

Product Data Sheet

Anti-CD172a Antibody

Catalog # Source Reactivity Applications

CPA2871 Rabbit H, Mk WB, IF/IC, IP

Description Rabbit polyclonal antibody to CD172a

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the C-term

region of human CD172a. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of CD172a protein.

Clonality Polyclonal

Form Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,

and 0.01% sodium azide.

Dilution WB (1/500 - 1/1000), IF/IC (1/100 - 1/500), IP (1/10 - 1/100)

Gene Symbol SIRPA

Alternative Names BIT; MFR; MYD1; PTPNS1; SHPS1; SIRP; Tyrosine-protein phosphatase non-receptor

type substrate 1; SHP substrate 1; SHPS-1; Brain Ig-like molecule with tyrosine-based

activation motifs; Bit; CD172 antigen-like family member A; Inhibitory receptor

SHPS-1; Macrophage fusion receptor; MyD-1 antigen; Signal-regulatory protein

alpha-1; Sirp-alpha-1; Signal-regulatory protein alpha-2; Sirp-alpha-2;

Signal-regulatory protein alpha-3; Sirp-alpha-3; p84; CD172a

Entrez Gene 140885 (Human)

SwissProt P78324 (Human)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

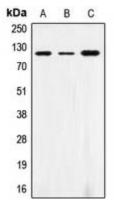
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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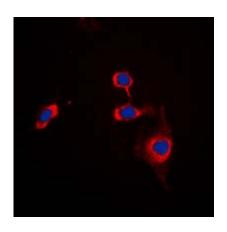
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Western blot analysis of CD172a expression in THP1 (A), AML193 (B), HL60 (C) whole cell lysates.



Immunofluorescent analysis of CD172a staining in THP1 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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