

# TransIT®-mRNA Transfection Kit for CRISPR/Cas9 mRNA and gRNA Delivery



Instructions for use with MIR 2225, 2250, 2251, 2255, 2256

## SPECIFICATIONS

Storage	Store both TransIT®-mRNA Reagent and mRNA Boost Reagent tightly capped at -20°C. <b>Before each use</b> , warm to room temperature and vortex gently.
Product Guarantee	1 year from the date of purchase, when properly stored and handled.

## ► CAS9 mRNA + gRNA 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 (per well). Most cell types should be ~80% confluent on day of transfection.

**For adherent cells:** Plate cells at a density of 0.8—3.0 x 10<sup>5</sup> cells/ml.

**For suspension cells:** Plate cells at a density of 2.5—5.0 x 10<sup>5</sup> cells/ml.

2. Culture cells overnight.

### B. Prepare TransIT®-mRNA Reagent:mRNA Boost:RNA complexes

1. Warm TransIT®-mRNA and Boost Reagent to room temperature and vortex gently.
2. Place \_\_\_µl of OptiMEM® I Reduced-Serum Medium in a sterile tube.
3. Add \_\_\_µl of guide RNA (50 uM stock solution; 50 nM final concentration per well). Mix gently by pipetting. NOTE: If using 2-part crRNA + tracrRNA, first combine at a 1:1 molar ratio and incubate for 10 minutes at room temperature to anneal. Then add to tube containing OptiMEM®.
4. Add \_\_\_µl of mRNA encoding Cas9. Mix gently by pipetting.
5. Add \_\_\_µl of mRNA Boost Reagent. Mix gently by pipetting.
5. Add \_\_\_µl TransIT®-mRNA Transfection Reagent. Mix gently by pipetting.
6. Incubate at room temperature for 2-5 minutes to allow complexes to form.

### C. Distribute transfection complexes to cells

1. Add the complexes (prepared in Step B) drop-wise to different areas of the wells.
2. Gently rock the culture vessel to evenly distribute the complexes.
3. Incubate 24-72 hours. NOTE: A post-transfection media exchange is not necessary.
4. 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 <sup>2</sup>	3.8 cm <sup>2</sup>	9.6 cm <sup>2</sup>
Complete growth medium	0.5 ml	1 ml	2.5 ml
Serum-free medium	50 µl	100 µl	250 µl
guide RNA (50 µM stock, 50 nM final)	0.5 µl	1 µl	2.5 µl
Cas9 mRNA (1 mg/ml stock)	0.5 µl	1 µl	2.5 µl
mRNA Boost Reagent	0.5 µl	1 µl	2.5 µl
TransIT®-mRNA Transfection Reagent	0.5 µl	1 µl	2.5 µl

### ► Transfection Optimization:

Cell type, cell confluency, reagent volume, and post-transfection incubation time are a few of the key parameters that affect the outcome of transfection experiments. For more on transfection optimization, see the TransIT®-mRNA [full protocol \(PDF\)](#) or [Tips from the Bench](#).



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