Enquiries to Ministry of Health 0800 855 066

## APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

Page 1 **Form SA1744** 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	r:		Fax Number:		
Omalizun	nab				
Application	Patient has a diagnosis of severe asthma  and  Past or current evidence of atopy, documented by skin prick testing or RAST  and  Total serum human immunoglobulin E (IgE) between 76 IU/mL and 1300 IU/ml at baseline  and  Proven adherence with optimal inhaled therapy including high dose inhaled corticosteroid (budesonide 1,600 mcg per day or fluticasone propionate 1,000 mcg per day or equivalent), plus long-acting beta-2 agonist therapy (at least salmeterol 50 mcg bd or eformoterol 12 mcg bd) for at least 12 months, unless contraindicated or not tolerated				
	or contraindicated or not tolerat	es of systemic corticosteroids equivalent to at least 28 days treatment in the past 12 months, unless rated  exacerbations needing systemic corticosteroids in the previous 12 months, where an exacerbation is sted use of oral corticosteroids for at least 3 days or parenteral steroids			
and [ and		t (ACT) score of 10 or less ent's asthma control using the ACT and oral corticos weeks after the first dose to assess response to tre			

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

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Omalizur	nab - continued					
Initial application — severe chronic spontaneous urticaria Applications only from a clinical immunologist or dermatologist. Approvals valid for 6 months.  Prerequisites(tick boxes where appropriate)  Patient must be aged 12 years or older and						
and	Patient is symptomatic with Urticaria Activity Score 7 (UAS7) of 20 or above  and Patient has a Dermatology life quality index (DLQI) of 10 or greater  or Patient has a Urticaria Control Test (UCT) of 8 or less  and					
	Patient has been taking high dose antihistamines (e.g. 4 times standard dose) and ciclosporin (> 3 mg/kg day) for at least 6 weeks  Patient has been taking high dose antihistamines (e.g. 4 times standard dose) and at least 3 courses of systemic corticoster (> 20 mg prednisone per day for at least 5 days) in the previous 6 months  Patient has developed significant adverse effects whilst on corticosteroids or ciclosporin					
and	or Treatment to be stopped if in Complete response* to 6 do	nadequate response* following 4 doses				
Renewal — severe asthma						
Current approval Number (if known):						
An increase in the Asthma Control Test (ACT) score of at least 5 from baseline  and  A reduction in the maintenance oral corticosteroid dose or number of exacerbations of at least 50% from baseline						

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Address:	DOB:	Address:					
	Address:						
Fax Number:		Fax Number:					
Omalizumab - continued							
Renewal — severe chronic spontaneous urticaria							
Current approval Number (if known):							
Applications only from a clinical immunologist or dermatologist. Approvals valid for 6 months.  Prerequisites(tick boxes where appropriate)							
Patient has previously adequately or	Patient has previously adequately responded* to 6 doses of omalizumab						
	Patient has previously had a complete response* to 6 doses of omalizumab						
	essation of omalizumab therapy						

Note: \*Inadequate response defined as less than 50% reduction in baseline UAS7 and DLQI score, or an increase in Urticaria Control Test (UCT) score of less than 4 from baseline. Patient is to be reassessed for response after 4 doses of omalizumab. Complete response is defined as UAS7 less than or equal to 6 and DLQI less than or equal to 5; or UCT of 16. Relapse of chronic urticaria on stopping prednisone/ciclosporin does not justify the funding of omalizumab.

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