

## RyR-2 (phospho Ser2808) Polyclonal Antibody

YP0967 Catalog No:

Human; Mouse; Rat Reactivity:

**Applications:** IHC;IF;ELISA

Target: RyR-2

Fields: >>Calcium signaling pathway;>>cAMP signaling pathway;>>Cardiac muscle

> contraction:>>Adrenergic signaling in cardiomyocytes:>>Apelin signaling pathway;>>Circadian entrainment;>>Insulin secretion;>>Oxytocin signaling

pathway;>>Pancreatic secretion;>>Prion disease;>>Pathways of

neurodegeneration - multiple diseases;>>Hypertrophic

cardiomyopathy;>>Arrhythmogenic right ventricular cardiomyopathy;>>Dilated

cardiomyopathy;>>Diabetic cardiomyopathy

Gene Name: RYR2

Ryanodine receptor 2 **Protein Name:** 

E9Q401

**Human Gene Id:** 6262

**Human Swiss Prot** 

Q92736

No:

Mouse Gene Id: 20191

**Mouse Swiss Prot** 

No:

Rat Gene Id: 689560

B0LPN4 Rat Swiss Prot No:

The antiserum was produced against synthesized peptide derived from human Immunogen:

RyR2 around the phosphorylation site of Ser2808. AA range:2774-2823

**Specificity:** Phospho-RyR-2 (S2808) Polyclonal Antibody detects endogenous levels of

RyR-2 protein only when phosphorylated at S2808.

Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.



**Sormedation:** Polyclonal, Rabbit, IgG

**Dilution :** IHC 1:100 - 1:300. ELISA: 1:10000.. IF 1:50-200

**Purification:** The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Molecularweight: 564kD

**Cell Pathway:** Calcium; Cardiac muscle contraction; Hypertrophic cardiomyopathy

(HCM); Arrhythmogenic right ventricular cardiomyopathy (ARVC); Dilated

cardiomyopathy;

**Background :** This gene encodes a ryanodine receptor found in cardiac muscle sarcoplasmic

reticulum. The encoded protein is one of the components of a calcium channel, composed of a tetramer of the ryanodine receptor proteins and a tetramer of FK506 binding protein 1B proteins, that supplies calcium to cardiac muscle. Mutations in this gene are associated with stress-induced polymorphic ventricular tachycardia and arrhythmogenic right ventricular dysplasia. [provided by RefSeq,

Jul 2008],

**Function:** developmental stage:Expressed in myometrium during

pregnancy., disease: Defects in RYR2 are the cause of catecholaminergic polymorphic ventricular tachycardia type 1 (CPVT1) [MIM:604772]; also known as stress-induced polymorphic ventricular tachycardia (VTSIP). CPVT1 is an autosomal dominant form of arrhythmogenic disorder characterized by stress-induced, bidirectional ventricular tachycardia that may degenerate into cardiac arrest and cause sudden death., disease: Defects in RYR2 are the cause of familial arrhythmogenic right ventricular dysplasia 2 (ARVD2) [MIM:600996]; also known as arrhythmogenic right ventricular cardiomyopathy 2 (ARVC2). ARVD is an autosomal dominant disease characterized by partial degeneration of the myocardium of the right ventricle, electrical instability, and sudden death. It is clinically defined by electrocardiographic and angiographic criteria; pathologic

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Subcellular Location:

Sarcoplasmic reticulum membrane; Multi-pass membrane protein. Membrane; Multi-pass membrane protein. Sarcoplasmic reticulum. The number of predicted transmembrane domains varies between orthologs, but both N-terminus and C-

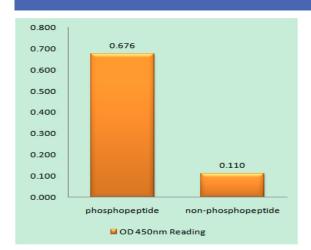
terminus seem to be cytoplasmic. .

**Expression:** Detected in heart muscle (at protein level). Heart muscle, brain (cerebellum and

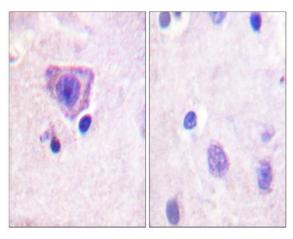
hippocampus) and placenta.



## **Products Images**



Enzyme-Linked Immunosorbent Assay (Phospho-ELISA) for Immunogen Phosphopeptide (Phospho-left) and Non-Phosphopeptide (Phospho-right), using RyR2 (Phospho-Ser2808) Antibody



Immunohistochemistry analysis of paraffin-embedded human brain, using RyR2 (Phospho-Ser2808) Antibody. The picture on the right is blocked with the phospho peptide.