Enquiries to Ministry of Health 0800 855 066

## APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

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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:	
Venetoclax			
Initial application — relapsed/refractory chroni Applications from any relevant practitioner. Approx Prerequisites(tick boxes where appropriate)			
Individual has chronic lymphocytic	leukaemia requiring treatment		
Individual has received at least one prior therapy for chronic lymphocytic leukaemia			
and Individual has not previously received funded venetoclax			
and The individual's disease has relapsed			
Venetoclax to be used in combination with six 28-day cycles of rituximab commencing after the 5-week dose titration schedule with venetoclax			
and Individual has an ECOG performance status of 0-2			
Renewal — relapsed/refractory chronic lymphocytic leukaemia  Current approval Number (if known):			
Initial application — previously untreated chronic lymphocytic leukaemia with 17p deletion or TP53 mutation* Applications from any relevant practitioner. Approvals valid for 6 months.  Prerequisites(tick boxes where appropriate)			
Individual has previously untreated	d chronic lymphocytic leukaemia		
There is documentation confirming that individual has 17p deletion by FISH testing or TP53 mutation by sequencing and			
Individual has an ECOG performation	nce status of 0-2		
Renewal — previously untreated chronic lymple Current approval Number (if known):		ntion*	
The treatment remains clinically appropri	iate and the patient is benefitting from and tolerating t des small lymphocytic lymphoma (SLL)* and B-cell p		

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

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Venetoclax - continued			
Initial application — previously untreated acute myeloid leukaemia Applications from any relevant practitioner. Approvals valid for 6 months.  Prerequisites(tick boxes where appropriate)  The individual is currently on treatment with venetoclax and met all remaining special authority criteria prior to commencing treatment or  Individual has previously untreated acute myeloid leukaemia (see note a), according to World Health Organization (WHO)  Classification  Venetoclax not to be used in combination with standard intensive remission induction chemotherapy  and  Venetoclax to be used in combination with azacitidine or low dose cytarabine			
Renewal — previously untreated acute myeloid Current approval Number (if known):	vals valid for 6 months.		
b) 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.