

Transfecting Plasmid DNA into NCI-H460 Cells Using Lipofectamine[™] LTX Reagent

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Introduction

Lipofectamine LTX[™] Reagent is a proprietary, animal-origin free formulation for the transfection of DNA into eukaryotic cells with low cytotoxicity. This reference provides a recommended procedure to transfect plasmid DNA into NCI-H460, human large cell lung cancer cells (ATCC No. HTB-177) using Lipofectamine LTX[™] Reagent.

Important Guidelines for Transfection

Follow these important guidelines when transfecting NCI-H460 cells using Lipofectamine LTX[™] Reagent:

- Maintain the same seeding conditions between experiments. Use low-passage cells; make sure cells are healthy and greater than 90% viable before transfection.
- Transfection can be performed both in the presence or absence of serum. Test serum-free media for compatibility with Lipofectamine LTX[™] Reagent.
- We recommend Opti-MEM[®] I Reduced Serum Medium (Cat. No. 31985-070) to dilute the DNA Lipofectamine LTX[™] Reagent before complexing.
- Using PLUS[™] Reagent (Cat. No. 11514-015) enhances transfection performance in NCI-H460 Cells
- Visit <u>www.invitrogen.com/genedelivery</u> or contact Technical Services for other specialized transfection protocols.
- Lipofectamine LTX[™] Reagent performs well with vector-based RNAi experiments. For siRNA and Stealth RNAi transfections, we recommend Lipofectamine RNAiMAX. Go to www.invitrogen.com/RNAi or contact Technical Service for more information.

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Materials Needed

Have the following reagents on-hand before beginning:

- NCI-H460 cells maintained in RPMI Medium 1640 (Cat. No. 21870-076) supplemented with 4mM L-Glutamine (Cat. No 25030-081), 10% fetal bovine serum (Cat No. 16000-044). Grow cells at 37° C with 5% CO₂.
- Plasmid DNA of interest.
- Lipofectamine LTX[™] Reagent
- Opti-MEM® I Reduced Serum Media
- Appropriate tissue culture plates and supplies

Transfecting NCI-H460 Cells

Use this procedure to transfect plasmid DNA into NCI-H460 cells in a 24-well format (for other formats, see Scaling Up or Down Transfections, below). All amounts and volumes are given on a per well basis.

- 1. The day before transfection, trypsinize and count the cells. Plate 7.5×10^4 cells per well in 0.5 ml of complete growth medium. Cell density should be 50-80% confluent on the day of transfection.
- 2. (Optional) The day of transfection, remove growth medium from cells and replace with 0.5 ml of complete growth medium.
- 3. For each well of cells to be transfected, dilute $0.35~\mu g$ of DNA in $100~\mu l$ of Opti-MEM® I Reduced Serum Media without serum.
- 4. If using PLUS[™] Reagent: Mix PLUS[™] Reagent gently before use, then add 0.35 μl PLUS[™] Reagent (a 1:1 ratio to DNA) directly to the diluted DNA. Mix gently and incubate 5-15 minutes at room temperature.
- 5. For each well of cells, add 1.75-2.75 μl of Lipofectamine LTX[™] Reagent into the above diluted Opti-MEM[®]:DNA solution, mix gently and incubate 30 minutes at room temperature to form DNA- Lipofectamine LTX[™] Reagent complexes.
- 6. After 30 minute incubation, add 100 µl of the DNA- Lipofectamine LTX[™] Reagent complexes directly to each well containing cells and mix gently by rocking the plate back and forth.
- 7. Complexes do not have to be removed following transfection. Incubate the cells at 37°C in a CO₂ incubator for 18-24 hours post-transfection before assaying for transgene expression.

Scaling Up or Down Transfections

Culture Vessel	Surface Area per well	Volume Plating Medium	Cells per well	Volume Dilution Medium	DNA	Lipofectamine LTX [™] Reagent	PLUS™ Reagent
96-well	0.3 cm^2	100 µl	1.5×10^4	20 µl	70 ng	$0.35 - 0.55 \mu l$	0.07 µl
48-well	1 cm ²	200 µl	3×10^4	40 µl	140 ng	$0.7 - 1.2 \mu l$	0.14 µl
24-well	2 cm ²	500 µl	7.5×10^4	100 µl	350 ng	1.75-2.75 μl	0.35 µl
12-well	4 cm ²	1 ml	1.5×10^5	200 µl	0.7 µg	$3.5 - 5.5 \mu l$	0.7 µl
6-well	10 cm ²	2 ml	3.75×10^5	500 µl	1.75 µg	8.75-13.75	1.75 µl

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