# abcam

## Product datasheet

# Recombinant Human Carbonic anhydrase 2/CA2 protein ab85484

1 References 1 图像

描述

产品名称 重组人Carbonic anhydrase 2/CA2蛋白

纯**度** > 95 % SDS-PAGE.

ab85484 was purified by using conventional chromatography.

表达系统 Escherichia coli

**蛋白长度** Full length protein

无动物成分 No

性质 Recombinant

种属 Human

序列 MSHHWGYGKH NGPEHWHKDF PIAKGERQSP

VDIDTHTAKY DPSLKPLSVS YDQATSLRIL
NNGHAFNVEF DDSQDKAVLK GGPLDGTYRL
IQFHFHWGSL DGQGSEHTVD KKKYAAELHL
VHWNTKYGDF GKAVQQPDGL AVLGIFLKVG
SAKPGLQKVV DVLDSIKTKG KSADFTNFDP
RGLLPESLDY WTYPGSLTTP PLLECVTWIV
LKEPISVSSE QVLKFRKLNF NGEGEPEELM

VDNWRPAQPL KNRQIKASFK

技术指标

Our **Abpromise guarantee** covers the use of **ab85484** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

应**用** SDS-PAGE

形式 Liquid

制备和贮存

稳定性和存储 Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw

cycles.

pH: 7.4

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#### 常规信息

### 功能 Essential for bone resorption and osteoclast differentiation (By similarity). Reversible hydration of

carbon dioxide. Can hydrates cyanamide to urea. Involved in the regulation of fluid secretion into

the anterior chamber of the eye.

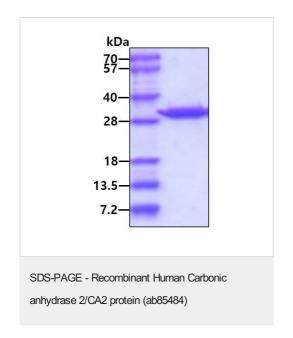
疾病相关 Defects in CA2 are the cause of osteopetrosis autosomal recessive type 3 (OPTB3)

[MIM:259730]; also known as osteopetrosis with renal tubular acidosis, carbonic anhydrase II deficiency syndrome, Guibaud-Vainsel syndrome or marble brain disease. Osteopetrosis is a rare genetic disease characterized by abnormally dense bone, due to defective resorption of immature bone. The disorder occurs in two forms: a severe autosomal recessive form occurring in utero, infancy, or childhood, and a benign autosomal dominant form occurring in adolescence or adulthood. Autosomal recessive osteopetrosis is usually associated with normal or elevated amount of non-functional osteoclasts. OPTB3 is associated with renal tubular acidosis, cerebral calcification (marble brain disease) and in some cases with mental retardation.

**序列相似性** Belongs to the alpha-carbonic anhydrase family.

细胞定位 Cytoplasm.

#### 图片



15% SDS-PAGE analysis of 3µg ab85484.

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