# abcam

### Product datasheet

## Recombinant Human BTD protein (His tag) ab219238

#### 1 Image

**Description** 

Product name Recombinant Human BTD protein (His tag)

Purity > 85 % SDS-PAGE.

Affinity purified

**Endotoxin level** < 1.000 Eu/µg

Expression system Baculovirus infected insect cells

Accession P43251-3

Protein length Full length protein

Animal free No

**Nature** Recombinant

**Species** Human

Sequence AHTGESVADHHEAEYYVAAVYEHPSILSLNPLALISRQEA

LELMNQNLD

**IYEQQVMTAAQKDVQIIVFPEDGIHGFNFTRTSIYPFLDFMP** 

**SPQVVRWN** 

PCLEPHRFNDTEVLQRLSCMAIRGDMFLVANLGTKEPCH

**SSDPRCPKDGR** 

YQFNTNVVFSNNGTLVDRYRKHNLYFEAAFDVPLKVDLITF

**DTPFAGRFG** 

IFTCFDILFFDPAIRVLRDYKVKHVVYPTAWMNQLPLLAAIEI

QKAFAVA

FGINVLAANVHHPVLGMTGSGIHTPLESFWYHDMENPKSH

LIIAQVAKNP

VGLIGAENATGETDPSHSKFLKILSGDPYCEKDAQEVHCD

**EATKWNVNAP** 

PTFHSEMMYDNFTLVPVWGKEGYLHVCSNGLCCYLLYER

PTLSKELYALG

VFDGLHTVHGTYYIQVCALVRCGGLGFDTCGQEITEATGIF

**EFHLWGNFS** 

TSYIFPLFLTSGMTLEVPDQLGWENDHYFLRKSRLSSGLV

TAALYGRLYE RDLEHHHHHH

Predicted molecular weight 58 kDa including tags

Amino acids 44 to 545

Tags His tag C-Terminus

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Additional sequence information This product is the mature full length protein from aa 44 to 545. The signal peptide is not included (NP 001268652).

#### **Specifications**

Our <u>Abpromise guarantee</u> covers the use of ab219238 in the following tested applications.

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

SDS-PAGE **Applications** 

**Form** Liquid

#### **Preparation and Storage**

Stability and Storage Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

pH: 7.40

Constituents: 10% Glycerol (glycerin, glycerine), 90% PBS

#### **General Info**

**Function** Catalytic release of biotin from biocytin, the product of biotin-dependent carboxylases

degradation.

Involvement in disease Defects in BTD are the cause of biotinidase deficiency (BTD deficiency) [MIM:253260]; also

> called late-onset multiple carboxylase deficiency. BTD deficiency is a juvenile form of multiple carboxylase deficiency, an autosomal recessive disorder of biotin metabolism, characterized by ketoacidosis, hyperammonemia, excretion of abnormal organic acid metabolites, and dermatitis. BTD deficiency is characterized by seizures, hypotonia, skin rash, alopecia, ataxia, hearing loss, and optic atrophy. If untreated, symptoms usually become progressively worse, and coma and

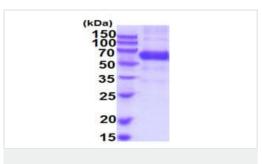
death may occur.

Sequence similarities Belongs to the CN hydrolase family. BTD/VNN subfamily.

Contains 1 CN hydrolase domain.

**Cellular localization** Secreted > extracellular space.

#### **Images**



SDS-PAGE - Recombinant Human BTD protein (His tag) (ab219238)

15% SDS-PAGE analysis of 3 µg ab219238.

Molecular weight: 50-70 kDa (SDS-PAGE under reducing conditions).

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