antibodies - online.com







anti-PRKAG2 antibody (N-Term)

Images



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100 μL	
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PRKAG2	
N-Term	
Human, Monkey	
Rabbit	
Polyclonal	
This PRKAG2 antibody is un-conjugated	
Western Blotting (WB), Immunohistochemistry (IHC), Immunofluorescence (IF),	
Immunochromatography (IC)	
KLH-conjugated synthetic peptide encompassing a sequence within the N-term region of	
human AMPK gamma 2.	
Recognizes endogenous levels of AMPK gamma 2 protein.	
Rabbit polyclonal antibody to AMPK gamma 2	
The antibody was purified by immunogen affinity chromatography.	
PRKAG2	
AMPK gamma 2 (PRKAG2 Products)	
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Target Details

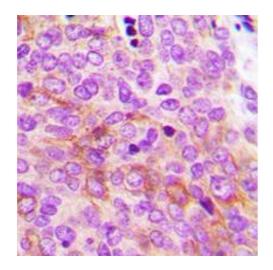
Background:	5'-AMP-activated protein kinase subunit gamma-2, AMPK gamma2, AMPK subunit gamma-2, H91620p
Gene ID:	51422, 108099
UniProt:	Q9UGJ0, Q91WG5
Pathways:	AMPK Signaling, Cellular Glucan Metabolic Process, Ribonucleoside Biosynthetic Process, Regulation of Carbohydrate Metabolic Process, Warburg Effect

Application Details

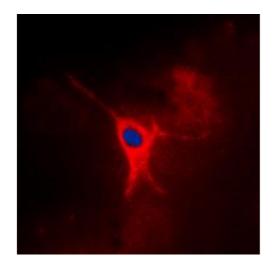
Application Notes:	WB (1:500 - 1:1000), IH (1:100 - 1:200), IF/IC (1:100 - 1:500)
Restrictions:	For Research Use only

Handling

Format:	Liquid
Buffer:	Liquid in 0.42 % Potassium phosphate, 0.87 % Sodium chloride, pH 7.3, 30 % glycerol, and 0.01 % sodium azide.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid freeze/thaw cycles.
Expiry Date:	12 months



kDa A B 200 140 100 80 60 50 40 30 20



Immunohistochemistry

Image 1. Immunohistochemical analysis of AMPK gamma 2 staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

Western Blotting

Image 2. Western blot analysis of AMPK gamma 2 expression in MCF7 (A), HeLa (B) whole cell lysates.

Immunofluorescence

Image 3. Immunofluorescent analysis of AMPK gamma 2 staining in MCF7 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).