

## 铜代谢结构域蛋白9 抗体

产品货号: mlR8180 英文名称: COMMD9 中文名称: 铜代谢结构域蛋白9抗体 别 名: COMD9\_HUMAN; COMM domain-containing protein 9; COMMD9; HSPC166. 研究领域: 细胞生物 通道蛋白 抗体来源: Rabbit 克隆类型: Polyclonal 交叉反应: Human, Mouse, Rat, Dog, Pig, Cow, Sheep, 产品应用: WB=1:500-2000 ELISA=1:500-1000 IHC-P=1:400-800 IHC-F=1:400-800 IF=1:100-500 (石蜡切片需

not yet tested in other applications.

做抗原修复)

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



Good elisakit producers	<b>州台州大里</b> 490

分子量: 22kDa

细胞定位: 细胞核 细胞浆

性 状: Lyophilized or Liquid

浓 度: 1mg/ml

免疫原: KLH conjugated synthetic peptide derived from human COMMD9:11-100/198

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 °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

产品介绍: COMMD9 (COMM domain containing 9) is a 198 amino acid ubiquitously expressed protein that contains one COMM domain and may be involved in signaling events throughout the cell. The gene encoding



COMMD9 maps to human chromosome 11, which houses over 1,400 genes and comprises nearly 4% of the human genome. Jervell and Lange-Nielsen syndrome, Jacobsen syndrome, Niemann-Pick disease, hereditary angioedema and Smith-Lemli-Opitz syndrome are associated with defects in genes that maps to chromosome 11.

Tissue Specificity:
Ubiquitous.
Similarity:
Contains 1 COMM domain.
SWISS:
Q9P000
Gene ID:
29099
Important Note:
This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic
applications.
产品图片



