ENT2/SLC29A2 Rabbit mAb

Catalog No.: A19691 Recombinant



Basic Information

Observed MW

Calculated MW

50kDa

Category

Primary antibody

Applications

ELISA,IHC-P,IF/ICC

Cross-Reactivity Human, Mouse, Rat

CloneNo number

ARC2234

Background

The uptake of nucleosides by transporters, such as SLC29A2, is essential for nucleotide synthesis by salvage pathways in cells that lack de novo biosynthetic pathways. Nucleoside transport also plays a key role in the regulation of many physiologic processes through its effect on adenosine concentration at the cell surface (Griffiths et al., 1997 [PubMed 9396714]).

Recommended Dilutions

IHC-P 1:50 - 1:200
IF/ICC 1:50 - 1:200

Immunogen Information

Gene ID3177

Swiss Prot
Q14542

Immunogen

A synthetic peptide corresponding to a sequence within amino acids 1-100 of human ENT2/SLC29A2 (Q14542).

Synonyms

ENT2; DER12; HNP36; ENT2/SLC29A2

Contact

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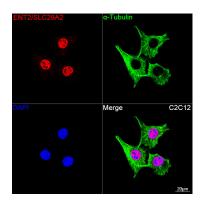
Product Information

SourceIsotypePurificationRabbitIgGAffinity purification

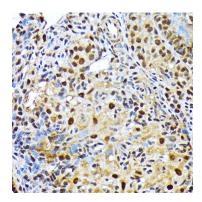
Storage

Store at -20°C. Avoid freeze / thaw cycles.

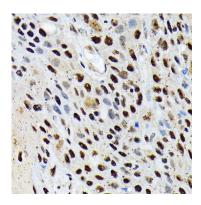
Buffer: PBS with 0.02% sodium azide, 0.05% BSA, 50% glycerol, pH7.3.



Confocal imaging of C2C12 cells using ENT2/SLC29A2 Rabbit mAb (A19691, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with $\alpha\text{-Tubulin}$ Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.



Immunohistochemistry analysis of paraffinembedded rat ovary using ENT2/SLC29A2 Rabbit mAb (A19691) at dilution of 1:100 (40x lens).Perform microwave antigen retrieval with 10 mM Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Immunohistochemistry analysis of paraffinembedded human esophageal cancer using ENT2/SLC29A2 Rabbit mAb (A19691) at dilution of 1:100 (40x lens).Perform microwave antigen retrieval with 10 mM Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.