

Prior Authorization Request

TARCEVA (erlotinib) and generics

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: Date of Birth (YYYY/MM/DD): Relationship: | Employee | Spouse | Dependent Language: English French Gender: | | Male | | Female 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								
TARCEVA (erlotinib) and generics		New request	Renewal request*					
Dose	Administration (ex: oral, IV, etc)	Frequency	Duration					
Site of drug administration:			<u> </u>					
☐ Home ☐ Physician's office/Infusion clinic ☐ Hospital (outpatient) ☐ Hospital (inpatient)								
* Please submit proof of prior coverage if available								
SECTION 2 – ELIGIBILITY CRITERIA								
Please indicate if the patient satisfies the below criteria:								
Non-Small Cell Lung Cancer – EGFR Expression Status Positive or Unknown								
For the treatment of locally advanced or metastatic non-small cell lung cancer (NSCLC) with epidermal growth factor receptor (EGFR) expression status positive or unknown in an adult, AND								
The patient has progressed after failure of at least one prior chemotherapy regimen (Please list prior therapies in the chart below), AND								
Erlotinib will be used as monotherapy								
Non-Small Cell Lung Cancer – EGFR Mutation, First-Line								
For the treatment of locally advanced or metastatic non-small cell lung cancer (NSCLC) with activating epidermal growth factor receptor (EGFR) mutations in an adult, AND								
The patient has not received prior systemic therapy, AND								
Erlotinib will be used as monotherapy								
Non-Small Cell Lung Cancer – E	GFR Mutation, Maintenance							
For the maintenance treatment of locally advanced or metastatic non-small cell lung cancer (NSCLC) with activating epidermal growth factor receptor (EGFR) mutations in an adult, AND								
The patient has received 4 cycles of standard platinum-based first-line chemotherapy (Please list prior therapies in the chart below), AND								
Erlotinib will be used a	s monotherapy							



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OR None of the above of	riteria applies.						
Relevant additional infor	mation:						
Please list previously tried therapies							
Drug	Dosage and	Duration of therapy		Reason for cessation			
	administration	From	То	Inadequate response	Allergy/ Intolerance		
			•				
SECTION 3 – PRESCRIBER INFORMATION							
Physician's Name:							
Address:							
Tel:		Fax:					
License No.:		Specialty:					
Physician Signature:		Date:					
Please fax or mail the Fax: Express Scripts Canada Clinical Services Mail: Express Scripts Canada Clinical Services							

completed form to **Express Scripts Canada®** 1 (855) 712-6329

5770 Hurontario Street, 10th Floor Mississauga, ON L5R 3G5