SZABO SCANDIC

Part of Europa Biosite

Produktinformation

- Forschungsprodukte & Biochemikalien
- Zellkultur & Verbrauchsmaterial
- Diagnostik & molekulare Diagnostik
- Laborgeräte & Service

Weitere Information auf den folgenden Seiten!
See the following pages for more information!

Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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FBXW4 polyclonal antibody (A01)

Catalog # : H00006468-A01 **規格 :** [50 uL]

List All

Specification

Product Description: Mouse polyclonal antibody raised against a partial recombinant FBXW4.

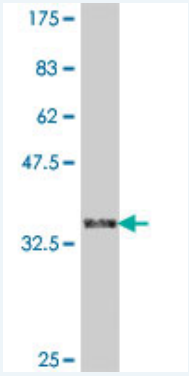
Immunogen: FBXW4 (NP_071322, 41 a.a. ~ 140 a.a) partial recombinant protein with GST tag.

Sequence: SYLDMRALGRLAQVCRWLRRFTSCDLLWRRIARASLNSGFTRLGTDLM TSVPVKERVKVSQNWRLGRCREGILLKWRCSQMPWMQLEDDSLYISQA NFIL

Host: Mouse

Reactivity: Human

Quality Control Testing: Antibody Reactive Against Recombinant Protein.



Western Blot detection against Immunogen (37.11 KDa) .

Storage Buffer: 50 % glycerol

Storage Instruction: Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

MSDS: Download

Datasheet: Download

Applications

Western Blot (Recombinant protein)

Protocol Download

ELISA

Gene Information

Entrez GeneID: 6468

GeneBank Accession#: NM_022039

Application Image

Western Blot (Recombinant protein)

ELISA

Protein Accession#:	NP_071322
Gene Name:	FBXW4
Gene Alias:	DAC,FBW4,FBWD4,SHFM3,SHSF3
Gene Description:	F-box and WD repeat domain containing 4
Omim ID:	600095, 608071
Gene Ontology:	Hyperlink
Gene Summary:	This gene is a member of the F-box/WD-40 gene family, which recruit specific target proteins through their WD-40 protein-protein binding domains for ubiquitin mediated degradation. In mouse, a highly similar protein is thought to be responsible for maintaining the apical ectodermal ridge of developing limb buds; disruption of the mouse gene results in the absence of central digits, underdeveloped or absent metacarpal/metatarsal bones and syndactyly. This phenotype is remarkably similar to split hand-split foot malformation in humans, a clinically heterogeneous condition with a variety of modes of transmission. An autosomal recessive form has been mapped to the chromosomal region where this gene is located, and complex rearrangements involving duplications of this gene and others have been associated with the condition. A pseudogene of this locus has been mapped to one of the introns of the BCR gene on chromosome 22. [provided by RefSeq
Other Designations:	F-box and WD-40 domain protein 4,F-box/WD repeat protein 4,OTTHUMP00000059175,dactylin