

LUSPATERCEPT SPECIAL AUTHORIZATION REQUEST FORM

Patients may or may not meet eligibility requirements as established by Alberta government sponsored drug programs.

Please complete all required sections to allow your request to be processed.

PATIENT INFORMATION					COVERAGE TYPE				
LAST NAME	FIRST NAME INITIAL			INITIAL	☐ Alberta Blue Cross				
BIRTH DATE (YYYY-MM-DD)	ALBERTA PERSONAL HEALTH NUMBER				☐ Alberta Human Services				
					Other				
ADDRESS	CITY PROV POSTAL CODE				ID/CLIENT/COVERAGE NUMBER				
PRESCRIBER INFORMATION									
PRESCRIBER LAST NAME FIRST NAME INITIAL PRESCRIBER PROFESSIONAL ASSOCIATION REGISTRATION							TION		
	☐ CPS	☐ CPSA ☐ ACO REGISTRATION NUMBER							
ADDRESS		CARNA DADA+C ACP Other							
CITY, PROVINCE			PHONE FAX						
POSTAL CODE	FAX NUMBE				PROVIDED WITH EACH REQUEST SUBMITTED				
Please provide the following information for ALL requests									
Diagnosis									
☐ Treatment of red blood cell (RBC) transfusion-dependent anemia associated with						١ ا	Weight (kg)		
□ beta-thalassemia → Complete Section I ONLY									
□ very low- to intermediate-risk MDS in patients with ring sideroblasts → Complete Section II ONLY □ Other, specify						-			
For patients new to coverage and already on the requested drug, specify start date (YYYY-MM-DD)									
Section I. Luspatercept for BETA-THALASSEMIA ASSOCIATED ANEMIA requests									
A. INITIAL requests for treatment-naïve and treatment-experienced BETA-THALASSEMIA patients									
Please indicate the patient's pre-treatment transfusion burden over the 24 weeks prior to treatment initiation (RBC units/8 weeks)									
Is the patient receiving regular transfusions, defined as the following in the 24 weeks prior to treatment initiation?									
Receiving 6 to 20 RBC units							☐ Yes	☐ No	
No transfusion-free period greater than 35 days							☐ Yes	□No	
B. RENEWAL requests and INITIAL requests for treatment-experienced BETA-THALASSEMIA patients									
Please indicate the patient's current transfusion burden over the last approval period (RBC units/8 weeks)									
Section II. Luspatercept for MYELODYSPLASTIC SYNDROME (MDS) ASSOCIATED ANEMIA requests									
A. INITIAL requests for treatment-naïve and treatment-experienced MDS patients									
Has the patient used erythropoietin-based the	rapy?								
☐ Yes, please indicate response									
□ No , please indicate why this therapy is not suitable									
B. RENEWAL requests and INITIAL request	ts for treatment-exper	ienced MC)S pati	ents					
Has the patient been RBC transfusion independent over a minimum of 16 consecutive weeks within the previous 24 weeks of treatment?					ious	☐ Yes	□No		
Additional information related to request									
PRESCRIBER'S SIGNATURE	DATE (YYYY-MM-DD)	Please forwa	rd this re	equest to					
T NESSKIBER S SIGNATURE	שאוב (דו דו - ואואויטט)	Alberta B 10009 108	Pease forward this request to Alberta Blue Cross, Clinical Drug Services 10009 108 Street NW, Edmonton, Alberta T5J 3C5 FAX 780-498-8384 in Edmonton • 1-877-828-4106 toll free all other areas						
ONCE VOUR REQUEST HAS SHO								01.10. G10G0	

The information on this form is being collected and pursuant to sections 20, 21 and 22 of the Health Information Act, and sections 33 and 34 of the Freedom of Information and Protection of Privacy Act, for the purposes of determining or verifying eligibility to participate in a program or receive a benefit, product or health service. If you have any questions regarding the collection or use of this information, please contact an Alberta Blue Cross privacy matters representative toll-free at 1-855-498-7302 or write to Privacy Matters, Alberta Blue Cross, 10009 - 108 Street, Edmonton AB T5J 3C5.



