

#### **Product Description**

Product Name: Nalm6-Fluc-Puro/HuCD19-Puro (Mid) Catalog Number: CL183 Lot Number: IMP036

Species:	Human ( <i>Homo sapiens</i> )
Tissues:	Peripheral blood
Cell type:	Lymphoma
Parental cells:	Nalm6 (ATCC® CRL-3273 <sup>™</sup> )*
Morphology:	Lymphocyte-like
Growth mode:	Suspension
Reporter gene:	Firefly luciferase (Fluc)
Transgene:	CD19
Selection gene:	Puromycin (Puro)
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This is a cell line derived from the human B cell precursor leukemia Nalm6 cell line (ATCC® CRL-3273<sup>TM</sup>). Imanis Nalm6-Fluc-Puro cells (Imanis #CL151) were transduced with a lentiviral vector encoding the CD19 cDNA and the puromycin resistance gene. A moderate CD19 expressing population was generated by selection using a semi-solid methyl-cellulose-based medium. The lentiviral vectors are self-inactivating (SIN) vectors in which the viral enhancer and promoter have been deleted. Transcription inactivation of the LTR in the SIN provirus increases biosafety by preventing mobilization by replication competent viruses and enables regulated expression of the genes from the internal promoters without *cis*-acting effects of the LTR (Miyoshi et al. J Virol. 1998. 72:8150-8157).

\* The ATCC trademark and any and all ATCC catalog numbers are trademarks of the American Type Culture Collection

## **Mycoplasma Testing**

This cell line has been tested for mycoplasma contamination and is mycoplasma free.

## **Cell Line Authentication**

Authentication of the parental Nalm6 cell line was confirmed by STR profiling.

#### **Recommended Uses**

These cells are suitable for in vitro and in vivo experimentation.

The Fluc transgene facilitates *in vivo* noninvasive bioluminescent imaging of implanted cells.

## References

<sup>1</sup>Miyoshi et al. J Virol. 1998. 72:8150-8157.

## **Biosafety Notice**

This cell line was generated by transduction with a lentiviral vector. Cell lines transduced with lentiviral vectors are classified as biosafety level 2 reagents and should be used under appropriate biosafety level for institutional guidelines.

#### Storage Instructions

Remove cells from the dry ice packaging and immediately store in the vapor phase above liquid nitrogen (below -130°C).

#### **Complete Growth Medium**

RPMI-1640 Medium (RPMI) supplemented with 10 mM HEPES 10% Fetal Bovine Serum (FBS) 1% Penicillin/Streptomycin 1 µg/mL Puromycin

Puromycin should <u>NOT</u> be added to the medium until a culture has been well established from the thawed cells (about 1 week). It is also recommended that a backup frozen cell stock be generated (see below) before adding puromycin to the growth medium.

Caution! Typical commercial puromycin stocks are provided at a concentration of 10 mg/mL or 10,000X.

## **Thawing Instructions**

- 1. Thaw cells by gently swirling in a 37°C water bath. To limit contamination, do not submerge the O-ring and cap.
- When cells are ~70% thawed (less than 1 min), remove the vial and wipe down with 70% ethanol. Allow tube to dry completely.
- In a biosafety cabinet, transfer the cells into a 15 mL conical tube containing 5 mL of complete growth medium. Centrifuge cells at ~300 x g for 3-5 min.
- 4. Remove supernatant and resuspend cells in 1 mL complete growth medium. Remove an aliquot for counting.
- Dilute the cells further with growth medium to achieve a final density of 1 x 10<sup>6</sup> cells/mL. Transfer the cells to a T25 or T75 flask based on volume.
- Incubate the culture at 37°C with 5% CO<sub>2</sub>. Cells should reach full confluency 3 days after thawing.
  - Note: For optimal cell health and viability, change media every  $\sim$  3 days even if the cells aren't confluent.

## **Subculturing Instructions**

The cells should be subcultured through dilution in fresh complete growth medium as needed to maintain a density between  $7 \times 10^5$  and  $3 \times 10^6$  cells/mL. As needed, passage using centrifugation as described below to limit the amount of debris in cultures.

- Pipet the cell suspension gently to dislodge any cells loosely attached to the culture flask. Transfer the desired volume (half, one fifth, etc.) of the cells to a conical tube.
- 2. Centrifuge at ~150 x g for 3 min. (Note: a short, low speed spin is recommended to limit the amount of cell debris in the pellet.)
- 3. Remove supernatant and resuspend cells in complete growth medium. Transfer to an appropriately sized flask.

## **Freezing Medium**

These cells can be amplified and used to generate additional frozen stocks. Preparation of low passage frozen stocks is highly recommended. Frozen stocks should be preserved in a designated cryopreservation medium or in complete growth medium without puromycin supplemented with 5-10% DMSO.



# Nalm6-Fluc-Puro/HuCD19-Puro (Mid)

#### **Certificate of Analysis**

Testing performed by Imanis Life Sciences

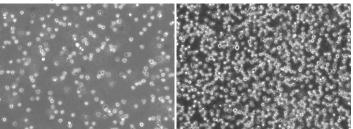
Test description	Result
Post thaw viable cell recovery	87%
Cells per vial	~ 8.5 x 10 <sup>6</sup>
Sterility	No contamination detected
Mycoplasma	No contamination detected
Puromycin selection	Pass QC
Luciferase expression	Pass QC
Human CD19 expression	Pass QC
Average doubling time	40.6 h*

\*Doubling time represents the average doubling time during <u>logarithmic growth</u>. This value should be used for general estimation only.

# Morphology

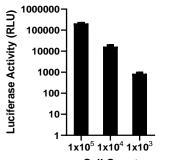
Low density, 200X

High density, 200X



Low- and high-density photos taken at different times after thawing.

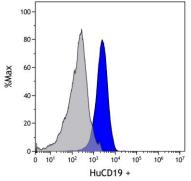
# Luciferase Expression



#### Cell Count

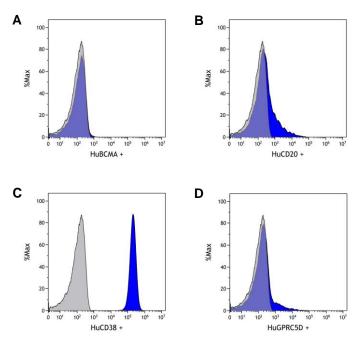
The indicated number of cells were placed in wells of a 96-well plate. After the addition of 15 mg/mL d-luciferin, bioluminescence was immediately read using a microplate reader.

## **Human CD19 Expression**



Nalm6-Fluc-Puro/HuCD19-Puro (Mid) cells were stained with an anti-HuCD19 antibody (blue) or isotype control antibody (grey) and analyzed by flow cytometry.

# **Expression Profiling of Surface Markers**



Nalm6-Fluc-Puro/HuCD19-Puro (Mid) (blue) cells were stained with an anti-HuBCMA antibody (A), anti-HuCD20 antibody (B), anti-HuCD38 antibody (C), anti-HuGPRC5D antibody (D), or isotype control antibody (grey; all panels) and analyzed by flow cytometry.

Quality control by: AWD/EMS Quality Assurance by: RLV Effective Date: 13-Nov-2024

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