

Product Sheet

H_MICA*001 Luciferase B16-F10 Cell Line

Catalog number: GM-C30325

Version 3.3.1.250108

Description	H_MICA*001 Luciferase B16-F10 Cell Line is a clonal stable B16-F10 cell line that constitutively expresses the human MICA*001 gene, constructed using lentiviral technology.
Quantity	5E6 Cells per vial, 1 mL
Product Format	3 vials of frozen cells
Shipping	Shipped on dry ice
Storage Conditions	Liquid nitrogen immediately upon receipt
Target	Human_MICA*001
Gene ID/Uniprot ID	Q29983
Host Cell	B16-F10
Recovery Medium	DMEM+10% FBS+1% P.S
Growth medium	DMEM+10% FBS+1% P.S+10 µg/mL Blasticidin+0.5 µg/mL Puromycin
Note	None
Freezing Medium	90% FBS+10% DMSO
Growth properties	Adherent
Growth Conditions	37°C, 5% CO ₂
Mycoplasma Testing	The cell line has been screened to confirm the absence of Mycoplasma species.
Safety considerations	Biosafety Level 2
Note	It is recommended to expand the cell culture and store a minimum of 10 vials at an early passage for potential future use.

Materials

Reagent	Manufacturer/Catalogue No.
DMEM	Gibco/C11995500BT
Fetal Bovine Serum	Cegrogen biotech/A0500-3010
Pen/Strep	Thermo/15140-122
Blasticidin	Genomeditech/GM-040404
Puromycin	Genomeditech/GM-040401
Anti-MICA/MICB hIgG1 Antibody(36 NF G236A)	Genomeditech/GM-48843AB
GMOne-Step Luciferase Reporter Gene Assay Kit	Genomeditech/GM-040503

Figures

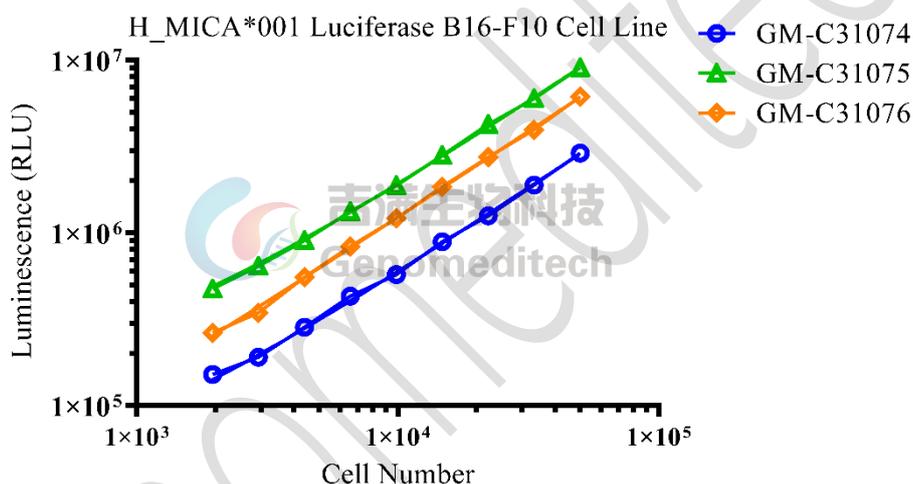


Figure 1 | Correlation between the number of cells and bioluminescence values. Serial dilutions of H_MICA*001 Luciferase B16-F10 Cell Line (Cat. GM-C30325) (96-well format). The firefly luciferase activity was measured using the GMOne-Step Luciferase Reporter Gene Assay Kit (Cat. [GM-040503](#)).

