

Product Datasheet

Anti-Beclin-1 Antibody (orb1473782)

Description Mouse monoclonal antibody to Beclin-1

Species/Host Mouse

Reactivity Human

Conjugation Unconjugated

Tested Applications IF, IH, WB

Immunogen Recombinant fusion protein of human Beclin-1. The exact sequence is

proprietary.

Target BECN1

Preservatives Mouse IgG3 kappa. Supplied in crude ascites with 0.01% sodium azide.

Form/Appearance Mouse IgG3 kappa. Supplied in crude ascites with 0.01% sodium azide.

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 Monoclonal

Antibody Type Primary Antibody

Source Mouse

Uniprot ID Q14457

Entrez 8678

Dilution Range WB (1/500 - 1/1000), IH (1/50 - 1/200), IF/IC (1/10 - 1/50)

Expiration Date 12 months from date of receipt.

Biorbyt Ltd.

7 Signet Court, Swann's Road, Cambridge, CB5 8LA, United Kingdom Email: info@biorbyt.com Phone: +44 (0) 1223 859-353 | Fax: +1 (415) 651-8558 **Biorbyt LLC.**

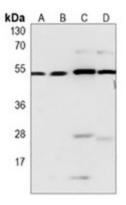
68 TW Alexander Drive,

Durham, NC, 27713, United States

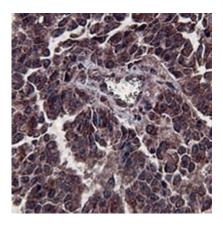
Email: $\underline{info@biorbyt.com}$, $\underline{support@biorbyt.com}$ Phone: $\underline{+1 (415) 906-5211}$ | Fax: $\underline{+1 (415) 651-8558}$



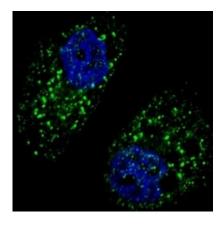




Western blot analysis of Beclin-1 expression in HepG2 (A), 293 (B), Jurkat (C), Hela (D) whole cell lysates. (Predicted band size: 51 kD; Observed band size: 51 kD)



Immunohistochemical analysis of Beclin-1 staining in human breast carcinoma 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 Beclin-1 staining in U251 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 AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).