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anti-NAT8B antibody (AA 221-227) (Alexa Fluor 555)



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| Quantity: | 100 μL |
|----------------------|--------------------------------------------------------------------------------------------------------------------------------|
| Target: | NAT8B |
| Binding Specificity: | AA 221-227 |
| Reactivity: | Human |
| Host: | Rabbit |
| Clonality: | Polyclonal |
| Conjugate: | This NAT8B antibody is conjugated to Alexa Fluor 555 |
| 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 NAT8B |
|-----------------------|-----------------------------------------------------------|
| Isotype: | IgG |
| Predicted Reactivity: | Human,Mouse,Rat,Sheep,Pig,Horse |
| Purification: | Purified by Protein A. |

Target Details

| Target: | NAT8B | |
|-----------------------------------------------------------------------------------------------------|------------------------|--|
| Alternative Name: | NAT8B (NAT8B Products) | |
| Background: Synonyms: Camello like protein 2, Camello-like protein 2, CML2, Hcml2, N acetyltransfer | | |

NAT8B, NAT8B_HUMAN, NAT8BP, Probable N acetyltransferase 8B, Probable N-acetyltransferase 8B.

Background: Acetyltransferases and deacetylases are protein groups most often associated with oncogenesis and cell cycle regulation. NAT-8B (N-acetyltransferase 8B), also known as CML2 (camello-like protein 2), is a 227 amino acid single-pass membrane protein that is implicated in gastrulation regulation. A member of the camello family, NAT-8B contains one N-acetyltransferase domain and is encoded by a gene that maps to human chromosome 2p13.2. The NAT-8B gene is susceptible to a nonsense mutation at Serine 16, which leads to a stop codon and subsequently, a non-functional protein that is truncated in length. Similarly, a nonsense mutation at Glutamine 168 is thought to lead to a non-functional protein, as it causes the N-acetyltransferase to become disrupted. Human chromosome 2 consists of 237 million bases, encodes over 1,400 genes and makes up approximately 8 % of the human genome. A number of genetic diseases are linked to genes on chromosome 2 including Harlequin icthyosis, sitosterolemia and Alstr syndrome.

Application Details

IF(IHC-P) 1:50-200

Application Notes:

| Application Notes. | Ir(InC-r) 1.30-200 | |
|--------------------|--------------------------------------------------------------------------------------------------------------------|--|
| | 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 | |