

## KV8.2 Rabbit pAb

CatalogNo: YT2515

### Key Features

#### Host Species

- Rabbit

#### Reactivity

- Human, Mouse

#### Applications

- WB, IHC

#### MW

- 62kD (Observed)

#### Isotype

- IgG

### Recommended Dilution Ratios

WB 1:500-2000

IHC 1:50-300

### Storage

**Storage\*** -15°C to -25°C/1 year (Do not lower than -25°C)

**Formulation** Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

### Basic Information

**Clonality** Polyclonal

### Immunogen Information

**Immunogen** The antiserum was produced against synthesized peptide derived from human KCNV2. AA range: 187-236

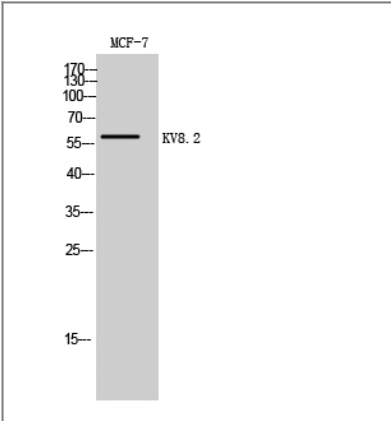
**Specificity** KV8.2 Polyclonal Antibody detects endogenous levels of KV8.2 protein.

### Target Information

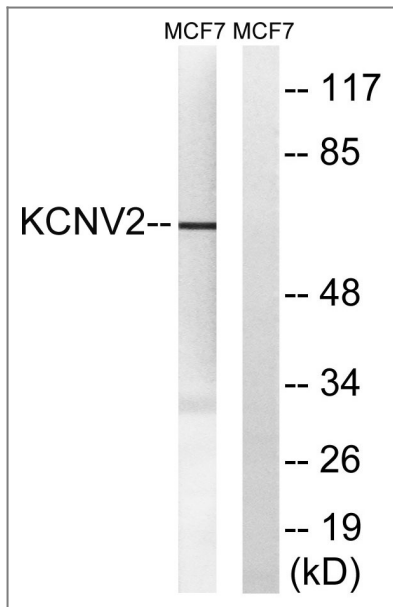
**Gene name** KCNV2

<b>Protein Name</b>	Potassium voltage-gated channel subfamily V member 2		
	<b>Organism</b>	<b>Gene ID</b>	<b>UniProt ID</b>
	Human	<a href="#">169522</a> ;	<a href="#">Q8TDN2</a> ;
	Mouse	<a href="#">240595</a> ;	<a href="#">Q8CFS6</a> ;
<b>Cellular Localization</b>	Cell membrane; Multi-pass membrane protein. Has to be associated with KCNB1 or possibly another partner to get inserted in the plasma membrane. Remains intracellular in the absence of KCNB1.		
<b>Tissue specificity</b>	Detected in lung, liver, kidney, pancreas, spleen, thymus, prostate, testis, ovary and colon.		
<b>Function</b>	<p>Disease:Defects in KCNV2 are the cause of cone dystrophy retinal type 3B (RCD3B) [MIM:610356]; also called cone dystrophy with night blindness and supernormal rod responses KCNV2-related. RCD3B is a rare form of cone dystrophy associated with supernormal rod responses. The disorder is characterized by reduced visual acuity, photoaversion, night blindness, and abnormal color vision. At an early age, the retina shows subtle depigmentation at the macula and, later, more obvious areas of atrophy.,Domain:The segment S4 is probably the voltage-sensor and is characterized by a series of positively charged amino acids at every third position.,Function:Potassium channel subunit. Modulates channel activity by shifting the threshold and the half-maximal activation to more negative values.,similarity:Belongs to the potassium channel family. V subfamily.,subcellular location:Has to be associated with KCNB1 or possibly another partner to get inserted in the plasma membrane. Remains intracellular in the absence of KCNB1.,subunit:Heteromultimer with KCNB1, KCNC1 and KCNF1. Does not form homomultimers.,tissue specificity:Detected in lung, liver, kidney, pancreas, spleen, thymus, prostate, testis, ovary and colon.,</p>		

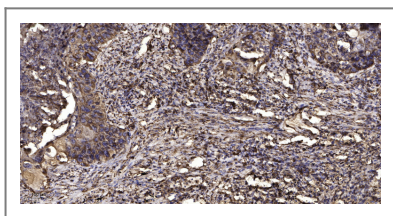
| Validation Data



Western Blot analysis of MCF-7 cells using KV8.2 Polyclonal Antibody



Western blot analysis of lysates from MCF-7 cells, using KCNV2 Antibody. The lane on the right is blocked with the synthesized peptide.



Immunohistochemical analysis of paraffin-embedded human Squamous cell carcinoma of lung. 1, Antibody was diluted at 1:200(4° overnight). 2, Tris-EDTA,pH9.0 was used for antigen retrieval. 3,Secondary antibody was diluted at 1:200(room temperature, 45min).

## | Contact information

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Please scan the QR code to access additional product information:  
**KV8.2 Rabbit pAb**

For Research Use Only. Not for Use in Diagnostic Procedures.

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