HOSPITAL MEDICINES LIST RESTRICTIONS CHECKLIST

Use this checklist to determine if a patient meets the restrictions for funding in the **hospital setting**. For more details, refer to Section H of the Pharmaceutical Schedule. For community funding, see the Special Authority Criteria.

PRESC	CRIBER	PATIENT:
Name:		
Ward:		NHI:
Меро	lizuma	
Re-as Prere	sessmer quisites Preso	evere eosinophilic asthma required after 12 months ick boxes where appropriate) ibed by, or recommended by a respiratory physician or clinical immunologist, or in accordance with a protocol or guideline that has been ed by the Health NZ Hospital.
	and or and or and or and or	Patient must be aged 12 years or older Patient must have a diagnosis of severe eosinophilic asthma documented by a respiratory physician or clinical immunologist Conditions that mimic asthma eg. vocal cord dysfunction, central airway obstruction, bronchiolitis etc. have been excluded 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 herapy regimen, unless contraindicated or not tolerated 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 Patient has received continuous oral corticosteroids of at least the equivalent of 10 mg per day over the previous 3 months Freatment is not to be used in combination 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 esponse to treatment
CONT Re-as	sessmer quisites	Patient has not previously received an anti-IL5 biological therapy for their severe eosinophilic asthma Patient was refractory or intolerant to previous anti-IL5 biological therapy Patient was not eligible to continue treatment with previous anti-IL5 biological therapy and discontinued within 12 months of commencing treatment P-Severe eosinophilic asthma required after 2 years ick boxes where appropriate) Severe eosinophilic asthma required after 2 years ick boxes where appropriate)
and		An increase in the Asthma Control 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

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PRESCRIBER	PATIENT:
Name:	Name:
Ward:	NHI:
Mepolizumab - continued	
INITIATION – eosinophilic granulomatosis with polyangiitis Re-assessment required after 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 contraindicated to all): azathioprine, cyclophosphamide, leflun	
	ee months and is unable to maintain disease control at doses below
CONTINUATION – eosinophilic granulomatosis with polyangiitis Re-assessment required after 12 months Prerequisites (tick box where appropriate) O Patient has no evidence of clinical disease progression	

ı	COMMITTI	mai	ure	above	details	are	correct.

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Signed.	Date:	
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