HOSPITAL MEDICINES LIST RESTRICTIONS CHECKLIST

July 2025

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.

PRESC	CRIBI	ER		PATIENT:
Name:				
Ward:				NHI:
/edol	izur	nab)	
Re-ass	sessr	nent	requ	n's disease - adults uired after 6 months poxes where appropriate)
	and)	Patie	ent has active Crohn's disease
		or	0	Patient has had an initial approval for prior biologic therapy and has experienced intolerable side effects or insufficient benefit to meet renewal criteria (unless contraindicated)
		or	0	Patient has a CDAI score of greater than or equal to 300, or HBI score of greater than or equal to 10
			0	Patient has extensive small intestine disease affecting more than 50 cm of the small intestine
		or	0	Patient has evidence of short gut syndrome or would be at risk of short gut syndrome with further bowel resection
		or	0	Patient has an ileostomy or colostomy, and has intestinal inflammation
í	and	_		
			\circ	Patient has tried but experienced an inadequate response to (including lack of initial response and/or loss of initial response) from prior therapy with immunomodulators and corticosteroids
		or	0	Patient has experienced intolerable side effects from immunomodulators and corticosteroids
		or	0	Immunomodulators and corticosteroids are contraindicated
	`			
Re-ass	sessr	nent	t requ	Crohn's disease - adults uired after 2 years poxes where appropriate)
		or	0	CDAI score has reduced by 100 points, or HBI score has reduced by 3 points, from when the patient was initiated on biologic therapy
		or	0	CDAI score is 150 or less, or HBI is 4 or less
		or	0	The patient has experienced an adequate response to treatment, but CDAI score and/or HBI score cannot be assessed
í	and (O	Vedo	olizumab to administered at a dose no greater than 300 mg every 8 weeks

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

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RESCRIE	BER		PATIENT:
ame:			Name:
/ard:			NHI:
edolizu	mal) - co	ontinued
Re-assess	smen	t requ	n's disease - children* uired after 6 months poxes where appropriate)
and	0	Paec	diatric patient has active Crohn's disease
		0	Patient has had an initial approval for prior biologic therapy and has experienced intolerable side effects or insufficient benefit to meet renewal criteria (unless contraindicated)
	or	0	Patient has a Paediatric Crohn's Disease Activity Index (PCDAI) score of greater than or equal to 30
		0	Patient has extensive small intestine disease
and		0	Patient has tried but experienced an inadequate response to (including lack of initial response and/or loss of initial response) from prior therapy with immunomodulators and corticosteroids
	or	0	Patient has experienced intolerable side effects from immunomodulators and corticosteroids
	OI OI	0	Immunomodulators and corticosteroids are contraindicated
ote: Indi	catio	n mar	ked with * is an unapproved indication.
e-assess	smen	t requ	Crohn's disease - children* uired after 2 years poxes where appropriate)
	or	0	PCDAI score has reduced by 10 points from when the patient was initiated on biologic therapy
	or	0	PCDAI score is 15 or less The patient has experienced an adequate response to treatment, but CDAI score cannot be assessed
and	0	Vedo	olizumab to administered at a dose no greater than 300mg every 8 weeks
Note: Indi	catio	n mar	ked with * is an unapproved indication.

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PRES	CRIB	ER			PATIENT:		
Name	:				Name:		
Ward:					NHI:		
Vedo	lizu	mak) - co	ontinued			
Re-as	ssess	men	t requ	ntive colitis nired after 6 months noxes where appropriate)			
	and	0	Patie	nt has active ulcerative colitis			
		or	0	Patient has had an initial approval for prior biologic thera meet renewal criteria (unless contraindicated)	apy and has experienced intolerable side effects or insufficient benefit to		
			0	Patient has a SCCAI score is greater than or equal to 4			
		or	0	Patient's PUCAI score is greater than or equal to 20*			
	and						
			0	Patient has tried but experienced an inadequate respon from prior therapy with immunomodulators and corticost	se to (including lack of initial response and/or loss of initial response) teroids		
		or	0	Patient has experienced intolerable side effects from im	munomodulators and corticosteroids		
		or	0	Immunomodulators and corticosteroids are contraindica	ited		
Note:	Indio	catio	n mar	ked with * is an unapproved indication.			
Re-as	ssess	men	t requ	Ilcerative colitis ired after 2 years poxes where appropriate)			
		or	0	The SCCAI score has reduced by 2 points or more from	the SCCAI score since initiation on biologic therapy		
			0	The PUCAI score has reduced by 10 points or more from	m the PUCAI score since initiation on biologic therapy *		
	and (0	Vedo	lizumab will be used at a dose no greater than 300 mg in	ntravenously every 8 weeks		
Note:	India	catio	n mar	ked with * is an unapproved indication.			

I confirm that the above details are correct:

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