

# Rabbit Anti-C3ORF31 antibody

### SL15173R

<b>Product Name:</b>	C3ORF31
Chinese Name:	3号染色体开放阅读框31抗体
Alias:	C3orf31; Chromosome 3 open reading frame 31; MGC16471; MMP37 like protein; MMP37 like protein mitochondrial; MMP37-like protein, mitochondrial; MMP37 HUMAN.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat, Dog, Pig, Cow, Sheep,
Applications:	WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800ICC=1:100-500IF=1:100-500 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	33kDa
Cellular localization:	cytoplasmicThe cell membrane
Form:	Lyophilized or Liquid
<b>Concentration:</b>	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human C3ORF31:101-200/316
Lsotype:	IgG
Purification:	affinity purified by Protein A
Storage Buffer:	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Storage:	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized antibody is stable at room temperature for at least one month and for greater than a year when kept at -20 °C. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.
PubMed:	<u>PubMed</u>
Product Detail:	C3orf31 (chromosome 3 open reading frame 31), also known as MGC16471 or DKFZp434E0519, is a 316 amino acid mitochondrial protein that belongs to the MMP37 family and may be involved in translocation of transit peptide-containing proteins across the mitochondrial inner membrane. C3orf24 is encoded by a gene that maps to human chromosome 3p25.2. Chromosome 3 is made up of approximately 214

million bases encoding over 1,100 genes. Notably, there is a chemokine receptor gene cluster and a variety of human cancer related loci on chromosome 3. Particular regions of the chromosome 3 short arm are deleted in many types of cancer cells. Key tumor suppressing genes on chromosome 3 encode apoptosis mediator RASSF1, cell migration regulator HYAL1 and angiogenesis suppressor SEMA3B. Marfan Syndrome, porphyria, von Hippel-Lindau syndrome, osteogenesis imperfecta and Charcot-Marie-Tooth disease are a few of the numerous genetic diseases associated with chromosome 3.

#### Function:

May be involved in the translocation of transit peptide-containing proteins across the mitochondrial inner membrane (By similarity).

#### **Subcellular Location:**

Mitochondrion (By similarity).

#### Similarity:

Belongs to the MMP37/TAM41 family.

## SWISS:

Q96BW9

#### Gene ID:

132001

#### Database links:

Entrez Gene: 514330Cow

Entrez Gene: 100725702 Guinea pig

Entrez Gene: 100051402Horse

Entrez Gene: 132001Human

Entrez Gene: 362419Rat

Entrez Gene: 641323Zebrafish

Omim: 614948Human

SwissProt: Q32L81Cow

SwissProt: Q96BW9Human

SwissProt: Q3TUH1Mouse

SwissProt: Q3B7H2Zebrafish

Important No
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This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

