

Recombinant Human PRPH2 293 Cell Lysate

Cat. No. PRPH2-2821HCL **Lot. No.** (See product label)

SPECIFICATION

Product Overview

Species	Human
Source	HEK293
Description	Antigen standard for peripherin 2 (retinal degeneration, slow) (PRPH2) is a lysate prepared from HEK293T cells transiently transfected with a TrueORF gene-carrying pCMV plasmid and then lysed in RIPA Buffer. Protein concentration was determined using a colorimetric assay. The antigen control carries a C-terminal Myc/DDK tag for detection.
Components	This product includes 3 vials: 1 vial of gene-specific cell lysate, 1 vial of control vector cell lysate, and 1 vial of loading buffer. Each lysate vial contains 0.1 mg lysate in 0.1 ml (1 mg/ml) of RIPA Buffer (50 mM Tris-HCl pH7.5, 250 mM NaCl, 5 mM EDTA, 50 mM NaF, 1% NP40). The loading buffer vial contains 0.5 ml 2X SDS Loading Buffer (125 mM Tris-Cl, pH6.8, 10% glycerol, 4% SDS, 0.002% Bromophenol blue, 5% beta-mercaptoethanol).
Size	0.1 mg
Storage Instruction	Store at -80°C. Minimize freeze-thaw cycles. After addition of 2X SDS Loading Buffer, the lysates can be stored at -20°C. Product is guaranteed 6 months from the date of shipment.

 Tel: 1-631-559-9269 1-516-512-3133

 Email: info@creative-biomart.com  Fax: 1-631-938-8127

 45-1 Ramsey Road, Shirley, NY 11967, USA

Applications

ELISA, WB, IP. WB: Mix equal volume of lysates with 2X SDS Loading Buffer. Boil the mixture for 10 min before loading (for membrane protein lysates, incubate the mixture at room temperature for 30 min). Load 5 ug lysate per lane.

GENE INFORMATION
Gene Name

PRPH2 peripherin 2 (retinal degeneration, slow) [Homo sapiens]

Official Symbol

PRPH2

Synonyms

PRPH2; peripherin 2 (retinal degeneration, slow); RDS, retinal degeneration, slow , retinal degeneration, slow (retinitis pigmentosa 7) , RP7; peripherin-2; CACD2; rd2; retinal peripherin; TSPAN22; tspan-22; tetraspanin-22; peripherin 2, homolog of mouse; peripherin, photoreceptor type; retinal degeneration slow protein; DS; RDS; RP7; AVMD; PRPH; AOFMD;

Gene ID

5961

mRNA Refseq

NM_000322

Protein Refseq

NP_000313

MIM

179605

UniProt ID

P23942

**Chromosome
Location**

6p21.2-cen

Pathway

Amyotrophic lateral sclerosis (ALS), organism-specific biosystem; Amyotrophic lateral sclerosis (ALS), conserved biosystem;

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