

## 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	
lsotype	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	GeneID: 55670 Human
	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]
Research Area	Controls and Markers antibody; Signaling Transduction antibody
Calculated Mw	34 kDa

## Images

250kDa 150kDa	ARG65269 anti-PEX26 antibody WB image
100kDa 75kDa	Western Blot: Human Kidney lysate (35 $\mu g$ protein in RIPA buffer) stained with ARG65269 anti-PEX26 antibody at 0.5 $\mu g/ml$ dilution.
50kDa	
37kDa	
25kDa	
20kDa	
15kDa	