

UM-SCC-128 Cells | 305726

General information

Description

UM-SCC-128 is a human HNSCC cell line from the University of Michigan UM-SCC panel, growing as an adherent monolayer with epithelial-like morphology. It is part of the higher-numbered UM-SCC series and provides a model for comparative preclinical HNSCC studies.

Applicable in HNSCC research, drug sensitivity and resistance studies, radiation response, and UM-SCC multi-line panel comparative experiments. Suitable for pharmacological screening and xenograft models.

Organism

Human

Tissue

Head and neck, oral cavity

Disease

Head and neck squamous cell carcinoma (HNSCC)

Metastatic site

Primary tumor site (head and neck)

Applications

HNSCC research; drug sensitivity and resistance; radiation response; UM-SCC panel studies; xenograft models

Characteristics

Age

Age unspecified

Gender

Sex unspecified

Ethnicity

Ethnicity unspecified

Morphology

Epithelial-like

Cell type

Epithelial cells

Growth properties

Adherent

Regulatory Data

Citation

UM-SCC-128 (Cytion catalog number 305726)

Biosafety level

1

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NCBI_TaxID 9606**CellosaurusAccession** Not assigned**GMO Status** No genetic modification; wildtype HNSCC cell line

Biomolecular Data

Handling

Culture Medium DMEM, w: 4.5 g/L Glucose, w: 4 mM L-Glutamine, w: 3.7 g/L NaHCO₃, w: 1.0 mM Sodium pyruvate (Cytion article number 820300a)**Supplements** Supplement the medium with 10% FBS and 1% non-essential amino acids (NEAA)**Dissociation Reagent** Accutase**Doubling time** approx. 24 to 48 hours**Subculturing** Remove medium, wash with PBS without calcium and magnesium, cover with Accutase, incubate 8–10 min at RT, resuspend in medium, centrifuge 300×g 3 min, discard supernatant, reseed in fresh medium.**Split ratio** 1 to 5**Seeding density** 1 to 3×10^4 cells/cm²**Fluid renewal** Every 2 to 3 days**Freeze medium** As a cryopreservation medium, we use complete growth medium + 10% DMSO for adequate post-thaw viability.

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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°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.

Quality Control & Molecular Analysis