

Datasheet for ABIN7319115 TPI1 Protein (His tag)



Overview

| Quantity: | 50 µg |
|-------------------------------|--|
| Target: | TPI1 |
| Origin: | Human |
| Source: | Escherichia coli (E. coli) |
| Protein Type: | Recombinant |
| Purification tag / Conjugate: | This TPI1 protein is labelled with His tag. |
| | |
| Product Details | |
| Purpose: | Recombinant Human TPI1/TIM Protein (His Tag) |

| Sequence: | Met 1-Gln249 |
|------------------|---|
| Characteristics: | Recombinant Human Triosephosphate Isomerase is produced by our E.coli expression system and the target gene encoding Met1-Gln249 is expressed with a 6His tag at the N-terminus. |
| Purity: | > 90 % as determined by reducing SDS-PAGE. |
| Endotoxin Level: | < 1.0 EU per μ g as determined by the LAL method. |

Target Details

| Target: | TPI1 |
|-------------------|--|
| Alternative Name: | TPI1/TIM (TPI1 Products) |
| Background: | Background: Triose-phosphate isomerase, also named Triose-phosphate isomerase, TPI and TIM, is an enzyme that catalyzes the reversible interconversion of the triose phosphate isomers |
| | dihydroxyacetone phosphate and D-glyceraldehyde 3-phosphate. TPI has been found in nearly |

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| | every organism searched for the enzyme, including animals such as mammals and insects as |
|---------------------|--|
| | well as in fungi, plants, and bacteria. However, some bacteria that do not perform glycolysis, |
| | like ureaplasmas, lack TPI. TPI plays an important role in glycolysis and is essential for efficient |
| | energy production. TPI deficiency is an autosomal recessive disorder and the most severe |
| | clinical disorder of glycolysis. Triose phosphate isomerase deficiency is associated with |
| | neonatal jaundice, chronic hemolytic anemia, progressive neuromuscular dysfunction, |
| | cardiomyopathy and increased susceptibility to infection and characterized by chronic |
| | hemolytic anemia. |
| | Synonym: Triosephosphate Isomerase, TIM, Triose-Phosphate Isomerase, TPI1, TPI |
| Molecular Weight: | 28.8 kDa |
| UniProt: | P60174 |
| Pathways: | Cell RedoxHomeostasis |
| Application Details | |
| Restrictions: | For Research Use only |
| Handling | |
| Format: | Frozen, Liquid |
| Buffer: | Supplied as a 0.2 μm filtered solution of 20 mM TrisHCl, 1 mM DTT, 10 % Glycerol, pH 8.0. |
| Storage: | -20 °C |
| Storage Comment: | Store at < -20°C, stable for 6 months. Please minimize freeze-thaw cycles. |
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