Enquiries to Ministry of Health 0800 855 066

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:		
Mepolizumab				
Initial application — Severe eosinophilic asth Applications only from a respiratory physician or Prerequisites(tick boxes where appropriate)	ma clinical immunologist. Approvals valid for 12 months.			
Patient must be aged 12 years of	rolder			
Patient must have a diagnosis of severe eosinophilic asthma documented by a respiratory physician or clinical immunologist				
and Conditions that mimic asthma eg. vocal cord dysfunction, central airway obstruction, bronchiolitis etc. have been excluded				
and Patient has a blood eosinophil count of greater than 0.5 × 10^9 cells/L in the last 12 months				
Patient must be adherent to optimised asthma therapy including inhaled corticosteroids (equivalent to at least 1000 mcg per day of fluticasone propionate) plus long acting beta-2 agonist, or budesonide/formoterol as part of the single maintenance and reliever therapy regimen, unless contraindicated or not tolerated				
and				
Patient has had at least 4 exacerbations needing systemic corticosteroids in the previous 12 months, where an exacerbation is defined as either documented use of oral corticosteroids for at least 3 days or parenteral corticosteroids or				
	Patient has received continuous oral corticosteroids of at least the equivalent of 10 mg per day over the previous 3 months			
and Treatment is not to be used in co	mbination with subsidised benralizumab			
Patient has an Asthma Control Test (ACT) score of 10 or less. Baseline measurements of the patient's asthma control using the ACT and oral corticosteroid dose must be made at the time of application, and again at around 52 weeks after the first dose to assess response to treatment				
	received an anti-IL5 biological therapy for their severe	eosinophilic asthma		
	ry or intolerant to previous anti-IL5 biological therapy			
	ble to continue treatment with previous anti-IL5 biologi ment	cal therapy and discontinued within 12 months		
Renewal — Severe eosinophilic asthma				
Current approval Number (if known):				
Applications only from a respiratory physician or Prerequisites (tick boxes where appropriate)	clinical immunologist. Approvals valid for 2 years.			
An increase in the Asthma Contr	ol Test (ACT) score of at least 5 from baseline			
	Exacerbations have been reduced from baseline by 50% as a result of treatment with mepolizumab			
Reduction in continuous oral corticosteroid use by 50% or by 10 mg/day while maintaining or improving asthma control				
<u> </u>				

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

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Mepolizumab - continued			
Initial application — eosinophilic granulomatosis with polyangiitis Applications only from a relevant specialist or any relevant practitioner on the recommendation of a relevant specialist. Approvals valid for 12 months. Prerequisites(tick boxes where appropriate) The patient has eosinophilic granulomatosis with polyangiitis and The patient has trialled and not received adequate benefit from at least one of the following for at least three months (unless contraindicated to all): azathioprine, cyclophosphamide, leflunomide, methotrexate, mycophenolate, or rituximab The patient has trialled prednisone for a minimum of three months and is unable to maintain disease control at doses below 7.5 mg per day Corticosteroids are contraindicated			
Renewal — eosinophilic granulomatosis with p Current approval Number (if known): Applications only from a relevant specialist or any r Prerequisites(tick box where appropriate) Patient has no evidence of clinical disease	relevant practitioner on the recommendation of a rele	vant specialist. Approvals valid for 12 months.	