

做抗原修复)

## KIAA1712 蛋白抗体

产品货号: mlR17024 英文名称: KIAA1712 中文名称: KIAA1712 蛋白抗体 别 名: Centrosomal protein of 44 kDa; Cep44; CEP44\_HUMAN; HBV PreS1-transactivated protein 3; PS1TP3. 研究领域: 细胞生物 免疫学 抗体来源: Rabbit 克隆类型: Polyclonal 交叉反应: Human, Mouse, Dog, Horse, Rabbit, 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



not yet tested in other applications.

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

分子量: 44kDa

细胞定位: 细胞浆

性 状: Lyophilized or Liquid

浓 度: 1mg/ml

免疫原: KLH conjugated synthetic peptide derived from human KIAA1712:301-390/390

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 Representing approximately 6% of the human genome, chromosome 4 contains nearly 900 genes. Notably, the Huntingtin gene, which is found to encode an expanded glutamine tract in cases of Huntington's disease, is on chromosome 4. FGFR-3 is also encoded on chromosome 4 and has been associated with thanatophoric dwarfism, achondroplasia, Muenke syndrome and bladder cancer. Chromosome 4 is also tied to Ellis-van Creveld syndrome, methylmalonic acidemia and polycystic kidney disease. Chromosome 4 reportedly contains the largest gene deserts (regions of the genome with no protein encoding genes) and has one of the two lowest recombination frequencies of the human chromosomes. The KIAA1712 gene product has been provisionally designated KIAA1712 pending further characterization.

## **Subcellular Location:**

Cytoplasm; cytoskeleton; centrosome. Cytoplasm; cytoskeleton; spindle pole. Midbody. Localizes to centrosomes, spindle poles, and weakly in midbody.

**SWISS:** 

Q9C0F1

Gene ID:

80817

## **Important Note:**

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