abcam

Product datasheet

Recombinant Human Calreticulin 3 protein ab185419

Description

Product name Recombinant Human Calreticulin 3 protein

Purity > 95 % SDS-PAGE.

Purity is greater than 95% as determined by SEC-HPLC and reducing SDS-PAGE.

Endotoxin level < 1.000 Eu/μg
Expression system Escherichia coli

Accession Q96L12

Protein length Full length protein

Animal free No

Nature Recombinant

Species Human

Sequence MTVYFQEEFLDGEHWRNRWLQSTNDSRFGHFRLSSGKF

YGHKEKDKGLQT

TQNGRFYAISARFKPFSNKGKTLVIQYTVKHEQKMDCGGG

YIKVFPADID

QKNLNGKSQYYIMFGPDICGFDIKKVHVILHFKNKYHENKKL

IRCKVDGF

THLYTLILRPDLSYDVKIDGQSIESGSIEYDWNLTSLKKETS

PAESKDWE

QTKDNKAQDWEKHFLDASTSKQSDWNGDLDGDWPAPM

LQKPPYQDGLKPE

GIHKDVWLHRKMKNTDYLTQYDLSEFENIGAIGLELWQVR

SGTIFDNFLI

TDDEEYADNFGKATWGETKGPEREMDAIQAKEEMKKAR

EEEEEELLSGKINRHEHYFNQFHRRNEL

Predicted molecular weight 43 kDa

Amino acids 20 to 384

Specifications

Our <u>Abpromise guarantee</u> covers the use of ab185419 in the following tested applications.

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

Applications SDS-PAGE

HPLC

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Form Lyophilized

Preparation and Storage

Stability and Storage Shipped at 4°C. Store at -80°C. Avoid freeze / thaw cycle.

pH: 7.40

Constituent: 100% PBS

Reconstitution Always centrifuge tubes before opening. Do not mix by vortex or pipetting. It is not recommended

to reconstitute to a concentration less than 100 μ g/ml. Dissolve the lyophilised protein in 3X PBS. Please aliquot the reconstituted solution and store at < -20 degrees, avoid freeze/thaw cycles.

General Info

Function During spermatogenesis, may act as a lectin-independent chaperone for specific client proteins

such as ADAM3. Required for sperm fertility (By similarity). CALR3 capacity for calcium-binding

may be absent or much lower than that of CALR.

Tissue specificity Testis specific.

Involvement in diseaseDefects in CALR3 are the cause of familial hypertrophic cardiomyopathy type 19 (CMH19)

[MIM:613875]. CMH19 is a hereditary heart disorder characterized by ventricular hypertrophy, which is usually asymmetric and often involves the interventricular septum. The symptoms include dyspnea, syncope, collapse, palpitations, and chest pain. They can be readily provoked by exercise. The disorder has inter- and intrafamilial variability ranging from benign to malignant

forms with high risk of cardiac failure and sudden cardiac death.

Sequence similarities Belongs to the calreticulin family.

DomainCan be divided into a N-terminal globular domain, a proline-rich P-domain forming an elongated

arm-like structure and a C-terminal acidic domain. The P-domain binds one molecule of calcium with high affinity, whereas the acidic C-domain binds multiple calcium ions with low affinity. The interaction with glycans occurs through a binding site in the globular lectin domain.

The zinc binding sites are localized to the N-domain.

Cellular localization Endoplasmic reticulum lumen.

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