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**Research Use Only. Not for
diagnostic or therapeutic use.**

EB06971 - Goat Anti-NHERF2 (isoform a) Antibody

Size: 100µg specific antibody in 200µl



Target Protein

Principal Names: SLC9A3R2, solute carrier family 9 (sodium/hydrogen exchanger), isoform 3 regulator 2, HGNC:11076, E3KARP, NHE3RF2, NHERF-2, NHERF2, OCTS2, SIP-1, SIP1, TKA-1, sodium/hydrogen exchanger, solute carrier family 9 (sodium/hydrogen exchanger), isoform 3 regulatory factor 2, solute carrier family 9 isoform 3 regulator 2, solute carrier family 9 (sodium/hydrogen exchanger), member 3 regulator 2, MGC104639, solute carrier family 9 (sodium/hydrogen exchanger), isoform 3 regulatory factor 2

Official Symbol: SLC9A3R2

Accession Number(s): NP_001123484.1

Human GeneID(s): [9351](#)

Important Comments: Please note that this variant has an extra insertion of 11 residues compared to NP_004776.3. The immunizing peptide represents this insertion.

Immunogen

Peptide with sequence C-DGSAWKQDPFQ, from the internal region of the protein sequence according to NP_001123484.1.

Please note the [peptide](#) is available for sale.

Purification and Storage

Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity chromatography using the immunizing peptide.

Supplied at 0.5 mg/ml in Tris saline, 0.02% sodium azide, pH7.3 with 0.5% bovine serum albumin.

Aliquot and store at -20°C. Minimize freezing and thawing.

Applications Tested

Peptide ELISA: antibody detection limit dilution 1:32000.

Western blot: Approx 38kDa band observed in human kidney lysates (calculated MW of 37.4kDa according to BAA33216.1). Recommended concentration: 0.3-1µg/ml.

IHC: In paraffin embedded Human Kidney shows strong membranous staining in glomerulus-. Recommended concentration, 1-2µg/ml.

Species Reactivity

Tested: Human

Expected from sequence similarity: Human

EB06971 (3µg/ml) staining of paraffin embedded Human Kidney. Microwaved antigen retrieval with Tris/EDTA buffer pH9, HRP-staining.

EB06971 (0.3µg/ml) staining of human kidney lysate (35µg protein in RIPA buffer). Primary incubation was 1 hour. Detected by chemiluminescence.