

A2780-GFP | 305676

General information

Description

A2780-GFP is a genetically modified derivative of the human ovarian carcinoma cell line A2780, engineered to stably express green fluorescent protein (GFP). The parental A2780 cell line was established from an ovarian tumor in an adult patient and is widely used as a model for epithelial ovarian cancer, particularly in studies of chemotherapy response. It is known for its relative sensitivity to platinum-based agents such as cisplatin, making it a valuable system for investigating mechanisms of drug action and resistance. The GFP-expressing variant preserves these biological characteristics while incorporating a fluorescent reporter for enhanced experimental utility.

The stable expression of GFP enables real-time visualization and quantitative analysis of A2780-GFP cells in a variety of experimental settings. Fluorescence-based detection allows researchers to monitor cell proliferation, morphology, migration, and viability *in vitro*, as well as to track tumor growth and dissemination *in vivo*. This is particularly advantageous in xenograft and metastasis models, where GFP facilitates discrimination of tumor cells from surrounding host tissue. The fluorescent signal is generally stable across passages, although expression levels may vary depending on the transduction method and clonal selection.

A2780-GFP retains the core molecular and phenotypic features of the parental A2780 line, including pathways associated with DNA damage response, apoptosis, and chemotherapeutic sensitivity. As such, it is commonly used in high-content imaging assays, drug screening platforms, and co-culture systems where spatial and temporal resolution of tumor cell behavior is required. The addition of GFP significantly enhances the versatility of this model, supporting applications in ovarian cancer research, therapeutic evaluation, and studies of tumor cell dynamics.

Genetic modification: Stably modified by replication-incompetent lentiviral transduction to express the ZsGreen1 green fluorescent protein reporter; maintained as a polyclonal population under puromycin selection (1–5 µg/mL). S1/BSL-1 containment.

Organism Human

Tissue Ovary

Disease Ovarian endometrioid adenocarcinoma

Metastatic site Primary tumor site (ovary)

Applications Ovarian cancer imaging; GFP-based tumor tracking; *in vivo* fluorescence imaging; combination with parental A2780 for comparative studies; drug efficacy visualization

Characteristics

Age Age unspecified

Gender Female

A2780-GFP | 305676**Ethnicity** African American**Morphology** Epithelial-like**Cell type** Epithelial cells**Growth properties** Adherent**Regulatory Data****Citation** A2780-GFP (Cytion catalog number 305676)**Biosafety level** 1**NCBI_TaxID** 9606**CellosaurusAccession** Not assigned (A2780-GFP reporter derivative; parental A2780 CVCL_1099)**GMO Status** GMO-S1: This cell line contains a stably integrated ZsGreen1 green fluorescent protein reporter 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.**Biomolecular Data****Antigen expression** ZsGreen1 (green fluorescent protein)**Handling****Culture Medium** RPMI 1640, w: 2.0 mM stable Glutamine, w: 2.0 g/L NaHCO₃ (Cytion article number 820700a)**Supplements** Supplement the medium with 10% FBS**Dissociation Reagent** Accutase**Split ratio** 1 to 5

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Seeding density 1 to 3×10^4 cells/cm²

Fluid renewal 2 to 3 times per week

Freeze medium As a cryopreservation medium, we use complete growth medium + 10% DMSO for adequate post-thaw viability.

Thawing and Culturing Cells

1. Confirm that the vial remains deeply frozen upon delivery, as cells are shipped on dry ice to maintain optimal temperatures during transit.
2. 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.
3. 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.
4. Perform all subsequent steps under sterile conditions in a flow hood, disinfecting the cryovial with 70% ethanol before opening.
5. 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.
6. Centrifuge the mixture at 200 x g for 5 minutes, carefully discard the supernatant containing freezing medium.
7. Follow the procedure described under Post-Thaw Recovery

Incubation Atmosphere 37°C, 5% CO₂, humidified atmosphere.

Shipping Conditions Cryopreserved cell lines are shipped on dry ice in validated, insulated packaging with sufficient refrigerant to maintain approximately -78 °C throughout transit. On receipt, inspect the container immediately and transfer vials without delay to appropriate storage.

Storage Conditions For long-term preservation, place vials in vapor-phase liquid nitrogen at about -150 to -196 °C. Storage at -80 °C is acceptable only as a short interim step before transfer to liquid nitrogen.

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Quality Control & Molecular Analysis