQUEST TARGETED DMARDS FOR ANKYLOSING SPONDYLITIS RENEWAL / DOSING ADJUSTMENT

received in error.

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If you have received this fax in error, please write

MISDIRECTED across the front of the form and fax toll-free to 1-800-609-4884, then destroy the pages

For up-to-date criteria and forms, please check: www.gov.bc.ca/pharmacarespecialauthority

Fax requests to 1-800-609-4884 (toll free) OR mail requests to: PharmaCare, Box 9652 Stn Prov Govt, Victoria, BC V8W 9P4 This facsimile is Doctor privileged and contains confidential information intended only for PharmaCare. Any other distribution, copying or disclosure is strictly prohibited.

If PharmaCare approves this Special Authority request, approval is granted solely for the purpose of covering prescription costs. PharmaCare approval does not indicate that the requested medication is, or is not, suitable for any specific patient or condition.

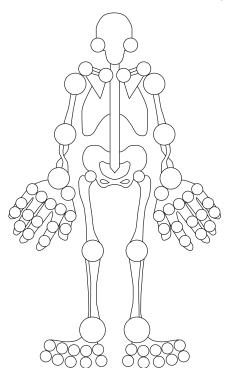
Forms with information missing will be returned for completion. If no prescriber fax or mailing address is provided, PharmaCare will be unable to return a response

SECTION 1 – SPECIALIST INFORMATION		SECTION 2 - PATIENT INFORMATION				
Rheumatologist's Name and Mailing Address		Patient (Family) Name				
		Patient (Given) Name(s)	Patient (Given) Name(s)			
College ID (use ONLY College ID	Phone Number (include area code)	Date of Birth (YYYY / MM / DI	D) Date of Ap	oplication (YYYY / MM / DD)		
CRITICAL FOR A TIMELY RESPONSE	 Rheumatologist's Fax Number	CRITICAL FOR PROCESSING	Personal Health Number (PHN)			
ECTION 3 – MEDICATI	ON REQUESTED					
Requested Dose and Interval	011 112 Q 013 112 D	Patient's Body Weight (if significa	antly changed)			
mg, every:		kg	kg			
ADALIMUMAB: 40 mg every two weeks  Indefinite coverage			GOLIMUMAB: 50 mg SC. once per month  Renewal of one year			
OR Renewal of three years OR Renewal of one year		INFLIXIMAB: 3-5 mg/k	INFLIXIMAB: 3-5 mg/kg every 8 weeks			
	IGEVITA®		O Indefinite coverage			
O HYRIMOZ® O IDA			OR			
_	mg every other week or 400 mg every 4 weeks	O AVSOLA® O INFLEC	○ AVSOLA® ○ INFLECTRA® ○ RENFLEXIS®			
○ Indefinite o		SECUKINUMAB: 150 m	SECUKINUMAB: 150 mg monthly			
OR O Renewal of		○ Renewal of o	Renewal of one year			
☐ ETANERCEPT: total d ☐ Indefinite c OR ☐ Renewal of OR ☐ Renewal of	overage three years		If approved, please note that claims with indefinite SA approvals will be monitored and any overuse or significant underuse will be subject to review.			
BRENZYS® 50 mg	) <b>ERELZI®</b> 25, 50 mg					
ECTION 4 - CURRENT	CLINICAL INFORMATION					
ESR or	CRP MORNING STIFFNESS (MINUTES)					
		PHYSICIAN GLOBAL ASSESSMENT OF IN (SCALE OF 0 - 10), 0 = REMISSION, 10 = S				
HARMACARE USE ON	ILY	Please comple	ete additional info	ormation on page 2		
TATUS	EF	FECTIVE DATE (YYYY / MM / DD)	DURATION OF APPR	OVAL		

## TARGETED DMARDS FOR ANKYLOSING SPONDYLITIS

		.,						
PATIENT NAME		F	PHN					
SECTION 5 - CURRENT MEDICATIONS (DMARDs, anti-inflammato	ries, corticosteroi	ds, opioids)						
MEDICATION DOSE			FREQUENCY					
SECTION 6 - CRITERIA FOR RENEWAL (For the criteria original	<b>ally specified</b> on	intial coverag	ge form, please provic	le current statu	s)			
Please complete all sections below RESP			ONSE TO THERAPY COMPARED TO BASELINE					
<b>Extra-articular manifestations</b> (specify even if resolved):		○ Worsened	d No Response	○ Improved	Resolved			
uveitis IBD psoriasis other:		O	. — () т.	Op. 12.12.	<b>O</b> 110201122			
AXIAL DISEASE								
☐ Spinal pain		○ Worsened	d O No Response	O Improved	○ Resolved			
PERIPHERAL DISEASE								
Active joints (complete homunculus below)		○ Worsened	d O No Response	○ Improved	○ Resolved			
		-	•	-	0			
<ul> <li>Active tenosynovitis and/or enthesitis         (indicate by arrow and "TS" or "E" on homunculus as applicable)     </li> </ul>		○ Worsened	d O No Response	○ Improved	Resolved			
☐ Copy of a current BASDAI attached. ☐ Copy of HAQ attached if predominantly peripheral disease.								
ADDITIONAL COMMENTS REGARDING PATIENT'S CURRENT MEDICAL STATUS								

## **SECTION 7 - HOMUNCULUS** *Indicate active joints, tenosynovitis and enthesitis.*



Report all adverse events to the post-market surveillance program, Canadian Vigilance, toll-free 1-866-234-2345 (health professionals only).

Personal information on this form is collected under the authority of, and in accordance with, the *British Columbia Pharmaceutical Services Act* 22(1) and *Freedom of Information and Protection of Privacy Act* 26 (a),(c),(e). The information is being collected for the purposes of (a) administering the PharmaCare program, (b) analyzing, planning and evaluating the Special Authority and other Ministry programs and (c) to manage and plan for the health system generally. If you have any questions about the collection of this information, call Health Insurance BC from Vancouver at 1-604-683-7151 or from elsewhere in BC toll free at 1-800-663-7100 and ask to consult a pharmacist concerning the Special Authority process.

I have discussed with the patient that the

purpose of releasing their information to PharmaCare is to obtain Special Authority for prescription coverage and for the purposes set out here.

Rheumatologist's Signature (Mandatory)

PharmaCare may request additional documentation to support this Special Authority request.

Actual reimbursement is subject to the rules of a patient's PharmaCare plan, including any annual deductible requirement, and to any other applicable PharmaCare pricing policy.