

DPF2 antibody

Cat. No. GTX31819

Host	Rabbit
Clonality	Polyclonal
Isotype	IgG
Applications	WB, IHC-P, ELISA
Reactivity	Human, Mouse

Package
100 µg

Applications

Application Note

*Optimal dilutions/concentrations should be determined by the researcher.

Suggested dilution	Recommended dilution
WB	1 µg/mL
IHC-P	5 µg/mL
ELISA	Assay dependent

Not tested in other applications.

Calculated MW 44 kDa. ([Note](#))

Product Note This antibody is predicted to not cross-react with other DPF protein family members.

Properties

Form	Liquid
Buffer	PBS
Preservative	0.02% Sodium azide
Storage	Store as concentrated solution. Centrifuge briefly prior to opening vial. For short-term storage (1-2 weeks), store at 4°C. For long-term storage, aliquot and store at -20°C or below. Avoid multiple freeze-thaw cycles.
Concentration	1 mg/ml (Please refer to the vial label for the specific concentration.)
Immunogen	DPF2 antibody was raised against an 18 amino acid synthetic peptide near the carboxy terminus of human DPF2. The immunogen is located within the last 50 amino acids of DPF2.
Purification	Purified by antigen-affinity chromatography
Conjugation	Unconjugated



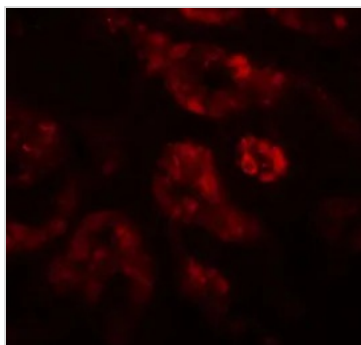
For full product information, images and publications, please visit our [website](#).

For laboratory research use only. Not for any clinical, therapeutic, or diagnostic use in humans or animals. Not for animal or human consumption.

Note

Purchasers shall not, and agree not to enable third parties to, analyze, copy, reverse engineer or otherwise attempt to determine the structure or sequence of the product.

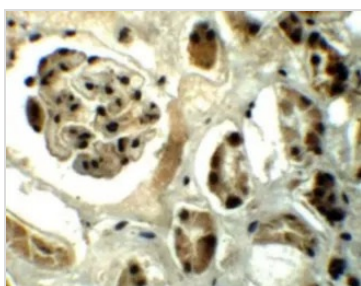
DATA IMAGES



GTX31819 IHC-P Image

IHC-P analysis of human kidney tissue using GTX31819 DPF2 antibody.

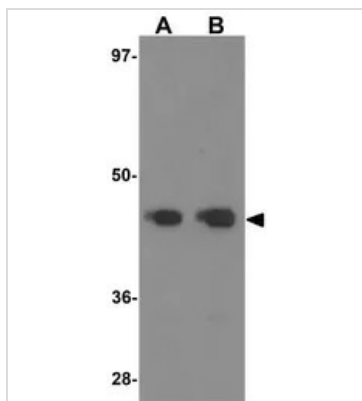
Working concentration : 20 µg/ml



GTX31819 IHC-P Image

IHC-P analysis of human kidney tissue using GTX31819 DPF2 antibody.

Working concentration : 5 µg/ml



GTX31819 WB Image

WB analysis of mouse kidney tissue lysate using GTX31819 DPF2 antibody.

Working concentration : 1 µg/ml



For full product information, images and publications, please visit our [website](#).