

Product Datasheet Anti-CD213a1 Antibody (orb393330)

Description Rabbit polyclonal antibody to IL13RA1.

Species/Host Rabbit

Reactivity Human, Mouse, Primate

Conjugation Unconjugated

Tested Applications IF, IH, WB

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

region of human CD213a1. The exact sequence is proprietary.

Target IL13RA1

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

glycerol, and 0.01% sodium azide.

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Storage Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -

20°C in small aliquots to prevent freeze-thaw cycles.

Note For research use only

Clonality Polyclonal

Antibody Type Primary Antibody

Source Rabbit

Uniprot ID P78552, 009030

Entrez 3597, 16164

Dilution Range WB: 1:500:1000, IHC-P: 1:100:200, IF/ICC: 1:100:500, IP: 1:10:100

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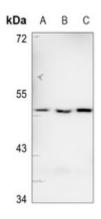
Email: <u>info@biorbyt.com</u>, <u>support@biorbyt.com</u> Phone: <u>+1 (415) 906-5211</u> | Fax: <u>+1 (415) 651-8558</u>



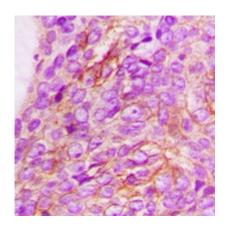


Expiration Date

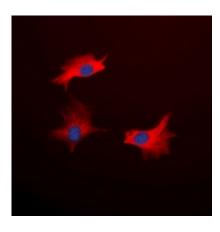
12 months from date of receipt.



Western blot analysis of CD213a1 expression in HEK293T (A), Hela (B), HGC27 (C) whole cell lysates. (Predicted band size: 48 kD; Observed band size: 49 kD)



Immunohistochemical analysis of CD213a1 staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of CD213a1 staining in MCF7 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 hidified 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).