RPMI 8226

300431

GENERAL INFORMATION

Description

RPMI 8226 cells are a human myeloma cell line that was established in 1966 from the peripheral blood of a 61-year-old male patient with multiple myeloma. This cell line was named after the Roswell Park Memorial Institute (RPMI) where it was developed, and the number 8226 denotes its specific catalog number in the cell bank.
The RPMI 8226 cell line is an important model system for studying multiple myeloma and related aspects of plasma cell biology, immunology research, and cancer therapy. RPMI 8226 cells are known to produce and secrete kappa light chains of immunoglobulins, a feature that is often exploited in research studies to investigate antibody production and secretion mechanisms.
RPMI 8226 cells exhibit numerous chromosomal abnormalities, which are typical of multiple myeloma cells. These include translocations, deletions, and amplifications that affect various oncogenes and tumor suppressor genes.
The human myeloma cell line RPMI 8226 are widely used in drug discovery and development research, and have been used to investigate drug resistance pathways and evaluating combination therapies.
In summary, RPMI 8226 cells provide a critical in vitro model for multiple myeloma research, enabling the investigation of the biological and molecular mechanisms underlying this disease and the development of therapeutic strategies.

Tissue	Peripheral blood
Disease	Multiple Myeloma

Synonyms

- RPMI-8226
- RPMI.8226
- RPMI8226
- RPMI no. 8226
- RPMI no 8226
- RPMI #8226
- 8226
- RPMI 8226/S
- RPMI-8226S
- RPMI8226/S
- 8226/S
- Roswell Park Memorial Institute 8226
- GM02132
- GM2132
- GM 2132
- GM02132C
- Simpson

CLASSIFICATION

Organism	Human
Sex	Male
Growth properties	Suspension
Biosafety level	1

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Banking information
Sparkasse Heidelberg
IBAN: DE35 6725 0020 0009 2076 86
BIC: SOLADES1HDB

CHARACTERISTICS

Morphology	Round cells

IDENTIFIERS

Citation

RPMI 8226 (Cytion catalog number 300431)

NCBI TaxID	9606
Cellosaurus	CVCL_0014

EXPRESSION AND SUSCEPTIBILITY

Antigen expression

HLA Aw19, B15, B37, Cw2

Isoenzymes

G6PD, A

Reverse transcriptase	Negative

Products

Immunoglobulin light chain

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 5 x 10^5 viable cells/ml. Subculture at 1-2 x 10^6 cells/ml. Maximum cell density is at 1-2 x 10^6 cell/ml.
Fluid renewal	2 to 3 times per week
Post-thaw recovery	After thawing allow the cells to recover from the freezing process for at least 24 hours.

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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	61 years

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