antibodies .- online.com







anti-FOXRED1 antibody (AA 251-350) (Cy7)



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| Quantity: | 100 μL |
|----------------------|--|
| Target: | FOXRED1 |
| Binding Specificity: | AA 251-350 |
| Reactivity: | Mouse, Rat |
| Host: | Rabbit |
| Clonality: | Polyclonal |
| Conjugate: | This FOXRED1 antibody is conjugated to Cy7 |
| Application: | Western Blotting (WB), Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p)) |

Product Details

| Immunogen: | KLH conjugated synthetic peptide derived from human FOXRED1 |
|-----------------------|---|
| Isotype: | IgG |
| Cross-Reactivity: | Mouse, Rat |
| Predicted Reactivity: | Human,Dog,Pig,Horse,Rabbit |
| Purification: | Purified by Protein A. |

Target Details

| Target: | FOXRED1 |
|-------------------|----------------------------|
| Alternative Name: | FOXRED1 (FOXRED1 Products) |

Target Details

| Background: | Synonyms: FAD dependent oxidoreductase domain containing 1, FAD dependent | |
|---------------------|--|--|
| | oxidoreductase domain containing protein 1, FAD-dependent oxidoreductase domain- | |
| | containing protein 1, FOXRED 1, FOXRED1, FP634, FXRD1_HUMAN, H17. | |
| | Background: FOXRED1 is a 486 amino acid single-pass membrane protein. Utilizing FAD as a | |
| | cofactor, FOXRED1 may act as a chaperone protein essential for the function of mitochondrial | |
| | complex I. Mutations to FOXRED1 may result in mitochondrial complex I deficiency (MT-C1D), | |
| | which results in a wide range of clinical maladies from lethal neonatal disease to adult onset | |
| | neurodegenerative disorders. Common phenotypes of MT-C1D include cardiomyopathy, liver | |
| | disease, Leigh syndrome, Leber hereditary optic neuropathy, and some forms of Parkinson | |
| | disease. FOXRED1 exists as three alternatively spliced isoforms and is encoded by a gene | |
| | mapping to human chromosome 11q24.2. With approximately 135 million base pairs and 1,400 | |
| | genes, chromosome 11 makes up around 4 % of human genomic DNA and is considered a | |
| | gene and disease association dense chromosome. | |
| Gene ID: | 55572 | |
| | | |
| 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: | -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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