

Rabbit Anti-C2orf27 antibody

SL15148R

Product Name: C2orf27 Chinese Name: 2号染色体开放阅读框27抗体 Alias: C2orf27A; Chromosome 2 open reading frame 27A; MGC50273; OTTHUMP00000162444; Uncharacterized protein C2orf27. Organism Species: Rabbit Clonality: Polyclonal		
Alias:C2orf27A; Chromosome 2 open reading frame 27A; MGC50273; OTTHUMP00000162444; Uncharacterized protein C2orf27.Organism Species:Rabbit		
Allas: OTTHUMP00000162444; Uncharacterized protein C2orf27. Organism Species: Rabbit		
OTTHUMP00000162444; Uncharacterized protein C2orf27. Organism Species: Rabbit		
Clonality: Polyclonal		
Cionanty. I orycional		
React Species: Human,		
WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800IC0	C=1:100-	
Applications: 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: 22kDa	22kDa	
Cellular localization: The nucleusExtracellular matrix	The nucleusExtracellular matrix	
Form: Lyophilized or Liquid	Lyophilized or Liquid	
Concentration: 1mg/ml	1mg/ml	
immunogen: KLH conjugated synthetic peptide derived from human C2orf27 :51-150/202	KLH conjugated synthetic peptide derived from human C2orf27 :51-150/203	
Lsotype: IgG		
Purification: affinity purified by Protein A	affinity purified by Protein A	
Storage Buffer: 0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.	
Store at -20 癈 for one year. Avoid repeated freeze/thaw cycles. The lyophil	ized	
antibody is stable at room temperature for at least one month and for greater	than a year	
Storage: when kept at -20癈. When reconstituted in sterile pH 7.4 0.01M PBS or dilu	ent of	
antibody the antibody is stable for at least two weeks at 2-4 癈.		
PubMed: PubMed		
The second largest human chromosome, 2 consists of 237 million bases enco	oding over	
1,400 genes and making up approximately 8% of the human genome. A num		
Product Detail: genetic diseases are linked to genes on chromosome 2. Harlequin icthyosis, a		
morbid skin deformity, is associated with mutations in the ABCA12 gene. T		
metabolic disorder sitosterolemia is associated with ABCG5 and ABCG8. A		

extremely rare recessive genetic disorder, Alstr鰉 syndrome is due to mutations in the ALMS1 gene. Interestingly, chromosome 2 contains what appears to be a vestigial second centromere and vestigial telomeres which gives credence to the hypothesis that human chromosome 2 is the result of an ancient fusion of two ancestral chromosomes seen in modern form today in apes. The C2orf27 gene product has been provisionally designated C2orf27 pending further characterization.
SWISS: Q580R0
Gene ID: 29798
Database links: UniProtKB/Swiss-Prot: Q580R0.1
Important Note: This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

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