

Product datasheet for **TP502773**

Ebp (NM_007898) Mouse Recombinant Protein

Product data:

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|---------------------------------------|--|
| Product Type: | Recombinant Proteins |
| Description: | Purified recombinant protein of Mouse phenylalkylamine Ca ²⁺ antagonist (emopamil) binding protein (Ebp), with C-terminal MYC/DDK tag, expressed in HEK293T cells, 20ug |
| Species: | Mouse |
| Expression Host: | HEK293T |
| Expression cDNA Clone or AA Sequence: | >MR202773 protein sequence Red =Cloning site Green =Tags(s) |
| | MTTNTVPLHPYWPRHLKLDNFVPNDLPTSHILVGLFSISGGLIVITWLLSSRASVPLGAGRRRLALCWFA VCTFIHLVIEGWFSLYNGILLEDQAFLSQLWKEYSKGDSRYILSDSFVVCMETVTACLWGPLSLWVVIAF LRQQPFRFVLQLVSMGQIYGDVLYFLTELHEGLQHGEIGHVPYFVWFVFLNAVWLVIP SILVLD AIKH LTSAQSVLDSKVMKIKSKHN TRTRPLEQKLISEEDLAANDILDYKDDDDKV |
| Tag: | C-MYC/DDK |
| Predicted MW: | 26.2 kDa |
| Concentration: | >0.05 µg/µL as determined by microplate BCA method |
| Purity: | > 80% as determined by SDS-PAGE and Coomassie blue staining |
| Buffer: | 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol |
| Note: | For testing in cell culture applications, please filter before use. Note that you may experience some loss of protein during the filtration process. |
| Storage: | Store at -80°C after receiving vials. |
| Stability: | Stable for 12 months from the date of receipt of the product under proper storage and handling conditions. Avoid repeated freeze-thaw cycles. |
| RefSeq: | NP_031924 |
| Locus ID: | 13595 |
| UniProt ID: | P70245 |
| RefSeq Size: | 1755 |



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Cytogenetics: X 3.7 cM

RefSeq ORF: 693

Synonyms: AI255399; m; mSl; P; Pabp; Sl; Td

Summary: This gene encodes a transmembrane protein that localizes to the endoplasmic reticulum. This protein catalyses the conversion of delta8 to delta7 sterols, an important step in sterol biosynthesis. Mutations in this gene are responsible for the mouse tattered mutant phenotype. Tattered males are embryonic lethal, while heterozygous females have developmental defects. Deficiency of the related gene in human causes X-linked dominant chondrodysplasia punctata. [provided by RefSeq, May 2015]