

Transfecting Plasmid DNA into Caco-2 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 Caco-2 Human Colorectal Adenocarcinoma Cells (ATCC Cat. No. HTB-37) using Lipofectamine LTX Reagent (Cat. No. 15338-100).

Important Guidelines for Transfection

Follow these important guidelines when transfecting DNA into Caco-2 cells using Lipofectamine™ LTX Reagent:

- The addition of antibiotics to media during transfection may result in cell death, and has not been tested for Caco-2 cells. If you wish to use antibiotics during transfection, test your conditions thoroughly.
- Maintain the same seeding conditions between experiments. Use low-passage cells; make sure that 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.
- Using PLUS™ Reagent (Cat. No. 11514-015) enhances transfection performance in Caco-2 cells.
- We recommend Opti-MEM® I Reduced Serum Medium (Cat. No. 31985-062) to dilute the DNA and Lipofectamine™ LTX Reagent before complexing.
- Visit www.invitrogen.com/transfection or contact Technical Service for other specialized transfection protocols (including cell-type specific advice on use of PLUS™ Reagent and antibiotics, and a protocol for vector-based RNAi).
- Lipofectamine[™] LTX Reagent performs well with vector-based RNAi experiments. For siRNA and Stealth[™] RNAi transfections, we recommend Lipofectamine[™] RNAiMAX (Cat. No. 13778-075). Go to www.invitrogen.com/RNAi or contact Technical Service for more information.

Part no.: 25-0930W Rev. Date: 9 June 2006

Materials Needed

Have the following reagents on hand before beginning:

- Caco-2 cells maintained in DMEM supplemented with L-glutamine (Cat. No. 11965-084), 0.1 mM MEM Non-Essential Amino Acids Solution (Cat. No. 11140-050), and 10% Fetal Bovine Serum (Cat. No. 26140-079).
 Grow cells at 37°C with 5% CO₂.
- Plasmid DNA of interest (100 ng/µl or higher)
- Lipofectamine™ LTX Reagent (store at +4°C until use), and PLUS™ Reagent (if desired; store at 4°C)
- Opti-MEM® I Reduced Serum Medium
- Appropriate tissue culture plates and supplies

Transfection of Caco-2 Cells

Use this procedure to transfect plasmid DNA into Caco-2 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 4×10^4 cells per well in 0.5 ml of complete growth medium. Cell density should be $50 \sim 80\%$ confluent on the day of transfection.
- 2. For each well of cells to be transfected, dilute 0.5 μg of DNA into 100 μl of Opti-MEM® I Reduced Serum Medium without serum.
- 3. If using PLUS[™] Reagent: Mix PLUS[™] Reagent gently before use, then add 0.5 µl PLUS[™] Reagent (a 1:1 ratio to DNA) directly to the diluted DNA. Mix gently and incubate for 5-15 minutes at room temperature.
- 4. For each well of cells, dilute $0.75-1.75 \mu l$ of LipofectamineTM LTX into the above diluted DNA solution, mix gently and incubate for 25 minutes at room temperature to form DNA-LipofectamineTM LTX complexes.
- 5. Remove growth medium from cells and replace with 0.5 ml of complete growth medium. Add 100 µl of the DNA-Lipofectamine™ LTX complexes directly to each well containing cells and mix gently by rocking the plate back and forth.
- 6. 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

To transfect Caco-2 cells in different tissue culture formats, vary the amounts of Lipofectamine[™] LTX Reagent, DNA, cells, medium and PLUS[™] Reagent used in proportion to the relative surface area, as shown in the table (amounts given on a per well basis).

Culture vessel	Surface area per well ¹	Volume plating medium	Cells per well	Volume dilution medium ²	DNA	Lipofectamine™ LTX Reagent	PLUS™ Reagent
96-well	0.3 cm^2	100 µl	8×10^{3}	20 µl	100 ng	0.15 - 0.35 µl	0.1 µl
48-well	1 cm^2	200 µl	2×10^{4}	40 µl	200 ng	$0.3 - 0.7 \mu l$	0.2 µl
24-well	2 cm^2	500 µl	4×10^{4}	100 µl	500 ng	0.75 – 1.75 µl	0.5 µl
12-well	4 cm^2	1 ml	8×10^{4}	200 µl	1 µg	$1.5 - 3.5 \mu l$	1.0 µl
6-well	10 cm^2	2 ml	2×10^{5}	500 µl	2.5 µg	$3.75 - 8.75 \mu$ l	2.5 µl

¹Surface areas may vary depending on the manufacturer.

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²If the volume of Lipofectamine[™] LTX Reagent is too small to dispense accurately, and you cannot pool dilutions, predilute Lipofectamine[™] LTX Reagent 10-fold in Opti-MEM[®] I Reduced Serum Medium, and dispense a 10-fold higher amount (should be at least 1.0 μ l per well). Discard any unused diluted Lipofectamine[™] LTX Reagent.