

**Anti-TRAP1 Picoband Antibody**  
**Catalog # ABO12520****Specification**

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**Anti-TRAP1 Picoband Antibody - Product Information**

Application	WB, IHC-P
Primary Accession	<a href="#">Q12931</a>
Host	Rabbit
Reactivity	Human, Rat
Clonality	Polyclonal
Format	Lyophilized

**Description**

Rabbit IgG polyclonal antibody for Heat shock protein 75 kDa, mitochondrial (TRAP1) detection. Tested with WB, IHC-P in Human; Rat.

**Reconstitution**

Add 0.2ml of distilled water will yield a concentration of 500ug/ml.

**Anti-TRAP1 Picoband Antibody - Additional Information**

**Gene ID** 10131

**Other Names**

Heat shock protein 75 kDa, mitochondrial, HSP 75, TNFR-associated protein 1, Tumor necrosis factor type 1 receptor-associated protein, TRAP-1, TRAP1, HSP75

**Calculated MW**

80110 MW KDa

**Application Details**

Immunohistochemistry (Paraffin-embedded Section), 0.5-1 µg/ml, Human, By Heat  
<br>Western blot, 0.1-0.5 µg/ml, Human, Rat<br>

**Subcellular Localization**

Mitochondrion . Mitochondrion inner membrane . Mitochondrion matrix .

**Tissue Specificity**

Found in skeletal muscle, liver, heart, brain, kidney, pancreas, lung, placenta and bladder. Expression is highly reduced in bladder cancer and renal cell carcinoma specimens compared to healthy tissues, but it is increased in other type of tumors. .

**Protein Name**

Heat shock protein 75 kDa, mitochondrial

**Contents**

Each vial contains 5mg BSA, 0.9mg NaCl, 0.2mg Na<sub>2</sub>HPO<sub>4</sub>, 0.05mg Na<sub>3</sub>.

**Immunogen**

E.coli-derived human TRAP1 recombinant protein (Position: A571-H704). Human TRAP1 shares

91.7% and 94% amino acid (aa) sequence identity with mouse and rat TRAP1, respectively.

**Purification**

Immunogen affinity purified.

**Cross Reactivity**

No cross reactivity with other proteins

**Storage**

**At -20°C for one year. After r°Constitution, at 4°C for one month. It°Can also be aliquotted and stored frozen at -20°C for a longer time.Avoid repeated freezing and thawing.**

**Anti-TRAP1 Picoband Antibody - Protein Information**

**Name** TRAP1

**Synonyms** HSP75, HSPC5 {ECO:0000303|PubMed:1866360

**Function**

Chaperone that expresses an ATPase activity. Involved in maintaining mitochondrial function and polarization, downstream of PINK1 and mitochondrial complex I. Is a negative regulator of mitochondrial respiration able to modulate the balance between oxidative phosphorylation and aerobic glycolysis. The impact of TRAP1 on mitochondrial respiration is probably mediated by modulation of mitochondrial SRC and inhibition of SDHA.

**Cellular Location**

Mitochondrion. Mitochondrion inner membrane Mitochondrion matrix

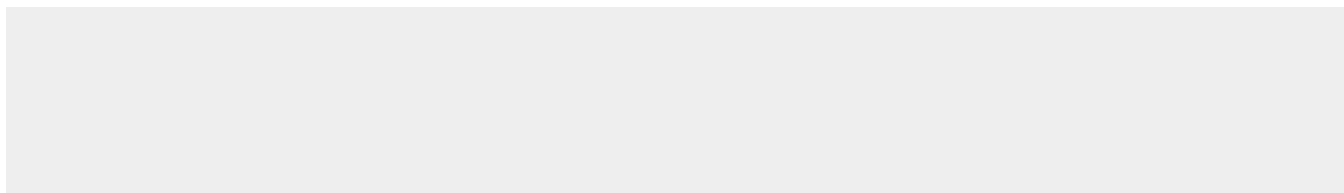
**Tissue Location**

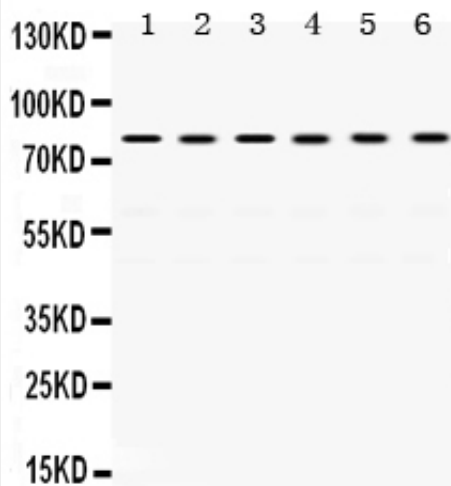
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**Anti-TRAP1 Picoband Antibody - Protocols**

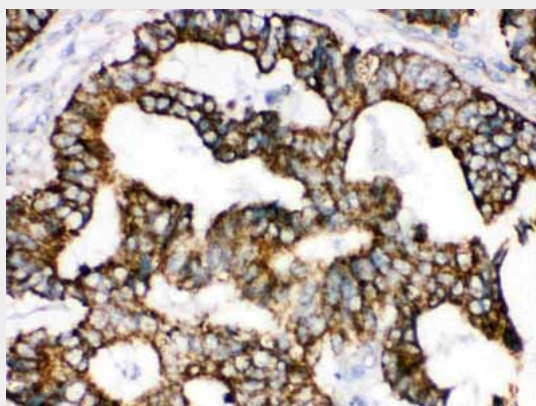
Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

**Anti-TRAP1 Picoband Antibody - Images**



Anti- TRAP1 Picoband antibody, ABO12520, Western blotting All lanes: Anti TRAP1 (ABO12520) at 0.5ug/ml  
Lane 1: Rat Cardiac Muscle Tissue Lysate at 50ug  
Lane 2: Rat Kidney Tissue Lysate at 50ug  
Lane 3: Rat Brain Tissue Lysate at 50ug  
Lane 4: SMMC Whole Cell Lysate at 40ug  
Lane 5: PANC Whole Cell Lysate at 40ug  
Lane 6: A549 Whole Cell Lysate at 40ug  
Predicted bind size: 80KD  
Observed bind size: 80KD



Anti- TRAP1 Picoband antibody, ABO12520, IHC(P) IHC(P): Human Intestinal Cancer Tissue

#### **Anti-TRAP1 Picoband Antibody - Background**

Heat shock protein 75 kDa, mitochondrial is a protein that in humans is encoded by the TRAP1 gene. It is mapped to 16p13.3. This gene encodes a mitochondrial chaperone protein that is member of the heat shock protein 90 (HSP90) family. The encoded protein has ATPase activity and interacts with tumor necrosis factor type I. And this protein may function in regulating cellular stress responses. In addition, it was found that TRAP1 interacted with the N-terminal half of TNFR1. Also, TRAP1 interacted with the C-terminal ends of the proteins encoded by both multiple exostoses-causing genes, EXT1 and EXT2, but not with EXTL1 or EXTL3.