



Rabbit Anti-FBXO11 antibody

SL8472R

Product Name:	FBXO11
Chinese Name:	F-box蛋白相关蛋白11抗体
Alias:	F box protein 11; F-box only protein 11; FBX11; FBX11_HUMAN; Fbxo11; Fbxo 11; Fbxo-11; PRMT9; VIT-1; VIT1; Vitiligo associated protein VIT 1; Vitiligo-associated protein 1.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human,Mouse,Rat,Chicken,Pig,Cow,Horse,Sheep,
Applications:	ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800IF=1:50-200 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	104kDa
Cellular localization:	The nucleus
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human FBXO11:65-160/927
Lsotype:	IgG
Purification:	affinity purified by Protein A
Storage Buffer:	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Storage:	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized antibody is stable at room temperature for at least one month and for greater than a year when kept at -20°C. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.
PubMed:	PubMed
Product Detail:	This gene encodes a member of the F-box protein family which is characterized by an approximately 40 amino acid motif, the F-box. The F-box proteins constitute one of the four subunits of ubiquitin protein ligase complex called SCFs (SKP1-cullin-F-box), which function in phosphorylation-dependent ubiquitination. The F-box proteins are divided into 3 classes: Fbws containing WD-40 domains, Fbls containing leucine-

rich repeats, and Fbxs containing either different protein-protein interaction modules or no recognizable motifs. The protein encoded by this gene belongs to the Fbxs class. It can function as an arginine methyltransferase that symmetrically dimethylates arginine residues, and it acts as an adaptor protein to mediate the neddylation of p53, which leads to the suppression of p53 function. This gene is known to be down-regulated in melanocytes from patients with vitiligo, a skin disorder that results in depigmentation. Polymorphisms in this gene are associated with chronic otitis media with effusion and recurrent otitis media (COME/ROM), a hearing loss disorder, and the knockout of the homologous mouse gene results in the deaf mouse mutant Jeff (Jf), a single gene model of otitis media. Alternatively spliced transcript variants encoding distinct isoforms have been identified for this gene. [provided by RefSeq, Jun 2010].

Function:

Substrate recognition component of the SCF (SKP1-CUL1-F-box protein) E3 ubiquitin-protein ligase complex which mediates the ubiquitination and subsequent proteasomal degradation of target proteins. Probably recognizes and binds to phosphorylated target proteins. Binds to and neddylates phosphorylated p53/TP53, inhibiting its transcriptional activity. SCF(FBXO11) does not seem to direct ubiquitination of TP53.

Subunit:

Component of the probable SCF(FBXO11) complex consisting of CUL1, RBX1, SKP1 and FBXO11. Interacts with TP53.

Subcellular Location:

Nucleus

Tissue Specificity:

Isoform 5 is expressed in keratinocytes, fibroblasts and melanocytes.

Similarity:

Contains 1 F-box domain.

Contains 19 PbH1 repeats.

Contains 1 UBR-type zinc finger.

SWISS:

Q86XK2

Gene ID:

80204

Database links:

[Entrez Gene: 80204](#)Human

[Entrez Gene: 225055](#)Mouse

[Entrez Gene: 301674](#)Rat

[Omim: 607871](#)Human
[SwissProt: Q86XK2](#)Human
[SwissProt: Q7TPD1](#)Mouse
[SwissProt: Q7TSL3](#)Rat
[Unigene: 352677](#)Human
[Unigene: 386857](#)Mouse
[Unigene: 36585](#)Rat

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