MDA-MB-231-Luc

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□□□ □□ □□	GMO-S1

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GMO-S1: This cell line contains a stably integrated firefly luciferase reporter cassette (Luc2, codon-optimized) introduced via replication-incompetent lentiviral transduction. The resulting polyclonal cell population was maintained under puromycin selection (1–5 µg/mL). S1 containment is required. This classification applies only within Germany and may differ elsewhere.

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MDA-MB-231-Luc (□□□□ □□□□ □□ 305693)

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Luc2 (□□□□, □□ □□□)

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□□□□: p.Gly464Val, □□□□; □□□□: p.Gly13Asp, □□□□; □□□□: p.Arg280Lys, □□□□

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DMEM:□□ F12(1:1), w: 3.1 g/L □□□, w: 1.6 mM L-□□□□, w: 15 mM HEPES, w: 1.0 mM □□□□ □□□, w: 1.2 g/L NaHCO3 (Cytion 820400a)

□□ □□	37°C, 5%CO2, □□ □□.

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10% FBS□ □□□ □□□□□

□□ □□	□□□□ 5□, 37°C□□

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Confirm that the vial remains deeply frozen upon delivery, as cells are shipped on dry ice to maintain optimal temperatures during transit.
Upon receipt, either store the cryovial immediately at temperatures below -150°C to ensure the preservation of cellular integrity, or proceed to step 3 if immediate culturing is required.
For immediate culturing, swiftly thaw the vial by immersing it in a 37°C water bath with clean water and an antimicrobial agent, agitating gently for 40-60 seconds until a small ice clump remains.
Perform all subsequent steps under sterile conditions in a flow hood, disinfecting the cryovial with 70% ethanol before opening.
Carefully open the disinfected vial and transfer the cell suspension into a 15 ml centrifuge tube containing 8 ml of room-temperature culture medium, mixing gently.
Centrifuge the mixture at 200 x g for 5 minutes, carefully discard the supernatant containing freezing medium.
Follow the procedure described under Post-Thaw Recovery

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Cytion GmbH
Nikola-Tesla-Str. 3
69124 Heidelberg
Tel: +49 (0)6221 405780
info@cytion.com
cytion.com

Registered office Heidelberg
VAT ID: DE283292917
Local court Mannheim - HRB 714394
CEO / Geschäftsführer: Jonathan Steubing

Banking information
Sparkasse Heidelberg
IBAN: DE35 6725 0020 0009 2076 86
BIC: SOLADES1HDB