

Human TMPRSS2 knockout Caco-2 cell line ab273737

画像数 2

製品の概要

製品名	Human TMPRSS2 knockout Caco-2 cell line
Parental Cell Line	Caco 2
Organism	Human
Mutation description	Knockout achieved by using CRISPR/Cas9, Homozygous: 77bp deletion in exon 3
Passage number	<20
Knockout validation	Sanger Sequencing, Western Blot (WB)
Biosafety level	1
特記事項	<p>Cryopreservation cell medium: Cell Freezing Medium-DMSO Serum free media, contains 8.7% DMSO in MEM supplemented with methyl cellulose.</p> <p>Culture medium: EMEM + 20% FBS</p> <p>Initial handling guidelines: Upon arrival, the vial should be stored in liquid nitrogen vapor phase and not at -80°C. Storage at -80°C may result in loss of viability.</p> <ol style="list-style-type: none"> 1. Thaw the vial in 37°C water bath for approximately 1-2 minutes. 2. Transfer the cell suspension (0.8 mL) to a 15 mL/50 mL conical sterile polypropylene centrifuge tube containing 8.4 mL pre-warmed culture medium, wash vial with an additional 0.8 mL culture medium (total volume 10 mL) to collect remaining cells, and centrifuge at 201 x g (rcf) for 5 minutes at room temperature. 10 mL represents minimum recommended dilution. 20 mL represents maximum recommended dilution. 3. Resuspend the cell pellet in 5 mL pre-warmed culture medium and count using a haemocytometer or alternative cell counting method. Based on cell count, seed cells in an appropriate cell culture flask at a density of 1×10^4 cells/cm². Seeding density is given as a guide only and should be scaled to align with individual lab schedules. 4. Incubate the culture at 37°C incubator with 5% CO₂. Cultures should be monitored daily. <p>Subculture guidelines:</p> <p>All seeding densities should be based on cell counts gained by established methods. A guide seeding density of 1×10^4 cells/cm² is recommended.</p> <p>A partial media change 24 hours prior to subculture may be helpful to encourage growth, if required.</p> <p>Cells should be passaged when they have achieved 80-90% confluence (approx 8×10^4-1×10^5 cells/cm²).</p>

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We will provide viable cells that proliferate on revival.

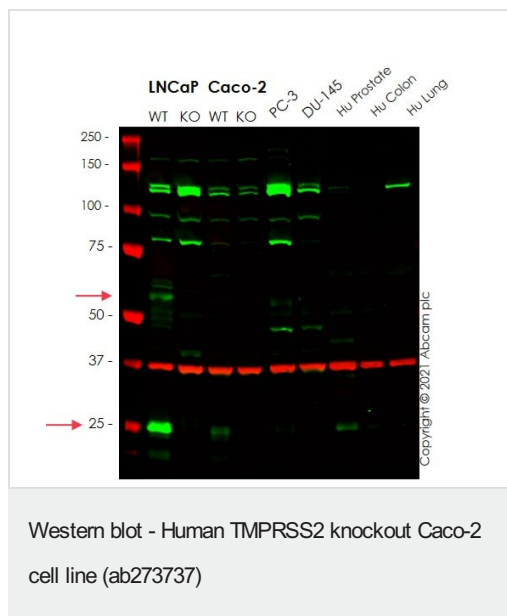
製品の特性

Number of cells	1 x 10 ⁶ cells/vial, 1 mL
Adherent /Suspension	Adherent
Tissue	Colon
Cell type	epithelial
Disease	Adenocarcinoma
Gender	Male
Mycoplasma free	Yes
保存方法	Shipped on Dry Ice. Store in liquid nitrogen.
バッファー	Constituents: 8.7% Dimethylsulfoxide, 2% Cellulose, methyl ether

ターゲット情報

組織特異性	Expressed strongly in small intestine. Also expressed in prostate, colon, stomach and salivary gland.
配列類似性	Belongs to the peptidase S1 family. Contains 1 LDL-receptor class A domain. Contains 1 peptidase S1 domain. Contains 1 SRCR domain.
細胞内局在	Cell membrane and Secreted. Activated by cleavage and secreted.

画像



All lanes : Anti-TMPRSS2 antibody [EPR3862] ([ab109131](#)) at 1/2000 dilution

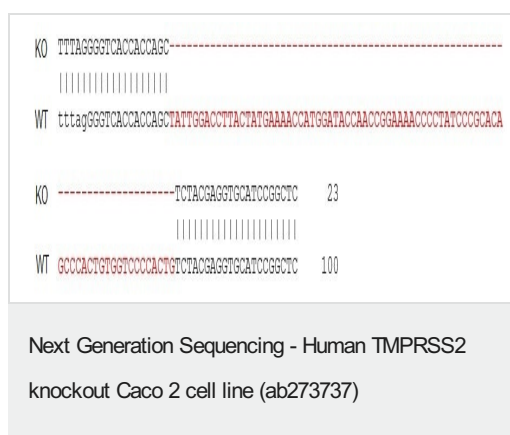
- Lane 1 :** Wild-type LNCaP cell lysate
- Lane 2 :** TMPRSS2 knockout LNCaP cell lysate
- Lane 3 :** Wild-type Caco-2 cell lysate
- Lane 4 :** TMPRSS2 knockout Caco-2 cell lysate
- Lane 5 :** PC-3 cell lysate
- Lane 6 :** DU 145 cell lysate
- Lane 7 :** Human Prostate cell lysate
- Lane 8 :** Human Colon cell lysate
- Lane 9 :** Human Lung cell lysate

Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Observed band size: 55,25 kDa

False colour image of Western blot: Anti-TMPRSS2 antibody [EPR3862] staining at 1/2000 dilution, shown in green; Mouse anti-GAPDH antibody [6C5] ([ab8245](#)) loading control staining at 1/20000 dilution, shown in red. In Western blot, [ab109131](#) was shown to bind specifically to TMPRSS2. A band was observed at 55, 25 kDa in wild-type LNCaP and at 25 kDa in Caco-2 cell lysates with no signal observed at this size in TMPRSS2 knockout LNCaP cell line [ab273745](#) (knockout LNCaP cell lysate [ab275499](#)) and TMPRSS2 knockout Caco-2 cell line [ab273737](#) (knockout Caco-2 cell lysate [ab277340](#)). To generate this image, wild-type and TMPRSS2 knockout cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 5 % milk in TBS-0.1 % Tween® 20 (TBS-T) before incubation with primary antibodies overnight at 4 °C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged. Secondary antibodies used were Goat anti-Rabbit IgG H&L (IRDye® 800CW) preabsorbed ([ab216773](#)) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed ([ab216776](#)) at 1/20000 dilution.



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