



General information

Organism	Human
Tissue	Blood
Disease	Plasma Cell Leukemia
Applications	3D cell culture, Immune system disorder research, Immunology
Synonyms	ARH 77, ARH77

Characteristics

Age	33 years
Gender	Female
Ethnicity	European
Morphology	Lymphoblast
Cell type	B lymphoblast
Growth properties	Suspension

Identifiers / Biosafety / Citation

Citation	ARH-77 (Cytion catalog number 300306)
Biosafety level	2

Expression / Mutation

Antigen expression	CD11a +, CD19 +, CD20 +, CD28 +, CD38 -, CD49e, +CD3 -, CD10 -, CD13 -, CD19 +, CD20 +, CD34 -, CD37 +, CD71 +, cyCD79 +, CD80 +, CD138 -, HLA-DR +, sm/cylgG +, sm/cylgM -, sm/cykappa +, sm/cylambda -
Viruses	EBV + (transformant), HBV -, HCV -, HIV-1 -, HIV-2 -, HTLV-1/2 -, MLV -, SMRV



ARH-77 Cells | 300306

Karyotype	human near diploid karyotype with 8% polyploidy - $46(44-48)xx$, +9, $del(1)(q23)$, $add(2)(q21)$, $add(3)(p11)$, $der(3)t(2,3)(q23,q26)$, $del(6)(p21)$, $der(9)t(9,17)(q10,q10)$ - $sideline$ with $der(x)t(x,1)(q23,p32)$, $del(16)(p13.2)$ and $unresolved$ $der(3)$ and $der(9)hsr$ markers - no IGH translocations detected
Handling	
Culture Medium	RPMI 1640, w: 2.1 mM stable Glutamine, w: 2.0 g/L NaHCO3 (Cytion article number 820700a)
Medium	Supplement the medium with 10% FBS

Subculturing

supplements

Gently homogenize the cell suspension in the flask by pipetting up and down, then take a representative sample to determine the cell density per ml. Dilute the suspension to achieve a cell concentration of 1×10^5 cells/ml with fresh culture medium, and aliquot the adjusted suspension into new flasks for further cultivation.

Freeze medium CM-1 (Cytion catalog number 800100) or CM-ACF (Cytion catalog number 806100)



ARH-77 Cells | 300306

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: 6,10 **D13S317**: 11,13 **D16S539**: 9,13 **D5S818**: 10,13 **D7S820**: 7,12 **TH01**: 8,9.3 **TPOX**: 8 **vWA**: 17 **D3S1358**: 16 **D21S11**: 29,30 **D18S51**: 14,16 Penta E: 12 **Penta D**: 10 **D8S1179**: 14,15 **FGA**: 20,21 **D6S1043**: 11,19 **D2S1338**: 17 **D12S391**: 19,20 **D19S433**: 14,15