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WBSCR17 Rabbit pAb

Catalog No.: A3481

Basic Information

Observed MW
68kDa

Calculated MW
68kDa

Category
Primary antibody

Applications
WB

Cross-Reactivity
Human, Mouse

Background

This gene encodes an N-acetylgalactosaminyltransferase. This gene is located centromeric to the common deleted region in Williams-Beuren syndrome (WBS), a multisystem developmental disorder caused by the deletion of contiguous genes at 7q11.23. This protein may play a role in membrane trafficking.

Recommended Dilutions

WB	1:500 - 1:2000

Immunogen Information

Gene ID
64409

Swiss Prot
Q6IS24

Immunogen
A synthetic peptide of human WBSCR17

Synonyms
GALNT16; GALNT20; GALNTL3; WBSCR17; GALNACT17; GalNAc-T17; GalNAc-T19; GalNAc-T5L

Contact

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Product Information

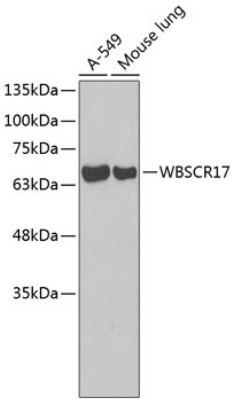
Source
Rabbit

Isotype
IgG

Purification
Affinity purification

Storage
Store at 4℃. Avoid freeze / thaw cycles.
Buffer: PBS with 0.02% sodium azide,pH7.3.

Validation Data



Western blot analysis of extracts of various cell lines, using WBSCR17 antibody (A3481).
Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.
Lysates/proteins: 25µg per lane.
Blocking buffer: 3% nonfat dry milk in TBST.