

**KANK1 Antibody**  
Catalog # ASC11646**Specification****KANK1 Antibody - Product Information**

Application	WB, IHC, IF
Primary Accession	<a href="#">Q14678</a>
Other Accession	<a href="#">NP_055973</a> , <a href="#">64464726</a>
Reactivity	Human, Mouse
Host	Rabbit
Clonality	Polyclonal
Isotype	IgG
Calculated MW	Predicted: 149 kDa
Application Notes	Observed: 150 kDa KDa KANK1 Antibody can be used for detection of KANK1 by Western blot at 1 µg/mL.

**KANK1 Antibody - Additional Information**

Gene ID 23189

**Target/Specificity**

KANK1; Two alternatively spliced transcript variants encoding different isoforms have been identified. The lower molecular weight band seen in the immunoblot is thought to be non-specific.

**Reconstitution & Storage**

KANK1 antibody can be stored at 4°C for three months and -20°C, stable for up to one year.

**Precautions**

KANK1 Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

**KANK1 Antibody - Protein Information**Name KANK1 ([HGNC:19309](#))**Function**

Adapter protein that links structural and signaling protein complexes positioned to guide microtubule and actin cytoskeleton dynamics during cell morphogenesis (PubMed:[22084092](http://www.uniprot.org/citations/22084092)), PubMed:[24120883](http://www.uniprot.org/citations/24120883)). At focal adhesions (FAs) rims, organizes cortical microtubule stabilizing complexes (CMSCs) and directly interacts with major FA component TLN1, forming macromolecular assemblies positioned to control microtubule-actin crosstalk at the cell edge (PubMed:[24120883](http://www.uniprot.org/citations/24120883)), PubMed:[27410476](http://www.uniprot.org/citations/27410476)). Recruits KIF21A in CMSCs at axonal growth cones and regulates axon guidance by suppressing microtubule growth without inducing microtubule disassembly once it reaches the cell cortex (PubMed:[24120883](http://www.uniprot.org/citations/24120883)). Interacts

with ARFGEF1 and participates in establishing microtubule-organizing center (MTOC) orientation and directed cell movement in wound healing (PubMed:<a href="http://www.uniprot.org/citations/22084092" target="\_blank">22084092</a>). Regulates actin stress fiber formation and cell migration by inhibiting RHOA activation in response to growth factors; this function involves phosphorylation through PI3K/Akt signaling and may depend on the competitive interaction with 14-3-3 adapter proteins to sequester them from active complexes (PubMed:<a href="http://www.uniprot.org/citations/18458160" target="\_blank">18458160</a>, PubMed:<a href="http://www.uniprot.org/citations/25961457" target="\_blank">25961457</a>). Inhibits the formation of lamellipodia but not of filopodia; this function may depend on the competitive interaction with BAIAP2 to block its association with activated RAC1. Inhibits fibronectin-mediated cell spreading; this function is partially mediated by BAIAP2 (PubMed:<a href="http://www.uniprot.org/citations/19171758" target="\_blank">19171758</a>). In the nucleus, is involved in beta-catenin- dependent activation of transcription (PubMed:<a href="http://www.uniprot.org/citations/16968744" target="\_blank">16968744</a>). During cell division, may regulate DAAM1-dependent RHOA activation that signals centrosome maturation and chromosomal segregation. May also be involved in contractile ring formation during cytokinesis (By similarity). Potential tumor suppressor for renal cell carcinoma (Probable).

#### Cellular Location

Cytoplasm, cell cortex. Cell projection, ruffle membrane; Peripheral membrane protein. Cytoplasm. Nucleus. Note=Shuttles between the cytoplasm and nucleus (PubMed:16968744). Colocalizes with CMSC components at focal adhesion rims. Colocalizes with KIF21A in membrane ruffles (PubMed:19559006, PubMed:27410476). Colocalizes with RHOA at the contractile ring. Colocalizes with RHOA and DAAM1 around centrosomes {ECO:0000250|UniProtKB:E9Q238, ECO:0000269|PubMed:16968744, ECO:0000269|PubMed:19559006, ECO:0000269|PubMed:27410476} [Isoform 2]: Cytoplasm. Nucleus Note=Shuttles between the cytoplasm and nucleus

#### Tissue Location

Widely expressed. Isoform 1 is predominantly expressed in heart and kidney. Isoform 2 probably is widely expressed at basic levels.

