

AMMR1 rabbit pAb

Catalog No: YT6949

Reactivity: Human; Mouse

Applications: WB

Target: AMMR1

Gene Name: AMMECR1

Protein Name: AMMR1

Human Gene Id: 9949

Human Swiss Prot

No:

Mouse Gene Id: 56068

Mouse Swiss Prot

No:

Immunogen: Synthesized peptide derived from human AMMR1 AA range: 12-62

Specificity: This antibody detects endogenous levels of AMMR1 at Human/Mouse

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

Source: Polyclonal, Rabbit, IgG

Q9Y4X0

Q9JHT5

Dilution: WB 1 ?500-2000

Purification: The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

1/2

Molecularweight: 37kD

Background: The exact function of this gene is not known, however, submicroscopic deletion

of the X chromosome including this gene, COL4A5, and FACL4 genes, result in a contiguous gene deletion syndrome, the AMME complex (Alport syndrome, mental retardation, midface hypoplasia, and elliptocytosis). Alternatively spliced transcript variants encoding different isoforms have been found for this gene.

[provided by RefSeq, Jan 2010],

Function: disease:Defects in AMMECR1 may be a cause of AMME complex

[MIM:300194]; also known as Alport syndrome with mental retardation, midface hypoplasia and elliptocytosis. The AMME complex is a contiguous gene deletion

syndrome., similarity: Contains 1 AMMECR1 domain.,

Subcellular Location:

Nucleus.

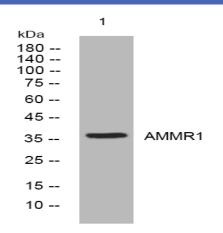
Sort: 1949

No4:

Host: Rabbit

Modifications: Unmodified

Products Images



Western blot analysis of lysates from SH-SY5Y cells, primary antibody was diluted at 1:1000, 4° over night