Enquiries to Ministry of Health 0800 855 066

## APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

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Initial application — relapsed/refractory chronic lymphocytic leukaemia Applications from any relevant practitioner. Approvals valid for 7 months.  Prerequisites(tick boxes where appropriate)  Individual has chronic lymphocytic leukaemia requiring treatment Individual has received at least one prior therapy for chronic lymphocytic leukaemia Individual has not previously received funded venetoclax Individual has not previously received funded venetoclax Individual's disease has relapsed Individual's disease has relapsed Individual has an ECOG performance status of 0-2    Renewal — relapsed/refractory chronic lymphocytic leukaemia   Current approval Number (if known):					
Address:					
Address:					
Fax Number: Fax Nu					
Venetoclax  Initial application — relapsed/refractory chronic lymphocytic leukaemia Applications from any relevant practitioner. Approvals valid for 7 months.  Prerequisites(tick boxes where appropriate)  Individual has chronic lymphocytic leukaemia requiring treatment Individual has received at least one prior therapy for chronic lymphocytic leukaemia Individual has not previously received funded venetoclax  The individual's disease has relapsed Individual's disease has relapsed Individual has an ECOG performance status of 0-2  Renewal — relapsed/refractory chronic lymphocytic leukaemia Current approval Number (if known):	 				
Initial application — relapsed/refractory chronic lymphocytic leukaemia Applications from any relevant practitioner. Approvals valid for 7 months.  Prerequisites(tick boxes where appropriate)  Individual has chronic lymphocytic leukaemia requiring treatment Individual has received at least one prior therapy for chronic lymphocytic leukaemia Individual has not previously received funded venetoclax 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 Individual has an ECOG performance status of 0-2  Renewal — relapsed/refractory chronic lymphocytic leukaemia Current approval Number (if known):					
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and					
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Venetoclax to be used in combination with six 28-day cycles of rituximab commencing after the 5-week dose titration schedule with venetoclax  Individual has an ECOG performance status of 0-2  Renewal — relapsed/refractory chronic lymphocytic leukaemia  Current approval Number (if known):	The individual's disease has relapsed				
Individual has an ECOG performance status of 0-2  Renewal — relapsed/refractory chronic lymphocytic leukaemia  Current approval Number (if known):	Venetoclax to be used in combination with six 28-day cycles of rituximab commencing after the 5-week dose titration schedule with venetoclax				
Current approval Number (if known):					
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 chronic lymphocytic leukaemia					
There is documentation confirming that individual has 17p deletion by FISH testing or TP53 mutation by sequencing and	7				
Individual has an ECOG performance status of 0-2					
Renewal — previously untreated chronic lymphocytic leukaemia with 17p deletion or TP53 mutation*  Current approval Number (if known):					
Applications from any relevant practitioner. 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.

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:		
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  Venetoclax to be used in combination with azacitidine or low dose cytarabine				
Renewal — previously untreated acute myeloid leukaemia  Current approval Number (if known):				
Applications from any relevant practitioner. Approvals valid for 6 months.  Prerequisites(tick box where appropriate)				
There is no evidence of disease progression Note:				
a) 'Acute myeloid leukaemia' includes myeloid sarcoma*				
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.