# abcam

# Product datasheet

# Human LAMB1 (Laminin beta 1) knockout HeLa cell lysate ab257499

## 画像数3

#### 製品の概要

製品名 Human LAMB1 (Laminin beta 1) knockout HeLa cell lysate

製品の概要

Knockout cell lysate achieved by CRISPR/Cas9.

Parental Cell Line HeLa

**Organism** Human

Mutation description Knockout achieved by using CRISPR/Cas9, Insertion of the selection cassette in exon3.

Passage number <20

**Knockout validation** Sanger Sequencing, Western Blot (WB)

 $\label{eq:Reconstitution notes} \textbf{To use as WB control, resuspend the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the lyophilizate in 50 $\mu$L of LDS* Sample Buffer to have a final labeled and the labele$ 

concentration of 2 mg/ml. For reducing conditions, we recommend a final concentration of 0.1 M

DTT.

\*Usage of SDS sample buffer is not recommended with these lyophilized lysates.

特記事項

**Lysate preparation:** Our lysates are made using RIPA buffer to which we add a protease inhibitor cocktail and phosphatase inhibitor cocktail (ratio: 300:100:10). *This means that the protein of interest is denatured.* If you require a native form of the protein please use the live cell version - found **here**. Please refer to our lysis protocol for further details on how our lysates are prepared.

**User storage instructions:** Lyophilizate may be stored at 4°C. After reconstitution, store at -20°C for short-term storage or -80°C for long-term storage.

Access thousands of knockout cell lysates, generated from commonly used cancer cell lines. **See here for more information on knockout cell lysates.** 

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アプリケーション **適用あり**: WB

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#### 製品の特性

#### 保存方法

Store at -80°C. Please refer to protocols.

内容	1 kit
ab262952 - Human LAMB1 knockout HeLa cell lysate	1 x 100μg
ab255929 - Human wild-type HeLa cell lysate	1 x 100µg

Cell type epithelial

**Disease** Adenocarcinoma

**Gender** Female

**STR Analysis** Amelogenin X D5S818: 11, 12 D13S317: 12, 13.3 D7S820: 8, 12 D16S539: 9, 10 vWA: 16, 18

TH01: 7 TPOX: 8,12 CSF1PO: 9, 10

#### ターゲット情報

機能 Binding to cells via a high affinity receptor, laminin is thought to mediate the attachment, migration

and organization of cells into tissues during embryonic development by interacting with other

extracellular matrix components.

配列類似性 Contains 13 Iaminin EGF-like domains.

Contains 1 Iaminin IV type B domain. Contains 1 Iaminin N-terminal domain.

ドメイン The alpha-helical domains I and II are thought to interact with other laminin chains to form a coiled

coil structure.

Domains VI and IV are globular.

細胞内局在 Secreted > extracellular space > extracellular matrix > basement membrane. Major component.

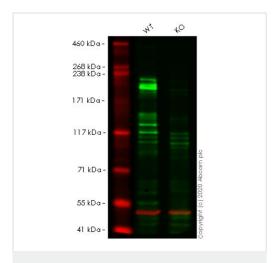
## アプリケーション

The Abpromise guarantee Abpromise保証は、次のテスト済みアプリケーションにおけるab257499の使用に適用されます

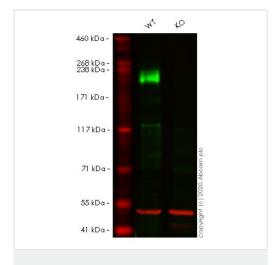
アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
WB		Use at an assay dependent concentration. Predicted molecular weight: 198 kDa.

#### 画像



Western blot - Human LAMB1 (Laminin beta 1) knockout HeLa cell lysate (ab257499)



Western blot - Human LAMB1 (Laminin beta 1) knockout HeLa cell lysate (ab257499)

Lane 1: Wild-type HeLa cell lysate (20µg)

Lane 2: LAMB1 knockout HeLa cell lysate (20µg)

**Lanes 1-2:** Merged signal (red and green). Green - <u>ab109293</u> observed at 200 kDa. Red - loading control <u>ab7291</u> observed at 50 kDa.

ab109293 Anti-Laminin beta 1 antibody [EPR3189(2)] was shown to specifically react with Laminin beta 1 in wild-type HeLa cells in western blot. Loss of signal was observed when knockout cell line ab265414 (knockout cell lysate ab257499) was used. Wild-type and Laminin beta 1 knockout samples were subjected to SDS-PAGE. Membrane was blocked for 1 hour at room temperature in 0.1% TBST with 3% non-fat dried milk. ab109293 and Anti-alpha Tubulin antibody [DM1A] - Loading Control (ab7291) were incubated overnight at 4°C at 1 in 1000 and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.

Lane 1: Wild-type HeLa cell lysate (20µg)

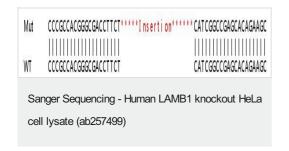
Lane 2: LAMB1 knockout HeLa cell lysate (20µg)

**Lanes 1-2:** Merged signal (red and green). Green - <u>ab108536</u> observed at 200 kDa. Red - loading control <u>ab7291</u> observed at 50 kDa.

ab108536 Anti-Laminin beta 1 antibody [EPR3189(2)] was shown to specifically react with Laminin beta 1 in wild-type HeLa cells in western blot. Loss of signal was observed when knockout cell line ab265414 (knockout cell lysate ab257499) was used. Wild-type and Laminin beta 1 knockout samples were subjected to SDS-PAGE. Membrane was blocked for 1 hour at room temperature in 0.1% TBST with 3% non-fat dried milk. ab108536 and Anti-alpha Tubulin antibody [DM1A] - Loading Control (ab7291) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit

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lgG H&L (IRDye<sup>®</sup> 800CW) preadsorbed (<u>ab216773</u>) and Goat anti-Mouse lgG H&L (IRDye<sup>®</sup> 680RD) preadsorbed (<u>ab216776</u>) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Homozygous: Insertion of the selection cassette in exon3

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