

## Rabbit Anti-C5orf60 antibody

## SL15212R

<b>Product Name:</b>	C5orf60
Chinese Name:	5号染色体开放阅读框60抗体
Alias:	C5orf60; CE060_HUMAN; Chromosome 5 open reading frame 60; FLJ35723; LOC285679; OTTHUMP00000223359; Putative uncharacterized protein C5orf60.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human,
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:	39kDa
Cellular localization:	The cell membrane
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human C5orf60:101-200/353
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:	With 181 million base pairs encoding around 1,000 genes, chromosome 5 is about 6% of human genomic DNA. It is associated with Cockayne syndrome through the ERCC8 gene and familial adenomatous polyposis through the adenomatous polyposis coli (APC) tumor suppressor gene. Treacher Collins syndrome is also chromosome 5 associated and is caused by insertions or deletions within the TCOF1 gene. Deletion of the p arm of chromosome 5 leads to Cri du chat syndrome. Deletion of 5q or

chromosome 5 altogether is common in therapy-related acute myelogenous leukemias and myelodysplastic syndrome. The C5orf60 gene product has been provisionally designated C5orf60 pending further characterization.

## **Subcellular Location:**

Membrane; Single-pass membrane protein (Potential).

SWISS: A6NFR6

**Gene ID:** 285679

Database links:

Entrez Gene: 285679 Human

SwissProt: A6NFR6 Human

Unigene: 558748 Human

## Important Note:

This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.