

Product datasheet

Recombinant Human NPHS2 protein ab188448

1 Image

Description

Product name	Recombinant Human NPHS2 protein	
Purity	> 95 % Densitometry.	
Endotoxin level	< 1.000 Eu/μg	
Expression system	Escherichia coli	
Accession	Q9NP85	
Protein length	Protein fragment	
Animal free	No	
Nature	Recombinant	
Species	Human	
Sequence	MKHHHHHHASVKVVQEYERVIIIFRLGHLLPGRAKGPGL FFFLPCLDTHK VDLRLQTLQLEIPFHEIVTKDMFIMEIDAICYRMENASLLLS SLAHVSKAV QFLVQTTMKRLLAHRSLTEILLERKSIAQDAKVALDSVT CIWGKVERIE IKDVRLPAGLQHS�AVEAEAQRQAKVRMIAAEAEKAA SESLRMAAEILSG TPAAVQLRYLHTLQSLSTKPSVTVLPLPFDLLNCLSS PSNRTQGSLPFP SPSKPVEPLNPKKKDSPML	
Predicted molecular weight	30 kDa including tags	
Amino acids	125 to 383	
Tags	His tag N-Terminus	

Specifications

Our [Abpromise guarantee](#) covers the use of **ab188448** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications	SDS-PAGE
	Mass Spectrometry
	Western blot

	ELISA
Mass spectrometry	LC-MS/MS
Form	Lyophilised

Preparation and Storage

Stability and Storage Shipped at 4°C. Upon delivery aliquot. Store at -80°C. Avoid freeze / thaw cycle.
pH: 4.00
Constituents: 0.02% DTT, 0.25% Sodium acetate

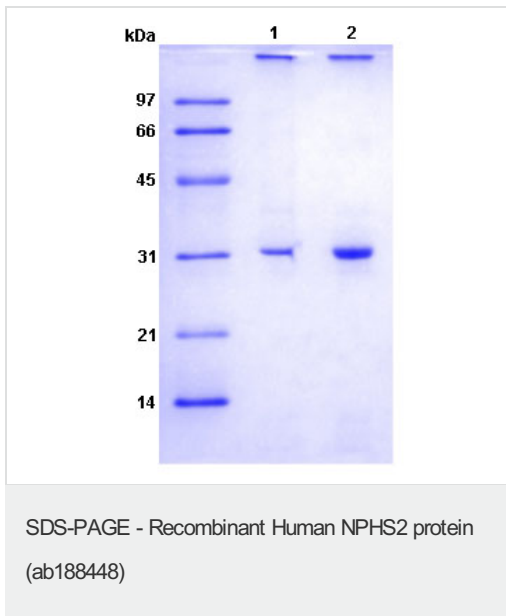
Lyophilized in 30 mM acetate buffer.

Reconstitution Add 0.1 M Acetate buffer pH4 to prepare a working stock solution of approximately 0.5 mg/mL and let the lyophilized pellet dissolve completely. For conversion into higher pH value, we recommend intensive dilution by relevant buffer to a concentration of 10 µg/mL. In higher concentrations the solubility of this antigen is limited.

General Info

Function	Plays a role in the regulation of glomerular permeability, acting probably as a linker between the plasma membrane and the cytoskeleton.
Tissue specificity	Almost exclusively expressed in the podocytes of fetal and mature kidney glomeruli.
Involvement in disease	Defects in NPHS2 are the cause of nephrotic syndrome type 2 (NPHS2) [MIM:600995]. It is a renal disorder characterized clinically by childhood onset of proteinuria, hypoalbuminemia, hyperlipidemia, and edema. Kidney biopsies show non-specific histologic changes such as focal segmental glomerulosclerosis and diffuse mesangial proliferation. The disorder is resistant to steroid treatment and progresses to end-stage renal failure in the first or second decades. Some patients show later onset of the disorder.
Sequence similarities	Belongs to the band 7/mec-2 family.
Cellular localization	Cell membrane.

Images



14% SDS-PAGE analysis of ab188448.

Lane 1: Reduced and boiled sample, 2.5 µg

Lane 2: Non-reduced and non-boiled sample, 2.5 µg

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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