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Recombinant PSPH Monoclonal Antibody

catalog number: AN300269P

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description

Reactivity Human

Immunogen Recombinant Human PSPH protein

Host Rabbit Isotype lgG B187 Clone **Purification** Protein A

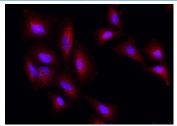
Buffer 0.2 µm filtered solution in PBS

Applications Recommended Dilution

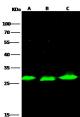
1:500-1:1000 **WB** 1:20-1:100 ICC/IF

ΙP 0.2-1 µL/mg of lysate

Data

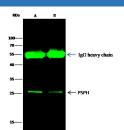


Immunofluorescence analysis of Human PSPH in A549 cells. Cells were fixed with 4% PFA, permeabilzed with 0.3% Monoclonal Antibody and 15 µl of 50 % Protein G agarose. Triton X-100 in PBS, blocked with 10% serum, and incubated with rabbit anti-Human PSPH Monoclonal Antibody (dilution ratio 1:60) at 4°C overnight. Then cells were stained with the Alexa Fluor® 594-conjugated Goat Anti-rabbit IgG secondary antibody (red) and counterstained with DAPI for nuclear staining (blue).



Western Blot with PSPH Monoclonal Antibody at dilution of 1:500. Lane A: MCF7 Whole Cell Lysate, Lane B: K562 Whole Cell Lysate, Lane C: HepG2 Whole Cell Lysate, Lysates/proteins at 30 µg per lane.

> Observed-MW:25 kDa Calculated-MW:25 kDa



Immunoprecipitation analysis using 0.5 µL anti-PSPH Western blot was performed from the immunoprecipitate using PSPH Monoclonal Antibody at a dilution of 1:1000. Lane A:0.5 mg K562 Whole Cell Lysate, Lane B:0.5 mg

> Observed-MW:25 kDa Calculated-MW:25 kDa

HepG2 Whole Cell Lysate

Preparation & Storage

For Research Use Only

Toll-free: 1-888-852-8623 Fax: 1-832-243-6017 Tel: 1-832-243-6086 Web: www.elabscience.com Email: techsupport@elabscience.com



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Storage This antibody can be stored at 2°C-8°C for one month without detectable loss of

activity. Antibody products are stable for twelve months from date of receipt when stored at -20°C to -80°C. Preservative-Free. Avoid repeated freeze-thaw cycles.

Shipping lce bag

Background

The protein encoded by this gene belongs to a subfamily of the phosphotransferases. This encoded enzyme is responsible for the third and last step in L-serine formation. It catalyzes magnesium-dependent hydrolysis of L-phosphoserine and is also involved in an exchange reaction between L-serine and L-phosphoserine. Deficiency of this protein is thought to be linked to Williams syndrome.

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