<u>TransIT-X2®</u> Dynamic Delivery System for CRISPR/Cas9 Ribonucleoprotein (RNP) Delivery

Mirus.

Instructions for use with MIR 6000, 6003, 6004, 6005, 6006, 6010

SPECIFICATIONS

Storage	Store <i>Trans</i> IT-X2® Dynamic Delivery System tightly capped at -20°C. **Before each use, warm to room temperature and vortex gently.
Product Guarantee	1 year from the date of purchase, when properly stored and handled.

▶ CRISPR RIBONUCLEOPROTEIN (RNP) TRANSFECTION PROTOCOL

Fill in volumes below based on culture vessel used for transfection (Table 1).

A. Plate cells

1. Approximately 18-24 hours before transfection, plate cells in ___ml complete growth medium. Most cell types should be ~80% confluent at the time of transfection.

For adherent cells: Plate cells at a density of $0.8-3.0 \times 10^5$ cells/ml. For suspension cells: Plate cells at a density of $2.5-5.0 \times 10^5$ cells/ml.

2. Culture overnight.

B. Prepare TransIT-X2®:RNP complexes (Immediately before transfection)

- 1. Warm TransIT-X2® to room temperature and vortex gently before using.
- 2. Place μl of OptiMEM® I Reduced-Serum Medium in a sterile tube.
- 3. Add ____µl of a 50 μM guide RNA stock solution (12 nM final concentration per well). Mix gently by pipetting. NOTE: If using 2-part crRNA + tracrRNA, combine at a 1:1 molar ratio and incubate for 10 minutes at room temperature to anneal. Then add to tube containing OptiMEM®.
- Add ____µl of a 30 µM Cas9 protein stock solution (6 nM final concentration per well).
 Mix gently by pipetting.
- 5. Incubate at room temperature for 10 minutes.
- 6. Add ul TransIT-X2® to the RNP mixture. Mix gently by pipetting.
- 7. Incubate at room temperature for 15 minutes.

C. Distribute complexes to cells

- 1. Add the *Trans*IT-X2®:RNP complexes (prepared in Step B) drop-wise to different areas of the well. Gently rock plate for even distribution of complexes.
- 2. Incubate 24-72 hours.
- 3. Harvest cells and assay as required.

Table 1. Recommended starting conditions

Culture vessel	24-well plate	12-well plate	6-well plate
Surface area	1.9 cm ²	3.8 cm ²	9.6 cm ²
Complete growth medium	0.5 ml	1 ml	2.5 ml
Serum-free medium	50 μl	100 μΙ	250 μΙ
guide RNA (50 μM stock, 12 nM final in well)	0.12 μΙ	0.24 μΙ	0.6 μΙ
Cas9 Protein (30 µM stock, 6 nM final in well)	0.1 μΙ	0.2 μΙ	0.5 μΙ
TransIT-X2® Reagent	1 μΙ	2 μΙ	5 μΙ

▶ Transfection Optimization:

The 2:1 ratio of guide RNA to Cas9 protein (12 nM gRNA:6 nM Cas9; final concentration per well) used in this protocol is a starting point for RNP transfection. Further ratio optimization may be required for some cell types.

For more on transfection optimization, see the TransIT-X2® <u>full protocol (PDF)</u> or <u>Tips</u> <u>from the Bench</u>.



Reagent Agent *is an online tool designed to help determine the best solution for nucleic acid delivery based on in-house data, customer feedback and citations.

Learn more at: mirusbio.com/ra

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Rev.A 051817