GLUT-1 (ABT197R) Rabbit mAb (Ready to Use)

CatalogNo: YM7128R Recombinant

Key Features

Host Species
- Rabbit

Reactivity
- Human,Mouse,Rat,

Applications
- IHC

Isotype
- IgG1,Kappa

Recommended Dilution Ratios

Ready to use for IHC

Storage

Storage*	2°C to 8°C/1 year
Formulation	The prediluted ready-to-use antibody is diluted in phosphate buffer saline containing stabilizing protein and 0.05% Proclin 300

Basic Information

Clonality	Monoclonal
Clone Number	ABT197R

Immunogen Information

Immunogen	Synthesized peptide derived from human GLUT-1 AA range:400-492
Specificity	This antibody detects endogenous levels of GLUT-1

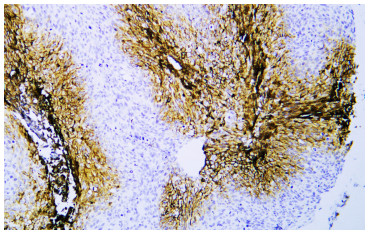
Target Information

Gene name	SLC2A1
Protein Name	Solute carrier family 2, facilitated glucose transporter member 1 (Glucose transporter type 1, erythrocyte/brain) (GLUT-1) (HepG2 glucose transporter)

Organism	Gene ID	UniProt ID
Human	6513;	P11166;

Cellular Localization	Membranous
Tissue specificity	Detected in erythrocytes (at protein level). Expressed at variable levels in many human tissues.
Function	Disease:Defects in SLC2A1 are the cause of autosomal dominant GLUT1 deficiency syndrome [MIM:606777]; also called blood-brain barrier glucose transport defect. This disease causes a defect in glucose transport across the blood-brain barrier. It is characterized by infantile seizures, delayed development, and acquired microcephaly.,Disease:Defects in SLC2A1 are the cause of dystonia type 18 (DYT18) [MIM:612126]. DYT18 is an exercise-induced paroxysmal dystonia/dyskinesia. Dystonia is defined by the presence of sustained involuntary muscle contraction, often leading to abnormal postures. DYT18 is characterized by attacks of involuntary movements triggered by certain stimuli such as sudden movement or prolonged exercise. In some patients involuntary exertion-induced dystonic, choreoathetotic, and ballistic movements may be associated with macrocytic hemolytic anemia.,Function:Facilitative glucose transporter. This isoform may be responsible for constitutive or basal glucose uptake. Has a very broad substrate specificity; can transport a wide range of aldoses including both pentoses and hexoses.,online information:GLUT1 entry,PTM:Phosphorylated upon DNA damage, probably by ATM or ATR.,similarity:Belongs to the major facilitator superfamily. Sugar transporter (TC 2.A.1.1) family. Glucose transporter subfamily.,subcellular location:Localizes primarily at the cell surface (By similarity). Identified by mass spectrometry in melanosome fractions from stage I to stage IV.,tissue specificity:Expressed at variable levels in many human tissues.,

Validation Data



Human cervical squamous cell carcinoma was stained with anti-GLUT-1 (ABT197R) rabbit mAb

Contact information

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Please scan the QR code to access additional product information:
GLUT-1 (ABT197R) Rabbit mAb (Ready to Use)