

Product datasheet

Recombinant Human CA8 protein ab123192

1 Image

Overview

Product name	Recombinant Human CA8 protein
Protein length	Full length protein

Description

Nature	Recombinant
Source	Escherichia coli

Amino Acid Sequence

Accession [P35219](#)

Species Human

Sequence

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MGSSHHHHHH SSGLVPRGSH MGSHMADLSF
IEDTVAFPEK EEDEEEEEEG VEWGYEEGVE
WGLVFPDANG EYQSPINLNS REARYDPSLL
DVR LSPNYVV CRDCEVTNDG HTIQVILKSK
SVLSGGPLPQ GHEFELYEVR FHWGRENQRG
SEHTVNFKAF PMELHLIHWN STLFGSIDEA
VGKPHGIAII ALFVQIGKEH VGLKAVTEIL
QDIQYK GKSK TIPCFNPNTL LPDPLLRDYW
VYEGSLTIPP CSEGVTWILF RYPLTISQLQ
IEEFRRLRTH VKGAELVEGC DGILGDNFRP
TQPLSDRVIR AAFQ
    
```

Molecular weight	36 kDa including tags
Amino acids	1 to 290
Tags	His tag N-Terminus

Specifications

Our [Abpromise guarantee](#) covers the use of **ab123192** in the following tested applications.

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

Applications	Mass Spectrometry SDS-PAGE
Mass spectrometry	MALDI-TOF
Purity	> 90 % SDS-PAGE.

ab123192 is purified using conventional chromatography techniques.

Form Liquid

Preparation and Storage

Stability and Storage Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw cycles.

pH: 8.00

Constituents: 0.02% DTT, 0.32% Tris HCl, 20% Glycerol

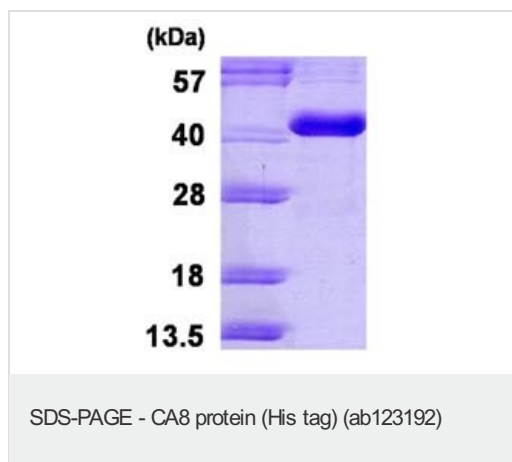
General Info

Function Does not have a carbonic anhydrase catalytic activity.

Involvement in disease Defects in CA8 are the cause of cerebellar ataxia mental retardation and dysequilibrium syndrome type 3 (CMARQ3) [MIM:613227]. CMARQ3 is a congenital cerebellar ataxia associated with dysarthria, quadripedal gait and mild mental retardation.

Sequence similarities Belongs to the alpha-carbonic anhydrase family.

Recombinant Human CA8 protein images



15% SDS-PAGE showing ab123192 (3µg).

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