antibodies -online.com





anti-FIBIN antibody (AA 101-200) (Cy5.5)



Go to Product page

| \sim | | | |
|--------|-----|-----|-----|
| | N/P | r\/ | i⊢₩ |

| Quantity: | 100 μL | |
|----------------------|--|--|
| Target: | FIBIN | |
| Binding Specificity: | AA 101-200 | |
| Reactivity: | Mouse | |
| Host: | Rabbit | |
| Clonality: | Polyclonal | |
| Conjugate: | This FIBIN antibody is conjugated to Cy5.5 | |
| Application: | Western Blotting (WB), Immunofluorescence (Paraffin-embedded Sections) (IF (p)), Immunofluorescence (Cultured Cells) (IF (cc)) | |

Product Details

| Immunogen: | KLH conjugated synthetic peptide derived from human FIBIN |
|-----------------------|---|
| Isotype: | IgG |
| Cross-Reactivity: | Mouse |
| Predicted Reactivity: | Human,Rat,Cow,Sheep,Pig,Horse,Rabbit |
| Purification: | Purified by Protein A. |

Target Details

| Target: | FIBIN |
|-------------------|------------------------|
| Alternative Name: | FIBIN (FIBIN Products) |

Target Details

Expiry Date:

| Background: | Synonyms: Fin bud initiation factor homolog, FIBIN, PSEC0235 | |
|---------------------|---|--|
| | Background: FIBIN (Fin bud initiation factor homolog) is a 211 amino acid protein involved in fin | |
| | initiation in zebrafish. The human homolog is encoded by a gene that maps to chromosome 11, | |
| | which makes up around 4 % of human genomic DNA and is considered a gene and disease | |
| | association dense chromosome. The chromosome 11 encoded Atm gene is important for | |
| | regulation of cell cycle arrest and apoptosis following double strand DNA breaks. Atm mutation | |
| | leads to the disorder known as ataxia-telangiectasia. The blood disorders Sickle cell anemia | |
| | and $_$ thalassemia are caused by HBB gene mutations. Wilms' tumors, WAGR syndrome and | |
| | Denys-Drash syndrome are associated with mutations of the WT1 gene. Jervell and Lange- | |
| | Nielsen syndrome, Jacobsen syndrome, Niemann-Pick disease, hereditary angioedema and | |
| | Smith-Lemli-Opitz syndrome are also associated with defects in chromosome 11. | |
| Gene ID: | 387758 | |
| UniProt: | Q8TAL6 | |
| Application Details | | |
| Application Notes: | IF(IHC-P) 1:50-200 | |
| | IF(IHC-F) 1:50-200 | |
| | IF(ICC) 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: | handled by trained staff only. -20 °C | |

12 months