

KIAA1429 蛋白抗体

产品货号: mlR17003 英文名称: KIAA1429 中文名称: KIAA1429 蛋白抗体 别 名: DKFZp434l116; DKFZp781B2117; fSAP121; KIAA1429; MGC138493; MGC141940; MSTP054; Protein virilizer homolog; VIR_HUMAN. 研究领域: 细胞生物 信号转导 转录调节因子 表观遗传学 抗体来源: Rabbit 克隆类型: Polyclonal

产品应用 : ELISA=1:500-1000 IHC-P=1:400-800 IHC-F=1:400-800 ICC=1:100-500 IF=1:100-500 (石蜡切片需

交叉反应: Human, Mouse, Rat, Cow, Horse, Rabbit, Sheep, Rhesus monkey, Gorilla, Orangutan, Platypus

做抗原修复)



not yet tested in other applications.

optimal dilutions/concentrations should be determined by the end user.

分子量: 202kDa

细胞定位: 细胞核

性 状: Lyophilized or Liquid

浓 度: 1mg/ml

免疫原: KLH conjugated synthetic peptide derived from human KIAA1429:1661-1760/1812

亚 型: IgG

纯化方法: affinity purified by Protein A

储存液: 0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.

保存条件: Store at -20 $^{\circ}$ 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 $^{\circ}$ 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 $^{\circ}$ C.



PubMed: PubMed

产品介绍: Made up of nearly 146 million bases, chromosome 8 encodes about 800 genes. Translocation of portions of chromosome 8 with amplifications of the c-Myc gene are found in some leukemias and lymphomas, and typically associated with a poor prognosis. Portions of chromosome 8 have been linked to schizophrenia and bipolar disorder. Trisomy 8, also known as Warkany syndrome 2, most often results in early miscarriage but is occasionally seen in a mosaic form in surviving patients who suffer to a varying degree from a number of symptoms including retarded mental and motor development, and certain facial and developmental defects. WRN is a DNA helicase encoded by chromosome 8 and shown defective in those with the early aging disorder Werner syndrome. Chromosome 8 is also associated with Pfeiffer syndrome, congenital hypothyroidism and Waardenburg syndrome. The KIAA1429 gene product has been provisionally designated KIAA1429 pending further characterization.

unction:	
May be involved in mRNA splicing regulation.	
Subcellular Location:	
Nucleus.	

Similarity:

Belongs to the vir family.

SWISS:

Q69YN4

Gene ID:

25962



Important Note:

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