

# Transfecting Plasmid DNA into ACHN Cells Using Lipofectamine<sup>™</sup> LTX Reagent

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#### Introduction

Lipofectamine LTX<sup>™</sup> 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 ACHN, human renal cancer cells (ATCC No. CRL-1611) using Lipofectamine LTX<sup>™</sup> Reagent.

## **Important Guidelines for Transfection**

Follow these important guidelines when transfecting ACHN cells using Lipofectamine LTX<sup>™</sup> 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<sup>™</sup> Reagent.
- We recommend Opti-MEM<sup>®</sup> I Reduced Serum Medium (Cat. No. 31985-070) to dilute the DNA Lipofectamine LTX<sup>™</sup> Reagent before complexing.
- Using PLUS<sup>™</sup> Reagent (Cat. No. 11514-015) enhances transfection performance in ACHN 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 <a href="https://www.invitrogen.com/RNAi">www.invitrogen.com/RNAi</a> or contact Technical Service for more information.

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### **Materials Needed**

Have the following reagents on-hand before beginning:

- ACHN cells maintained in Minimum Essential Medium (MEM) (Cat. No. 11090-081) supplemented with 4 mM L-Glutamine (Cat. No. 25030-081), 10% fetal bovine serum (Cat No.16000-044). Grow cells at 37° C with 5% CO<sub>2</sub>.
- Plasmid DNA of interest.
- Lipofectamine LTX™ Reagent (store at +4°C until ready to use) and PLUS™ Reagent (if desired; store at 4°C)
- Opti-MEM® I Reduced Serum Media
- Appropriate tissue culture plates and supplies

## **Transfecting ACHN Cells**

Use this procedure to transfect plasmid DNA into ACHN 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 1.25x10<sup>5</sup> 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  $0.5~\mu g$  of DNA in  $100~\mu l$  of Opti-MEM® I Reduced Serum Media without serum.
- 4. If using PLUS<sup>™</sup> Reagent: Mix PLUS<sup>™</sup> Reagent gently before use, then add 0.5 μl PLUS<sup>™</sup> 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 2.75-3.75 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<sup>™</sup> 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 CO<sub>2</sub> 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 <sup>™</sup> Reagent
96-well	$0.3 \text{ cm}^2$	100 µl	$2.5 \times 10^4$	20 µl	100 ng	$0.55 - 0.75 \mu l$	0.1 µl
48-well	1 cm <sup>2</sup>	200 µl	$5 \times 10^4$	40 µl	200 ng	1.1 – 1.5 µl	0.2 µl
24-well	2 cm <sup>2</sup>	500 µl	$1.25 \times 10^5$	100 µl	500 ng	2.75-3.75 µl	0.5 µl
12-well	4 cm <sup>2</sup>	1 ml	$2.5 \times 10^5$	200 µl	1 μg	$5.5 - 7.5 \mu l$	1.0 µl
6-well	10 cm <sup>2</sup>	2 ml	$6.25 \times 10^5$	500 µl	2.5 µg	13.75 – 18.75 μl	2.5 µl

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