

Rabbit Anti-C2orf15 antibody

SL15146R

Product Name:	C2orf15
Chinese Name:	2号染色体开放阅
Alias:	Uncharacterized protein C2orf15; CB015 HUMAN; C2orf15.
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:	17kDa
Cellular localization:	cytoplasmic
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human C2orf15:1-100/125
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 癈 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癈. 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 癈.
PubMed:	PubMed
Product Detail:	C2orf15 is a 125 amino acid protein encoded by a gene that maps to human
	chromosome 2q11.2. As the second largest human chromosome, chromosome 2 makes
	up approximately 8% of the human genome and contains 237 million bases encoding
	over 1,400 genes. A number of genetic diseases are linked to genes on chromosome 2.
	Harlequin icthyosis, a rare skin deformity, is associated with mutations in the ABCA12
	gene. The lipid metabolic disorder sitosterolemia is associated with ABCG5 and

ABCG8. An extremely rare recessive genetic disorder, Alstr鰉 syndrome, is related to mutations in the ALMS1 gene. Chromosome 2 contains a probable vestigial second centromere as well as vestigial telomeres, which gives credence to the hypothesis that human chromosome 2 formed as a result of an ancient fusion of two ancestral chromosomes, which are still present in modern day apes.

SWISS: Q8WU43

Gene ID: 150590

Database links:

Entrez Gene: 150590 Human

SwissProt: Q8WU43 Human

Unigene: 352211 Human

Important Note:

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