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.

PRES	CRIB	ER	PATIENT:				
Name	:						
Ward:			NHI:				
Benr	alizu	ıma	ab				
INITIA Re-as	assessmerequisites Presended and and and and or and	or	Revere eosinophilic asthma I required after 12 months (tick boxes where appropriate)  ribed by, or recommended by a respiratory physician or clinical immunologist, or in accordance with a protocol or guideline that has been sed by the Health NZ Hospital.  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 must be adherent to optimised asthma therapy including inhaled corticosteroids (equivalent to at least 1000 mcg per day of fluitoasone propionate) plus long-acting beta-2 agonist, or budesonide/formoterol as part of the anti-inflammatory reliever therapy plus maintenance 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  Treatment is not to be used in combination with subsidised mepolizumab  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				
		or	O Patient has not previously received an anti-IL5 biological therapy for their severe eosinophilic asthma  O Patient was refractory or intolerant to previous anti-IL5 biological therapy  and O Patient was not eligible to continue treatment with previous anti-IL5 biological therapy and discontinued within 12 months of commencing treatment				
CONTINUATION – Severe eosinophilic asthma Re-assessment required after 2 years Prerequisites (tick boxes where appropriate)  Prescribed by, or recommended by a respiratory physician or clinical immunologist, or in accordance with a protocol or guideline endorsed by the Health NZ Hospital.  An increase in the Asthma Control Test (ACT) score of at least 5 from baseline							
		or	Exacerbations have been reduced from baseline by 50% as a result of treatment with benralizumab  Reduction in continuous oral corticosteroid use by 50% or by 10 mg/day while maintaining or improving asthma control				

I confirm that the above details are correct:

Cianad.	Data.	
Signeg	 Date	