

# FreeStyle™ MAX Reagent

|   |                             |   |  |
|---|-----------------------------|---|--|
|    | <b>Package Contents</b>     | <b>Catalog Number</b><br>16447-100<br>16447-500<br>16447-750  | <b>Size</b><br>1.0 mL<br>15.0 mL<br>10 × 15.0 mL |
|    | <b>Storage Conditions</b>   | <ul style="list-style-type: none"> <li>Store at 4°C.</li> <li>Do not freeze.</li> </ul>   |  |
|    | <b>Required Materials</b>   | <ul style="list-style-type: none"> <li>FreeStyle™ 293-F Cells, FreeStyle™ CHO-S® Cells, or DG44 Cells</li> <li>FreeStyle™ 293 Expression Medium, FreeStyle™ CHO Expression Medium, or DG44 Medium</li> <li>Erlenmeyer flasks with vented caps</li> <li>Orbital shaker in temperature and CO<sub>2</sub> controlled incubator</li> <li>Plasmid DNA</li> <li>OptiPRO™ SFM</li> </ul>            |  |
|    | <b>Timing</b>               | Cell Preparation: 1 day<br>Transfection: 10–20 minutes  |  |
|    | <b>Selection Guide</b>      | <a href="#">Protein Expression Systems</a><br>Go online to view related products.   |  |
|  | <b>Product Description</b>  | <ul style="list-style-type: none"> <li>FreeStyle™ MAX Reagent is a proprietary, animal origin-free formulation for transfecting plasmid DNA into eukaryotic cells, which can be easily scaled up to produce large amounts of recombinant proteins.</li> <li>This transfection reagent is formulated specifically for use with FreeStyle™ 293-F, FreeStyle™ CHO-S®, and DG44 cells.</li> </ul> |  |
|  | <b>Important Guidelines</b> | <ul style="list-style-type: none"> <li>DNA-FreeStyle™ MAX complexes must be made in OptiPRO™ SFM and can be added directly to cells in culture medium.</li> <li>Cultivate FreeStyle™ 293-F and FreeStyle™ CHO-S® Cells, or DG44 Cells, in a humidified, 37°C, 8% CO<sub>2</sub> environment in suspension on an orbital shaker.</li> </ul>  |  |
|  | <b>Online Resources</b>     | Visit our <a href="#">product page</a> for additional information and protocols. For support, visit <a href="http://www.lifetechnologies.com/support">www.lifetechnologies.com/support</a> .  |  |



## Protocol Outline

- Culture cells at least three passages after thawing.
- Prepare and add DNA-lipid complexes to cells.
- Incubate cells for 1–7 days.
- Harvest.

## Transfection Protocol

-  See page 2 to view a typical procedure for transfecting FreeStyle™ 293-F and FreeStyle™ CHO-S® Cells for protein expression.
-  See page 3 to view a typical procedure for transfecting DG44 cells to generate stable cell lines.

## Transfection Conditions for FreeStyle™ Cells

**Final transfection volume:** 30 mL

**Number of cells to transfect:**  $3 \times 10^7$

**Amount of plasmid DNA:** 37.5 µg

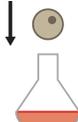
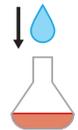
**Amount of FreeStyle™ MAX Reagent:** 37.5 µL

## Scaling Up or Down Transfections

## Limited Product Warranty and Disclaimer Details

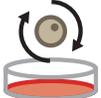
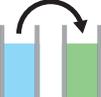
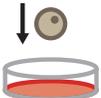
## Transfecting FreeStyle™ 293-F or FreeStyle™ CHO-S® Cells

Use the following protocol to transfect suspension cells. All amounts are given on a per-flask basis for 30-mL cultures in 125-mL shake flasks.

