

**SW1271 Cells | 305880****Renseignements généraux****Description**

The SW1271 cell line is a human small cell lung carcinoma (SCLC) model derived from an adult patient. It is characterized by its neuroendocrine phenotype, which is typical of SCLC, and displays molecular features relevant to therapeutic sensitivity and resistance. In a comprehensive epigenome-wide methylation analysis of SCLC cell lines, including SW1271, the line exhibited specific DNA methylation patterns that correlated with chemosensitivity to several classes of anticancer drugs. These included Aurora kinase inhibitors, CDK inhibitors, and DNA-damaging agents. The methylation status of key genes such as *TREX1*, *SLFN11*, *CEP350*, and *KDM1A* in SW1271 and other SCLC models has been associated with altered drug response, implicating epigenetic modulation as a determinant of therapeutic efficacy.

Furthermore, SW1271 has been used in integrated genomic and epigenomic studies to understand subtype-specific vulnerabilities in SCLC. This cell line, along with others representing different SCLC transcriptional subtypes (*ASCL1*, *NEUROD1*, *POU2F3*, and *YAP1*), helps delineate the heterogeneity within the disease. The methylation profile of SW1271 contributes to our understanding of the regulatory mechanisms affecting gene expression and drug response, including suppression of tumor suppressor genes and dysregulation of lineage-specific transcription factors. These insights position SW1271 as a valuable model for investigating epigenetically driven pathways in SCLC and for identifying potential biomarkers and therapeutic targets.

**Organism** Human**Tissue** Lung**Disease** Lung small cell carcinoma**Synonyms** SW-1271, SW 1271**Caractéristiques****Age** 69 years**Gender** Male**Ethnicity** Caucasian**Morphology** Epithelial**Cell type** Epithelial cell**Growth properties** Adherent

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## Données réglementaires

<b>Citation</b>	SW1271 (Cytion catalog number 305880)
<b>Biosafety level</b>	1
<b>NCBI_TaxID</b>	9606
<b>CellosaurusAccession</b>	CVCL_1716

## Données biomoléculaires

<b>Antigen expression</b>	Blood Type A; Rh +
<b>Mutational profile</b>	Mutation: NRAS, Simple, p.Gln61Arg (c.182A>G), Homozygous, SMARCA4, Simple, p.Asn774Lys (c.2322C>A), Homozygous.Mutation, TP53, Simple, p.Cys277Phe (c.830G>T), Homozygous

## Manipulation

<b>Culture Medium</b>	DMEM:Ham's F12 (1:1), w: 3.1 g/L Glucose, w: 2.5 mM L-Glutamine, w: 15 mM HEPES, w: 0.5 mM Sodium pyruvate, w: 1.2 g/L NaHCO3 (Cytion article number 820400a)
<b>Supplements</b>	Supplement the medium with 10% FBS, AB, 5µg/mL Insulin
<b>Dissociation Reagent</b>	Accutase
<b>Fluid renewal</b>	2 to 3 times per week
<b>Freeze medium</b>	As a cryopreservation medium, we use complete growth medium (including FBS) + 10% DMSO for adequate post-thaw viability, or CM-1 (Cytion catalog number 800100), which includes optimized osmoprotectants and metabolic stabilizers to enhance recovery and reduce cryo-induced stress.

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### 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  $-150^{\circ}\text{C}$  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  $37^{\circ}\text{C}$  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  $300 \times g$  for 3 minutes to separate the cells and carefully discard the supernatant containing residual freezing medium.
7. Gently resuspend the cell pellet in 10 ml of fresh culture medium. For adherent cells, divide the suspension between two T25 culture flasks; for suspension cultures, transfer all the medium into one T25 flask to promote effective cell interaction and growth.
8. Adhere to established subculture protocols for continued growth and maintenance of the cell line, ensuring reliable experimental outcomes.

### Incubation Atmosphere

$37^{\circ}\text{C}$ , 5%  $\text{CO}_2$ , humidified atmosphere.

### Shipping Conditions

Cryopreserved cell lines are shipped on dry ice in validated, insulated packaging with sufficient refrigerant to maintain approximately  $-78^{\circ}\text{C}$  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  $-196^{\circ}\text{C}$ . Storage at  $-80^{\circ}\text{C}$  is acceptable only as a short interim step before transfer to liquid nitrogen.

## Contrôle de la qualité et analyse moléculaire

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**Sterility**

Mycoplasma contamination is excluded using both PCR-based assays and luminescence-based mycoplasma detection methods.

To ensure there is no bacterial, fungal, or yeast contamination, cell cultures are subjected to daily visual inspections.