SA2483 - Upadacitinib

Crohn's disease - adult - Initial application	3
Crohn's disease - adult - Renewal	4
Crohn's disease - children* - Initial application	4
Crohn's disease - children* - Renewal	4
Rheumatoid Arthritis - Renewal	2
Rheumatoid Arthritis (previously treated with adalimumab or etanercept) - Initial application	2
Atopic dermatitis - Initial application	3
Atopic dermatitis - Renewal	
Ulcerative colitis - Initial application	5
Ulcerative colitis - Renewal	

APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

Page 2 Form SA2483 November 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:	
Upadacitinib			
Initial application — Rheumatoid Arthritis (previously treated with adalimumab or etanercept) Applications from any relevant practitioner. Approvals valid for 6 months. Prerequisites(tick boxes where appropriate) The individual has had an initial Special Authority approval for adalimumab and/or etanercept for rheumatoid arthritis The individual has experienced intolerable side effects with adalimumab and/or etanercept The individual has received insufficient benefit from at least a three-month trial of adalimumab and/or etanercept such that they do not meet the renewal criteria for rheumatoid arthritis And Rituximab is not clinically appropriate The individual is seronegative for both anti-cyclic citrullinated peptide (CCP) antibodies and rheumatoid factor The individual has been started on rituximab for rheumatoid arthritis in a Health NZ Hospital At four months following the initial course of rituximab the individual has received insufficient benefit such that they do not meet the renewal criteria for rheumatoid arthritis			
Renewal — Rheumatoid Arthritis			
or			

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

APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

Page 3 Form SA2483 November 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:	
Upadacitinib - continued			
Initial application — atopic dermatitis Applications from any relevant practitioner. Approvals valid for 6 months. Prerequisites(tick boxes where appropriate)			
Individual is currently on treatment	with upadacitinib for atopic dermatitis and met all re	maining criteria prior to commencing treatment	
Individual has moderate to severe atopic dermatitis, severity as defined by an Eczema Area and Severity Index (EASI) score of greater than or equal to 16 or a Dermatology Life Quality Index (DLQI) score of greater than or equal to 10 and Individual has received insufficient benefit from topical therapy (including topical corticosteroids or topical calcineurin inhibitors) for a 28-day trial within the last 6 months, unless contraindicated to all			
Individual has trialled and re ciclosporin, azathioprine, me	ceived insufficient benefit from at least one systemic ethotrexate or mycophenolate mofetil), unless contrain	ndicated to all	
while still on treatment but n	QI assessment has been completed for at least the m o longer than 1 month following cessation of each pri QLI assessment is no more than 1 month old at the tir	or treatment course	
Renewal — atopic dermatitis Current approval Number (if known):			
Individual has received a 75% or gupadacitinib	greater reduction in EASI score (EASI 75) as compare	ed to baseline EASI prior to commencing	
or	provement of 4 or more as compared to baseline DLC	QI prior to commencing upadacitinib	
Initial application — Crohn's disease - adult Applications from any relevant practitioner. Approvals valid for 6 months. Prerequisites(tick boxes where appropriate)			
Individual is currently on treatment	with upadacitinib for Crohn's disease and met all rer	naining criteria prior to commencing treatment	
Individual has active Crohn's	s disease		
Individual has had an benefit to meet renew	initial approval for prior biologic therapy and has expo al criteria	erienced intolerable side effects or insufficient	
Individual meets	s the initiation criteria for prior biologic therapies for C	rohn's disease	
Other biologic tl	nerapies for Crohn's disease are contraindicated		

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

APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

Page 4 Form SA2483 November 2025

APPL	ICANT (stamp or sticker acceptable)	PATIENT NHI:	REFERRER Reg No:
Reg N	lo:	First Names:	First Names:
Name	:	Surname:	Surname:
Addre	ss:	DOB:	Address:
		Address:	
Fax N	umber:		Fax Number:
Upac	dacitinib - continued		
Rene	ewal — Crohn's disease - adult		
Curre	ent approval Number (if known):		
	cations from any relevant practitioner. Appro- equisites(tick boxes where appropriate)	vals valid for 2 years.	
		pints from the CDAI score when the individual was in	itiated on biologic therapy
		from when individual was initiated on biologic therap	ру
	CDAI score is 150 or less		
	or HBI score is 4 or less		
	or	adequate response to treatment, but CDAI score car	anot be assessed
	The marriagar has experienced an	adoquate respense to rearmon, sat 02/11 cont out	
Initial application — Crohn's disease - children* Applications from any relevant practitioner. Approvals valid for 6 months. Prerequisites(tick boxes where appropriate)			
	Individual is currently on treatment	with upadacitinib for Crohn's disease and met all rer	naining criteria prior to commencing treatment
	Child has active Crohn's dis	ease	
		al approval for prior biologic therapy for Crohn's disea penefit to meet renewal criteria	se and has experienced intolerable side
		initiation criteria for prior biologic therapies for Crohr	n's disease
	and Other biologic t	nerapies for Crohn's disease are contraindicated	
Rene	ewal — Crohn's disease - children*		
Current approval Number (if known):			
Applications from any relevant practitioner. Approvals valid for 2 years.			
Prere	equisites(tick boxes where appropriate)		
	PCDAI score has reduced by 10 p	oints from the child was initiated on treatment	
	PCDAI score is 15 or less		
	The child has experienced an ade	quate response to treatment, but PCDAI score canno	t be assessed
Note	: Indications marked with * are unapproved ir	dications.	

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

APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

Page 5 Form SA2483 November 2025

APPL	ICANT (stamp or sticker acceptable)	PATIENT NHI:	REFERRER Reg No:
Reg N	lo:	First Names:	First Names:
Name	:	Surname:	Surname:
Addre	ss:	DOB:	Address:
		Address:	
Fax N	umber:		Fax Number:
Upac	lacitinib - continued		
App	Individual has active ulcerate and Individual has had an effects or insufficient to and Individual meets	with upadacitinib for ulcerative colitis and met all rem	colitis and has experienced intolerable side
Renewal — ulcerative colitis			
	ent approval Number (if known):		
	cations from any relevant practitioner. Approvequisites(tick boxes where appropriate)	als valid for 2 years.	
1.51	(
	The SCCAI score has reduced by 2 points or more from the SCCAI score when the individual was initiated on treatment or		
		oints or more from the PUCAI score when the individu	ual was initiated on treatment