

Product sheet

Ca AN3 | 300119

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Description Ca AN3 is a cell line derived from a patient with acute myeloid leukemia (AML). It is characterized by a high degree of heterogeneity and is used for studying drug response and resistance in AML. The cell line is maintained in suspension culture in RPMI 1640 medium supplemented with 10% fetal bovine serum (FBS) and 1% penicillin-streptomycin. The cell line is characterized by a high degree of heterogeneity and is used for studying drug response and resistance in AML. The cell line is maintained in suspension culture in RPMI 1640 medium supplemented with 10% fetal bovine serum (FBS) and 1% penicillin-streptomycin.

Organism Human

Tissue Blood, Bone Marrow

Disease Acute Myeloid Leukemia (AML)

Synonyms AN3_CA, AN3-CA, AN3 Ca, AN3CA, AN-3, AN3, Ca AN3, AN3 Ca (ATCC CCL-253) - ATCC CCL-253

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Age 55 years

Gender Male

Ethnicity Caucasian

Morphology Granulocytic

Cell type Myeloid

Growth properties Suspension

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Citation AN3 Ca (ATCC CCL-253) Cytion 300119

Biosafety level 1

NCBI_TaxID 9606

CellosaurusAccession CVCL_0028

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Thawing and Culturing Cells

1. Thaw the cells in a water bath at 37°C. Do not shake the vial. Transfer the cells to a centrifuge tube and centrifuge at 300 x g for 5 minutes. Remove the supernatant and resuspend the cells in 100 µl of culture medium.
2. Seed the cells into a 96-well plate at a density of 100,000 cells per well. Incubate at 37°C with 5% CO₂ for 24 hours.
3. After 24 hours, the cells should be at approximately 70% confluency. Remove the medium and replace with fresh culture medium.
4. Seed the cells into a 96-well plate at a density of 100,000 cells per well. Incubate at 37°C with 5% CO₂ for 24 hours.
5. After 24 hours, the cells should be at approximately 70% confluency. Remove the medium and replace with fresh culture medium.
6. Seed the cells into a 96-well plate at a density of 100,000 cells per well. Incubate at 37°C with 5% CO₂ for 24 hours.
7. After 24 hours, the cells should be at approximately 70% confluency. Remove the medium and replace with fresh culture medium.
8. Seed the cells into a 96-well plate at a density of 100,000 cells per well. Incubate at 37°C with 5% CO₂ for 24 hours.

Incubation Atmosphere 37°C, 5% CO₂, humidified

Flask Coating Cell culture medium

Freezing Procedure Freeze cells in a freezing medium at -80°C

Shipping Conditions Ship cells at -80°C

Storage Conditions Store cells at -150°C for 196 days

HLA

Sterility Sterilized by gamma irradiation

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STR

Amelogenin: x,x
CSF1PO: 12,14,15
D13S317: 12,14
D16S539: 10,14,15
D5S818: 11,14
D7S820: 7.1,10
TH01: 9.3,10
TPOX: 8,1
vWA: 14,19,20,21
D3S1358: 17
D21S11: 29,3
D18S51: 15,17,18
Penta E: 9,16
Penta D: 9,16
D8S1179: 12,14
FGA: 23
D1S1656: 13,18.3
D6S1043: 12,13,14,15,18
D2S1338: 20,23
D12S391: 20,21,23,24,25
D19S433: 14

HLA

A*: 03:01:01
B*: '44:02:01, '57:01:01
C*: '05:01:01, '06:02:01
DRB1*: '04:01:01G, '16:01:01
DQA1*: '01:02:02, '03:01:01
DQB1*: 03:02:01, 05:02:01
DPB1*: '05:01:01G, '13:01:01G
E: 01:03:02