

Recombinant *E. coli* Carbonic anhydrase 2/CA2 protein ab87351

1 图像

描述

产品名称	重组 <i>E. coli</i> Carbonic anhydrase 2/CA2蛋白		
纯度	> 95 % SDS-PAGE. ab87351 is purified using conventional chromatography techniques.		
表达系统	Escherichia coli		
蛋白长度	Full length protein		
无动物成分	No		
性质	Recombinant		
种属	Escherichia coli		
序列	MGSSHHHHHH	SSGLVPRGSH	MKDIDTLISN
	NALWSKMLVE	EDPGFFEKLA	QAQKPRFLWI
	GCSDSRVPAE	RLTGLEPGEL	FVHRNVANLV
	IHTDLNCLSV	VQYAVDVLEV	EHIICGHYG
	CGGVQAAVEN	PELGLINNL	LHIRDIWFKH
	SLLGEMPQE	RRLDTLCELN	VMEQVYNLGH
	STIMQSAWKR	GQKVTIHGWA	YGIHDGLLRD
	LDVTATNRET	LEQRYRHGIS	NLKLKHANHK

描述

重组*E. coli* Carbonic anhydrase 2/CA2蛋白

技术指标

Our **Abpromise guarantee** covers the use of **ab87351** 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

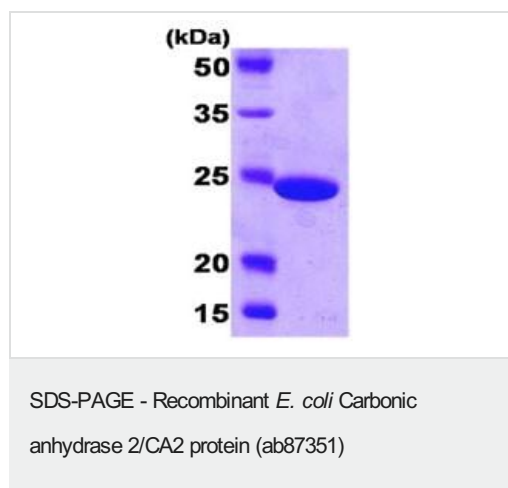
制备和贮存

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: 8.00 Constituents: 0.0154% DTT, 0.242% Tris
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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



ab87351 on 15% SDS-PAGE (3 μ g)

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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