

Product datasheet for AP17415PU-N

LRRC32 Rabbit Polyclonal Antibody

Product data:

OriGene Technologies, Inc.

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Product Type:	Primary Antibodies
Applications:	IHC, WB
Recommended Dilution:	Western Blot: 1/1000. Immunohistochemistry on Paraffin Sections: 1/10-1/50.
Reactivity:	Human, Mouse
Host:	Rabbit
Clonality:	Polyclonal
Immunogen:	KLH conjugated synthetic peptide between 234~260 amino acids from the Center region of Human GARP.
Specificity:	This antibody detects GARP at the center region. <i>Cellular Location</i> : Membrane; Single-pass type I membrane protein. <i>Tissue Location</i> : Preferentially expressed in regulatory T-cells (T(regs)).
Formulation:	PBS State: Purified State: Liquid purified Ig fraction Preservative: 0.09% Sodium Azide
Concentration:	lot specific
Purification:	Prepared by Saturated Ammonium Sulfate (SAS) precipitation followed by dialysis against PBS
Conjugation:	Unconjugated
Storage:	Store undiluted at 2-8°C for one month or (in aliquots) at -20 °C for longer. Avoid repeated freezing and thawing.
Stability:	Shelf life: one year from despatch.
Predicted Protein Size:	71979 Da
Gene Name:	leucine rich repeat containing 32
Database Link:	<u>Entrez Gene 434215 MouseEntrez Gene 2615 Human</u> <u>Q14392</u>



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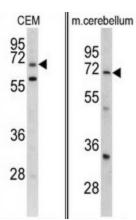
ORIGENE	LRRC32 Rabbit Polyclonal Antibody – AP17415PU-N
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Background:

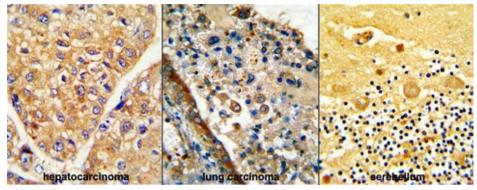
GARP, also known as leucine rich repeat containing 32, is a type I membrane protein with twenty-two leucine-rich repeats and a short intracytoplasmic region. GARP was identified based on the analysis of gene expression profiling of Treg and Th cells following TCR stimulation.

Synonyms:	Garpin, D11S833E
Protein Families:	Transmembrane

Product images:



Western blot analysis of GARP antibody (Center)) in CEM cell line (Left) and Mouse cerebellum tissue (Right) lysates (35 ug/lane). GARP (arrow) was detected using the purified Pab.



Formalin-fixed and paraffin-embedded (from Left to Right) human hepatocarcinoma, lung carcinoma and cerebellum reacted with GARP Antibody (Center), which was peroxidaseconjugated to the secondary antibody, followed by DAB staining. This data demonstrates the use of this antibody for immunohistochemistry; clinical relevance has not been evaluated.

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