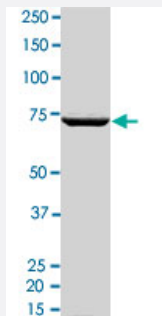


Bioactive

## EPHA2 (Human) Recombinant Protein

Catalog # P4677      Size 100 ug

### Applications



### Result of activity analysis

Result of activity analysis

□

### Specification

<b>Product Description</b>	Human EPHA2 (NM_004431.2, 585 a.a. - 976 a.a.) partial recombinant protein with GST-His tag expressed in Sf9 cells.
<b>Host</b>	insect
<b>Theoretical MW (kDa)</b>	73.691
<b>Form</b>	Liquid
<b>Preparation Method</b>	Insect cell (Sf9) expression system
<b>Purification</b>	One-step affinity purification using GSH-agarose
<b>Concentration</b>	0.505 ug/uL

Activity	30 pmol/ug x min
Quality Control Testing	2 ug/lane SDS-PAGE Stained with Coomassie Blue
Storage Buffer	In 50 mM Tris-HCl, 100 mM NaCl, pH 8.0. (5 mM DTT, 4 mM reduced glutathione, 20% glycerol)
Storage Instruction	Store at -80°C. Aliquot to avoid repeated freezing and thawing
Note	Result of activity analysis Result of activity analysis

## Applications

- Functional Study
- SDS-PAGE

## Gene Info — EPHA2

Entrez GeneID	<a href="#">1969</a>
Protein Accession#	<a href="#">NM_004431.2</a>
Gene Name	EPHA2
Gene Alias	ECK
Gene Description	EPH receptor A2
Omim ID	<a href="#">176946</a>
Gene Ontology	<a href="#">Hyperlink</a>
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 nervous 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. This gene encodes a protein that binds ephrin-A ligands. [provided by RefSeq]
Other Designations	ephrin receptor EphA2 epithelial cell receptor protein tyrosine kinase protein tyrosine kinase receptor protein tyrosine kinase regulated by p53 and E2F-1 soluble EPHA2 variant 1

## Pathway

- [Axon guidance](#)

## Disease

- [Cataract](#)
- [Genetic Predisposition to Disease](#)
- [Hearing Loss](#)