

DESCRIPTION

Species Reactivity	Human
Specificity	Detects human IL-4I1 in direct ELISAs.
Source	Monoclonal Rat IgG _{2B} Clone # 1006202
Purification	Protein A or G purified from hybridoma culture supernatant
Immunogen	Chinese Hamster Ovary cell line, CHO-derived human IL-4I1 Met1-His567 Accession # Q96RQ9
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied either lyophilized or as a 0.2 µm filtered solution in PBS.

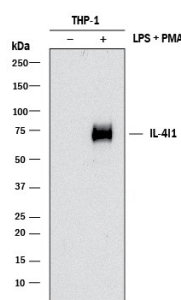
APPLICATIONS

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
Western Blot	2 µg/mL	See Below
Immunocytochemistry	5-25 µg/mL	See Below
Immunohistochemistry	8-25 µg/mL	See Below

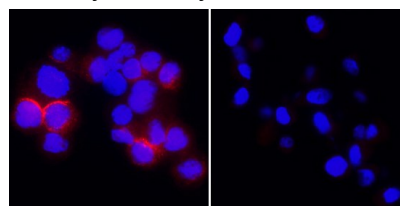
DATA

Western Blot



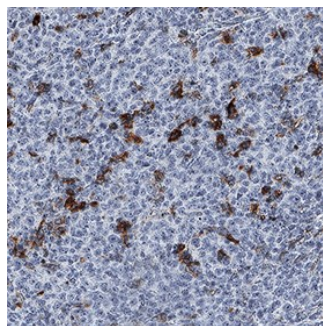
Detection of Human IL-4I1 by Western Blot. Western blot shows lysates of THP-1 human acute monocytic leukemia cell line untreated (-) or treated (+) with 200 nM PMA for 24 hours and 10 µg/mL LPS for 3 hours. PVDF membrane was probed with 2 µg/mL of Rat Anti-Human IL-4I1 Monoclonal Antibody (Catalog # MAB5684) followed by HRP-conjugated Anti-Rat IgG Secondary Antibody (Catalog # Catalog # HAF005). A specific band was detected for IL-4I1 at approximately 75 kDa (as indicated). This experiment was conducted under reducing conditions and using Immunoblot Buffer Group 1.

Immunocytochemistry



IL-4I1 in HDLM-2 Human Cell Line. IL-4I1 was detected in immersion fixed HDLM-2 human Hodgkin's lymphoma cell line using Rat Anti-Human IL-4I1 Monoclonal Antibody (Catalog # MAB5684) at 8 µg/mL for 3 hours at room temperature. Cells were stained using the NorthernLights™ 557-conjugated Anti-Rat IgG Secondary Antibody (red; Catalog # Catalog # NL013) and counterstained with DAPI (blue). Specific staining was localized to cytoplasm in lysosomes. View our protocol for [Fluorescent ICC Staining of Non-adherent Cells](#).

Immunohistochemistry



IL-4I1 in Human B Cell Lymphoma. IL-4I1 was detected in immersion fixed paraffin-embedded sections of human B cell lymphoma using Rat Anti-Human IL-4I1 Monoclonal Antibody (Catalog # MAB5684) at 5 µg/mL for 1 hour at room temperature followed by incubation with the Anti-Rat IgG VisUCyte™ HRP Polymer Antibody (Catalog # Catalog # VC005). Before incubation with the primary antibody, tissue was subjected to heat-induced epitope retrieval using Antigen Retrieval Reagent-Basic (Catalog # Catalog # CTS013). Tissue was stained using DAB (brown) and counterstained with hematoxylin (blue). Specific staining was localized to cytoplasm in lymphocytes. View our protocol for [IHC Staining with VisUCyte HRP Polymer Detection Reagents](#).

PREPARATION AND STORAGE

Reconstitution	Reconstitute at 0.5 mg/mL in sterile PBS.
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. • 12 months from date of receipt, -20 to -70 °C as supplied. • 1 month, 2 to 8 °C under sterile conditions after reconstitution. • 6 months, -20 to -70 °C under sterile conditions after reconstitution.

BACKGROUND

Interleukin 4 induced protein 1 (IL-4I1), also known as protein FIG-1 and L-amino acid oxidase, is encoded by a B-cell IL-4-inducible gene, FIG1, and is highly expressed in primary metastatic B-cell lymphomas (1-4). It belongs to the flavin monoamine oxidase family, FIG1 subfamily. Enzymological characterization reveals that IL-4I1 has L-amino acid oxidase activity with preference toward aromatic amino acids. Studies have shown that hIL-4I1 inhibited the proliferation of CD3-stimulated T lymphocytes with a similar effect on CD4(+) and CD8(+) T cells (5). Its inhibitory effect was dependent on enzymatic activity and H₂O₂ production. Its restricted expression to lymphoid tissues indicates that it may play an important function in the immune system (1, 4).

References:

1. Chu, C.C. and W.E. Paul. (1997) Proc. Natl. Acad. Sci. USA **94**:2507.
2. Mason, J.M. *et al.* (2004) J. Immunol. **173**:4561.
3. Chavan, S.S. *et al.* (2002) Biochim. Biophys. Acta. **1576**:70.
4. Copie-Bergman, C. *et al.* (2003) Blood **101**:2756.
5. Boulland, M.L. *et al.* (2007) Blood **110**:220.