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Rabbit Anti-FHAD1 antibody

SL13171R

Product Name:	FHAD1
Chinese Name:	叉头相关结构域包含蛋白1抗体
Alias:	FHA domain-containing protein 1; Forkhead-associated (FHA) phosphopeptide binding domain; Forkhead-associated domain-containing protein 1; KIAA1937; RP3-467K16.1; FHAD1_HUMAN.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human,Mouse,Rat,Cow,Rabbit,Sheep,
Applications:	ELISA=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:	162kDa
Cellular localization:	The nucleuscytoplasmic
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human FHAD1:401-500/1412
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 FHAD1 gene encodes a 1,420 amino acid protein and maps to human chromosome 1, the largest human chromosome which spans about 260 million base pairs and makes up 8% of the human genome. Other notable genes located on chromosome 1 include LMNA, which is associated with the rare aging disease Hutchinson-Gilford progeria, and the MUTYH gene, which is partially responsible for familial adenomatous

	polyposis. Stickler syndrome, Parkinsons, Gaucher disease and Usher syndrome. A breakpoint has been identified in 1q which disrupts the DISC1 gene and is linked to schizophrenia. Aberrations in chromosome 1 are found in a variety of cancers including head and neck cancer, malignant melanoma and multiple myeloma. **Similarity:** Contains 1 FHA domain. **SWISS:** B1AJZ9 **Gene ID:** 114827 **Database links:** Entrez Gene: 114827Human SwissProt: B1AJZ9Human Unigene: 659997Human **Important Note:** This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.