P3x63Ag8.653

400118

GENERAL INFORMATION

Description

The cells are resistant to 8-azaguanine and are HAT sensitive. They can be used as fusion partners for producing hybridomas. The cells do not secrete immunoglobulin. The cells have been reported to be cholesterol auxotroph due to a deficiency in 3-ketosteroid reductase activity.

Tissue	Hematopoietic
Disease	Myeloma

Synonyms

- P3-x63-Ag8.653
- P3-x63-Ag8-653
- P3-x63-Ag8 653
- P3-x63-Ag 8.653
- P3-x63Ag8.653
- P3-x63.Ag8.653
- P3/x63/Ag8.653
- P3x63 Ag8.653
- P3x63 AG8-653
- P3x63-Ag8.653
- P3x63-Ag8.653
- P3x63 AG 8.653
- P3x63Ag8653
- P3-x63-Ag8-6-5-3
- P3x63Ag8-6-5-3
- P3.times.63 Ag8.653
- P3.653
- x63-Ag 8.6.5.3
- x63-AG 8.653
- x63-Ag8-653
- x63-Ag8.653
- x63.Ag8.653
- x63Ag8-653
- x63Ag8.653
- x63AG8.653
- P3-653
- GM03570
- GM3570
- GM03570E
- NS653

CLASSIFICATION

Organism	Mouse
Sex	Female

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Banking information
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BIC: SOLADES1HDB

Growth properties	Adherent and suspension
Biosafety level	1

CHARACTERISTICS

Morphology	Round cells

IDENTIFIERS

Citation

P3x63Ag8.653 (Cytion catalog number 400118)

NCBI TaxID	10090
Cellosaurus	CVCL_4032

EXPRESSION AND SUSCEPTIBILITY

Viruses

Tested negative for ectromelia virus (mouse pox).

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 10% FBS

Dissociation reagent	Accutase

Subculturing

Gather the suspension cells in a 15 ml tube and gently wash the adherent cells with PBS lacking calcium and magnesium (use 3-5 ml for T25 flasks and 5-10 ml for T75 flasks). Apply Accutase (1-2 ml for T25 flasks, 2.5 ml for T75 flasks) ensuring full coverage of the cell layer. Allow the cells to incubate at room temperature for 10 minutes. Following incubation, combine and centrifuge both the suspension and adherent cells. After centrifugation, carefully resuspend the cell pellet and transfer the cell suspension into new flasks containing fresh medium.

Seeding density	Start new cultures at 4 x 10^5 cells/ml. The cell density should not exceed 2 x 10^6 cells/ml.
Fluid renewal	Every 3 to 4 days. Collect floating cells, centrifuge and add to the flask together with fresh medium.
Post-thaw recovery	After thawing, plate the cells at 5 x 10^4 cells/cm^2 and allow the cells to recover from the freezing process and to adhere for at least 48 hours.

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

Breed / subspecies	BALB/c

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.

Disclaimer

Our cells are provided for in vitro laboratory research purposes exclusively and are not intended for clinical or diagnostic use, nor are they to be administered to humans or used for veterinary purposes. Users must adhere to all applicable guidelines and regulations for the handling and use of these cells in a research setting.

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

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