

Human IL-33 Alexa Fluor® 350-conjugated Antibody

Recombinant Monoclonal Goat IgG Clone # 40015C Catalog Number: IC36253U

100 µg

DESCRIPTION			
Species Reactivity	Human		
Specificity	Detects human IL-33 in direct ELISAs and Western blots.		
Source	Recombinant Monoclonal Goat IgG Clone # 40015C		
Purification	Protein A or G purified from cell culture supernatant		
Immunogen	E.coli-derived recombinant human IL-33 Ser112-Thr270 Accession # 095760		
Conjugate	Alexa Fluor 350 Excitation Wavelength: 346 nm Emission Wavelength: 442 nm		
Formulation	Supplied 0.2 mg/mL in a saline solution containing BSA and Sodium Azide. See Certificate of Analysis for details.		
	*Contains <0.1% Sodium Azide, which is not hazardous at this concentration according to GHS classifications. Refer to the Safety Data Sheet (SDS) for additional information and handling instructions.		

APPLICATIONS				
Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.				
	Recommended Concentration	Sample		
Intracellular Staining by Flow Cytometry	0.25-1 μg/10 ⁶ cells	Human PBMCs fixed with Flow Cytometry Fixation Buffer (Catalog # FC004) and permeabilized with ice-cold methanol		

PREPARATION AND STORAGE		
Shipping	The product is shipped with polar packs. Upon receipt, store it immediately at the temperature recommended below.	
Stability & Storage Protect from light. Do not freeze.		
	 12 months from date of receipt, 2 to 8 °C as supplied. 	

BACKGROUND

IL-33, also known as NF-HEV and DVS 27, is a 30 kDa proinflammatory protein that may also regulate gene transcription (1-3). DVS 27 was identifed as a gene that is up-regulated in vasospastic cerebral arteries (1). NF-HEV was described as a nuclear factor that is preferentially expressed in the endothelial cells of high endothelial venules relative to endothelial cells from other tissues (2). IL-33 was identified based on sequence and structural homology with IL-1 family cytokines (3). DVS 27, NF-HEV, and IL-33 share 100% amino acid sequence identity. IL-33 is constitutively expressed in smooth muscle and airway epithelia. It is up-regulated in arterial smooth muscle, dermal fibroblasts, and keratinocytes following IL-1α or IL-1β stimulation (1, 3). Similar to IL-1, IL-33 can be cleaved *in vitro* by caspase-1, generating an N-terminal fragment that is slightly shorter than the C-terminal fragment (3, 4). The N-terminal portion of full length IL-33 contains a predicted bipartite nuclear localization sequence and a homeodomain-like helix-turn-helix DNA binding domain. By immunofluorescence, full length IL-33 localizes to the nucleus in HUVECs and transfectants (2). The C-terminal fragment, corresponding to mature IL-33, binds and triggers signaling through mast cell IL-1 R4/ST2L, a longtime orphan receptor involved in the augmentation of Th2 cell responses (3, 5-7). A ternary signaling complex is formed by the subsequent association of IL-33 and ST2L with IL-1R AcP (8). Stimulation of Th2 polarized lymphocytes with mature IL-33 *in vitro* induces IL-5 and IL-13 secretion (3). *In vitro* administration of mature IL-33 promotes increased production of IL-5, IL-13, IgE, and IgA, as well as splenomegaly and inflammatory infiltration of mucosal tissues (3). Full length and mature human IL-33 share 52-58% as sequence identity with mouse and rat IL-33. Human IL-33 shares less than 20% as sequence identity with other IL-1 family proteins.

References:

- 1. Onda, H. et al. (1999) J. Cereb. Blood Flow Metab. 19:1279.
- 2. Baekkevold, E.S. et al. (2003) Am. J. Pathol. 163:69.
- 3. Schmitz, J. et al. (2005) Immunity 23:479.
- 4. Black, R.A. et al. (1989) J. Biol. Chem. 264:5323.
- 5. Xu, D. et al. (1998) J. Exp. Med. 187:787.
- 6. Lohning, M. et al. (1998) Proc. Natl. Acad. Sci. USA 95:6930.
- 7. Dinarello, C.A. (2005) Immunity 23:461.
- 8. Chackerian, A.A. et al. (2007) J. Immunol. 179:2551.





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