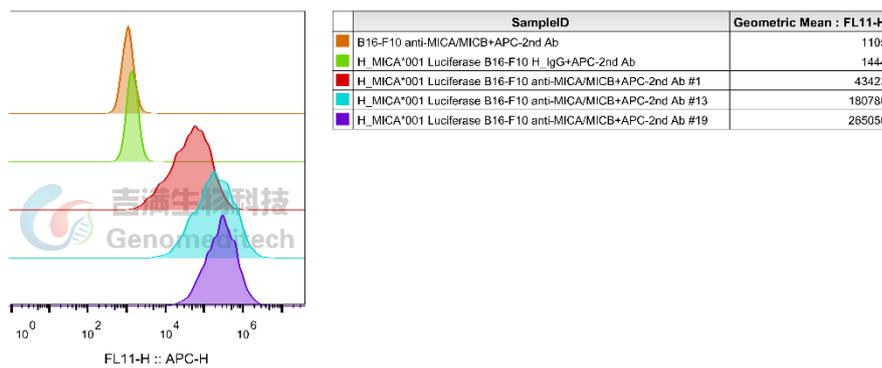


Figure 2 | H_MICA*001 Luciferase B16-F10 Cell Line (Cat. GM-C30325) was determined by flow cytometry using Anti-MICA/MICB hlgG1 Antibody(36 NF G236A) (Cat. [GM-48843AB](#)).

Cell Recovery

Recovery Medium: DMEM+10% FBS+1% P.S

To insure the highest level of viability, thaw the vial and initiate the culture as soon as possible upon receipt. If upon arrival, continued storage of the frozen culture is necessary, it should be stored in liquid nitrogen vapor phase and not at -70°C . Storage at -70°C will result in loss of viability.

- Thaw the vial by gentle agitation in a 37°C water bath. To reduce the possibility of contamination, keep the O-ring and cap out of the water. Thawing should be rapid (approximately 2 - 3 minutes).
- Remove the vial from the water bath as soon as the contents are thawed, and decontaminate by dipping in or spraying with 70% ethanol. All of the operations from this point on should be carried out under strict aseptic conditions.
- Transfer the vial contents to a centrifuge tube containing 5.0 mL complete culture medium and spin at approximately $176 \times g$ for 5 minutes. Discard supernatant.
- Resuspend cell pellet with the recommended recovery medium. And dispense into appropriate culture dishes.
- Incubate the culture at 37°C in a suitable incubator. A 5% CO_2 in air atmosphere is recommended if using the medium described on this product sheet.

Cell Freezing

Freezing Medium: 90% FBS+10% DMSO

- Centrifuge at $176 \times g$ for 3 minutes to collect cells.
- Resuspend the cells in pre-cooled freezing medium and adjust the cell density to 5×10^6 cells/mL.
- Aliquot 1 mL into each vial.
- Place the vial in a controlled-rate freezing container and store at -80°C for at least 1 day, then transfer to liquid nitrogen as soon as possible.

Cell passage

Growth medium: DMEM+10% FBS+1% P.S+10 µg/mL Blasticidin+0.5 µg/mL Puromycin

For the first 1 to 2 passages post-resuscitation, use the recovery medium. Once the cells have stabilized, switch to a growth medium.

- Remove and discard culture medium.
- Briefly rinse the cell layer with PBS to remove all traces of serum that contains trypsin inhibitor.
- Add 1.0 mL of 0.25% (w/v) Trypsin-EDTA solution to dish and observe cells under an inverted microscope until cell layer is dispersed (usually within 30 to 60 seconds at 37°C).
- Note: To avoid clumping do not agitate the cells by hitting or shaking the flask while waiting for the cells to detach. Cells that are difficult to detach may be placed at 37°C to facilitate dispersal.
- Add 2.0 mL of growth medium to mix well and aspirate cells by gently pipetting.
- After centrifugation, resuspend the pellet and add appropriate aliquots of the cell suspension to new culture vessels.
- Incubate cultures at 37°C.

Subcultivation Ratio: A subcultivation ratio of 1:3 - 1:4 is recommended

Medium Renewal: Every 2 to 3 days

Notes

- If small black dots appear inside the cells, particles are present in the intercellular spaces, or a few dead cells are floating in the medium during the cultivation process, these are all common phenomena in cell culture and do not affect cell proliferation. The cell pellet after centrifugation may also appear black.

Sequence

MICA*001 [Q29983-1](#)

MGLGPVFLLLAGIFPFAPPGAAAEPHSLRYNLTVLSWDGVSQSGFLTEVHLDGQPFLRCRQKCRAPQGG
 WAEDVLGNKTWDRETRDLTGNGKDLRMTLAHIKDKQEGHLSLQEIRVCEIHEDNSTRSSQHFYDYGELFLS
 QNLETKEWTMPQSSRAQTLAMNVRNFLKEDAMKTKTHYHAMHADCLQELRRYLKSGVVLRRTPPMVNV
 TRSEASEGNITVTCRASGFYPWNITLSWRQDGVLSHDTQQWGDVLPDGNQTYQTWVATRICQGEEQRFTC
 YMEHSGNHSTHPVPSGKVLVLQSHWQTFHVSAAVAIAAIFVIIIIFYVRCCKKKTSAAEGPELVSLQVLDQHPV
 GTSDHRDATQLGFQPLMSDLGSTGSTEGA

Related Products

MICA;MICB	
Cynomolgus_MICA(AAO24115) CHO-K1 Cell Line	Cynomolgus_MICA(Q2MGE0-1) CHO-K1 Cell Line
Cynomolgus_MICB CHO-K1 Cell Line	H_MICA CHO-K1 Cell Line
H_MICA HEK-293 Cell Line	H_MICA*001 MC38 Cell Line
H_MICA*008 CHO-K1 Cell Line	H_MICB CHO-K1 Cell Line
H_MICB HEK-293 Cell Line	

Anti-MICA/MICB hIgG1 Antibody(36 NF G236A)	Anti-MICA/MICB mIgG2a Antibody(7C6)
Anti-MICA/MICB mIgG2a Antibody(PDI-01)	
NKG2D	
H_NKG2D Blockade Reporter Jurkat Cell Line	Cynomolgus_NKG2D CHO-K1 Cell Line
H_NKG2D CHO-K1 Cell Line	H_NKG2D HEK-293 Cell Line
Anti-H_KLRK1(NKG2D) hIgG4 Antibody(Tesnatilimab)	Anti-NKG2D hIgG1 Antibody(A49MI)
In Vivo MAb Isotype Controls	
Human IgG1 Isotype Control(Anti-HEL)	Human IgG1 Isotype Control(Anti-MOPC-21)
Human IgG1 Isotype Control(Anti-RSV)	Human IgG1(LALA) Isotype Control(Anti-HEL)
Human IgG1(LALAPG) Isotype Control(Anti-HEL)	Human IgG1(N297A) Isotype Control(Anti-HEL)
Human IgG4(S228P) Isotype Control(Anti-HEL)	Mouse IgG1 Isotype Control(Anti-HEL)
Mouse IgG2a Isotype Control(Anti-HEL)	Mouse IgG2a Isotype Control(Anti-RSV)
Mouse IgG2a(D265A) Isotype Control(Anti-HEL)	

Limited Use License Agreement

Genomeditech (Shanghai) Co., Ltd grants to the Licensee all intellectual property rights, exclusive, non-transferable, and non-sublicensable rights of the Licensed Materials; Genomeditech (Shanghai) Co., Ltd will retain ownership of the Licensed Materials, cell line history packages, progeny, and the Licensed Materials including modified materials.

Between Genomeditech (Shanghai) Co., Ltd, and Licensee, Licensee is not permitted to modify cell lines in any way. The Licensee shall not share, distribute, sell, sublicense, or otherwise provide the Licensed Materials, or progenitors to third parties such as laboratories, departments, research institutions, hospitals, universities, or biotechnology companies for use other than for the purpose of outsourcing the Licensee's research.

Please refer to the Genomeditech Cell Line License Agreement for details.