

IPLB-Sf21 Insect Cells

Catalog	No(s).
631411	

Amount 1 Vial Lot Number Specified on product label.

Description

IPLB-Sf21 insect cells may be used as a host for propagating the *Autographa californica* multiple-enveloped nuclear polyhedrosis virus (AcMNPV) and its expression vector derivatives generated from our BacPAKTM system. The IPLB-Sf21 cell line is derived from pupal ovaries of the fall armyworm, *Spodoptera frugiperda* (1). Exponentially growing IPLB-Sf21 cells are concentrated by centrifugation and frozen in insect cell complete medium (2) containing 10% dimethysulfoxide (DMSO).

Package Contents

• 1 x 1 ml IPLB-Sf21 insect cells (2x10⁶ cells/vial)

Storage Conditions

• Liquid nitrogen vapor phase

Shelf Life

• 1 year from date of receipt under proper storage conditions.

Shipping Conditions

• Dry ice $(-70^{\circ}C)$

Product Documents

Documents for Clontech® products are available for download at www.clontech.com/manuals

References

- 1. Vaughn, J. L., *et al.* (1977) *In Vitro* **13**(4):213-217.
- 2. Hink, W. F. (1970) *Nature* **266**(5244):466–467.

Quality Control Data

- Cell viability before freezing: $\geq 80\%$
- Cell viability after thawing: ≥ 60% (as determined by trypan blue dye exclusion).

Establishing the IPLB-Sf21 Cell Line:

- 1. Add 5 ml TNM-FH (2) supplemented with 10% FBS, to a 25-cm2 tissue culture flask and warm to 27°C.
- 2. Remove the vial of cells from liquid nitrogen (wear cryogenic gloves and a face shield).
- 3. Thaw rapidly in a 37°C water bath with gentle agitation until the suspension is almost thawed.
- 4. Decontaminate the outside of the vial with 70% ethanol.
- 5. In a laminar flow hood, transfer the cell suspension to the prewarmed flask containing 5 ml TNM-FH/FBS medium (refer to step 1).
- 6. Incubate the flask at 27°C for 1–3 hr (no more than 12 hr) to allow the healthy cells to adhere to the inside surface.
- 7. After confirming that a significant portion of the cells has attached, gently replace medium with 5 ml prewarmed TNM-FH/FBS medium.
- 8. Incubate at 27°C until the cells form a 80–90% confluent layer (about 2–3 days)



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