

SNU-719 Cells | 305636**General information****Description**

The SNU-719 cell line is a human gastric carcinoma model established from the primary gastric tumor tissue of an adult male patient in Korea. It belongs to a collection of gastric cancer lines developed to support cancer research in East Asia, where gastric cancer prevalence is particularly high. SNU-719 was derived from a moderately differentiated adenocarcinoma and has demonstrated strong attachment to plastic culture surfaces, growing as a diffuse monolayer. The line was maintained in RPMI-1640 medium supplemented with 10% heat-inactivated fetal bovine serum.

Comprehensive biochemical and genetic profiling of SNU-719 revealed expression of carcinoembryonic antigen (CEA) and high levels of tissue polypeptide antigen (TPA) in both supernatant and cell lysate. However, alpha-fetoprotein (aFP) was not detected. Mutation analysis identified alterations in the TP53 gene, although the c-Ki-ras oncogene remained unmutated in this line. These features make SNU-719 a suitable model for studying the molecular mechanisms of gastric adenocarcinoma and for evaluating biomarker expression and therapeutic interventions. Additionally, STR and SNP profiling have confirmed its identity and uniqueness, ensuring the cell line's reliability for in vitro experimentation.

Organism

Human

Tissue

Stomach

Disease

tubular adenocarcinoma

Synonyms

SNU719, NCI-SNU-719

Characteristics**Age**

53 years

Gender

Male

Ethnicity

Korean

Morphology

Epithelial-like

Cell type

Epithelial

Growth properties

Adherent, monolayer

Regulatory Data

SNU-719 Cells | 305636**Citation** SNU-719 (Cytion catalog number 305636)**Biosafety level** 2**NCBI_TaxID** 9606**CellosaurusAccession** CVCL_5086**Biomolecular Data****Mutational profile** Mutation: CTNNB1, Simple, p.Gly34Val (c.101G>T), Heterozygous; Mutation: MET, Simple, p.Asp153Ala (c.458A>C), Heterozygous; Mutation: NRAS, Simple, p.Gln61Leu (c.182A>T), Homozygous; Mutation: PIK3CA, Simple, p.Pro104Arg (c.311C>G), Heterozygous**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**Doubling time** 43 hours**Subculturing** Remove medium, add fresh 0.25 % trypsin 0.02 % EDTA solution, stand culture flask at 37°C for 3 to 5 minutes, add culture medium and collect the cells, transfer the medium into 15ml tube, centrifuge, aspirate the medium, resuspend the pellets with culture medium and dispense into the culture flask**Fluid renewal** 2 to 3 times per week**Freeze medium** 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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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 300 x 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°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

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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.