## APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

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

[		The p	patient has had an initial Special Authority approval for adalimumab and/or etanercept for rheumatoid arthritis
and	The patient has experienced intolerable side effects from adalimumab and/or etanercept		
	or		The patient has received insufficient benefit from at least a three-month trial of adalimumab and/or etanercept such that they on not meet the renewal criteria for rheumatoid arthritis
and			
	or		The patient is seronegative for both anti-cyclic citrullinated peptide (CCP) antibodies and rheumatoid factor
		an	The patient has been started on rituximab for rheumatoid arthritis in a Health NZ Hospital d
			The patient has experienced intolerable side effects from rituximab
			At four months following the initial course of rituximab the patient has received insufficient benefit such that they do not meet the renewal criteria for rheumatoid arthritis

## **Renewal — Rheumatoid Arthritis**

Current approval Number (if known):.....

Applications only from a rheumatologist or Practitioner on the recommendation of a rheumatologist. Approvals valid for 6 months. Prerequisites(tick boxes where appropriate)

or

Following 6 months' initial treatment, the patient has at least a 50% decrease in active joint count from baseline and a clinically significant response to treatment in the opinion of the physician

On subsequent reapplications, the patient demonstrates at least a continuing 30% improvement in active joint count from baseline and a clinically significant response to treatment in the opinion of the physician

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

Signed: ..... Date: .....