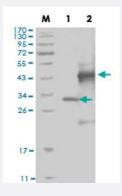


EPHA4 monoclonal antibody, clone 7D3D4

Catalog # MAB17687 Size 100 uL

Applications



Western Blot (Recombinant protein)

Western blot analysis of truncated (1) Trx-EphA4 recombinant protein and (2) GST-EPHA4 recombinant protein with EPHA4 monoclonal antibody.

Specification	
Product Description	Mouse monoclonal antibody raised against recombinant human EPHA4.
Immunogen	Recombinant protein corresponding to amino acids 777-986 of human EPHA4 from E. coli.
Host	Mouse
Reactivity	Human
Form	Liquid
Isotype	lgG1
Recommend Usage	ELISA (1:10000) Flow Cytometry Immunocytochemistry Immunohistochemistry Western Blot (1:500-1:2000) The optimal working dilution should be determined by the end user.
Storage Buffer	In ascites (0.03% sodium azide).



Product Information

Storage Instruction	Store at 4°C. For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.
Note	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which shoul d be handled by trained staff only.

Applications

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Gene Info — EPHA4	
Entrez GeneID	2043
Gene Name	EPHA4
Gene Alias	HEK8, SEK, TYRO1
Gene Description	EPH receptor A4
Omim ID	<u>602188</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene belongs to the ephrin receptor subfamily of the protein-tyrosine kinase family. EPH and EPH-related receptors have been implicated in mediating developmental events, particularly in the enervous system. Receptors in the EPH subfamily typically have a single kinase domain and an extracellular region containing a Cys-rich domain and 2 fibronectin type III repeats. The ephrin receptors are divided into 2 groups based on the similarity of their extracellular domain sequences and their affinities for binding ephrin-A and ephrin-B ligands. [provided by RefSeq
Other Designations	OTTHUMP00000164185 TYRO1 protein tyrosine kinase ephrin receptor EphA4 ephrin type-A receptor 4 receptor protein-tyrosine kinase HEK8 tyrosine-protein kinase receptor SEK

Pathway

Axon guidance

Disease



- Alzheimer Disease
- Cognition Disorders
- Genetic Predisposition to Disease
- Parkinson disease