

## **Product datasheet for TP761181**

## 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

## PRKAG2 (NM\_016203) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Purified recombinant protein of Human protein kinase, AMP-activated, gamma 2 non-catalytic

subunit (PRKAG2), transcript variant a, full length, with N-terminal HIS tag, expressed in E. coli,

50ug

Species: Human

**Expression Host:** E. coli

**Expression cDNA Clone** 

or AA Sequence:

A DNA sequence encoding human full-length PRKAG2

Tag: N-His

**Predicted MW:** 62.9 kDa

**Concentration:** >0.05 μg/μL as determined by microplate BCA method

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

**Buffer:** 50 mM Tris-HCl, pH 8.0, 8 M urea

**Note:** For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and

handling conditions. Avoid repeated freeze-thaw cycles.

**RefSeq:** NP 057287

**Locus ID:** 51422

**UniProt ID:** Q9UGI0, A0A090N8Q6

RefSeq Size: 2062 Cytogenetics: 7q36.1

RefSeq ORF: 1704

Synonyms: AAKG; AAKG2; CMH6; H91620p; WPWS





**Summary:** 

AMP-activated protein kinase (AMPK) is a heterotrimeric protein composed of a catalytic alpha subunit, a noncatalytic beta subunit, and a noncatalytic regulatory gamma subunit. Various forms of each of these subunits exist, encoded by different genes. AMPK is an important energy-sensing enzyme that monitors cellular energy status and functions by inactivating key enzymes involved in regulating de novo biosynthesis of fatty acid and cholesterol. This gene is a member of the AMPK gamma subunit family. Mutations in this gene have been associated with Wolff-Parkinson-White syndrome, familial hypertrophic cardiomyopathy, and glycogen storage disease of the heart. Alternate transcriptional splice variants, encoding different isoforms, have been characterized. [provided by RefSeq, Jan 2015]

**Protein Families:** Druggable Genome

Protein Pathways: Adipocytokine signaling pathway, Hypertrophic cardiomyopathy (HCM), Insulin signaling

pathway

## **Product images:**

