SLC19A2 Antibody



PACO14998

Reactivity:

Product Information

Size: Protein Background:

This gene encodes the thiamin transporter protein. Mutations in this gene cause

thiamin-responsive megaloblastic anemia syndrome (TRMA), which is an autosomal

 $recessive\ disorder\ characterized\ by\ diabetes\ mellitus,\ megaloblastic\ anemia\ and$

Human sensorineural deafness.

Source: Gene ID:

Rabbit SLC19A2

Isotype: Uniprot

IgG O60779

Applications: Synonyms:

ELISA, IHC solute carrier family 19 (thiamine transporter), member 2

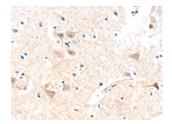
Recommended dilutions: Immunogen:

ELISA:1:1000-1:2000, IHC:1:15-1:50 Fusion protein of human SLC19A2.

Storage:

-20° C, pH7.4 PBS, 0.05% NaN3, 40% Glycerol

Product Images



The image on the left is immunohistochemistry of paraffin-embedded Human brain tissue using PACO14998(SLC19A2 Antibody) at dilution 1/15, on the right is treated with fusion protein. (Original magnification: x—200).