

Peroxin 26 antibody

Catalog No: #22179

Package Size: #22179 100ul

Description

Product Name	Peroxin 26 antibody
Host Species	Rabbit
Clonality	Polyclonal
Purification	Purified by antigen-affinity chromatography.
Applications	WB IHC IF
Species Reactivity	Hu
Immunogen Type	Recombinant protein
Immunogen Description	Recombinant protein fragment contain a sequence corresponding to a region within amino acids 1 and 265 of Human PEX26
Conjugates	Unconjugated
Target Name	Peroxin 26
Accession No.	Swiss-Prot:Q7Z412Gene ID:55670
Concentration	1mg/ml
Formulation	Supplied in 0.1M Tris-buffered saline with 10% Glycerol (pH7.0). 0.01% Thimerosal was added as a preservative.
Storage	Store at -20°C for long term preservation (recommended). Store at 4°C for short term use.

Application Details

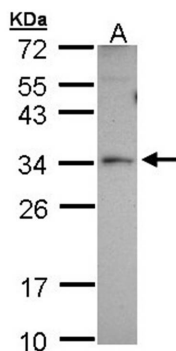
Predicted MW: 34kd

Western blotting: 1:500-1:3000

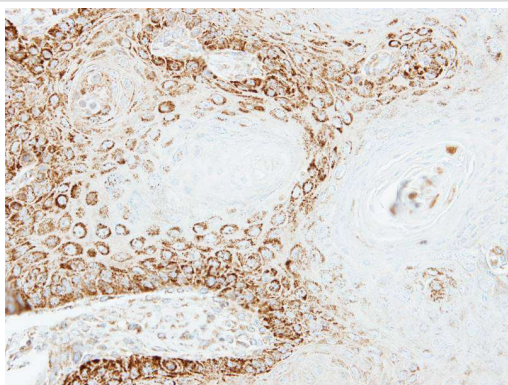
Immunohistochemistry: 1:50-1:500

Immunofluorescence: 1:100-1:200

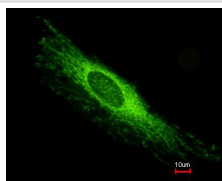
Images



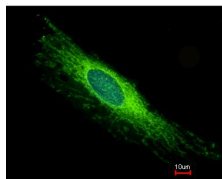
Sample (30 ug of whole cell lysate)
A: A431
12% SDS PAGE
Primary antibody diluted at 1: 1000



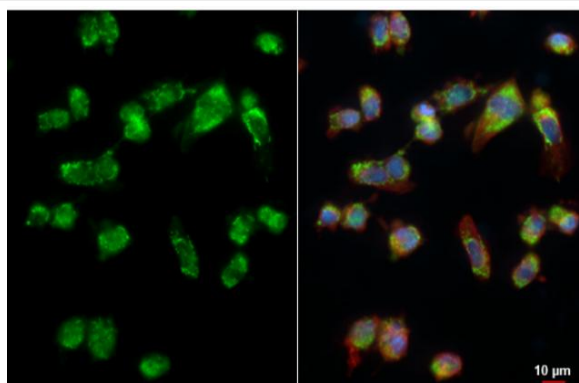
Immunohistochemical analysis of paraffin-embedded Cal27 xenograft, using PEX26 antibody at 1: 500 dilution.



Costained with Hoechst 33342



Immunofluorescence analysis of methanol-fixed HeLa, using PEX26 antibody at 1: 500 dilution.



Sample: ice-cold MeOH fixed HT-1080 cells Green: 1:500
Red: alpha Tubulin 1:1000 Blue: DAPI

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). Two transcript variants encoding the same protein have been identified for this gene. [provided by RefSeq]

Note: This product is for in vitro research use only and is not intended for use in humans or animals.