

Prior Authorization Request

TAFINLAR (dabrafenib)

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	Has the patient applied for reimbursement under a provincial plan? Yes No N/A				
Coverage	What is the coverage decision of the drug? Approved Denied *Attach decision letter*				
Primary	Has the patient applied for reimbursement under a primary plan?				
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.

Plan Member Signature

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

TAFINLAR (dabrafe	enib)		New request	Renewal request*	
Dose	Administration (ex: or	ral, IV, etc)	Frequency	Duration	
Site of drug administration:					
Home	e Physician's office/Infusion clinic		pital (outpatient)	Hospital (inpatient)	
* Please submit proof of prior coverage if available					

SECTION 2 – ELIGIBILITY CRITERIA

1. Please indicate if the patient satisfies the below criteria:
Melanoma – Unresectable or Metastatic
For the treatment of unresectable or metastatic melanoma with a BRAF V600 mutation in an adult, AND
TAFINLAR will be used as monotherapy and the patient has not been treated with BRAF or mitogen-activated protein kinase kinase (MEK) inhibitors, OR
TAFINLAR will be used in combination with MEKINIST (trametinib) and the patient has not previously progressed on a BRAF inhibitor
Melanoma – Resectable, Adjuvant
For the adjuvant treatment of melanoma with a BRAF V600 mutation and involvement of lymph node(s), following complete resection in an adult, AND
TAFINLAR will be used in combination with MEKINIST (trametinib)
Non-Small Cell Lung Cancer
For the treatment of metastatic non-small cell lung cancer (NSCLC) with a BRAF V600 mutation in an adult, AND
TAFINLAR will be used in combination with MEKINIST (trametinib)
Low-Grade Glioma
For the treatment of low-grade glioma (LGG) with a BRAF V600E mutation, AND
The patient is 1 year of age or older, AND
The patient has not received prior systemic therapy, AND
TAFINLAR will be used in combination with MEKINIST (trametinib)



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High-Grade Glioma					
For the treatment of high-g	grade glioma (HGG) with a BR	AF V600E mutati	ion, AND		
The patient is 1 year of ag	e or older, AND				
 The patient has relapsed, progressed, or failed to respond to radiation or chemotherapy (Please list prior therapies in the chart below), AND 					
TAFINLAR will be used in c	ombination with MEKINIST (t	rametinib)			
OR					
None of the above criteria	applies.				
Relevant additional information	n:				
2. Please list previously tried ther	apies				
Decore and		Duration of therapy		Reason for cessation	
	Dosage and	Duration o	of therapy		
Drug	Dosage and administration	Duration of From		Inadequate	r cessation Allergy/ Intolerance
Drug			of therapy To		Allergy/
Drug				Inadequate	Allergy/
Drug				Inadequate	Allergy/
Drug				Inadequate	Allergy/
Drug				Inadequate	Allergy/
Drug				Inadequate	Allergy/
Drug				Inadequate	Allergy/
	administration			Inadequate	Allergy/
	administration			Inadequate	Allergy/
	administration			Inadequate	Allergy/
SECTION 3 - PRESCRIBER INF	administration			Inadequate	Allergy/
SECTION 3 – PRESCRIBER INF Physician's Name:	administration			Inadequate	Allergy/

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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 Clir 1 (855) 712-6329	nical Services	Mail: Express Scripts Canada Clinical Services 5770 Hurontario Street, 10 th Floor Mississauga, ON L5R 3G5	