

Recombinant Human SGCA 293 Cell Lysate

Cat. No. SGCA-1890HCL **Lot. No.** (See product label)

SPECIFICATION

Species	Human
Source	HEK293
Description	Antigen standard for sarcoglycan, alpha (50kDa dystrophin-associated glycoprotein) (SGCA), transcript variant 1 is a lysate prepared from HEK293T cells transiently transfected with a TrueORF gene-carrying pCMV plasmid and then lysed in RIPA Buffer. Protein concentration was determined using a colorimetric assay. The antigen control carries a C-terminal Myc/DDK tag for detection.
Components	This product includes 3 vials: 1 vial of gene-specific cell lysate, 1 vial of control vector cell lysate, and 1 vial of loading buffer. Each lysate vial contains 0.1 mg lysate in 0.1 ml (1 mg/ml) of RIPA Buffer (50 mM Tris-HCl pH7.5, 250 mM NaCl, 5 mM EDTA, 50 mM NaF, 1% NP40). The loading buffer vial contains 0.5 ml 2X SDS Loading Buffer (125 mM Tris-Cl, pH6.8, 10% glycerol, 4% SDS, 0.002% Bromophenol blue, 5% beta-mercaptoethanol).
Size	0.1 mg
Storage Instruction	Store at -80°C. Minimize freeze-thaw cycles. After addition of 2X SDS Loading Buffer, the lysates can be stored at -20°C. Product is guaranteed 6 months from the date of shipment.
Applications	ELISA, WB, IP. WB: Mix equal volume of lysates with 2X SDS Loading Buffer. Boil the mixture for 10 min before loading (for membrane protein lysates, incubate the

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mixture at room temperature for 30 min). Load 5 ug lysate per lane.

GENE INFORMATION

Gene Name	SGCA sarcoglycan, alpha (50kDa dystrophin-associated glycoprotein) [Homo sapiens]
Official Symbol	SGCA
Synonyms	SGCA; sarcoglycan, alpha (50kDa dystrophin-associated glycoprotein); ADL, sarcoglycan, alpha (50kD dystrophin associated glycoprotein); alpha-sarcoglycan; 50kD DAG; A2; adhalin; adhalin (limb girdle muscular dystrophy 2D); DMDA2; LGMD2D; Sarcoglycan; alpha (50kD dystrophin associated glycoprotein; adhalin); SCARMD1; 50DAG; alpha-SG; dystroglycan-2; 50 kDa dystrophin-associated glycoprotein; ADL; DAG2; 50-DAG;
Gene ID	6442
mRNA Refseq	NM_000023
Protein Refseq	NP_000014
MIM	600119
UniProt ID	Q16586
Chromosome Location	17q21
Pathway	Arrhythmogenic right ventricular cardiomyopathy (ARVC), organism-specific biosystem; Arrhythmogenic right ventricular cardiomyopathy (ARVC), conserved

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biosystem; Dilated cardiomyopathy, organism-specific biosystem; Dilated cardiomyopathy, conserved biosystem; Hypertrophic cardiomyopathy (HCM), organism-specific biosystem; Hypertrophic cardiomyopathy (HCM), conserved biosystem; Viral myocarditis, organism-specific biosystem;

Function

calcium ion binding;

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