



# Product Information Sheet

# **Polyclonal Anti- Podocin**

Catalogue No. PA1322

Lot No. 09L01

Ig type rabbit IgG

Size 100µg/vial

## Specificity

Rat (Kidney), mouse. No cross reactivity with other proteins.

Recommended application
Western blot



#### Lane 1-4 : Rat Kidney tissue Lysate

### Immunogen

A synthetic peptide corresponding to a sequence at the middle region of human Podocin, different from the mouse sequence by one amino acid.

## Purity

Immunogen affinity purified.

## Application

	Concen- tration	Tested Species	Concluded Species	Antigen Retrieval
WB	1µg/ml	Rat	Ms	-
IHC-P	-	-	-	-
IHC-F	-	-	-	-
ICC	-	-	-	-

Other applications have not been tested.

Optimal dilutions should be determined by end user.

## Contents

Each vial contains 5mg BSA, 0.9mg NaCl, 0.2mg Na $_2$ HPO $_4$ , 0.05mg Thimerosal, 0.05mg NaN $_3$ .

## Reconstitution

0.2ml of distilled water will yield a concentration of 500µg/ml.

# To reorder contact us at:

Antagene, Inc. Storage

Toll Free: 1(866)964-2589 email: Info@antageneinc.com

At -20°C for one year. After reconstitution, at 4°C for one month. It can also be aliquotted and stored frozen at -20°C for longer time.

FOR RESEARCH USE ONLY. NOT FOR DIAGNOSTIC AND CLINICAL USE.

#### BACKGROUND

Podocin (PDCN) is a protein which lines the podocytes and assists in maintaining the barrier at the glomerular basement membrane. NPHS2 is a causative gene for Familial idiopathic nephrotic syndromes, which represents a heterogeneous group of kidney disorders, and include autosomal recessive steroid-resistant nephrotic syndrome, which is characterized by early childhood onset of proteinuria, rapid progression to end-stage renal disease and focal segmental glomerulosclerosis. By positional cloning, NPHS2 was mapped to 1q25-31. It is almost exclusively expressed in the podocytes of fetal and mature kidney glomeruli, and encodes a new integral membrane protein, podocin, belonging to the stomatin protein family. Boute et al. (2000) found ten different NPHS2 mutations, comprising nonsense, frameshift and missense mutations, to segregate with the disease, demonstrating a crucial role for podocin in the function of the glomerular filtration barrier.<sup>1</sup>

## REFERENCE

1. Boute, N.; Gribouval, O.; Roselli, S.; Benessy, F.; Lee, H.; Fuchshuber, A.; Dahan, K.; Gubler, M.-C.; Niaudet, P.; Antignac, C. : NPHS2, encoding the glomerular protein podocin, is mutated in autosomal recessive steroid-resistant nephrotic syndrome. Nature Genet. 24: 349-354, 2000. Note: Erratum: Nature Genet. 25: 125 only, 2000.