



# FBXO7 Polyclonal Antibody

<b>Catalog No</b>	BYab-10625
<b>Isotype</b>	IgG
<b>Reactivity</b>	Human;Rat;Mouse;
<b>Applications</b>	WB;ELISA
<b>Gene Name</b>	FBXO7 FBX7
<b>Protein Name</b>	F-box protein 7
<b>Immunogen</b>	Synthetic peptide from human protein at AA range: 371-420
<b>Specificity</b>	The antibody detects endogenous FBXO7 protein
<b>Formulation</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Source</b>	Polyclonal, Rabbit,IgG
<b>Purification</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Dilution</b>	WB 1:500-2000, ELISA 1:10000-20000
<b>Concentration</b>	1 mg/ml
<b>Purity</b>	≥90%
<b>Storage Stability</b>	-20°C/1 year
<b>Synonyms</b>	FBXO7 FBX7
<b>Observed Band</b>	58kD
<b>Cell Pathway</b>	Cytoplasm . Nucleus . Mitochondrion . Cytoplasm, cytosol . Predominantly cytoplasmic (PubMed:16096642). A minor proportion is detected in the nucleus (PubMed:16096642). Relocates from the cytosol to depolarized mitochondria (PubMed:23933751) . .
<b>Tissue Specificity</b>	Fetal kidney,Pancreas,
<b>Function</b>	disease:Defects in FBXO7 may be the cause of parkinsonian-pyramidal syndrome (PKPS) [MIM:260300]. PKPS is a hypokinetic rigid disorder, the most common example of which is Parkinson disease. PKPS is a rare disorder that exhibits both Parkinsonian and pyramidal-associated signs. Symptoms, which may be vague in the beginning, start in young adulthood, progress relatively slowly, and may culminate in severe movement incapacity. Response to levodopa is usually dramatic and sustained for many years. Most, but not all, reported cases have been familial and associated with parental consanguinity, suggesting autosomal-recessive inheritance. .function:Substrate recognition component of a (SKP1-CUL1-F-box protein) E3 ubiquitin-protein ligase complex which mediates

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the ubiquitination and subsequent proteasomal degradation of target proteins.  
Recognizes BIRC2 and DLGAP5.,pathway:Protein modification;

### Background

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 the 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 and it may play a role in regulation of hematopoiesis. Alternatively spliced transcript variants of this gene have been identified with the full-length natures of only some variants being determined. [provided by RefSeq, Jul 2008],

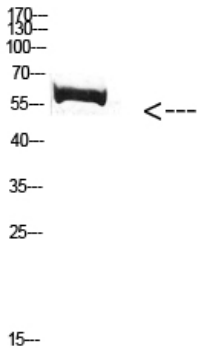
### matters needing attention

Avoid repeated freezing and thawing!

### Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

## Products Images



Western Blot analysis of mouse-kidney cells using Antibody diluted at 1000. Secondary antibody(catalog#:RS0002) was diluted at 1:20000

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