

Prior Authorization Request

COSENTYX (secukinumab)

Instructions

Plan Member Signature

Please complete Part A and have your physician complete Part B. Completion and submission is not a guarantee of approval. Any fees related to the completion of this form are the responsibility of the plan member. Drugs in the Prior Authorization Program may be eligible for reimbursement if the patient does not qualify for coverage under a primary plan or a government program. Drugs used for indications not approved by Health Canada may be denied. For Quebec plan members, RAMQ exception drug criteria may apply. The decision for approval versus denial is based on pre-defined clinical criteria, primarily based on Health Canada approved indication(s) and on supporting evidence-based clinical protocols. The plan member will be notified whether their request has been approved or denied. Please note that you have the right to appeal the decision made by Express Scripts Canada.

Part A - Patient Patient information First Name: Last Name: Insurance Carrier Name/Number: Group Number: Client ID: Relationship: Employee Spouse Dependent Date of Birth (YYYY/MM/DD): Gender: Male Female Language: | English | French Address: City: Province: Postal Code: Email address: Telephone (home): Telephone (cell): Telephone (work): Coordination of benefits **Patient** Is the patient enrolled in any patient assistance program? Yes No **Assistance Program** Contact Name: _ Has the patient applied for reimbursement under a provincial plan? Yes No N/A **Provincial** Coverage What is the coverage decision of the drug? Approved Denied *Attach decision letter* Has the patient applied for reimbursement under a primary plan? | Yes | No | N/A **Primary** Coverage What is the coverage decision of the drug? Approved Denied *Attach decision letter* **Authorization** On behalf of myself and my eligible dependents, I authorize my group benefit provider, and its agents, to exchange the personal information contained on this form. I give my consent on the understanding that the information will be used solely for purposes of administration and management of my group benefit plan. This consent shall continue so long as my dependents and I are covered by, or are claiming benefits under the present group contract, or any modification, renewal, or reinstatement thereof.

Date



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Part B - Prescriber

Please see instructions on page 1 and complete all sections below. <u>Incomplete forms may result in automatic denial</u>. Please do **not** provide genetic test information or results.

SECTION 1 - DRUG REQUESTED

SECTION I - DIVOG NEQUE	51LD					
COSENTYX (secukinumab)		New request Renewal request*				
Dose	Administration (ex: oral, IV, etc)	Frequency	Duration			
Site of drug administration:			_			
Home Physician	n's office/Infusion clinic	Hospital (outpatient)	Hospital (inpatient)			
* Please submit proof of prior of	coverage if available					
SECTION 2 – ELIGIBILITY C	RITERIA					
1. Please indicate if the par	tient satisfies the below criteria:					
Plaque Psoriasis						
For the treatment of m	oderate to severe plaque psorias	is, AND				
The patient is 6 years of	of age or older, AND					
The patient has an affe hands, feet or genital r	ected body surface area (BSA) of a region, AND	10% or greater, or there is involv	vement of the patient's face,			
The patient has a Psor	iasis Area and Severity Index (PAS	SI) score of 10 or greater, AND				
The patient has had ar inaccessible, AND	n inadequate response or has a d	ocumented intolerance to photo	otherapy, unless it is			
The patient has had an inadequate response or has a documented intolerance to conventional systemic therapy, or to another biologic response modifier (<i>Please list prior therapies in the chart below</i>)						
Psoriatic Arthritis						
For the treatment of ps	soriatic arthritis in an adult, AND					
	n inadequate response or has a de RDs), or to another biologic respo					
Ankylosing Spondylitis						
For the treatment of ar	nkylosing spondylitis in an adult, A	AND				
The patient has a Bath scale, AND	Ankylosing Spondylitis Disease A	ctivity Index (BASDAI) score of 4	or greater on a 10-point			
inflammatory drugs (N	n inadequate response or has a do SAIDs) for a minimum of 2 weeks minimum of 3 months, or to anoth	each, or to at least 2 disease m	nodifying anti-rheumatic			



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Non-Radiographic Axial Spondyloarthriti	s				
For the treatment of non-radiog		nritis in an adult	, AND		
The patient has objective signs resonance imaging (MRI), AND				tein (CRP) and/or	r magnetic
The patient has had an inadeque inflammatory drugs (NSAIDs) for prior therapies in the chart below	or a minimum of 2 weeks				
OR None of the above criteria appl	ies.				
Relevant additional information:					
Please list previously tried therap	pies				
		Duration of therapy		Reason for cessation Inadequate Allergy/	
Drug	Dosage and administration			Inadequate	Allergy/
Drug		From	of therapy To		
Drug				Inadequate	Allergy/
Drug				Inadequate	Allergy/
Drug				Inadequate	Allergy/
Drug				Inadequate	Allergy/
Drug				Inadequate	Allergy/
Drug				Inadequate	Allergy/
SECTION 3 – PRESCRIBER INFORI	administration			Inadequate	Allergy/
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	administration			Inadequate	Allergy/
SECTION 3 – PRESCRIBER INFORI	administration			Inadequate	Allergy/
SECTION 3 – PRESCRIBER INFORI Physician's Name: Address:	administration	From		Inadequate	Allergy/

Please fax or mail the completed form to Express Scripts Canada®

Fax: Express Scripts Canada Clinical Services 1 (855) 712-6329

Mail: Express Scripts Canada Clinical Services 5770 Hurontario Street, 10th Floor Mississauga, ON L5R 3G5