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

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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		
and The individual has experience or The individual has received i do not meet the renewal crite. and Rituximab is not clinically apport or The individual is seronegative or The individual has been and The individual has been and At four months for	vals valid for 6 months. secial Authority approval for adalimumab and/or etangled intolerable side effects with adalimumab and/or ensufficient benefit from at least a three-month trial of eria for rheumatoid arthritis	adalimumab and/or etanercept such that they odies and rheumatoid factor ealth NZ Hospital
Renewal — Rheumatoid Arthritis		
or		

APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

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APPLICANT (stamp or sticker acceptable)		PATIENT NHI:	REFERRER Reg No:	
Reg No:		First Names:	First Names:	
Name	2:	Surname:	Surname:	
Addre	ess:	DOB:	Address:	
		Address:		
Fax N	lumber:		Fax Number:	
Upad	dacitinib - continued			
App	al application — atopic dermatitis lications from any relevant practitioner. Appro equisites(tick boxes where appropriate)	vals valid for 6 months.		
	Individual is currently on treatment	t with upadacitinib for atopic dermatitis and met all re	maining criteria prior to commencing treatment	
	Individual has moderate to s greater than or equal to 16 o	severe atopic dermatitis, severity as defined by an Ecor a Dermatology Life Quality Index (DLQI) score of g	reater than or equal to 10	
	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 received insufficient benefit from at least one systemic therapy for a minimum of three months (eg ciclosporin, azathioprine, methotrexate or mycophenolate mofetil), unless contraindicated to all			
		QI assessment has been completed for at least the moon longer than 1 month following cessation of each pri		
	The most recent EASI or DO	QLI assessment is no more than 1 month old at the tir	me of application	
	-			
Rene	ewal — atopic dermatitis			
Curre	ent approval Number (if known):			
	ications from any relevant practitioner. Approrequisites(tick boxes where appropriate)	vals valid for 12 months.		
	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	
App	al application — Crohn's disease - adult lications from any relevant practitioner. Appro equisites(tick boxes where appropriate)	vals valid for 6 months.		
	Individual is currently on treatment	t with upadacitinib for Crohn's disease and met all rer	maining 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	
		s the initiation criteria for prior biologic therapies for C	crohn's disease	
		herapies 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

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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:	
Upad	lacitinib - continued			
Rene	ewal — Crohn's disease - adult			
Curre	ent approval Number (if known):			
	cations from any relevant practitioner. Appro-			
Prere	equisites(tick boxes where appropriate)			
	CDAI score has reduced by 100 pe	pints from the CDAI score when the individual was ini	tiated on biologic therapy	
	or HBI score has reduced by 3 points	from when individual was initiated on biologic therap	y	
	or CDAI score is 150 or less			
	or			
	HBI score is 4 or less			
	The individual has experienced an	adequate response to treatment, but CDAI score car	nnot be assessed	
Appl	Child has active Crohn's dis	vals valid for 6 months. with upadacitinib for Crohn's disease and met all ren	se and has experienced intolerable side	
Rene	ewal — Crohn's disease - children*			
Curre	ent 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 points 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:	ote: Indications marked with * are unapproved indications.			

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APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

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						January 2020
APPLICANT (stamp or sticker acceptable)		sticker acceptable)	PATIENT NHI:	REFERRER Reg No:		
Reg N	lo:				First Names:	First Names:
Name	:				Surname:	Surname:
Addre	ss:				DOB:	Address:
					Address:	
Fax N	umbei	·:				Fax Number:
Upac	laciti	nib -	- cont	inued		
App	ication	ns from tes(tic	n any k box ndivid	Individual has active ulceration limited in the lim	with upadacitinib for ulcerative colitis and met all ren	colitis and has experienced intolerable side
Renewal — ulcerative colitis						
Current approval Number (if known):						
Applications from any relevant practitioner. Approvals valid for 2 years. Prerequisites(tick boxes where appropriate)						
	The SCCAI score has reduced by 2 points or more from the SCCAI score when the individual was initiated on treatment or			vidual was initiated on treatment		
	j. [P	UCAI	score has reduced by 10 po	pints or more from the PUCAI score when the individ	ual was initiated on treatment

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