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Datasheet for ABIN4997569 anti-BPGM antibody (Alexa Fluor 750)



Overview

| Quantity: | 100 µL |
|--------------|--------------------------------------------------------------------------------------------------------|
| Target: | BPGM |
| Reactivity: | Human, Mouse, Rat |
| Host: | Rabbit |
| Clonality: | Polyclonal |
| Conjugate: | This BPGM antibody is conjugated to Alexa Fluor 750 |
| Application: | Western Blotting (WB), Immunofluorescence (Paraffin-embedded Sections) (IF (p)), Flow Cytometry (FACS) |

Product Details

| Immunogen: | KLH conjugated synthetic peptide derived from human BPGM |
|-------------------|----------------------------------------------------------|
| Isotype: | lgG |
| Cross-Reactivity: | Human, Mouse, Rat |
| Purification: | Purified by Protein A. |

Target Details

| Target: | BPGM |
|-------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Alternative Name: | BPGM (BPGM Products) |
| Background: | Synonyms: 2,3-bisphosphoglycerate mutase, 2,3-bisphosphoglycerate synthase, 3- bisphosphoglycerate mutase, 3-bisphosphoglycerate synthase, 3-diphosphoglycerate mutase, |
| | Ab2 098, Al323730, AL022789, Bisphosphoglycerate mutase, BPG dependent PGAM, BPG- |

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| Gene ID: | 669 |
|----------|---------------------------------------------------------------------------------------------------|
| | hemolytic anemia, splenomegaly, cholelithiasis and cholecystitis. |
| | the cause of bisphosphoglycerate mutase deficiency (BPGMD) . A disease characterized by |
| | Citrullinemia and Shwachman-Diamond syndrome. Involvement in disease:Defects in BPGM are |
| | Osteogenesis imperfecta, Williams-Beuren syndrome, Pendred syndrome, Lissencephaly, |
| | genome. Defects in some of the genes localized to chromosome 7 have been linked to |
| | chromosome 7, which houses over 1,000 genes and comprises nearly 5 % of the human |
| | the concentration of 2,3-BPG within the cell. The gene encoding BPGM maps to human |
| | glyceroyl phosphate to 2,3-bisD-glycerate (2,3-BPG), a reaction that is essential for controlling |
| | the regulation of hemoglobin oxygen. Specifically, BPGM catalyzes the conversion of 3-D- |
| | to the phosphoglycerate mutase family and exists as a homodimer that plays a crucial role in |
| | Background: BPGM (2,3-bisphosphoglycerate mutase) is a 259 amino acid protein that belongs |
| | PMGE_HUMAN, 2,3-bisphosphoglycerate mutase, erythrocyte, erythrocyte, 2 antibody. |
| | dependent PGAM , Bpgm, BPGM, C86192, DPGM, Erythrocyte 2,3 bisphosphoglycerate mutase, |

Application Details

| Application Notes: | IF(IHC-P) 1:50-200 |
|--------------------|--------------------------------------------------------------------------------------------------------------------|
| Restrictions: | For Research Use only |
| Handling | |
| Format: | Liquid |
| Concentration: | 1 μg/μL |
| Buffer: | Aqueous buffered solution containing 0.01M TBS (pH 7.4) with 1 % BSA, 0.03 % Proclin300 and 50 % Glycerol. |
| Preservative: | ProClin |
| Precaution of Use: | This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE, which should be handled by trained staff only. |
| Storage: | -20 °C |
| Storage Comment: | Store at -20°C. Aliquot into multiple vials to avoid repeated freeze-thaw cycles. |
| Expiry Date: | 12 months |

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