Enquiries to Ministry of Health 0800 855 066

APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

Page 1 Form SA2289 December 2025

APPLICANT (stamp or sticker acceptable)	PATIENT NHI:	REFERRER Reg No:		
Reg No:	First Names:	First Names:		
Name:	Surname:	Surname:		
Address:	DOB:	Address:		
	Address:			
Fax Number:		Fax Number:		
Brentuximab				
Initial application — relapsed/refractory Hodgl Applications from any relevant practitioner. Appro Prerequisites(tick boxes where appropriate)				
and	refractory CD30-positive Hodgkin lymphoma after two	o or more lines of chemotherapy		
or				
Patient has relapsed/refractory CD30-positive Hodgkin lymphoma				
Patient has previously undergone autologous stem cell transplant				
and Patient has not previously received funded brentuximab vedotin and Response to brentuximab vedotin treatment is to be reviewed after a maximum of 6 treatment cycles				
Brentuximab vedotin to be administered at doses no greater than 1.8 mg/kg every 3 weeks				
Renewal — relapsed/refractory Hodgkin lymphoma				
Current approval Number (if known):				
Applications from any relevant practitioner. Approvals valid for 9 months. Prerequisites(tick boxes where appropriate)				
Patient has achieved a partial or complete response to brentuximab vedotin after 6 treatment cycles				
and Treatment remains clinically appropriate and the patient is benefitting from treatment and treatment is being tolerated				
Patient is to receive a maximum o	f 16 total cycles of brentuximab vedotin treatment			
Initial application — anaplastic large cell lymphoma Applications from any relevant practitioner. Approvals valid for 6 months. Prerequisites(tick boxes where appropriate)				
Patient has relapsed/refractory CI	030-positive systemic anaplastic large cell lymphoma			
	Patient has an ECOG performance status of 0-1			
Patient has not previously received brentuximab vedotin				
	Response to brentuximab vedotin treatment is to be reviewed after a maximum of 6 treatment cycles			
	Brentuximab vedotin to be administered at doses no greater than 1.8 mg/kg every 3 weeks			

I confirm the above details are correct and that in signing this form I understand I may be audited.

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APPLICANT (stamp or sticker acceptable)	PATIENT NHI:	REFERRER Reg No:	
Reg No:	First Names:	First Names:	
Name:	Surname:	Surname:	
Address:	DOB:	Address:	
	Address:		
Fax Number:		Fax Number:	
Brentuximab - continued			
Renewal — anaplastic large cell lymphoma			
Current approval Number (if known):			
Applications from any relevant practitioner. Approvals valid for 9 months.			
Prerequisites(tick boxes where appropriate)			
Patient has achieved a partial or complete response to brentuximab vedotin after 6 treatment cycles			
Treatment remains clinically appropriate and the patient is benefitting from treatment and treatment is being tolerated and			
Patient is to receive a maximum of 16 total cycles of brentuximab vedotin treatment			

I confirm the above details are correct and that in signing this form I understand I may be audited.