# abcam

## Product datasheet

# Recombinant Human CANT1 protein ab123189

### 1 Image

**Description** 

Product name Recombinant Human CANT1 protein

Purity > 90 % SDS-PAGE.

ab123189 is purified using conventional chromatography techniques.

**Expression system** Escherichia coli

Accession Q8WVQ1

Protein length Protein fragment

Animal free No

**Nature** Recombinant

**Species** Human

**Sequence** MGSSHHHHHH SSGLVPRGSH MGSHMRPAPG

RPPTHNAHNW RLGQAPANWY NDTYPLSPPQ

RTPAGIRYRI AVIADLDTES RAQEENTWFS YLKKGYLTLS

DSGDKVAVEW DKDHGVLESH LAEKGRGMEL
SDLIVFNGKL YSVDDRTGVV YQIEGSKAVP
WVILSDGDGT VEKGFKAEWL AVKDERLYVG
GLGKEWTTTT GDVVNENPEW VKVVGYKGSV
DHENWVSNYN ALRAAAGIQP PGYLIHESAC
WSDTLQRWFF LPRRASQERY SEKDDERKGA

NLLLSASPDF GDIAVSHVGA VVPTHGFSSF KFIPNTDDQI IVALKSEEDS GRVASYIMAF TLDGRFLLPE TKIGSVKYEG

IEFI

Predicted molecular weight 41 kDa including tags

Amino acids 63 to 401

Tags His tag N-Terminus

#### **Specifications**

Our Abpromise guarantee covers the use of ab123189 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

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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, 10% Glycerol (glycerin, glycerine), 0.29% Sodium

chloride

#### **General Info**

Function Calcium-dependent nucleotidase with a preference for UDP. The order of activity with different

substrates is UDP > GDP > UTP > GTP. Has very low activity towards ADP and even lower

activity towards ATP. Does not hydrolyze AMP and GMP.

**Tissue specificity** Widely expressed.

Involvement in disease Defects in CANT1 are the cause of Desbuquois dysplasia (DBQD) [MIM:251450]. A

chondrodysplasia characterized by severe prenatal and postnatal growth retardation (less than -5 SD), joint laxity, short extremities, progressive scoliosis, round face, midface hypoplasia, prominent bulging eyes. The main radiologic features are short long bones with metaphyseal splay, a 'Swedish key' appearance of the proximal femur (exaggerated trochanter), and advance

carpal and tarsal bone age. Two forms of Desbuquois dysplasia are distinguished on the basis of the presence (type 1) or absence (type 2) of characteristic hand anomalies: an extra ossification center distal to the second metacarpal, delta phalanx, bifid distal thumb phalanx, and phalangeal

dislocations.

**Sequence similarities** Belongs to the apyrase family.

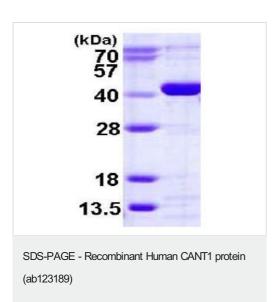
Post-translational modifications

N-glycosylated.

**Cellular localization** Endoplasmic reticulum membrane. Golgi apparatus > Golgi stack membrane. Processed form:

Secreted.

# **Images**



15% SDS-PAGE showing ab123189 (3µg).

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

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- · We investigate all quality concerns to ensure our products perform to the highest standards

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