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.

PRESCRIBER	PATIENT:			
Name:	Name:			
Ward:	NHI:			
Benralizumab				
INITIATION – Severe eosinophilic asthma Re-assessment required after 12 months Prerequisites (tick boxes where appropriate) Prescribed by, or recommended by a respirate endorsed by the Health NZ Hospital.	ory physician or clinical immunologist, or in accordance with a protocol or guideline that has been			
	e 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 and Patient has a blood eosinophil count of greater than 0.5 × 10°9 cells/L in the last 12 months and				
O Patient must be adherent to optimised a	asthma therapy including inhaled corticosteroids (equivalent to at least 1000 mcg per day of beta-2 agonist, or budesonide/formoterol as part of the anti-inflammatory reliever therapy plus icated or not tolerated			
or O Patient has received continuous of	pations needing systemic corticosteroids in the previous 12 months, where an exacerbation is e of oral corticosteroids for at least 3 days or parenteral corticosteroids 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 combinate and Patient has an Asthma Control Test (AC and oral corticosteroid dose must be mare response to treatment and	ion with subsidised mepolizumab CT) score of 10 or less. Baseline measurements of the patient's asthma control using the ACT ade at the time of application, and again at around 52 weeks after the first dose to assess			
O Patient was refractory or int	ord an anti-IL5 biological therapy for their severe eosinophilic asthma colerant to previous anti-IL5 biological therapy continue treatment with previous anti-IL5 biological therapy and discontinued within treatment			
CONTINUATION – Severe eosinophilic asthma Re-assessment required after 2 years Prerequisites (tick boxes where appropriate)				
Prescribed by, or recommended by a respirate endorsed by the Health NZ Hospital.	ory physician or clinical immunologist, or in accordance with a protocol or guideline that has been			
An increase in the Asthma Control Test	(ACT) score of at least 5 from baseline d from baseline by 50% as a result of treatment with benralizumab			
or	costeroid use by 50% or by 10 mg/day while maintaining or improving asthma control			

I confirm that the above details are correct:

Cianad.	Data.	
Siurieu.	 Date.	