

Rabbit Anti-TRIM29 (acetyl K116) antibody

SL16731R

Product Name:	TRIM29 (acetyl K116)	
Chinese Name:	乙酰化TRIM29蛋白抗体	
Alias:	Ataxia telangiectasia group D associated protein; Ataxia telangiectasia group D- associated protein; ATDC; FLJ36085; TRI29_HUMAN; TRIM 29; TRIM29; Tripartite motif containing 29; Tripartite motif containing protein 29; Tripartite motif protein 29; Tripartite motif protein TRIM29; Tripartite motif-containing protein 29.	
Organism Species:	Rabbit	
Clonality:	Polyclonal	
React Species:	Human,Pig,Cow,Rabbit,	
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:	66kDa	
Cellular localization:	cytoplasmic	
Form:	Lyophilized or Liquid	
Concentration:	1mg/ml	
immunogen:	KLH conjugated Synthesised acetylpeptide derived from human TRIM29 around the acetylation site of K116:AK(Ac-K)PP	
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:	PubMed	
Product Detail:	The protein encoded by this gene belongs to the TRIM protein family. It has multiple zinc finger motifs and a leucine zipper motif. It has been proposed to form homo- or	

heterodimers which are involved in nucleic acid binding. Thus, it may act as a transcriptional regulatory factor involved in carcinogenesis and/or differentiation. It may also function in the suppression of radiosensitivity since it is associated with ataxia telangiectasia phenotype. [provided by RefSeq, Jul 2008]

Function:

It is able to complement the radiosensitivity defect of an ataxia telangiectasia (AT) fibroblast cell line.

Subcellular Location: Cytoplasm. Colocalizes with intermediate filaments.

Tissue Specificity: Expressed in placenta, prostate and thymus.

Post-translational modifications: Constitutively phosphorylated by PKC on serine/threonine in A431 cells.

Similarity: Contains 1 B box-type zinc finger.

SWISS: Q14134

Gene ID: 23650

Database links:

Entrez Gene: 23650 Human

Entrez Gene: 525062 Cow

Entrez Gene: 72169 Mouse

Entrez Gene: 300656 Rat

<u>Omim: 610658</u> Human

SwissProt: Q14134 Human

SwissProt: Q8R2Q0 Mouse

Unigene: 504115 Human

Unigene: 273277 Mouse

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

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