Panc 03.27

305381

GENERAL INFORMATION

Description

Panc 03.27 (PL11) is a human pancreatic adenocarcinoma cell line derived from a primary tumor. This cell line is a valuable resource for studying the molecular and genetic alterations characteristic of pancreatic cancer, a disease with notoriously poor prognosis and limited treatment options. Panc 03.27 is particularly utilized in research focused on genomic instability, allelic loss, and tumor suppressor gene alterations.

This cell line is characterized by significant chromosomal abnormalities, including regions of allelic loss and homozygous deletions, as revealed through high-resolution SNP arrays. Such deletions frequently implicate loci associated with tumor suppressor genes, providing insight into the genetic mechanisms driving pancreatic tumorigenesis. Panc 03.27 has been instrumental in understanding the impact of chromosomal instability patterns on the progression and heterogeneity of pancreatic cancers, highlighting genomic ""holes"" or regions prone to homozygous deletions.

Panc 03.27 is used in preclinical research to evaluate novel therapeutic approaches, including those targeting genetic vulnerabilities specific to pancreatic cancer. Its genomic profile makes it a key model for studies of somatic mutations, gene loss, and tumor progression, helping to identify potential biomarkers and treatment targets. The cell line's ability to provide consistent and reproducible results ensures its ongoing importance in pancreatic cancer research and drug development.

Tissue	Pancreas
Disease	Adenocarcinoma

Synonyms

- Panc 3.27
- Panc-03.27
- PANC-03-27
- Panc_03_27
- Panc-3_27
- PANC3.27
- Panc3.27
- Panc3_27
- Panc-327
- PANC 327
- Panc327
- PANC0327
- Panc0327
- PL11
- PL-11

CLASSIFICATION

Organism	Human
Sex	Female
Growth properties	Adherent
Biosafety level	1

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

CHARACTERISTICS

Morphology	Epithelial-like
Cell type	Ephitelial

IDENTIFIERS

Citation

Panc 03.27 (Cytion catalog number 305381)

NCBI TaxID	9606
Cellosaurus	CVCL_1635

EXPRESSION AND SUSCEPTIBILITY

Antigen expression

MHC class I +; MHC class II -; Blood type A; Rh+

Oncogenes

KRAS mutant (G12V)

Mutational profile

Mutation: KRAS, Simple, p.Gly12Val (c.35G>T), Heterozygous; Mutation: TP53, Simple, c.375+5G>T, Homozygous

HANDLING AND CULTURE

Culture medium

RPMI 1640, w: 2.0 mM stable Glutamine, w: 2.0 g/L NaHCO3 (Cytion article number 820700a)

Incubation atmosphere	37°C, 5% CO_2, humidified atmosphere.

Supplements

Supplement the medium with 15% FBS and 0,01 mg/ml human recombinant insulin

Dissociation reagent	Accutase

Subculturing

Remove the old medium from the adherent cells and wash them with PBS that lacks calcium and magnesium. For T25 flasks, use 3-5 ml of PBS, and for T75 flasks, use 5-10 ml. Then, cover the cells completely with Accutase, using 1-2 ml for T25 flasks and 2.5 ml for T75 flasks. Let the cells incubate at room temperature for 8-10 minutes to detach them. After incubation, gently mix the cells with 10 ml of medium to resuspend them, then centrifuge at 300xg for 3 minutes. Discard the supernatant, resuspend the cells in fresh medium, and transfer them into new flasks that already contain fresh medium.

Seeding density	2-4*10^4/cm²
Fluid renewal	2 to 3 times per week

Thawing and culturing

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 300 x g for 3 minutes to separate the cells and carefully discard the supernatant containing residual freezing medium.
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.
Adhere to established subculture protocols for continued growth and maintenance of the cell line, ensuring reliable experimental outcomes.

QUALITY CONTROL AND CRYOPRESERVATION

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.

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.

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.

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.

DONOR

Donor age	65 years
Ethnicity	Caucasian

LEGAL AND SAFETY

Safety precautions

When planning to store a cryovial in liquid nitrogen for future thawing, it is mandatory to adhere to stringent safety measures. Appropriate protective gloves and clothing are essential, and the use of a face mask or safety goggles is required during the transfer of frozen samples to or from the liquid nitrogen tank. This is to mitigate the risk of injury from potential cryovial explosions upon removal, which can result in the projection of sharp fragments.

Warranty

At Cytion, we stand behind the quality and viability of our products.
We guarantee that our cells will be viable at the start of culture for
60 days after shipment from our facilities.
We assure that all materials conform to the specifications described in the
Certificate of Analysis and/or Product Sheet.
For full details, please refer to our
Warranty Statement.

Material Transfer Agreement required	Yes