

## 公司产品仅供科学研究实验, 不得用于临床!

### 商品详情:

英文名称: C10orf62

中文名称: **10号染色体开放阅读框62抗体**

别名: bA548K23.1; Hypothetical protein LOC414157; MERIT40; chromosome 10 open reading frame 62; NBA1; Uncharacterized protein C10orf62; C1062\_HUMAN.

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: (predicted: Human, Mouse, Rat, Pig, )

产品应用: ELISA=1:5000-10000

not yet tested in other applications.

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

理论分子量: 25kDa

细胞定位: 细胞核 细胞浆

性状: Liquid

浓度: 1mg/ml

免疫原: KLH conjugated synthetic peptide derived from human C10orf62: 22-130/223

亚型: IgG

纯化方法: affinity purified by Protein A

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

保存条件: Shipped at 4°C. Store at -20 °C for one year. Avoid repeated freeze/thaw cycles.

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

**10号染色体开放阅读框62抗体**产品介绍: C10orf62 is a 223 amino acid protein encoded by a gene that maps to human chromosome 10q24.1. Spanning nearly 135 million base pairs, chromosome 10 makes up approximately 4.5% of total DNA in cells and encodes nearly 1,200 genes. Several protein-coding genes, including those that encode for chemokines, cadherins, excision repair proteins, early growth response factors (Egrs) and fibroblast growth receptors (FGFRs), are located on chromosome 10. Defects in some of the genes that map to chromosome 10 are associated with Charcot-Marie Tooth disease, Jackson-Weiss syndrome, Usher syndrome, nonsyndromic deafness, Wolman's syndrome, Cowden syndrome, multiple endocrine neoplasia type 2 and porphyria.

