

ABHD11 polyclonal antibody

Catalog: BS65323

Host: Rabbit

Reactivity: Human

BackGround:

abhydrolase domain containing 11(ABHD11) Homo sapiens This gene encodes a protein containing an alpha/beta hydrolase fold domain. This gene is deleted in Williams syndrome, a multisystem developmental disorder caused by the deletion of contiguous genes at 7q11.23. [provided by RefSeq, Mar 2016],

Product:

Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Molecular Weight:

~ 32 kDa

Swiss-Prot:

Q8NFV4

Purification&Purity:

The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.

Applications:

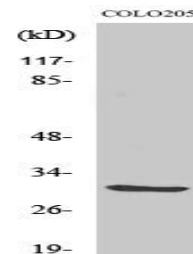
Western Blot: 1/500 - 1/2000. ELISA: 1/20000. Not yet tested in other applications.

Storage&Stability:

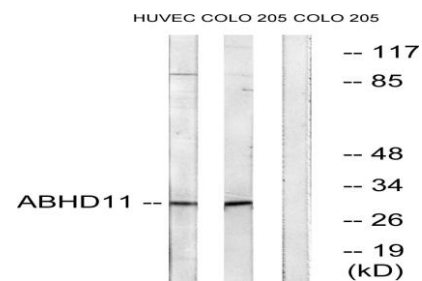
Store at 4 °C short term. Aliquot and store at -20 °C long term. Avoid freeze-thaw cycles.

Specificity:

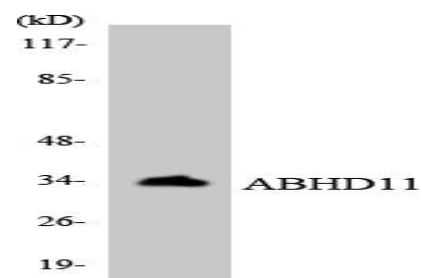
ABHD11 Polyclonal Antibody detects endogenous levels of ABHD11 protein.

DATA:

Western Blot analysis of various cells using ABHD11 Polyclonal Antibody



Western blot analysis of lysates from COLO and HUVEC cells, using ABHD11 Antibody. The lane on the right is blocked with the synthesized peptide.



Western blot analysis of the lysates from HeLa cells using ABHD11 antibody.

Note:

For research use only, not for use in diagnostic procedure.

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