

**APPLICANT** (stamp or sticker acceptable) **PATIENT NHI:** ..... **REFERRER** Reg No: .....

Reg No: ..... First Names: ..... First Names: .....

Name: ..... Surname: ..... Surname: .....

Address: ..... DOB: ..... Address: .....

..... Address: ..... .....

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Fax Number: ..... Fax Number: .....

## 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  
**and** ☐ 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  
**and** ☐ 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):.....

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

**Prerequisites**(tick boxes where appropriate)

- ☐ Treatment remains clinically appropriate and the individual is benefitting from and tolerating treatment  
**and** ☐ 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)

- ☐ Individual has previously untreated chronic lymphocytic leukaemia  
**and** ☐ There is documentation confirming that individual has 17p deletion by FISH testing or TP53 mutation by sequencing  
**and** ☐ 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.

Signed: ..... Date: .....

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

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Fax Number: .....	.....	Fax Number: .....

**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

and

☐ 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 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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