

DLL3 monoclonal antibody

Catalog: MB23355

Host: Mouse

Reactivity: Human

BackGround:

This gene encodes a member of the delta protein ligand family. This family functions as Notch ligands that are characterized by a DSL domain, EGF repeats, and a transmembrane domain. Mutations in this gene cause autosomal recessive spondylocostal dysostosis 1. Two transcript variants encoding distinct isoforms have been identified for this gene.

Product:

Purified antibody in PBS with 0.05% sodium azide

Molecular Weight:

65KDa

Swiss-Prot:

Q9NYJ7

Purification&Purity:

The antibody was affinity-purified from mouse ascites by affinity-chromatography using epitope-specific immunogen and the purity is > 95% (by SDS-PAGE).

Applications:

IHC:1/200 - 1/1000 FC:1/200 - 1/400

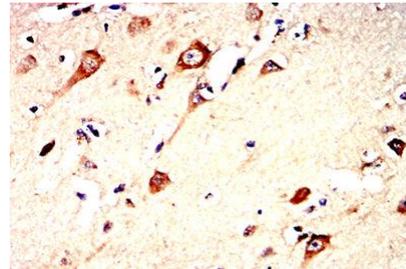
Storage&Stability:

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

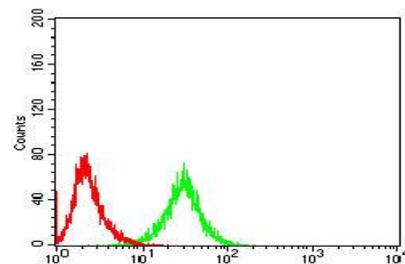
Isotype:

Mouse IgG1

DATA:



Immunohistochemical analysis of paraffin-embedded human brain tissues using DLL3 mouse mAb with DAB staining.



Flow cytometric analysis of HEK293 cells using DLL3 mouse mAb (green) and negative control (red).

Note:

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

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