

## Product description

Eighth generation cell line from ipsilateral brachial lymph node metastasis of seventh generation.

**Name:** NBF0LN8-1205BL Cell Line

**Disease:** Cancer

**Cancers detailed:** Melanoma

**Tissue:** Skin (lymph node metastasis)

**Organism:** Mouse

**Gender:** Female

**Growth properties:** Adherent

**Model:** Tumorigenic cell line; syngeneic

**Conditional:** No

**Parental cell line:** B16-F0

**Production details:** Eighth generation cell line derived from an ipsilateral brachial lymph node metastasis of a seventh generation (LN7) after one month of growth. A luciferase transgene was integrated into the B16-F0 parental line but is no longer expressed in LN metastatic progeny. WES suggests the presence of BRAF and PTEN mutations. These lines are syngeneic in the C57Bl/6 mouse background.

**Morphology:** Mixed morphology, predominantly spindle-shaped

**CRISPR Edited:** No

**Biosafety level:** 1

**Application:** Syngeneic transplantable models of melanoma metastasis

**Mycoplasma free:** Yes

## Contributor(s)

**Inventor:** Nathan E. Reticker-Flynn

**Institute:** Stanford University

## Properties

**Product format:** Frozen

**Unpacking and storage:**

1. Check all containers for leakage or breakage.

2. Remove the frozen cells from the dry ice packaging and immediately place the cells at a temperature below  $-130^{\circ}\text{C}$ , preferably in liquid nitrogen vapor, until ready for use.

**Recommended medium:** DMEM (high glucose) + 10% FBS + 4mM L-glutamine (or GlutaMAX) + 1% Pen/Strep (optional).

**Subculture:** Split confluent cultures 1:6 to 1:10 using 0.25% Trypsin-EDTA for 1 min at  $37.0^{\circ}\text{C}$ .

**Culture conditions:**  $37.0^{\circ}\text{C} \pm 1.0^{\circ}\text{C}$  incubator with  $5.0\% \pm 1.0\%$   $\text{CO}_2$

**Cryopreservation medium:** DMEM, 50% FBS, 10% DMSO

### Handling instructions

1. Please ensure that vials are frozen when received, and store at  **$<-130^{\circ}\text{C}$  long term**. When removing frozen cells from storage, it is important to minimize exposure to room temperature ( $15 - 25^{\circ}\text{C}$ ). If not proceeding directly to thawing, place the cells on dry ice or in a liquid nitrogen container.
2. **Do not thaw at room temperature.** To thaw, swirl the vial quickly in a  $37^{\circ}\text{C}$  water bath with O-ring and cap above the water to avoid contamination. Remove from the water bath with a small ice pellet remaining (this should not take more than 2 minutes) and wipe the exterior with 70% ethanol or isopropanol before transferring to a biosafety cabinet. Further steps should be conducted under aseptic conditions.
3. We strongly recommend that the volume of cell suspension is measured at this point, and a 20  $\mu\text{L}$  aliquot be removed for a **viable cell count** using trypan blue or similar dye. This ensures that provided cells are viable, and the cell count can be used to determine volume of growth medium to be added to the cell suspension.
4. Transfer the remaining cell suspension to a centrifuge tube using a pipette.
5. Rinse the vial with 1 mL of medium and add it dropwise to the cells. Wash by adding 10 mL of medium dropwise, while gently swirling the tube.
6. Centrifuge the cell suspension at  $140 \times g$  for 5 minutes at room temperature. Carefully remove the supernatant with a pipette, leaving a small amount of medium to ensure the cell pellet is not disturbed. Resuspend the cell pellet by gently flicking the tube.
7. Gently add required volume of pre-warmed culture medium, transfer to a suitable cell culture flask and incubate.

### Notes:

1. Cell death will be observed post-thaw. It is recommended that cells be thawed in the morning and growth medium is replaced at the end of the day, if necessary.
2. Always culture for a minimum of two passages after thawing before experimental use.
3. Accutase™ is recommended to prepare cells for injections, but Trypsin also works. When preparing cells for injection, use DMEM (without serum or phenol red) and keep the cells at room temperature. Inject  $200\mu\text{L}$  of cell suspension at a concentration of  $10^6$  cells/mL.

### References

- Reticker-Flynn, N.E., et al. (2022). 185(11):1924-1942.e23. PMID: 35525247

### Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner:  
NBF0LN8-1205BL Cell Line was invented by Nathan E. Reticker-Flynn (CancerTools.org #162233).

PRODUCTS ARE FOR RESEARCH USE ONLY AND NOT INTENDED FOR HUMAN OR ANIMAL DIAGNOSTIC OR THERAPEUTIC USES UNLESS OTHERWISE STATED.

While CancerTools.org has made all reasonable efforts to ensure that the information provided by CancerTools.org and its suppliers is correct, it makes no warranties or representations as to the accuracy or completeness of such information.