

# NIR1 Polyclonal Antibody

Purified Rabbit Polyclonal Antibody (Pab)

Catalog # AP59023

## Product Information

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| <b>Application</b>             | WB, IF, E  |
| <b>Primary Accession</b>       | <a href="#">Q9BZ71</a>   |
| <b>Reactivity</b>              | Rat, Pig, Bovine   |
| <b>Host</b>                    | Rabbit   |
| <b>Clonality</b>               | Polyclonal   |
| <b>Calculated MW</b>           | 106781   |
| <b>Physical State</b>          | Liquid   |
| <b>Immunogen</b>               | KLH conjugated synthetic peptide derived from human NIR1/RDGBA3  |
| <b>Epitope Specificity</b>     | 131-250/974  |
| <b>Isotype</b>                 | IgG  |
| <b>Purity</b>                  | affinity purified by Protein A   |
| <b>Buffer</b>                  | 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.  |
| <b>SUBCELLULAR LOCATION</b>    | Endomembrane system; Peripheral membrane protein   |
| <b>SIMILARITY</b>              | Belongs to the PtdIns transfer protein family. PI transfer class IIA subfamily. Contains 1 DDHD domain.  |
| <b>SUBUNIT</b>                 | Interacts with PTK2B via its C-terminus.   |
| <b>DISEASE</b>                 | Defects in PITPNM3 are the cause of cone-rod dystrophy type 5 (CORD5) [MIM:600977]. CORDs are inherited retinal dystrophies belonging to the group of pigmentary retinopathies. CORDs are characterized by retinal pigment deposits visible on fundus examination, predominantly in the macular region, and initial loss of cone photoreceptors followed by rod degeneration. This leads to decreased visual acuity and sensitivity in the central visual field, followed by loss of peripheral vision. Severe loss of vision occurs earlier than in retinitis pigmentosa. |
| <b>Important Note</b>          | This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.  |
| <b>Background Descriptions</b> | Catalyzes the transfer of phosphatidylinositol and phosphatidylcholine between membranes (in vitro) (By similarity). Binds calcium ions. Involvement in disease:Defects in PITPNM3 are the cause of cone-rod dystrophy type 5 (CORD5) . CORDs are inherited retinal dystrophies belonging to the group of pigmentary retinopathies. CORDs are characterized by retinal pigment deposits visible on fundus examination, predominantly in the macular region, and initial loss of cone photoreceptors followed by rod degeneration.  |

## Additional Information

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| <b>Gene ID</b>     | 83394  |
| <b>Other Names</b> | Membrane-associated phosphatidylinositol transfer protein 3, Phosphatidylinositol transfer protein, membrane-associated 3, PITPnm 3, Pyk2 N-terminal domain-interacting receptor 1, NIR-1, PITPNM3, NIR1 |

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| <b>Target/Specificity</b> | Detected in brain and spleen, and at low levels in ovary.   |
| <b>Dilution</b>           | WB=1:500-2000,IF=1:50-200,ELISA=1:5000-10000  |
| <b>Format</b>             | 0.01M TBS(pH7.4) with 1% BSA, 0.09% (W/V) sodium azide and 50% Glyce  |
| <b>Storage</b>            | Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C. |

## Protein Information

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| <b>Name</b>              | PITPNM3  |
| <b>Synonyms</b>          | NIR1   |
| <b>Function</b>          | Catalyzes the transfer of phosphatidylinositol and phosphatidylcholine between membranes (in vitro) (By similarity). Binds calcium ions. |
| <b>Cellular Location</b> | Endomembrane system; Peripheral membrane protein   |
| <b>Tissue Location</b>   | Detected in brain and spleen, and at low levels in ovary.  |

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.