

ULK2 Antibody (Internal)
Rabbit Polyclonal Antibody
Catalog # ALS16022**Specification**

ULK2 Antibody (Internal) - Product Information

Application	WB, IF, IHC
Primary Accession	Q8IYT8
Reactivity	Human
Host	Rabbit
Clonality	Polyclonal
Calculated MW	113kDa KDa

ULK2 Antibody (Internal) - Additional Information**Gene ID** 9706**Other Names**

Serine/threonine-protein kinase ULK2, 2.7.11.1, Unc-51-like kinase 2, ULK2, KIAA0623

Target/Specificity

At least two isoforms of ULK2 are known to exist; this antibody will detect both isoforms. ULK2 antibody is predicted to not cross-react with ULK1.

Reconstitution & Storage

Long term: -20°C; Short term: +4°C. Avoid repeat freeze-thaw cycles.

Precautions

ULK2 Antibody (Internal) is for research use only and not for use in diagnostic or therapeutic procedures.

ULK2 Antibody (Internal) - Protein Information**Name** ULK2**Synonyms** KIAA0623**Function**

Serine/threonine-protein kinase involved in autophagy in response to starvation. Acts upstream of phosphatidylinositol 3-kinase PIK3C3 to regulate the formation of autophagophores, the precursors of autophagosomes. Part of regulatory feedback loops in autophagy: acts both as a downstream effector and a negative regulator of mammalian target of rapamycin complex 1 (mTORC1) via interaction with RPTOR. Activated via phosphorylation by AMPK, also acts as a negative regulator of AMPK through phosphorylation of the AMPK subunits PRKAA1, PRKAB2 and PRKAG1. May phosphorylate ATG13/KIAA0652, FRS2, FRS3 and RPTOR; however such data need additional evidences. Not involved in ammonia-induced autophagy or in autophagic response of cerebellar granule neurons (CGN) to low potassium concentration. Plays a role early in neuronal differentiation and is required for granule cell axon formation: may govern axon formation via

Ras-like GTPase signaling and through regulation of the Rab5-mediated endocytic pathways within developing axons.

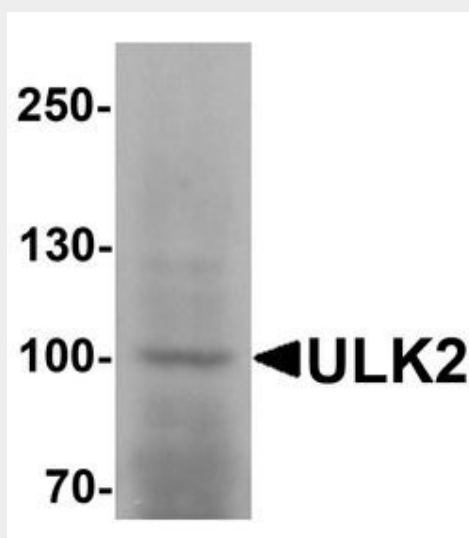
Cellular Location

Cytoplasmic vesicle membrane; Peripheral membrane protein. Note=Localizes to pre-autophagosomal membrane

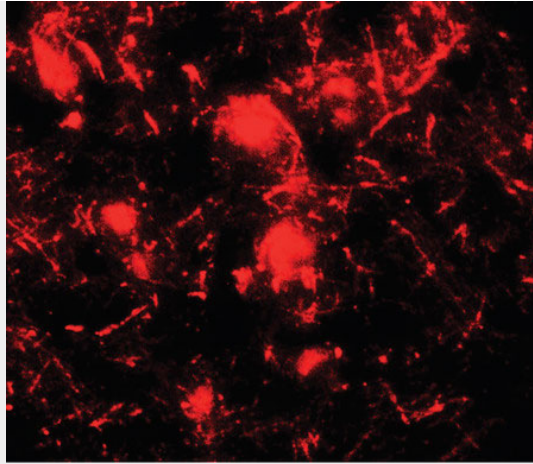
ULK2 Antibody (Internal) - Protocols

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

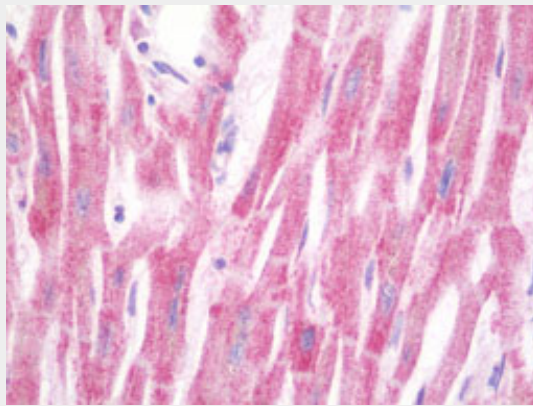
- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

ULK2 Antibody (Internal) - Images

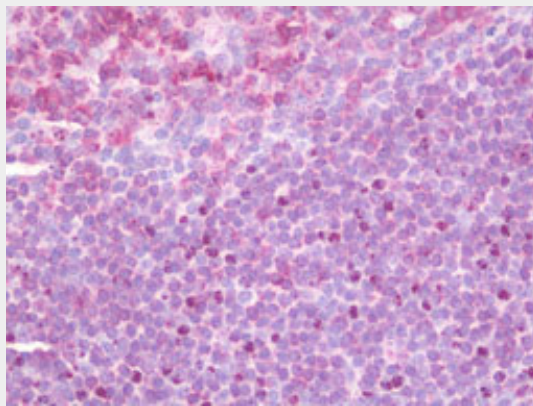
Western blot analysis of ULK2 in human brain tissue lysate with ULK2 antibody at 1 ug/ml.



Immunofluorescence of ULK2 in human brain tissue with ULK2 antibody at 20 ug/ml.



Anti-ULK2 antibody IHC staining of human heart.



Anti-ULK2 antibody IHC staining of human tonsil.

ULK2 Antibody (Internal) - Background

Serine/threonine-protein kinase involved in autophagy in response to starvation. Acts upstream of phosphatidylinositol 3- kinase PIK3C3 to regulate the formation of autophagophores, the precursors of autophagosomes. Part of regulatory feedback loops in autophagy: acts both as a downstream effector and a negative regulator of mammalian target of rapamycin complex 1 (mTORC1) via interaction with RPTOR. Activated via phosphorylation by AMPK, also acts as a negative regulator of AMPK through phosphorylation of the AMPK subunits PRKAA1, PRKAB2 and PRKAG1. May phosphorylate ATG13/KIAA0652, FRS2, FRS3 and RPTOR; however such data need additional evidences. Not involved in ammonia-induced autophagy or in autophagic response of cerebellar

granule neurons (CGN) to low potassium concentration. Plays a role early in neuronal differentiation and is required for granule cell axon formation: may govern axon formation via Ras-like GTPase signaling and through regulation of the Rab5-mediated endocytic pathways within developing axons.

ULK2 Antibody (Internal) - References

Ishikawa K.,et al.DNA Res. 5:169-176(1998).
Zody M.C.,et al.Nature 440:1045-1049(2006).
Chan E.Y.W.,et al.Mol. Cell. Biol. 29:157-171(2009).
Lee E.J.,et al.Autophagy 7:689-695(2011).
Loffler A.S.,et al.Autophagy 7:696-706(2011).