

**Product Name:** alpha Sarcoglycan Rabbit Monoclonal Antibody**Catalog #:** AMRe87007

For research use only.

**Summary**

<b>Description</b>	Recombinant rabbit monoclonal antibody
<b>Host</b>	Rabbit
<b>Application</b>	WB,IHC,IP
<b>Reactivity</b>	Human,Mouse,Rat
<b>Conjugation</b>	Unconjugated
<b>Modification</b>	Unmodified
<b>Isotype</b>	IgG
<b>Clonality</b>	Monoclonal
<b>Form</b>	Liquid
<b>Concentration</b>	0.15mg/ml. The concentration of this product may be batch-dependent.
<b>Storage</b>	Aliquot and store at -20°C (valid for 12 months). Avoid freeze/thaw cycles.
<b>Shipping</b>	Ice bags
<b>Buffer</b>	Supplied in 50mM Tris-Glycine(pH 7.4), 0.15M NaCl, 40% Glycerol, 0.01% sodium azide and 0.05% protective protein. Stable for 12 months from date of receipt.
<b>Purification</b>	Affinity Purification

**Application**

<b>Dilution Ratio</b>	WB 1:1000-1:5000,IHC 1:500-1:2000,IP 1:20-1:50
<b>Molecular Weight</b>	Calculated MW:43 kDa; Observed MW:50 kDa

**Antigen Information**

<b>Gene Name</b>	alpha Sarcoglycan
<b>Alternative Names</b>	ADL; DAG2; 50DAG; DMDA2; LGMD2D; SCARMD1; adhalin
<b>Gene ID</b>	6442
<b>SwissProt ID</b>	Q16586
<b>Immunogen</b>	A synthetic peptide of human alpha Sarcoglycan

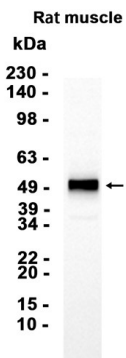
**Background**

This gene encodes a component of the dystrophin-glycoprotein complex (DGC), which is critical to the stability of muscle fiber membranes and to the linking of the actin cytoskeleton to the extracellular matrix. Its expression is thought to be restricted to

striated muscle. Mutations in this gene result in type 2D autosomal recessive limb-girdle muscular dystrophy. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Oct 2008]

## Research Area

## Image Data



Western blot analysis of extracts from Rat muscle tissue using alpha Sarcoglycan Rabbit Monoclonal Antibody at 1:1000.