

Prior Authorization Request

LENVIMA (lenvatinib)

Instructions

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

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 Assistance	Is the patient enrolled in any patient assistance program?			
Program	Contact Name: Fax:			
Provincial Coverage	Has the patient applied for reimbursement under a provincial plan? Yes No N/A			
	What is the coverage decision of the drug? Approved Denied *Attach decision letter*			
Primary Coverage	Has the patient applied for reimbursement under a primary plan?			
	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

LENVIMA (lenvatinib)		New request	Renewal request*		
Dose	Administration (ex: oral, IV, etc)	Frequency	Duration		
Site of drug administration:					
Home Physician's office/Infusion clinic Hospital (outpatient) Hospital (inpatient)			Hospital (inpatient)		
* Please submit proof of prior coverage if available					

SECTION 2 - ELIGIBILITY CRITERIA

1. Please indicate if the patient satisfies the below criteria:
Differentiated Thyroid Cancer
For the treatment of locally recurrent or metastatic, progressive, radioactive iodine-refractory differentiated thyroid cancer in an adult
Renal Cell Carcinoma – First-Line
For the treatment of advanced or metastatic renal cell carcinoma (RCC) in an adult, AND
The patient has not received prior systemic therapy for metastatic disease, AND
LENVIMA will be used in combination with KEYTRUDA (pembrolizumab)
Renal Cell Carcinoma – Second-Line
For the treatment of advanced renal cell carcinoma in an adult, AND
The patient has had an inadequate response or has a documented intolerance to 1 or more vascular endothelial growth factor (VEGF)-targeted therapy (<i>Please list prior therapies in the chart below</i>), AND
LENVIMA will be used in combination with everolimus
Hepatocellular Carcinoma
For the treatment of unresectable hepatocellular carcinoma (HCC) in an adult, AND
The patient has not received prior systemic therapy, AND
The patient has Child-Pugh Score Class A liver function



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Endometrial Cancer	
For the treatment of advanced endometrial carcinoma that is not microsatellite instability high (MSI-H) or mismatch repair deficient (dMMR) in an adult, AND	
The patient has progressed following treatment with platinum-based systemic therapy (<i>Please list prior therapies in the chart below</i>), AND	
LENVIMA will be used in combination with KEYTRUDA (pembrolizumab)	
OR None of the above criteria applies.	
Relevant additional information:	

2. Please list previously tried therapies

Decore and	Duration	of therapy	Reason for cessation		
administration	From	То	Inadequate response	Allergy/ Intolerance	
	Dosage and administration	Dosage and administration	administration	Dosage and Inadequate	

SECTION 3 - PRESCRIBER INFORMATION

Physician's Name:				
Address:				
Tel:		Fax:		
License No.:		Specialty:		
Physician Signature:		Date:		
Please fax or mail the completed form to Express Scripts Canada®	Fax: Express Scripts Canada Cl 1 (855) 712-6329	inical Services N	Mail:	Express Scripts Canada Clinical Services 5770 Hurontario Street, 10 th Floor Mississauga, ON L5R 3G5