

# HK2 (Hexokinase II) (Center) Antibody

Rabbit Polyclonal Antibody Catalog # ABV10116

### **Specification**

### HK2 (Hexokinase II) (Center) Antibody - Product Information

Application WB, IHC, E
Primary Accession P52789
Reactivity Human
Host Rabbit
Clonality Polyclonal
Isotype Rabbit IgG
Calculated MW 102380

# HK2 (Hexokinase II) (Center) Antibody - Additional Information

**Gene ID 3099** 

Positive Control Western Blot: A375 cell lysate

Immunohistochemistry: Human cancer

tissue

Application & Usage The antibody can be used for ELISA (0.25

 $\mu$ g/ml), Western blotting (0.5 - 2.5  $\mu$ g/ml) and Immunohistochemistry (2.5 - 5.0

μg/ml).

#### **Other Names**

Hexokinase type II, HK II, Muscle form hexokinase

Target/Specificity HK2 (Hexokinase II)

**Antibody Form** 

Liquid

**Appearance**Colorless liquid

#### **Formulation**

 $100~\mu g$  (0.25 mg/ml) purified rabbit Ig polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide.

#### **Handling**

The antibody solution should be gently mixed before use.

**Reconstitution & Storage** 

-20 °C

**Background Descriptions** 



#### **Precautions**

HK2 (Hexokinase II) (Center) Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

### HK2 (Hexokinase II) (Center) Antibody - Protein Information

# Name HK2 (<u>HGNC:4923</u>)

#### **Function**

Catalyzes the phosphorylation of hexose, such as D-glucose and D-fructose, to hexose 6-phosphate (D-glucose 6-phosphate and D- fructose 6-phosphate, respectively) (PubMed:<a href="http://www.uniprot.org/citations/23185017" target="\_blank">23185017</a>, PubMed:<a href="http://www.uniprot.org/citations/26985301" target="\_blank">26985301</a>, PubMed:<a href="http://www.uniprot.org/citations/29298880" target="\_blank">29298880</a>). Mediates the initial step of glycolysis by catalyzing phosphorylation of D-glucose to D-glucose 6-phosphate (PubMed:<a href="http://www.uniprot.org/citations/29298880" target="\_blank">29298880</a>). Plays a key role in maintaining the integrity of the outer mitochondrial membrane by preventing the release of apoptogenic molecules from the intermembrane space and subsequent apoptosis (PubMed:<a href="http://www.uniprot.org/citations/18350175" target="\_blank">18350175</a>).

#### **Cellular Location**

Mitochondrion outer membrane; Peripheral membrane protein. Cytoplasm, cytosol Note=The mitochondrial-binding peptide (MBP) region promotes association with the mitochondrial outer membrane (PubMed:29298880) The interaction with the mitochondrial outer membrane via the mitochondrial-binding peptide (MBP) region promotes higher stability of the protein (PubMed:29298880). Release from the mitochondrial outer membrane into the cytosol induces permeability transition pore (PTP) opening and apoptosis (PubMed:18350175).

#### **Tissue Location**

Predominant hexokinase isozyme expressed in insulin-responsive tissues such as skeletal muscle

### HK2 (Hexokinase II) (Center) Antibody - Protocols

Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- <u>Immunohistochemistry</u>
- Immunofluorescence
- <u>Immunoprecipitation</u>
- Flow Cytomety
- Cell Culture

## HK2 (Hexokinase II) (Center) Antibody - Images

### HK2 (Hexokinase II) (Center) Antibody - Background

In vertebrates there are four major glucose-phosphorylating isoenzymes, designated hexokinase I, II, III, and IV. Hexokinase is an allosteric enzyme inhibited by its product GLC-6-P. Hexokinase activity is involved in the first step in several metabolic pathways. HK3 is bound to the outer mitochondrial membrane. Its hydrophobic N-terminal sequence may be involved in membrane bindng. It is the predominant hexokinase isozyme expressed in insuline-responsive tissues such as skeletal muscle. The N- and C-terminal halves of this hexokinase show extensive sequence





similarity to each other. The catalytic activity is associated with the C-terminus while regulatory function is associated wiht the N-terminus. Altho µgh found in NIDDM patients, genetic variations of HK2 do not contribute to the disease.