

Bertilimumab

Cat. No.:	HY-P99474
CAS No.:	375348-49-5
Target:	CCR
Pathway:	GPCR/G Protein; Immunology/Inflammation
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.

BIOLOGICAL ACTIVITY

Description	Bertilimumab (CAT 213; iCo-008) is a human monoclonal antibody targeting eotaxin-1 (CCL11). Bertilimumab has the potential for allergic disorders research ^[1] .	
In Vitro	Bertilimumab (CAT 213) neutralizes the ability of eotaxin-1 to cause an increase in intracellular calcium signaling (with an IC ₅₀ value of 2.86 nM), migration of CCR3-expressing L1.2 cells (with an IC ₅₀ value of 0.48 nM), and inhibition of the eotaxin1-evoked shape change of human eosinophils in vitro (with an IC ₅₀ of 0.71 nM) ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
In Vivo	Bertilimumab (CAT 213) (0.01-10 mg/kg) administered i.v. 30 min before i.po. injection of human eotaxin1 (1 µg) causes a significant dose-dependent inhibition of eosinophil recruitment in IL-5-treated, ovalbumin-sensitized mice. Bertilimumab also significantly inhibits neutrophil and mononuclear cell influx into the air pouch, which resulted in a dose-related inhibition of total cell influx ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
	Animal Model:	Female BALB/c mice (17-21 g) injected human eotaxin1 ^[1] .
	Dosage:	0.01 mg/kg, 0.1 mg/kg, 1 mg/kg, 10 mg/kg
	Administration:	i.v.; once
	Result:	Caused a significant dose-dependent inhibition of eosinophil recruitment in IL-5-treated, ovalbumin-sensitized mice.

REFERENCES

[1]. Sarah Main, et al. A potent human anti-eotaxin1 antibody, CAT-213: isolation by phage display and in vitro and in vivo efficacy. J Pharmacol Exp Ther. 2006 Dec;319(3):1395-404.

Caution: Product has not been fully validated for medical applications. For research use only.

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