

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 from any relevant practitioner. Approvals valid for 8 months.

Prerequisites(tick boxes where appropriate)

<input type="checkbox"/>	Individual has chronic lymphocytic leukaemia requiring treatment
and	<input type="checkbox"/>
<input type="checkbox"/>	Individual has received at least one prior therapy for chronic lymphocytic leukaemia
and	<input type="checkbox"/>
<input type="checkbox"/>	Individual has not previously received funded venetoclax
and	<input type="checkbox"/>
<input type="checkbox"/>	The individual's disease has relapsed
and	<input type="checkbox"/>
<input type="checkbox"/>	Venetoclax to be used in combination with six 28-day cycles of rituximab commencing after the 5-week dose titration schedule with venetoclax
and	<input type="checkbox"/>
<input type="checkbox"/>	Individual has an ECOG performance status of 0-2

Renewal — relapsed/refractory chronic lymphocytic leukaemia

Current approval Number (if known):.....

Applications from any relevant practitioner. Approvals valid for 6 months.

Prerequisites(tick boxes where appropriate)

<input type="checkbox"/>	Treatment remains clinically appropriate and the individual is benefitting from and tolerating treatment
and	<input type="checkbox"/>
<input type="checkbox"/>	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 from any relevant practitioner. Approvals valid for 6 months.

Prerequisites(tick boxes where appropriate)

<input type="checkbox"/>	Individual has previously untreated chronic lymphocytic leukaemia
and	<input type="checkbox"/>
<input type="checkbox"/>	There is documentation confirming that individual has 17p deletion by FISH testing or TP53 mutation by sequencing
and	<input type="checkbox"/>
<input type="checkbox"/>	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)

<input type="checkbox"/>	The treatment remains clinically appropriate and the patient is benefitting from and tolerating treatment
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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.

Signed: Date:

Post application to Health New Zealand, Private Bag 3015, Wanganui – email: customerservice@health.govt.nz

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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)

or	<input type="checkbox"/> The individual is currently on treatment with venetoclax and met all remaining special authority criteria prior to commencing treatment
	<input type="checkbox"/> Individual has previously untreated acute myeloid leukaemia (see note a), according to World Health Organization (WHO) Classification
	and <input type="checkbox"/> Venetoclax not to be used in combination with standard intensive remission induction chemotherapy
	and <input type="checkbox"/> 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.

Signed: Date:

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