

HARA-B Cells | 300465**General information****Description**

The HARA-B cell line is derived from human squamous cell carcinoma of the lung, specifically established from metastatic bone tissue in a mouse model. This cell line is a secondary development from the original HARA cell line and is characterized by its high expression of parathyroid hormone-related protein (PTHrP), which plays a significant role in the extensive bone metastasis observed in these cells. The HARA-B line has been instrumental in studying the mechanisms of bone metastasis associated with lung cancer.

Scientific studies involving HARA-B often focus on its utility in modeling hypercalcemia, a common paraneoplastic syndrome associated with certain cancers, including lung cancer. The hypercalcemia in this model is induced by hypodermic transplantation of the cells, providing a valuable tool for understanding the interactions between cancer cells and bone cells, as well as the pathways that lead to bone degradation and calcium release. This cell line helps researchers investigate potential therapeutic strategies to mitigate bone metastasis and associated complications in lung cancer patients.

Organism

Human

Tissue

Lung

Disease

Lung squamous cell carcinoma

Metastatic site

Pleural effusion

Synonyms

HARAB

Characteristics**Age**

57 years

Gender

Male

Ethnicity

Japanese

Growth properties

Adherent

Regulatory Data**Citation**

HARA-B (Cytion catalog number 300465)

NCBI_TaxID

9606

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CellosaurusAccession CVCL_2915**Biomolecular Data****Protein expression** Produces high level of parathyroid hormone-related peptide (PTHrP).**Handling****Culture Medium** RPMI 1640, w: 2.0 mM stable Glutamine, w: 2.0 g/L NaHCO₃ (Cytion article number 820700a)**Supplements** Supplement the medium with 10% FBS**Dissociation Reagent** 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.**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.

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Thawing and Culturing Cells

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.
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.

Incubation Atmosphere

37°C, 5% CO₂, humidified atmosphere.

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.

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.

Quality Control & Molecular Analysis

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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.