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## anti-AGPS antibody (AA 31-130) (Cy7)



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| Quantity:            | 100 μL  |  |
|----------------------|---|--|
| Target:              | AGPS  |  |
| Binding Specificity: | AA 31-130   |  |
| Reactivity:          | Rat   |  |
| Host:                | Rabbit  |  |
| Clonality:           | Polyclonal  |  |
| Conjugate:           | This AGPS antibody is conjugated to Cy7   |  |
| Application:         | Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p)) |  |

#### **Product Details**

| Immunogen:            | KLH conjugated synthetic peptide derived from human AGPS/Alkyl-DHAP synthase |  |
|-----------------------|--|--|
| Isotype:              | IgG  |  |
| Cross-Reactivity:     | Rat  |  |
| Predicted Reactivity: | Human,Mouse,Cow,Pig,Horse  |  |
| Purification:         | Purified by Protein A.   |  |

#### Target Details

| Target:           | AGPS                                     |
|-------------------|--|
| Alternative Name: | AGPS/Alkyl-DHAP synthase (AGPS Products) |

#### Target Details

| Background:         | Synonyms: AAG5, ADAP-S, ADAS, ADAS_HUMAN, ADHAPS, ADPS, Aging associated gene 5               |
|---------------------|---|
|                     | protein, Aging-associated gene 5 protein, AGPS, ALDHPSY, Alkyl-DHAP synthase,                 |
|                     | Alkyldihydroxyacetonephosphate synthase, Alkyldihydroxyacetonephosphate synthase,             |
|                     | peroxisomal, Alkylglycerone phosphate synthase, Alkylglycerone-phosphate synthase,            |
|                     | peroxisomal.  |
|                     | Background: AGPS is a 658 amino acid enzyme that is required for glycerolipid metabolism and  |
|                     | ether lipid biosynthesis. Localized to the inner aspect of the peroxisomal membrane, AGPS is  |
|                     | likely part of a heterotrimeric complex that is also composed of GNPAT and a modified form of |
|                     | GNPAT. Containing one FAD-binding PCMH-type domain, AGPS utilizes FAD as a cofactor in the    |
|                     | synthesis of alkyl-glycerone 3-phophate and a long-chain acid anion from 1-acteyl-glyerone 3- |
|                     | phosphate and a long-chain alcohol. Defects in the gene encoding AGPS results in rhizomelic   |
|                     | chondrodysplasia punctata type 3, a disease characterized by vertebral disorders, severe      |
|                     | mental retardation, cutaneous lesions, cataracts and rhizomelic shortening of the humerus and |
|                     | femur.  |
| Gene ID:            | 8540  |
| Pathways:           | SARS-CoV-2 Protein Interactome  |
| 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  |
|                     |   |

### Handling

| Storage Comment: | Store at -20°C. Aliquot into multiple vials to avoid repeated freeze-thaw cycles. |
|------------------|---|
| Expiry Date:     | 12 months   |