



General information

Organism Human

Tissue Colon

Disease Adenocarcinoma

Synonyms SW403, SW 403

Characteristics

Age 51 years

Gender Female

Ethnicity Caucasian

Morphology Epithelial-like

Growth properties Adherent

Identifiers / Biosafety / Citation

Citation SW-403 (Cytion catalog number 300350)

Biosafety level

Expression / Mutation

Antigen Colon antigen 3, positive. The cells are positive for keratin by immunoperoxidase staining. CSAp negative (CSApexpression).

Isoenzymes G6PD, B, PGM1, 1, PGM3, 1-2, 6PGD, A, ES-D, 1, PEP-D, 1

Tumorigenic

Yes, in nude mice

Reverse Negative

transcriptase

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SW-403 Cells | 300350

Products	Carcinoembryonic antigen (CEA) 155 ng/10 exp6 cells/10 days, keratin
Mutational profile	SW-403 cells carry a heterozygous Kras mutation in codon12: GGT>GTT
Handling	
Culture Medium	Ham's F12, w: 1.0 mM stable Glutamine, w: 1.0 mM Sodium pyruvate, w: 1.1 g/L NaHCO3 (Cytion article number 820600a)
Medium supplements	Supplement the medium with 10% FBS
Passaging solution	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.
Split ratio	A ratio of 1:2 to 1:6 is recommended
Fluid renewal	1 to 2 times per week
Freeze medium	CM-1 (Cytion catalog number 800100) or CM-ACF (Cytion catalog number 806100)



SW-403 Cells | 300350

Handling of cryopreserved cultures

- 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. Optionally, skip centrifugation but remove any remaining freezing medium after 24 hours.
- 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.

Quality control / Genetic profile / HLA

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.





STR profile Amelogenin: x,x

CSF1PO: 10,13
D13S317: 13
D16S539: 10,12
D5S818: 11
D7S820: 8,9
THO1: 6
TPOX: 8,9
vWA: 14,18
D3S1358: 15
D21S11: 28,29
D18S51: 17
Penta E: 5
Penta D: 9
D8S1179: 11
FGA: 19

HLA alleles A*: 02:05:01, 03:01:01

B*: 07:02:01, 49:01:01 **C***: 07:01:01, 07:02:01 **DRB1***: 04:01:01, 04:05:01

DQA1*: 03:03:01

DQB1*: 03:01:01, 03:02:01

DPB1*: 04:01:01 **E**: 01:03:02, 01:03:05