

**FBX09 / FBX09 Antibody (aa431-447)**  
**Rabbit Polyclonal Antibody**  
**Catalog # ALS11326**

**Specification**

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**FBX09 / FBX09 Antibody (aa431-447) - Product Information**

Application	WB, IHC-P, IF, E
Primary Accession	<a href="#">O9UK97</a>
Reactivity	Human
Host	Rabbit
Clonality	Polyclonal
Calculated MW	52kDa KDa
Dilution	WB~~1:1000 IHC-P~~N/A IF~~1:50~200 E~~N/A

**FBX09 / FBX09 Antibody (aa431-447) - Additional Information**

**Gene ID** 26268

**Other Names**

F-box only protein 9, Cross-immune reaction antigen 1, Renal carcinoma antigen NY-REN-57, FBX09, FBX9, VCIA1

**Target/Specificity**

Amino acids 431-447 of human FBOX9 protein.

**Reconstitution & Storage**

+4°C or -20°C, Avoid repeated freezing and thawing.

**Precautions**

FBX09 / FBX09 Antibody (aa431-447) is for research use only and not for use in diagnostic or therapeutic procedures.

**FBX09 / FBX09 Antibody (aa431-447) - Protein Information**

**Name** FBX09

**Synonyms** FBX9, VCIA1

**Function**

Substrate recognition component of a SCF (SKP1-CUL1-F-box protein) E3 ubiquitin-protein ligase complex which mediates the ubiquitination and subsequent proteasomal degradation of target proteins and plays a role in several biological processes such as cell cycle, cell proliferation, or maintenance of chromosome stability (PubMed:<a href="http://www.uniprot.org/citations/23263282" target="\_blank">23263282</a>, PubMed:<a href="http://www.uniprot.org/citations/34480022" target="\_blank">34480022</a>). Ubiquitinates

mTORC1-bound TTI1 and TELO2 when they are phosphorylated by CK2 following growth factor deprivation, leading to their degradation. In contrast, does not mediate ubiquitination of TTI1 and TELO2 when they are part of the mTORC2 complex. As a consequence, mTORC1 is inactivated to restrain cell growth and protein translation, while mTORC2 is the activated due to the relief of feedback inhibition by mTORC1 (PubMed:<a href="http://www.uniprot.org/citations/23263282" target="\_blank">23263282</a>). Plays a role in maintaining epithelial cell survival by regulating the turn- over of chromatin modulator PRMT4 through ubiquitination and degradation by the proteasomal pathway (PubMed:<a href="http://www.uniprot.org/citations/34480022" target="\_blank">34480022</a>). Regulates also PPARgamma stability by facilitating PPARgamma/PPARG ubiquitination and thereby plays a role in adipocyte differentiation (By similarity).

#### Cellular Location

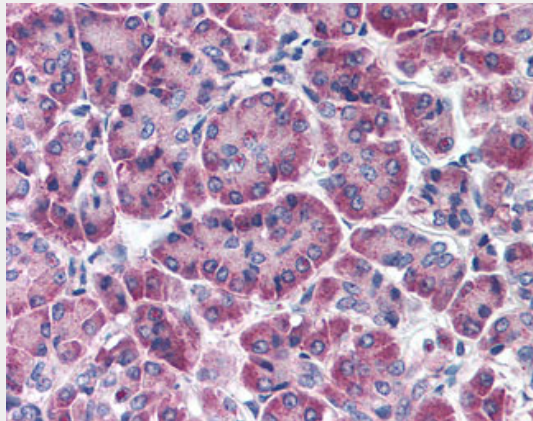
Cytoplasm.

#### FBX09 / FBX09 Antibody (aa431-447) - Protocols

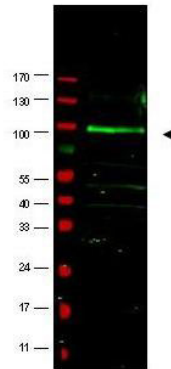
Provided below are standard protocols that you may find useful for product applications.

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

#### FBX09 / FBX09 Antibody (aa431-447) - Images



Anti-FBX09 antibody IHC of human pancreas.



Anti-FBOX9 Antibody - Western Blot.

### **FBX09 / FBX09 Antibody (aa431-447) - Background**

Substrate recognition component of a SCF (SKP1-CUL1-F-box protein) E3 ubiquitin-protein ligase complex which mediates the ubiquitination and subsequent proteasomal degradation of TTI1 and TELO2 in a CK2-dependent manner, thereby directly regulating mTOR signaling. SCF(FBX09) recognizes and binds mTORC1-bound TTI1 and TELO2 when they are phosphorylated by CK2 following growth factor deprivation, leading to their degradation. In contrast, the SCF(FBX09) does not mediate ubiquitination of TTI1 and TELO2 when they are part of the mTORC2 complex. As a consequence, mTORC1 is inactivated to restrain cell growth and protein translation, while mTORC2 is activated due to the relief of feedback inhibition by mTORC1.

### **FBX09 / FBX09 Antibody (aa431-447) - References**

- Winston J.T., et al. *Curr. Biol.* 9:1180-1182(1999).
- Deng H.-X., et al. Submitted (DEC-2004) to the EMBL/GenBank/DDBJ databases.
- Bechtel S., et al. *BMC Genomics* 8:399-399(2007).
- Ota T., et al. *Nat. Genet.* 36:40-45(2004).
- Totoki Y., et al. Submitted (MAR-2005) to the EMBL/GenBank/DDBJ databases.