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.

PRES	CRIB	ER			PATIENT:
Name	:				Name:
Ward:					NHI:
lbrut	inib				
Re-a	ssess	ment	t required a	nphocytic leukaemia (CLL) after 6 months where appropriate)	
	and	\mathcal{I}	Individual I	has chronic lymphocytic leukaemia (CLL) requiring	therapy
	and	C	Individual has not previously received funded ibrutinib		
	and	C	Ibrutinib is	to be used as monotherapy	
		or	and O	There is documentation confirming that the individual has experienced intolerable side effect	
			and O and	Individual has received at least one prior immuno Individual's CLL has relapsed Individual has experienced intolerable side effect	ts with venetoclax in combination with rituximab regimen
		or	O Indiv	vidual's CLL is refractory to or has relapsed following	ng a venetoclax regimen
Re-a	ssess equisi	meni ites	t required a (tick box wh	ic lymphocytic leukaemia (CLL) after 12 months here appropriate) clinical disease progression	
				tic leukaemia (CLL)' includes small lymphocytic lympocations marked with * are Unapproved indications.	phoma (SLL) and B-cell prolymphocytic

I confirm that the above details are correct:	
Signed:	Date: