

# Product Sheet

## Luciferase LNCaP Cell Line

Catalog number: GM-C44045

Version 3.3.1.260512

<b>Description</b>	Luciferase LNCaP Cell Line is a clonal stable LNCaP cell line that constitutively expresses the Luciferase gene, constructed using lentiviral technology.
<b>Quantity</b>	5E6 Cells per vial, 1 mL
<b>Product Format</b>	1 vials of frozen cells
<b>Shipping</b>	Shipped on dry ice
<b>Storage Conditions</b>	Liquid nitrogen immediately upon receipt
<b>Target</b>	/
<b>Gene ID/Uniprot ID</b>	/
<b>Host Cell</b>	LNCaP Cell Line
<b>Recovery Medium</b>	RPMI 1640+10% FBS+1% P.S
<b>Growth medium</b>	RPMI 1640+10% FBS+1% P.S+2.5 µg/mL Puromycin
<b>Note</b>	None
<b>Freezing Medium</b>	90% FBS+10% DMSO
<b>Growth properties</b>	Adherent
<b>Growth Conditions</b>	37°C, 5% CO <sub>2</sub>
<b>Mycoplasma Testing</b>	The cell line has been screened to confirm the absence of Mycoplasma species.
<b>Safety considerations</b>	Biosafety Level 2
<b>Note</b>	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.
Puromycin	Genomeditech/GM-040401
Pen/Strep	Thermo/15140-122
Fetal Bovine Serum	ExCell/FSP500
RPMI 1640	gibco/C11875500BT
GMOne-Step 2.0 Luciferase Reporter Gene Assay Kit	Genomeditech/GM-040513

## Figures

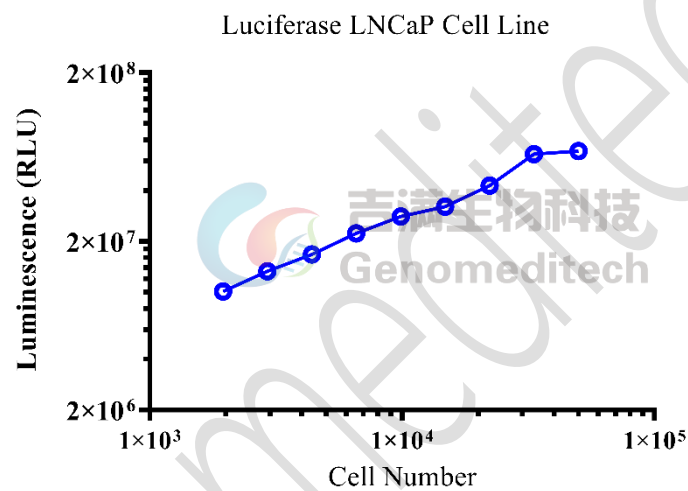


Figure 1 | Correlation between the number of cells and bioluminescence values. Serial dilutions of Luciferase LNCaP Cell Line (Cat. GM-C44045) (96-well format). The firefly luciferase activity was measured using the GMOne-Step 2.0 Luciferase Reporter Gene Assay Kit (Cat. GM-040513).

## Cell Recovery

Recovery Medium: RPMI 1640+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.

- c) Transfer the vial contents to a centrifuge tube containing 5.0 mL complete culture medium. And spin at approximately 176 x g for 5 minutes. Discard supernatant.
- d) Resuspend cell pellet with the recommended complete medium. And dispense the suspension into an appropriate culture flask and initially place the flask in an upright position after thawing.
- e) Incubate the culture at 37°C in a suitable incubator. A 5% CO<sub>2</sub> in air atmosphere is recommended if using the medium described on this product sheet.

## Cell Freezing

Freezing Medium: 90% FBS+10% DMSO

- a) Centrifuge at 176 x g for 3 minutes to collect cells.
- b) Resuspend the cells in pre-cooled freezing medium and adjust the cell density to 5E6 cells/mL.
- c) Aliquot 1 mL into each vial.
- d) 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: RPMI 1640+10% FBS+1% P.S+2.5 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.

- a) Subculturing is necessary when the cell density reaches 80%. It is recommended to perform subculturing at a ratio of 1:2 every 4-5 days. Ensure that the density does not exceed 90%, as overcrowding can lead to reduced viability due to compression.
- b) Remove and discard culture medium.
- c) Briefly rinse the cell layer with PBS to remove all traces of serum that contains trypsin inhibitor.
- d) 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 3 to 5 minutes at 37°C).
- e) 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.
- f) Add 2.0 mL of growth medium to mix well and aspirate cells by gently pipetting.
- g) After centrifugation, resuspend the pellet and add appropriate aliquots of the cell suspension to new culture vessels.
- h) Incubate cultures at 37°C.

**Subcultivation Ratio: A subcultivation ratio of 1:2 is recommended**

**Medium Renewal: Every 4 to 5 days**

## Notes

- a) Cells immediately after thawing typically contain a certain proportion of dead cells, which is a normal phenomenon. With appropriate adjustment and recovery, cell viability generally improves markedly. Once the culture stabilizes, the proportion of dead cells decreases after subsequent passages, and the cell growth rate becomes more consistent.

- b) Standard tissue culture (TC)-treated culture flasks or dishes may not provide optimal cell adhesion, which can increase the difficulty of cell culture. It is therefore recommended to pre-coat the culture surface with poly-L-lysine solution to enhance cell attachment.
- c) FBS should be heat-inactivated at 56 °C for 30 minutes in a water bath. This process inactivates complement and some viruses, while generally having minimal impact on the activity of most growth factors and cytokines.

## Related Products

Labeled Cells	
<a href="#">Luciferase-GFP MCF-7 Cell Line</a>	<a href="#">Luciferase A498 Cell Line</a>
<a href="#">Luciferase B16-F10 Cell Line</a>	<a href="#">Luciferase Daudi Cell Line</a>
<a href="#">Luciferase HEK-293 Cell Line</a>	<a href="#">Luciferase HL-60 Cell Line</a>
<a href="#">Luciferase Jurkat Cell Line</a>	<a href="#">Luciferase MIA PaCa-2 Cell Line</a>
<a href="#">Luciferase MM.1R Cell Line</a>	<a href="#">Luciferase NCI-H929 Cell Line</a>
<a href="#">Luciferase OVCAR3 Cell Line</a>	<a href="#">Luciferase RAMOS Cell Line</a>
<a href="#">Luciferase U-937 Cell Line</a>	<a href="#">Luciferase-ZsGreen1 K562 Cell Line</a>
<a href="#">Luciferase-ZsGreen1 Raji Cell Line</a>	
<a href="#">D-Luciferin, Potassium Salt</a>	<a href="#">D-Luciferin, Sodium Salt</a>

## License Agreement:

**By purchasing and using this cell line product, the user voluntarily agrees to accept and abide by the following policies:**

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