Enquiries to Ministry of Health 0800 855 066

APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

Page 1 Form SA1868 April 2024

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:
Venetoclax		
Initial application — relapsed/refractory chronic lymphocytic leukaemia Applications only from a relevant specialist or medical practitioner on the recommendation of a relevant specialist. Approvals valid for 7 months. Prerequisites(tick boxes where appropriate)		
Patient has chronic lymphocytic le	ukaemia requiring treatment	
Patient has received at least one prior therapy for chronic lymphocytic leukaemia		
Patient has not previously received funded venetoclax and		
The patient's disease has relapsed within 36 months of previous treatment		
Venetoclax to be used in combination with six 28-day cycles of rituximab commencing after the 5-week dose titration schedule with venetoclax		
Patient has an ECOG performance status of 0-2		
Renewal — relapsed/refractory chronic lymphocytic leukaemia Current approval Number (if known):		
Venetoclax is to be discontinued after a maximum of 24 months of treatment following the titration schedule unless earlier discontinuation is required due to disease progression or unacceptable toxicity		
Initial application — previously untreated chronic lymphocytic leukaemia with 17p deletion or TP53 mutation* Applications only from a relevant specialist or medical practitioner on the recommendation of a relevant specialist. Approvals valid for 6 months. Prerequisites(tick boxes where appropriate)		
Patient has previously untreated c	hronic lymphocytic leukaemia	
	There is documentation confirming that patient has 17p deletion by FISH testing or TP53 mutation by sequencing	
Patient has an ECOG performanc	e status of 0-2	
	hocytic leukaemia with 17p deletion or TP53 muta	ntion*
Current approval Number (if known):		
Applications only from a relevant specialist or medical practitioner on the recommendation of a relevant specialist. Approvals valid for 6 months. Prerequisites(tick box where appropriate)		
The treatment remains clinically appropriate and the patient is benefitting from and tolerating treatment		
Note: 'Chronic lymphocytic leukaemia (CLL)' includes small lymphocytic lymphoma (SLL)* and B-cell prolymphocytic leukaemia (B-PLL)*. Indications marked with * are Unapproved indications.		

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