

## NPHS1 Polyclonal Antibody

**Catalog No.** E-AB-70211

**Note:** Centrifuge before opening to ensure complete recovery of vial contents.

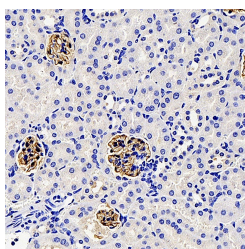
### Description

|                     |                                                                            |
|---------------------|----------------------------------------------------------------------------|
| <b>Reactivity</b>   | Mouse                                                                      |
| <b>Immunogen</b>    | KLH conjugated Synthetic peptide corresponding to Mouse Nephtrin           |
| <b>Host</b>         | Rabbit                                                                     |
| <b>Isotype</b>      | IgG                                                                        |
| <b>Purification</b> | Affinity purification                                                      |
| <b>Conjugation</b>  | Unconjugated                                                               |
| <b>Buffer</b>       | PBS with 0.02% sodium azide, 1% protective protein and 50% glycerol, pH7.4 |

### Applications Recommended Dilution

|            |               |
|------------|---------------|
| <b>IHC</b> | 1:1000-1:2000 |
|------------|---------------|

### Data



Immunohistochemistry analysis of paraffin-embedded mouse kidney using NPHS1 Polyclonal Antibody at dilution of 1:1000.

### Preparation & Storage

|                |                                             |
|----------------|---------------------------------------------|
| <b>Storage</b> | Store at -20°C. Avoid freeze / thaw cycles. |
|----------------|---------------------------------------------|

### Background

This gene encodes a member of the immunoglobulin family of cell adhesion molecules that functions in the glomerular filtration barrier in the kidney. The gene is primarily expressed in renal tissues, and the protein is a type-1 transmembrane protein found at the slit diaphragm of glomerular podocytes. The slit diaphragm is thought to function as an ultrafilter to exclude albumin and other plasma macromolecules in the formation of urine. Mutations in this gene result in Finnish-type congenital nephrosis 1, characterized by severe proteinuria and loss of the slit diaphragm and foot processes. In Western blots, nephrin antibodies generated against the two terminal extracellular Ig domains of recombinant human nephrin recognized a 180-kDa protein in lysates of human glomeruli and a 150-kDa protein in transfected COS-7 cell lysates.

### For Research Use Only