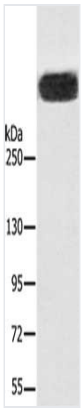
RYR1 Antibody – Rabbit Polyclonal

PRODUCT OVERVIEW

Product Name	RYR1, Polyclonal Antibody
Full Product Name	RYR1 Antibody
Product Gene Name	anti-RYR1 antibody
Product Synonym Names	ryanodine receptor 1 (skeletal)
Clonality	Polyclonal
Isotype	IgG
Host	Rabbit
Species Reactivity	Human, Mouse, Rat
Form/Format	Liquid; In Phosphate Buffered Saline, pH 7.4, 0.05% NaN3, and 40% Glycerol
Purity/Purification	Antigen Affinity Purification
Preparation and Storage	Upon receipt, store at -20 degree C or -80 degree C. Avoid repeated freeze.
Molecular Weight	565,176 Da
OMIM	117000
3D Structure	ModBase 3D Structure for P21817
Immunogen	Synthetic peptide of human RYR1
Immunogen Species	Human
Conjugate	Non-conjugated
Product Categories/Family for anti-RYR1 antibody	Neuroscience
ISO Certification	Manufactured in an ISO 9001:2015 Certified Laboratory.
Applications Tested/Suitable for anti-RYR1 antibody	ELISA (EIA), Western Blot (WB)
Application Notes for anti-RYR1 antibody	ELISA: 1:1000-1:2000 WB: 1:200-1:1000
Related Product Information for anti-RYR1 antibody	MBS7129659 is an antibody targeting RYR1. Also known as RYR1 Antibody. Reactive in Human, Mouse, Rat. Host: Rabbit.
Other Notes	Small volumes of anti-RYR1 antibody vial(s) may occasionally become entrapped in the seal of the product vial during shipment and storage. If necessary, briefly centrifuge the vial on a tabletop centrifuge to dislodge any liquid in the container`s cap. Certain products may require to ship with dry ice and additional dry ice fee may apply.



SDS-PAGE of anti-RYR1 antibody
Gel: 6%SDS-PAGE, Lysate: 40 ug, Lane: Mouse brain tissue, Primary antibody: MBS7129659(RYR1 Antibody) at dilution 1/400, Secondary antibody: Goat anti rabbit IgG at 1/8000 dilution, Exposure time: 2 minutes

ADDITIONAL DETAILS

NCBI/Uniprot data below describe general gene information for RYR1. It may not necessarily be applicable to this product.

NCBI GI #	108935904
NCBI GeneID	6261
NCBI Accession #	P21817.3
UniProt Primary Accession #	P21817
UniProt Secondary Accession #	Q16314; Q16368; Q9NPK1; Q9P1U4
UniProt Related Accession #	P21817
NCBI Official Full Name	Ryanodine receptor 1
NCBI Official Synonym Full Names	ryanodine receptor 1 (skeletal)
NCBI Official Symbol	RYR1
NCBI Official Synonym Symbols	CCO; MHS; RYR; MHS1; RYDR; SKRR; RYR-1; PPP1R137
UniProt Protein Name	Ryanodine receptor 1
UniProt Synonym Protein Names	Skeletal muscle calcium release channel; Skeletal muscle ryanodine receptor; Skeletal muscle-type ryanodine receptor; Type 1 ryanodine receptor
Protein Family	Ryanodine receptor
UniProt Gene Name	RYR1
UniProt Synonym Gene Names	RYDR; RYR-1; RyR1
UniProt Entry Name	RYR1_HUMAN
NCBI Protein Information	ryanodine receptor 1; central core disease of muscle; type 1-like ryanodine receptor; skeletal muscle ryanodine receptor; skeletal muscle calcium release channel; protein phosphatase 1, regulatory subunit 137; sarcoplasmic reticulum calcium release channe
NCBI Summary for RYR1	This gene encodes a ryanodine receptor found in skeletal muscle. The encoded protein functions as a calcium release channel in the sarcoplasmic reticulum but also serves to connect the sarcoplasmic reticulum and transverse tubule. Mutations in this gene are associated with malignant hyperthermia susceptibility, central core disease, and minicore myopathy with external ophthalmoplegia. Alternatively spliced transcripts encoding different isoforms have been described. [provided by RefSeq, Jul 2008]

UniProt Comments for RYR1	RYR1: Calcium channel that controls communication between transverse-tubules and sarcoplasmic reticulum. Contraction of skeletal muscle is triggered by release of calcium ions from SR following depolarization of T-tubules. Can mediate the release of Ca(2+) from intracellular stores in neurons, and may thereby promote prolonged Ca(2+) signaling in the brain. Required for normal development of muscle fibers, skeletal muscle, heart morphogenesis, and skin development and ossification during embryogenesis. Defects in RYR1 are the cause of malignant hyperthermia susceptibility type 1 (MHS1) and central core disease of muscle (CCD). CCD is an autosomal dominant congenital myopathy, but a severe autosomal recessive form also exists. Defects in RYR1 are the cause of multiminicore disease with external ophthalmoplegia (MMDO), congenital myopathy with fiber-type disproportion (CFTD). Defects in RYR1 may be a cause of Samaritan myopathy, a congenital myopathy with benign course. Belongs to the ryanodine receptor (TC 1.A.3.1) family. RYR1 subfamily. 3 isoforms of the human protein are produced by alternative splicing. Protein type: Transporter; Transporter, ion channel; Membrane protein, integral; Membrane protein, multi-pass; Channel, calcium Chromosomal Location of Human Ortholog: 19q13.1 Cellular Component: I band; sarcoplasmic reticulum membrane; smooth endoplasmic reticulum; sarcoplasmic reticulum; integral to plasma membrane; cytoplasm; junctional membrane complex; T-tubule; plasma membrane; cell cortex Molecular Function: voltage-gated calcium channel activity; calmodulin binding; protein binding; protease binding; calcium-release channel activity; calcium channel activity; ryanodine-sensitive calcium-release channel activity Biological Process: skin development; cytosolic calcium ion homeostasis; muscle contraction; calcium ion transport; release of sequestered calcium ion into cytosol; response to hypoxia; skeletal muscle fiber development; release of sequestered calcium ion by sarcoplasmic reticulum into cytosol; transmembrane transport; response to caffeine Disease: Minicore Myopathy With External Ophthalmoplegia; Malignant Hyperthermia, Susceptibility To, 1; Central Core Disease Of Muscle; Myopathy, Congenital, With Fiber-type Disproportion
Research Articles on RYR1	**1.** R2355W can be included in the list of causative mutations for malignant hyperthermia. V2354M might be a causative mutation.
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