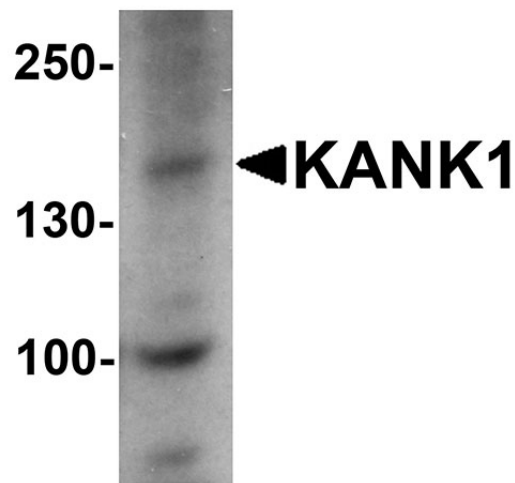
### KANK1 Antibody - Protocols

Provided below are standard protocols that you may find useful for product applications.

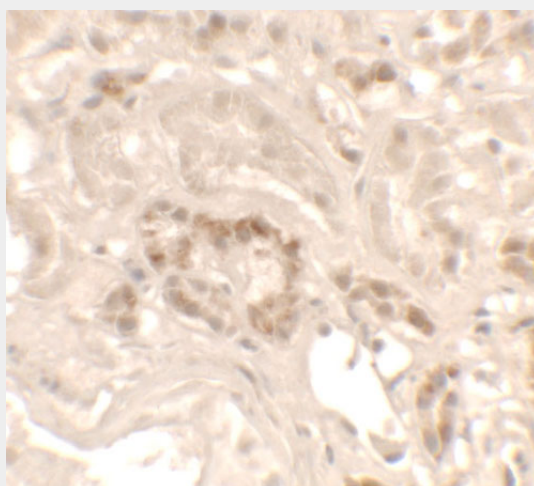
- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

### KANK1 Antibody - Images

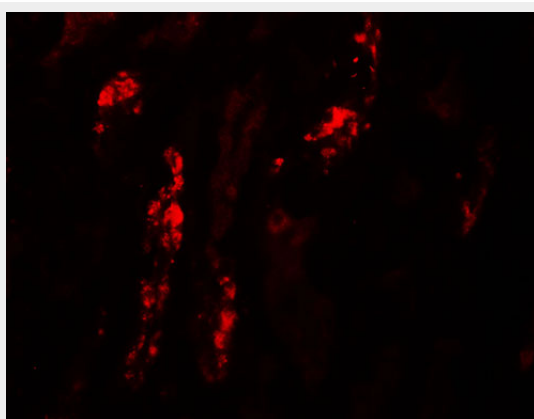




Western blot analysis of KANK1 in 3T3 cell lysate with KANK1 antibody at 1  $\mu$ g/mL.



Immunohistochemistry of KANK1 in human kidney tissue with KANK1 antibody at 2.5  $\mu$ g/ml.



Immunofluorescence of KANK1 in human kidney tissue with KANK1 antibody at 20  $\mu$ g/ml.

#### **KANK1 Antibody - Background**

KANK1 Antibody: Ankyrins are membrane adaptor molecules that play important roles in the control of cytoskeleton formation by regulating actin polymerization. KANK1 (KN motif and ankyrin repeat domain-containing protein 1), also known as ANKRD15, is a 1,352 amino acid protein that contains at least 12 exons and 5 ANK repeats. It binds to beta-catenin and regulates its subcellular

distribution. KANK1 is ubiquitously expressed and localizes to cytoplasm. It may function as a tumor suppressor for renal cell carcinoma. Mutations in this gene cause cerebral palsy spastic quadriplegic type 2, a central nervous system development disorder.

### **KANK1 Antibody - References**

Zhu Y, Kakinuma N, Wang Y, et al. Kank proteins: a new family of ankyrin-repeat domain containing proteins. *Biochim. Biophys. Acta* 2008; 1780:128-33.

Roy BC, Kakinuma N, Kiyama R. Kank attenuates actin remodeling by preventing interaction between IRSp53 and Rac1. *J. Cell Biol.* 2009; 184:253-67.

Sarkar S, Roy BC, Hatano N, et al. A novel ankyrin repeat-containing gene (Kank) located at 9p24 is a growth suppressor of renal cell carcinoma. *J. Biol. Chem.* 2002; 277:36585-91.

Lerer I, Sagi M, Meiner V, et al. Deletion of the ANKRD15 gene at 9p24.3 causes parent-of-origin-dependent inheritance of familial cerebral palsy. *Hum. Mol. Genet.* 2005; 14: 3911-20