

Use this checklist to determine if a patient meets the restrictions for funding in the **hospital setting**. For more details, refer to [Section H](#) of the Pharmaceutical Schedule. For community funding, see the [Special Authority Criteria](#).

PRESCRIBER

Name:

Ward:

PATIENT:

Name:

NHI:

Ibrutinib

INITIATION – chronic lymphocytic leukaemia (CLL)

Re-assessment required after 6 months

Prerequisites (tick boxes where appropriate)

Patient has chronic lymphocytic leukaemia (CLL) requiring therapy

and

Patient has not previously received funded ibrutinib

and

Ibrutinib is to be used as monotherapy

and

There is documentation confirming that patient has 17p deletion or TP53 mutation

and

Patient has experienced intolerable side effects with venetoclax monotherapy

or

Patient has received at least one prior immunochemotherapy for CLL

and

Patient's CLL has relapsed within 36 months of previous treatment

and

Patient has experienced intolerable side effects with venetoclax in combination with rituximab regimen

or

Patient's CLL is refractory to or has relapsed within 36 months of a venetoclax regimen

CONTINUATION – chronic lymphocytic leukaemia (CLL)

Re-assessment required after 12 months

Prerequisites (tick boxes where appropriate)

No evidence of clinical disease progression

and

The treatment remains appropriate and the patient is benefitting from 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 that the above details are correct:

Signed: Date: