

Product datasheet

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ARG65269 anti-PEX26 antibody

Package: 100 μg Store at: -20°C

Summary

Product Description Goat Polyclonal antibody recognizes PEX26

Tested Reactivity Hu
Tested Application WB

Specificity Reported variants represent identical protein: NP_060399.1, NP_001121121.1

Host Goat

Clonality Polyclonal

Isotype IgG

Target Name PEX26
Species Human

 Immunogen
 C-QKPNLEGSVSHK

 Conjugation
 Un-conjugated

Alternate Names PBD7B; PBD7A; PEX26M1T; Peroxisome assembly protein 26; Peroxin-26; Pex26pM1T

Application Instructions

Application table	Application	Dilution
	WB	0.5 - 1.5 μg/ml
Application Note	WB: Recommend incubate at RT for 1h. * The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	

Properties

Form Liquid

Purification Purified from goat serum by antigen affinity chromatography.

Buffer Tris saline (pH 7.3), 0.02% Sodium azide and 0.5% BSA.

Preservative 0.02% Sodium azide

Stabilizer 0.5% BSA

Concentration 0.5 mg/ml

Storage instruction For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot

and store at -20°C or below. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed

before use.

Note For laboratory research only, not for drug, diagnostic or other use.

Bioinformation

Database links <u>GeneID: 55670 Human</u>

Swiss-port # Q7Z412 Human

Background This gene belongs to the peroxin-26 gene family. It is probably required for protein import into

peroxisomes. It anchors PEX1 and PEX6 to peroxisome membranes, possibly to form heteromeric AAA ATPase complexes required for the import of proteins into peroxisomes. Defects in this gene are the cause of peroxisome biogenesis disorder complementation group 8 (PBD-CG8). PBD refers to a group of peroxisomal disorders arising from a failure of protein import into the peroxisomal membrane or matrix. The PBD group is comprised of four disorders: Zellweger syndrome (ZWS), neonatal adrenoleukodystrophy (NALD), infantile Refsum disease (IRD), and classical rhizomelic chondrodysplasia punctata (RCDP). Alternatively spliced transcript variants have been identified for this gene. [provided

by RefSeq, Dec 2010]

15kDa

Research Area Controls and Markers antibody; Signaling Transduction antibody

Calculated Mw 34 kDa

Images

250kDa
150kDa
100kDa
75kDa
Western Blot: Human Kidney lysate (35 μg protein in RIPA buffer)
stained with ARG65269 anti-PEX26 antibody at 0.5 μg/ml dilution.

50kDa
37kDa
25kDa
20kDa