

# C21orf2 Rabbit pAb

C21orf2 Rabbit pAb  
Catalog # AP59450

## Product Information

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<b>Application</b>	IHC-P, IHC-F, IF, E
<b>Primary Accession</b>	<a href="#">O43822</a>
<b>Reactivity</b>	Rat
<b>Predicted</b>	Human, Mouse
<b>Host</b>	Rabbit
<b>Clonality</b>	Polyclonal
<b>Calculated MW</b>	28340
<b>Physical State</b>	Liquid
<b>Immunogen</b>	KLH conjugated synthetic peptide derived from human C21orf2
<b>Epitope Specificity</b>	1-100/256
<b>Isotype</b>	IgG
<b>Purity</b>	affinity purified by Protein A
<b>Buffer</b>	0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.
<b>SIMILARITY</b>	Contains 3 LRR (leucine-rich) repeats. Contains 1 LRRCT domain. To <i>C.elegans</i> F09G8.5.
<b>Important Note</b>	This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.
<b>Background Descriptions</b>	The smallest of the human chromosomes, 21 makes up about 1.5% of the human genome. Chromosome 21 contains nearly 300 genes and 47 million base pairs. Down syndrome, also known as trisomy 21, is the disease most commonly associated with chromosome 21. Alzheimer's disease, Jervell and Lange-Nielsen syndrome and amyotrophic lateral sclerosis are also associated with chromosome 21. Translocations are found to occur between chromosome 21 and 8, and chromosome 21 and 12, in certain leukemias. The C21orf2 gene product has been provisionally designated C21orf2 pending further characterization.

## Additional Information

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<b>Gene ID</b>	755
<b>Other Names</b>	Cilia- and flagella-associated protein 410 {ECO:0000312 HGNC:HGNC:1260}, C21orf-HUMF09G8.5, Leucine-rich repeat-containing protein 76, YF5/A2, CFAP410 ( <a href="#">HGNC:1260</a> )
<b>Dilution</b>	IHC-P=1:100-500,IHC-F=1:100-500,ICC/IF=1:100-500,IF=1:100-500,ELISA=1:500 0-10000
<b>Storage</b>	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. 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.

## Protein Information

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<b>Name</b>	CFAP410 ( <a href="#">HGNC:1260</a> )
<b>Function</b>	Plays a role in cilia formation and/or maintenance (By similarity). Plays a role in the regulation of cell morphology and cytoskeletal organization (PubMed: <a href="#">21834987</a> ). Involved in DNA damage repair (PubMed: <a href="#">26290490</a> ).
<b>Cellular Location</b>	Mitochondrion. Cytoplasm, cytoskeleton, cilium basal body. Cell projection, cilium, photoreceptor outer segment. Cytoplasm Note=Colocalizes with NEK1 and SPATA7 at the basal body
<b>Tissue Location</b>	Widely expressed (PubMed:26974433, PubMed:9325172). Expressed in the retina (PubMed:26294103)

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Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.