# abcam

# Product datasheet

# Anti-LMBRD1 antibody ab105419

# 画像数3

#### 製品の概要

製品名 Anti-LMBRD1 antibody

製品の詳細 Rabbit polyclonal to LMBRD1

由来種 Rabbit

特異性 LMBRD1 antibody is predicted to not cross-react with other LMBRD1 family members.

アプリケーション 適用あり: ICC/IF, WB, IHC-P

種交差性 交差種: Human

免疫原 An 18 amino acid peptide from the C-terminus of Human LMBRD1 (NP\_060838)

ポジティブ・コントロール Human Brain Tissue Lysate

特記事項

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

#### 製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C.

**バッファー** pH: 7.2

Preservative: 0.02% Sodium azide

Constituent: PBS

精製度 Immunogen affinity purified

**ポリ/モノ** ポリクローナル

アイソタイプ IgG

アプリケーション

# アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
ICC/IF		Use a concentration of 20 µg/ml.
WB		Use a concentration of 1 - 2 µg/ml. Predicted molecular weight: 61 kDa.
IHC-P		Use at an assay dependent concentration.

#### ターゲット情報

機能 Probable lysosomal cobalamin transporter. Required to export cobalamin from lysosomes

allowing its conversion to cofactors. Isoform 3 may play a role in the assembly of hepatitis delta

virus (HDV).

組織特異性 Isoform 3 is expressed in liver.

関連疾患 Defects in LMBRD1 are the cause of methylmalonic aciduria and homocystinuria type cblF

(MMAFHC) [MIM:277380]; also known as homocystinuria-megaloblastic anemia

complementation type F. MMAFHC is a disorder of cobalamin metabolism characterized by decreased levels of the coenzymes adenosylcobalamin (AdoCbl) and methylcobalamin (MeCbl).

It is due to accumulation of free cobalamin in lysosomes, thus hindering its conversion to cofactors. Clinical features include developmental delay, stomatitis, glossitis, seizures and

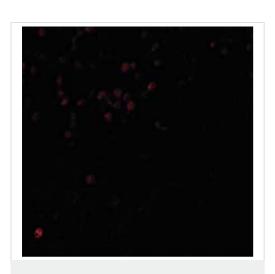
methylmalonic aciduria responsive to vitamin B12.

配列類似性 Belongs to the LIMR family. LMBRD1 subfamily.

**翻訳後修飾** N-glycosylated.

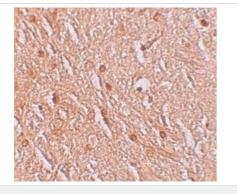
細胞内局在 Lysosome membrane.

### 画像



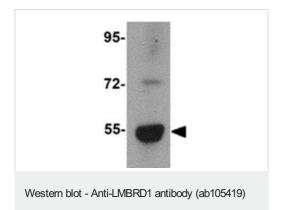
Immunocytochemistry/ Immunofluorescence - Anti-LMBRD1 antibody (ab105419)

Immunofluorescence of LMBRD1 in Human brain tissue with ab105419 at 20  $\,\mathrm{ug/mL}$ .



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-LMBRD1 antibody (ab105419)

Immunohistochemistry of LMBRD1 in Human brain tissue with ab105419 at 2.5  $\mu g/mL$ .



Anti-LMBRD1 antibody (ab105419) at 1  $\mu$ g/ml + Human brain tissue lysate at 15  $\mu$ g

Predicted band size: 61 kDa

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

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