# abcam

# Product datasheet

# Recombinant Human Abhd5/CGI-58 protein ab132557

# 1 Image

**Description** 

Product name Recombinant Human Abhd5/CGI-58 protein

Expression system Wheat germ
Accession Q8WTS1

Protein length Full length protein

Animal free No

Nature Recombinant

**Species** Human

Sequence MAAEEEVDSADTGERSGWLTGWLPTWCPTSISHLKEA

**EEKMLKCVPCTY** 

 ${\sf KKEPVRISNGNKIWTLKFSHNISNKTPLVLLHGFGGGLGLW}$ 

ALNFGDLCT

NRPVYAFDLLGFGRSSRPRFDSDAEEVENQFVESIEEWR

CALGLDKMILL

GHNLGGFLAAAYSLKYPSRVNHLILVEPWGFPERPDLADQ

**DRPIPVWIRA** 

LGAALTPFNPLAGLRIAGPFGLSLVQRLRPDFKRKYSSMF

**EDDTVTEYIY** 

HCNVQTPSGETAFKNMTIPYGWAKRPMLQRIGKMHPDIPV

**SVIFGARSCI** 

DGNSGTSIQSLRPHSYVKTIAILGAGHYVYADQPEEFNQKV

**KEICDTVD** 

Predicted molecular weight 64 kDa including tags

Amino acids 1 to 349

Tags GST tag N-Terminus

#### **Specifications**

Our Abpromise guarantee covers the use of ab132557 in the following tested applications.

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

**Applications** ELISA

SDS-PAGE

Western blot

Form Liquid

**Additional notes** This product was previously labelled as Abhd5.

### **Preparation and Storage**

Stability and Storage Shipped on dry ice. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles.

00.8 :Hg

Constituents: 0.31% Glutathione, 0.79% Tris HCI

#### **General Info**

Function Lysophosphatidic acid acyltransferase which functions in phosphatidic acid biosynthesis. May

regulate the cellular storage of triacylglycerol through activation of the phospholipase PNPLA2.

Involved in keratinocyte differentiation.

**Tissue specificity** Widely expressed in various tissues, including lymphocytes, liver, skeletal muscle and brain.

Expressed by upper epidermal layers and dermal fibroblasts in skin, hepatocytes and neurons (at

protein level).

Involvement in disease Defects in ABHD5 are the cause of Chanarin-Dorfman syndrome (CDS) [MIM:275630]; also

called triglyceride storage disease with impaired long-chain fatty acid oxidation or neutral lipid storage disease with ichthyosis. CDS is an autosomal recessive inborn error of lipid metabolism with multisystemic accumulation of triglycerides although plasma concentrations are normal.

Clinical characteristics are congenital generalized ichthyosis, vacuolated leukocytes,

hepatomegaly, myopathy, cataracts, neurosensory hearing loss and developmental delay. The disorder presents at birth with generalized, fine, white scaling of the skin and a variable degree of

erythema resembling non-bullous congenital ichthyosiform erythroderma.

Sequence similarities Belongs to the peptidase S33 family. ABHD4/ABHD5 subfamily.

**Developmental stage** Detected in fetal epidermis from 49 to 135 days estimated gestational age (at protein level).

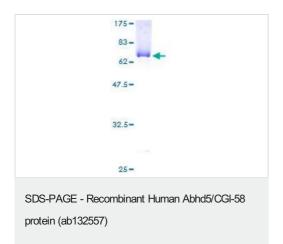
**Domain** The HXXXXD motif is essential for acyltransferase activity and may constitute the binding site for

the phosphate moiety of the glycerol-3-phosphate.

Cellular localization Cytoplasm. Lipid droplet. Colocalized with PLIN and ADRP on the surface of lipid droplets. The

localization is dependent upon the metabolic status of the adipocytes and the activity of PKA.

#### **Images**



12.5% SDS-PAGE analysis of ab132557 stained with Coomassie Blue.

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