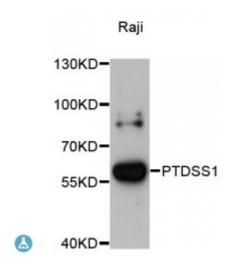


Anti-PTDSS1 Antibody



Description The protein encoded by this gene catalyzes the formation of

phosphatidylserine from either phosphatidylcholine or

phosphatidylethanolamine. Phosphatidylserine localizes to the

mitochondria-associated membrane of the endoplasmic reticulum, where it serves a structural role as well as a signaling role. Defects in this gene are a cause of Lenz-Majewski hyperostotic dwarfism. Two transcript variants

encoding different isoforms have been found for this gene.

Model STJ115032

Host Rabbit

Reactivity Human

Applications WB

Immunogen Recombinant fusion protein containing a sequence corresponding to amino

acids 1-70 of human PTDSS1 (NP_055569.1).

Gene ID 9791

Gene Symbol PTDSS1

Dilution range WB 1:500 - 1:2000

Purification Affinity purification

Note For Research Use Only (RUO).

Protein Name Phosphatidylserine synthase 1 PSS-1 PtdSer synthase 1

Molecular Weight 55.528 kDa

Clonality Polyclonal

Conjugation Unconjugated

Isotype IgG

Formulation PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

Storage Instruction Store at -20C. Avoid freeze / thaw cycles.

Database Links HGNC:95870MIM:151050Reactome:R-HSA-1483101

Alternative Names Phosphatidylserine synthase 1 PSS-1 PtdSer synthase 1

Function Catalyzes a base-exchange reaction in which the polar head group of

phosphatidylethanolamine (PE) or phosphatidylcholine (PC) is replaced by L-

serine, In membranes, PTDSS1 catalyzes mainly the conversion of phosphatidylcholine, Also converts, in vitro and to a lesser extent,

phosphatidylethanolamine

Cellular Localization Endoplasmic reticulum membrane,

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