

FuGENE® HD Transfection Reagent

INSTRUCTIONS FOR USE OF PRODUCTS E2311 AND E2312.

Quick
PROTOCOL

Transfection Protocol

Preparing the FuGENE® HD Transfection Reagent

1. Before use, allow the vial of FuGENE® HD Transfection Reagent to reach room temperature.
2. Mix by inverting or vortexing briefly. If a precipitate is visible, briefly warm at 37°C, then cool to room temperature.

General Transfection Protocol

1. To a sterile tube or U- or V-bottom plate, add 90–98µl of medium prewarmed to room temperature so that the final volume after adding the DNA is 100µl. Add 2µg of plasmid DNA (0.2–1µg/µl), and vortex. For a 3:1 FuGENE® HD Transfection Reagent:DNA ratio, add 6µl of FuGENE® HD Transfection Reagent directly to medium, and mix immediately. For other ratios, consult Table 1.

Note: Do not allow undiluted FuGENE® HD Transfection Reagent to contact the sides of the tube or U- or V-bottom plate.

Table 1. Volumes of FuGENE® HD Transfection Reagent and DNA for Various Reagent:DNA Ratios.

	Ratio of FuGENE® HD Transfection Reagent to DNA					
	4:1	3.5:1	3:1	2.5:1	2:1	1.5:1
Medium to a final volume of	100µl	100µl	100µl	100µl	100µl	100µl
DNA amount	2µg	2µg	2µg	2µg	2µg	2µg
Volume of FuGENE® HD Transfection Reagent	8µl	7µl	6µl	5µl	4µl	3µl

2. Incubate the FuGENE® HD Transfection Reagent/DNA mixture for 0–15 minutes at room temperature.
3. Add 2–10µl of the FuGENE® HD Transfection Reagent/DNA mixture per well to a 96-well plate containing 100µl of cells in growth medium. Mix by pipetting or using a plate shaker. Return cells to the incubator for 24–48 hours.
4. Measure transfection efficiency using an assay appropriate for the reporter gene. For transient transfection, cells are typically assayed 24–48 hours after transfection.

See additional protocol information in Technical Manual #TM328, available online at: www.promega.com/tbs/

For a list of conditions that have been used to transfect various cell types, visit our FuGENE® HD Protocol Database at: www.promega.com/techserv/tools/FugeneHD/

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ORDERING/TECHNICAL INFORMATION:

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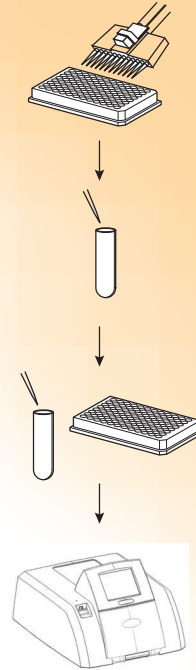


Plate cells one day prior to transfection.

Dilute DNA in medium.

Add FuGENE® HD Transfection Reagent. Incubate for 0–15 minutes.

Add FuGENE® HD Transfection Reagent:DNA mixture to cells, and mix gently.

Measure experimental results.

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