abcam

Product datasheet

Human RAB20 knockout HeLa cell line ab265924

3 Images

Overview

Product name Human RAB20 knockout HeLa cell line

Parental Cell Line HeLa
Organism Human

Mutation description Knockout achieved by using CRISPR/Cas9, 22 bp deletion in exon 1 and 28 bp deletion in exon 1

and 34 bp deletion in exon 1

Passage number <20

Knockout validation Sanger Sequencing

Biosafety level 2

General notes Recommended control: Human wild-type HeLa cell line (ab255928). Please note a wild-type

cell line is not automatically included with a knockout cell line order, if required please add recommended wild-type cell line at no additional cost using the code WILDTYPE-TMTK1.

Cryopreservation cell medium: Cell Freezing Medium-DMSO Serum free media, contains 8.7% DMSO in MEM supplemented with methyl cellulose.

Culture medium: DMEM (High Glucose) + 10% FBS

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.

- 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 2x10⁴ 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% $\rm CO_2$. Cultures should be monitored daily.

Subculture guidelines:

All seeding densities should be based on cell counts gained by established methods. A guide seeding density of 2x10⁴ cells/cm² is recommended.

A partial media change 24 hours prior to subculture may be helpful to encourage growth, if required.

1

Cells should be passaged when they have achieved 80-90% confluence.

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We will provide viable cells that proliferate on revival.

Properties

Number of cells 1 x 10⁶ cells/vial, 1 mL

Adherent /Suspension Adherent
Tissue Cervix
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

Mycoplasma free Yes

Storage instructions Shipped on Dry Ice. Store in liquid nitrogen.

Storage buffer Constituents: 8.7% Dimethylsulfoxide, 2% Cellulose, methyl ether

Target

Function Plays a role in apical endocytosis/recycling. Plays a role in the maturation and acidification of

phagosomes that engulf pathogens, such as S.aureus and M.tuberculosis. Plays a role in the

fusion of phagosomes with lysosomes.

Tissue specificity Low or absent expression in normal pancreas and stronger expression in 15 of 18 exocrine

pancreatic adenocarcinomas (at protein level).

Sequence similaritiesBelongs to the small GTPase superfamily. Rab family.

Cellular localization Golgi apparatus. Cytoplasmic vesicle, phagosome. Cytoplasmic vesicle, phagosome membrane.

Highly enriched on apical endocytic structures in polarized epithelial cells of kidney proximal tubules (By similarity). Recruited to phagosomes containing S.aureus or M.tuberculosis

(PubMed:21255211).

Images

Allele-1: 28 bp deletion in exon 1.

Mut	GGGAAGACGTCGCTGCTGCAGCGG	ACACGGT CAGCACG
WT	GGGAAGACGTCGCTGCTGCAGCGGTATATGGAGC	CGGCGCTTCCCGGACACGGTCAGCACG
Sanger Sequencing - Human RAB20 knockout HeLa		
Sanger Sequencing - Human RABZO knockout HeLa		
cell line (ab265924)		

Allele-2: 22 bp deletion in exon 1.

Mut	GGGAAGACGTCGCTGCTGCAGCGGCG	
WT	GGGAAGACGTCGCTGCTGCAGCGGTATATGGAGCGGCGCTTCCCGGACACGGTCAGCACG	
Sanger Sequencing - Human RAB20 knockout HeLa		
cell line (ab265924)		

Allele-3: 34 bp deletion in exon 1.

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