Enquiries to Ministry of Health 0800 855 066

## APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

Page 1 Form SA2182 August 2025

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:		
Ustekinumab				
at the time of commencing treatment or Patient has active Crohn's did and Patient has had an initive effects or insufficient by Patient meets the and Pati	th ustekinumab commenced prior to 1 February 2023 ent	ease and has experienced intolerable side		
Renewal — Crohn's disease - adults  Current approval Number (if known):				
CDAI score has reduced by therapy  or  CDAI score is 150 or less, or	100 points, or HBI score has reduced by 3 points, from HBI is 4 or less	om when the patient was initiated on biologic		
The patient has experienced	an adequate response to treatment, but CDAI score	and/or HBI score cannot be assessed		
and Ustekinumab to be administered at	t a dose no greater than 90 mg every 8 weeks			

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	Address:			
Fax Number:		Fax Number:		
Initial application — Crohn's disease - children* Applications from any relevant practitioner. Approvals valid for 6 months.  Prerequisites(tick boxes where appropriate)  Patient is currently on treatment with ustekinumab commenced prior to 1 February 2023 and met all remaining criteria (criterion 2) below at the time of commencing treatment  Patient has active Crohn's disease  and  Patient has had an initial approval for prior biologic therapy and has experienced intolerable side effects or insufficient benefit to meet renewal criteria  Patient meets the initiation criteria for prior biologic therapies for Crohn's disease  and  Other biologics for Crohn's disease are contraindicated				
Note: Indication marked with * is an unapproved in	dication.			
Renewal — Crohn's disease - children*  Current approval Number (if known):				
or PCDAI score is 15 or less or The patient has experienced	an adequate response to treatment, but CDAI score			

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Name:		Surname:	Surname:		
Address:		DOB:	Address:		
		Address:			
	per:		Fax Number:		
Ustekin	umab - continued				
Application	at the time of commencing treatments  Patient has active ulcerative and  Patient has had an initive effects or insufficient by and  Patient meets the and	ith ustekinumab commenced prior to 1 February 2023 ent	litis and has experienced intolerable side		
Renewal — ulcerative colitis  Current approval Number (if known):					
	or PUCAI score has reduced by	ed by 2 points or more from the SCCAI score since in y 10 points or more from the PUCAI score since initia			
an	Ustekinumab will be used at a dos	e no greater than 90 mg intravenously every 8 weeks			
Note: Cr	Note: Criterion marked with * is for an unapproved indication.				

I confirm the above details are correct and that in signing this form I understand I may be audited.