

## Product datasheet for **TP760061**

### SLC25A38 (NM\_017875) Human Recombinant Protein

#### Product data:

|                                       |  |
|---------------------------------------|--|
| Product Type:                         | Recombinant Proteins   |
| Description:                          | Recombinant protein of human solute carrier family 25, member 38 (SLC25A38), full length, with N-terminal HIS tag, expressed in E.Coli, 50ug         |
| Species:                              | Human  |
| Expression Host:                      | E. coli  |
| Expression cDNA Clone or AA Sequence: | A DNA sequence encoding human full-length SLC25A38   |
| Tag:                                  | N-His  |
| Predicted MW:                         | 33.6 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, pH 8.0, 150 mM NaCl, 1% sarkosyl, 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.  |
| 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:                               | <a href="#">NP_060345</a>  |
| Locus ID:                             | 54977  |
| UniProt ID:                           | <a href="#">Q96DW6</a>   |
| RefSeq Size:                          | 2124   |
| Cytogenetics:                         | 3p22.1   |
| RefSeq ORF:                           | 912  |
| Synonyms:                             | SIDBA2   |


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**Summary:**

This gene is a member of the mitochondrial carrier family. The encoded protein is required during erythropoiesis and is important for the biosynthesis of heme. Mutations in this gene are the cause of autosomal congenital sideroblastic anemia (anemia, sideroblastic, 2, pyridoxine-refractory). A related pseudogene is found on chromosome 1. [provided by RefSeq, Aug 2017]

**Product images:**