| Timeline |   | Steps   | Procedure Details   |  |                                    |
|----------|---|---|---|--|------------------------------------|
| Day -1   | 1 |  Expand cells                        | For each 30-mL transfection, you will need $3 \times 10^7$ cells in 30 mL of FreeStyle™ 293 Expression Medium or FreeStyle™ CHO Expression Medium.<br><b>For FreeStyle™ 293-F Cells:</b> One day prior to transfection, passage at $6-7 \times 10^5$ cells/mL; shake at 120–135 rpm.<br><b>For FreeStyle™ CHO-S® Cells:</b> One day prior to transfection, passage at $5-6 \times 10^5$ cells/mL; shake at 120–135 rpm.   |  |                                    |
|          | 2 |  Count cells and determine viability | Use the trypan blue dye exclusion method to determine cell viability and clumping in a small aliquot of cells. Use an automated cell counter or a hemocytometer to determine cell counts. On the day of transfection, your cells should have a density of $1.2-1.5 \times 10^6$ cells/mL at >95% viability.   |  |                                    |
|          | 3 |  Seed cells in flask                 | Dilute cells to $1 \times 10^6$ cells/mL. You will need $3 \times 10^7$ cells for each 30-mL transfection.<br>Use fresh, pre-warmed FreeStyle™ 293 Expression Medium or FreeStyle™ CHO Expression Medium to a total volume of 30 mL for each 30-mL transfection.  |  |                                    |
| Day 0    | 4 |  Prepare DNA-lipid complexes        | Prepare DNA-lipid complexes as follows: <ol style="list-style-type: none"> <li>Dilute 37.5 µg of plasmid DNA in OptiPRO™ SFM reduced serum medium to a total volume of 0.6 mL. Mix gently.</li> <li>Dilute 37.5 µL of FreeStyle™ MAX Reagent in OptiPRO™ SFM reduced serum medium to a total volume of 0.6 mL. Mix gently and incubate for 5 minutes at room temperature. Incubation times longer than five minutes may result in decreased activity.</li> <li>After the 5-minute incubation, add the diluted DNA to the diluted reagent to obtain a total volume of 1.3 mL. Mix gently.</li> <li>Incubate for 20–30 minutes at room temperature to allow the DNA-lipid complexes to form.</li> </ol> |  |                                    |
|          | 5 |  Add DNA-lipid complex to cells    | Add 1.2 mL of complex to each cell suspension flask. Each flask should have a total volume of 30 mL, and contain approximately $1 \times 10^6$ viable cells/mL.<br>To the negative control flask, add 2 mL of reduced serum medium instead of complex.  |  |                                    |
| Days 1-7 | 6 |  Incubate                          | Temperature<br>37°C   | Humidified Atmosphere<br>8% CO <sub>2</sub> in air | Orbital Shaker Platform<br>125 rpm |
|          | 7 |  Harvest cells or media            | Assay for recombinant protein expression. Perform this step 1–7 days post-transfection. Harvest media instead of cells if recombinant protein is secreted.  |  |                                    |

## Transfecting DG44 Cells to Generate Stable Cell Lines

Use this procedure to transfect linearized DNA into DG44 cells. All amounts are on a per-flask basis for 30-mL cultures in 125-mL shake flasks.

| Timeline |   | Steps                              | Procedure Details   |   |   |
|----------|---|------------------------------------|---|---|---|
| Day 0    | 1    | Prepare and culture the DG44 cells | a. Passage the cells at $3 \times 10^5$ cell/mL.<br>b. Shake at 130–135 rpm at 37°C, 8% CO <sub>2</sub> .<br>c. Culture in CD DG44 Medium (Cat. No. 12610-010) with 8 mM L-glutamine (Cat. No. 25030-081) and 18 mL/L of 10% Pluronic® F-68 (Cat. No. 24040-032). |   |   |
| Day 1    | 2    | Passage the DG44 cells again       | Passage cells again at $3 \times 10^5$ cell/mL.   |   |   |
|          | 3    | Prepare the cells                  | Count the cells. Cell viability should be >95%.<br>In each flask, add $1.5 \times 10^7$ cells in a total volume of 30 mL CD DG44 Medium.  |   |   |
|          | 4    | Combine lipid and linearized DNA   | Gently invert the tube to mix the reagent. Then, add 18 µg of linearized DNA and 15 µg of FreeStyle™ MAX Reagent into 1.2 mL of OptiPRO™ SFM (at room temperature), and gently invert to mix.   |   |   |
| Day 2    | 5   | Incubate the DNA-lipid mixture     | Incubate for 10 minutes at room temperature, but no longer than 20 minutes.   |   |   |
|          | 6  | Add DNA-lipid mixture to cells     | Slowly add 1.2 mL of mixture into the 125-mL flask containing the cells while slowly swirling the flask.  |   |   |
|          | 7  | Incubate                           | <b>Temperature</b><br>37°C  | <b>Humidified Atmosphere</b><br>8% CO <sub>2</sub> in air | <b>Orbital Shaker Platform</b><br>130–135 rpm |
| Day 4    | 8  | Place cells on a selective medium  | Place cells on a selective medium (for example, CD OptiCHO™ Medium, Cat. No. 12681-011).  |   |   |