# abcam

### Product datasheet

# Human IFNGR1 knockout HeLa cell lysate ab257477

#### 画像数 2

#### 製品の概要

製品名 Human IFNGR1 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, 1 bp insertion in exon1.

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 $\mu$L of LDS* Sample$ 

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
ab262099 - Human IFNGR1 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

Receptor for interferon gamma. Two receptors bind one interferon gamma dimer.

関連疾患

Defects in IFNGR1 are a cause of mendelian susceptibility to mycobacterial disease (MSMD) [MIM:209950]; also known as familial disseminated atypical mycobacterial infection. This rare condition confers predisposition to illness caused by moderately virulent mycobacterial species,

such as Bacillus Calmette-Guerin (BCG) vaccine and environmental non-tuberculous

mycobacteria, and by the more virulent Mycobacterium tuberculosis. Other microorganisms rarely cause severe clinical disease in individuals with susceptibility to mycobacterial infections, with the

exception of Salmonella which infects less than 50% of these individuals. The pathogenic mechanism underlying MSMD is the impairment of interferon-gamma mediated immunity whose severity determines the clinical outcome. Some patients die of overwhelming mycobacterial disease with lepromatous-like lesions in early childhood, whereas others develop, later in life, disseminated but curable infections with tuberculoid granulomas. MSMD is a genetically

heterogeneous disease with autosomal recessive, autosomal dominant or X-linked inheritance.

配列類似性 Belongs to the type II cytokine receptor family.

Contains 2 fibronectin type-III domains.

Contains 2 lg-like C2-type (immunoglobulin-like) domains.

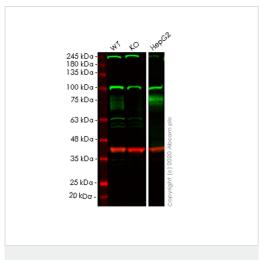
翻訳後修飾 Phosphorylated at Ser/Thr residues.

細胞内局在 Membrane.

#### アプリケーション

**The Abpromise guarantee** <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab257477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: 54 kDa.



Western blot - Human IFNGR1 knockout HeLa cell lysate (ab257477) Lane 1: Wild-type HeLa cell lysate (20 µg)

Lane 2: IFNGR1 knockout HeLa cell lysate (20 µg)

Lane 3: HepG2 cell lysate (20 µg)

**Lanes 1-3:** Merged signal (red and green). Green - <u>ab134070</u> observed at 70-95 kDa. Red - loading control <u>ab8245</u> observed at 36 kDa.

ab134070 Anti-IFNGR1 antibody [EPR7866] was shown to specifically react with IFNGR1 in wild-type HeLa cells. Loss of signal was observed when knockout cell line ab265111 (knockout cell lysate ab257477) was used. Wild-type and IFNGR1 knockout samples were subjected to SDS-PAGE. ab134070 and Anti-GAPDH antibody [6C5] - Loading Control (ab8245) were incubated overnight at 4°C at 1 in 1000 dilution 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.



Homozygous: 1 bp insertion in exon1

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