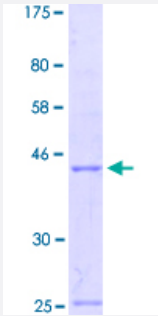


# EN2 (Human) Recombinant Protein (Q01)

Catalog # H00002020-Q01      Size 25 ug, 10 ug

## Applications



Specification	
Product Description	Human EN2 partial ORF (NP_001418.2, 86 a.a. - 210 a.a.) recombinant protein with GST-tag at N-terminal.
Sequence	GTCCAGAGGGRGGGAGGEGGASGAEGGGGAGGSEQLLGSGSREPRQNPPCAPGAGGPLPAA GSDSPGDGEGGSKTSLHGGAKKGGDPGGPLDGSLKARGLGGGDLVSSDSDSSQAGANLGA QP
Host	Wheat Germ (in vitro)
Theoretical MW (kDa)	39.38
Interspecies Antigen Sequence	Mouse (79); Rat (78)
Preparation Method	<a href="#">in vitro wheat germ expression system</a>
Purification	Glutathione Sepharose 4 Fast Flow
Quality Control Testing	12.5% SDS-PAGE Stained with Coomassie Blue
Storage Buffer	50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.
Storage Instruction	Store at -80°C. Aliquot to avoid repeated freezing and thawing.
Note	Best use within three months from the date of receipt of this protein.

# Applications

- Enzyme-linked Immunoabsorbent Assay
- Western Blot (Recombinant protein)
- Antibody Production
- Protein Array

# Gene Info — EN2

Entrez GeneID	<a href="#">2020</a>
GeneBank Accession#	<a href="#">NM_001427</a>
Protein Accession#	<a href="#">NP_001418.2</a>
Gene Name	EN2
Gene Alias	AUTS1, AUTS10
Gene Description	engrailed homeobox 2
Omim ID	<a href="#">131310 611016</a>
Gene Ontology	<a href="#">Hyperlink</a>
Gene Summary	Homeobox-containing genes are thought to have a role in controlling development. In Drosophila, the 'engrailed' (en) gene plays an important role during development in segmentation, where it is required for the formation of posterior compartments. Different mutations in the mouse homologs, En1 and En2, produced different developmental defects that frequently are lethal. The human engrailed homologs 1 and 2 encode homeodomain-containing proteins and have been implicated in the control of pattern formation during development of the central nervous system. [provided by RefSeq]
Other Designations	engrailed homolog 2 engrailed-2

# Disease

- [Autistic Disorder](#)
- [Child Development Disorders](#)

- [Genetic Predisposition to Disease](#)
- [Parkinson disease](#)