



## Rabbit Anti-Synaptopodin 2 antibody

SL8743R

<b>Product Name:</b>	Synaptopodin 2
<b>Chinese Name:</b>	突触足蛋白2抗体
<b>Alias:</b>	DKFZp686G051; Genethonin 2; Genethonin-2; Myopodin; striated muscle specific; Synaptopodin-2; SYNP2 HUMAN; SYNPO 2; Synpo2.
<b>Organism Species:</b>	Rabbit
<b>Clonality:</b>	Polyclonal
<b>React Species:</b>	Human,Mouse,Rat,Pig,Cow,
<b>Applications:</b>	ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800ICC=1:100-500IF=1:100-500 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
<b>Molecular weight:</b>	117kDa
<b>Cellular localization:</b>	The nucleuscytoplasmic
<b>Form:</b>	Lyophilized or Liquid
<b>Concentration:</b>	1mg/ml
<b>immunogen:</b>	KLH conjugated synthetic peptide derived from human Synaptopodin 2:601-700/1093
<b>Lsotype:</b>	IgG
<b>Purification:</b>	affinity purified by Protein A
<b>Storage Buffer:</b>	Preservative: 15mM Sodium Azide, Constituents: 1% BSA, 0.01M PBS, pH 7.4
<b>Storage:</b>	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.
<b>PubMed:</b>	<a href="#">PubMed</a>
<b>Product Detail:</b>	SYNPO2 (Synaptopodin 2) is a Protein Coding gene. Diseases associated with SYNPO2 include Duchenne Muscular Dystrophy and Myopathy, Myofibrillar, 2. GO annotations related to this gene include actin binding and muscle alpha-actinin binding. An important paralog of this gene is SYNPO2L.  <b>Function:</b>

Has an actin-binding and actin-bundling activity. Can induce the formation of F-actin networks in an isoform-specific manner (PubMed:24005909, PubMed:23225103). At the sarcomeric Z lines is proposed to act as adapter protein that links nascent myofibers to the sarcolemma via ZYX and may play a role in early assembly and stabilization of the Z lines. Involved in autophagosome formation. May play a role in chaperone-assisted selective autophagy (CASA) involved in Z lines maintenance in striated muscle under mechanical tension; may link the client-processing CASA chaperone machinery to a membrane-tethering and fusion complex providing autophagosome membranes (By similarity). Involved in regulation of cell migration (PubMed:22915763, PubMed:25883213). May be a tumor suppressor (PubMed:16885336).

**Subcellular Location:**

Nucleus. Cytoplasm. Shuttles between the nucleus and the cytoplasm in a differentiation-dependent and stress-induced fashion. Localizes to the Z-disk in mature striated muscle. The nuclear export is XPO1-dependent (By similarity). Localized in a fiber-like pattern, partly overlapping with filamentous actin.

**Tissue Specificity:**

Skeletal muscle-specific.

**DISEASE:**

Down-regulated in muscle cell lines derived from patients with Duchenne muscular dystrophy (DMD).

**Similarity:**

Belongs to the synaptopodin family.  
Contains 1 PDZ (DHR) domain.

**SWISS:**

Q9UMS6

**Gene ID:**

171024

**Database links:**

[Entrez Gene: 171024](#) Human

[Entrez Gene: 118449](#) Mouse

[Entrez Gene: 499702](#) Rat

[SwissProt: Q9UMS6](#) Human

[SwissProt: Q91YE8](#) Mouse

[Unigene: 655519](#) Human

[Unigene: 317009](#) Mouse

[Unigene: 34359](#) Mouse

[Unigene: 474733](#) Mouse

**Important Note:**

This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

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