

## PD-L1-TF mRNA-LNP

Ready-to-use lipid nanoparticles



Order Information		
Catalog#	Size	GenPept No.
PM-LNP-0077	200uL	

#### Description

PD-L1 (programmed cell death protein ligand 1), also known as CD274, is a type 1 transmembrane protein mainly expressed on the cell surface of tumor cells or antigen-presenting cells. PD-1, (also known as CD274), is a membrane protein of the human immunoglobulin superfamily, expressed on the surface of T cells, primary B cells, and myeloid-derived dendritic cells. PD-1 and PD-L1 were also found to be co-expressed on the surface of SCLC cancer cells. PD-1/PD-L1 act as inhibitory checkpoint molecules in the immune system. PD-1 binds to PD-L1 to form a complex and transmit inhibitory signals to reduce the proliferation and activity of killer T lymphocytes. Blocking PD-1 with anti-PD-1 antibody, or blocking PD-L1 with anti-PD-L1 antibody, can inhibit the formation of PD-1/PD-L1 complex, thereby

## Composition

mRNA-LNPs suspended in PBS (-Ca, -Mg) (pH: 7.0-7.4).

## **Translated Protein sequence**

Available up request



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Product is delivered on blue ice. Store at 4°C for up to 3 months.

### **Application & Handling**

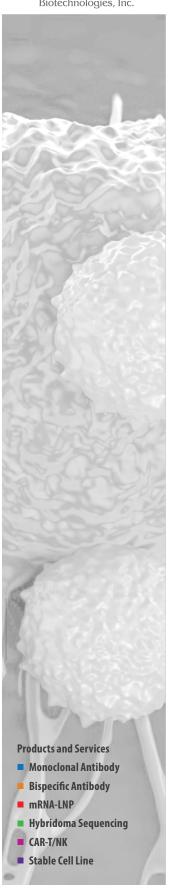
Upon receiving product, briefly pulse spin before opening to ensure product is at bottom of container. It is important not to spin for too long as this may rupture mRNA-LNPs. Do not vortex. Work with mRNA-LNPs on ice and minimize the time that the product spends at room temperature. After handling the product during experiments, return immediately to ice. mRNA-LNP products should only be handled with certified RNase-free reagents and consumables. Use of filtered pipette tips is highly recommended.

#### Safety & Research Disclosure

All ProMab mRNA lipid nanoparticle products are for in vitro research use only. Products are not FDA approved for human use.

#### **General Protocol**

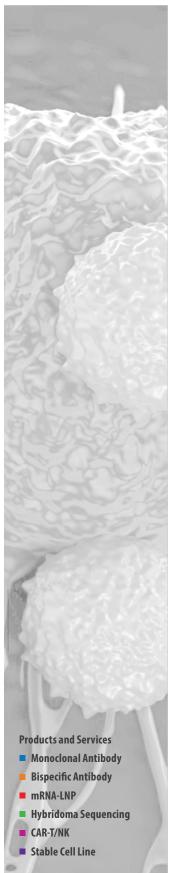
- 1. Prior to transfection: Plate 1ml of cells at a density of [1.0E6 cells/ml] in a single well of a 12-well culture plate. Ensure the cells you are using are viable and healthy. Try not to let your cells sit for longer than 5 minutes prior to transfection. Cell clumping at the time of transfection may reduce transfection efficiency. \*Note: If cell clumping occurs, gently pipette your culture up & down to ensure you have a single cell suspension before transfecting.
- 2. Briefly pipette mRNA-LNP mix up & down to resuspend. Add 20-40ul of the mRNA-LNP product dropwise directly to your 1ml culture. Gently tilt plate back and forth to mix (not necessary if you are using cells which will be immediately placed back into a shaker). Place your transfected cells back into their original culture conditions.
- 3. Check cell expression by FACS or by using other detection methods at 24hr intervals after transfection. \*Note: This is a generalized protocol for transfection using mammalian suspension culture cells. Transfection volume may be scaled up or down proportionately using the volumes given. HEK-293s cells were grown and transfected in FreeStyle™ F17 Expression Medium (Gibco, Cat#: A1383501), supplemented with GlutaMAX™ (Gibco, Cat#: 35050061), and Poloxamer 188 Non-ionic Surfactant (Gibco, Cat#: 24040032). T-cells were grown and transfected in a culture medium supplemented with 10% FBS (Omega Scientific, Cat#: FB-02). When transfecting cells using mRNA-LNPs, it is typically necessary for the cell culture medium to be supplemented with 10% FBS at the time of transfection. Without this supplement, transfection efficiency will drop significantly. For mRNA-LNP transfection of cells which cannot use FBS in the culture medium, please contact us at (510) 860-4615. Alternative transfection methods are available.





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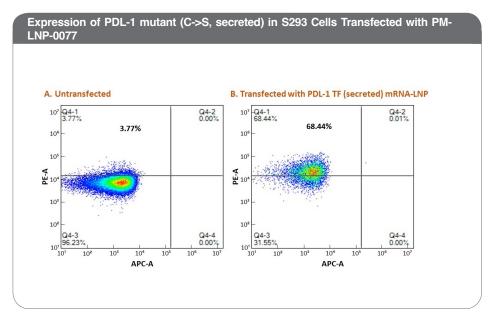


Figure 1. Flow Cytometry. PM-LNP-0077 nanoparticle-treated HEK293S cells secrete TF-tagged PD-L1 protein, detected with a labeled anti-CD274 antibody (ProMab, Cat# 31530).