

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Product datasheet for TA501146M

PKA R2 (PRKAR2A) Mouse Monoclonal Antibody [Clone ID: OTI1F8]

Product data:

Product Type:	Primary Antibodies
Clone Name:	OTI1F8
Applications:	FC, IF, IHC, WB
Recommended Dilution:	WB 1:2000, IHC 1:150, IF 1:100, FLOW 1:100
Reactivity:	Human, Mouse, Rat
Host:	Mouse
lsotype:	lgG1
Clonality:	Monoclonal
Immunogen:	Full length human recombinant protein of human PRKAR2A (NP_004148) produced in HEK293T cell.
Formulation:	PBS (pH 7.3) containing 1% BSA, 50% glycerol and 0.02% sodium azide.
Concentration:	0.74 mg/ml
Purification:	Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography (protein A/G)
Conjugation:	Unconjugated
Storage:	Store at -20°C as received.
Stability:	Stable for 12 months from date of receipt.
Predicted Protein Size:	45.3 kDa
Gene Name:	protein kinase cAMP-dependent type II regulatory subunit alpha
Database Link:	<u>NP_004148</u> <u>Entrez Gene 19087 MouseEntrez Gene 29699 RatEntrez Gene 5576 Human P13861</u>



This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2025 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US

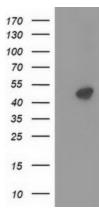
CRIGENE PKA R2 (PRKAR2A) Mouse Monoclonal Antibody [Clone ID: OTI1F8] – TA501146M

Background: CAMP is a signaling molecule important for a variety of cellular functions. cAMP exerts its effects by activating the cAMP-dependent protein kinase, which transduces the signal through phosphorylation of different target proteins. The inactive kinase holoenzyme is a tetramer composed of two regulatory and two catalytic subunits. cAMP causes the dissociation of the inactive holoenzyme into a dimer of regulatory subunits bound to four cAMP and two free monomeric catalytic subunits. Four different regulatory subunits and three catalytic subunits have been identified in humans. The protein encoded by this gene is one of the regulatory subunits. This subunit can be phosphorylated by the activated catalytic subunit. It may interact with various A-kinase anchoring proteins and determine the subcellular localization of cAMP-dependent protein kinase. This subunit has been shown to regulate protein transport from endosomes to the Golgi apparatus and further to the endoplasmic reticulum (ER). [provided by RefSeq]

Synonyms: PKR2; PRKAR2

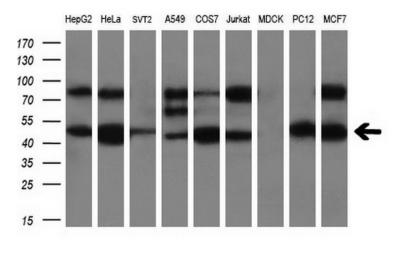
Protein Families:Druggable GenomeProtein Pathways:Apoptosis, Insulin signaling pathway

Product images:

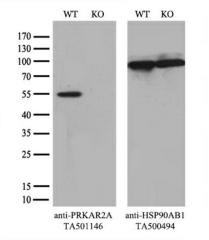


HEK293T cells were transfected with the pCMV6-ENTRY control (Cat# [PS100001], Left lane) or pCMV6-ENTRY PRKAR2A (Cat# [RC220376], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-PRKAR2A(Cat# [TA501146]). Positive lysates [LY401337] (100ug) and [LC401337] (20ug) can be purchased separately from OriGene.

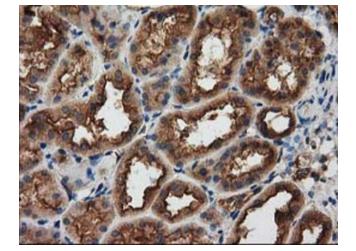
This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2025 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US



Western blot analysis of extracts (35ug) from 9 different cell lines by using anti-PRKAR2A monoclonal antibody at 1:200 dilution. (HepG2: human; HeLa: human; SVT2: mouse; A549: human; COS7: monkey; Jurkat: human; MDCK: canine; PC12: rat; MCF7: human)

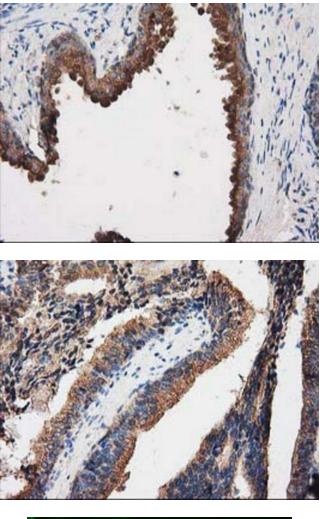


Equivalent amounts of cell lysates (10 ug per lane) of wild-type HeLa cells (WT, Cat# LC810HELA) and PRKAR2A-Knockout HeLa cells (KO, Cat# [LC812853]) were separated by SDS-PAGE and immunoblotted with anti-PRKAR2A monoclonal antibody [TA501146] (1:2000`). Then the blotted membrane was stripped and reprobed with anti-HSP90 antibody as a loading control.



Immunohistochemical staining of paraffinembedded Human Kidney tissue within the normal limits using anti-PRKAR2A mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.

This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2025 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US

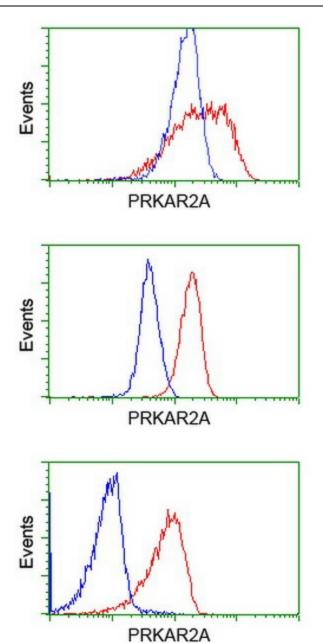


Immunohistochemical staining of paraffinembedded Human prostate tissue within the normal limits using anti-PRKAR2A mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.

Immunohistochemical staining of paraffinembedded Carcinoma of Human prostate tissue using anti-PRKAR2A mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.

Anti-PRKAR2A mouse monoclonal antibody ([TA501146]) immunofluorescent staining of COS7 cells transiently transfected by pCMV6-ENTRY PRKAR2A ([RC220376]).

This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2025 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US



HEK293T cells transfected with either [RC220376] overexpress plasmid (Red) or empty vector control plasmid (Blue) were immunostained by anti-PRKAR2A antibody ([TA501146]), and then analyzed by flow cytometry.

Flow cytometric Analysis of Hela cells, using anti-PRKAR2A antibody ([TA501146]), (Red), compared to a nonspecific negative control antibody (TA50011), (Blue).

Flow cytometric Analysis of Jurkat cells, using anti-PRKAR2A antibody ([TA501146]), (Red), compared to a nonspecific negative control antibody (TA50011), (Blue).

This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2025 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US