

HK EGFP-alpha-tubulin/H2B-mCherry Cells | 300670

General information

Description

The HK EGFP-alpha-tubulin/H2B-mCherry HeLa Kyoto cell line is a meticulously engineered model designed for detailed visualization of cellular processes. This clonal line has been stably transfected to express two fluorescent protein fusions that enable real-time imaging of both chromatin and the microtubular network. The red fluorescent protein mCherry is fused to the core histone protein H2B, creating H2B-mCherry. This fusion protein is expressed from the pH2B-mCherry-IRES-neo3 plasmid and serves as a chromatin marker, highlighting the nuclear DNA in live-cell imaging and facilitating studies on chromatin dynamics and nuclear architecture.

Additionally, this cell line expresses monomeric enhanced GFP (Green Fluorescent Protein) fused to α -tubulin, introduced via the pmEGFP- α -tubulin-IRES-puro2b plasmid. The GFP- α -tubulin fusion provides a vivid green fluorescence that outlines the microtubule structures within the cell. This feature is crucial for studying microtubule organization, dynamics, and their role in cell division and intracellular transport. The stable integration of these constructs allows for continuous, long-term observation of these cellular components without the need for repeated transfection, thus reducing variability and enhancing the reliability of experimental results. Drug resistance selection following transfection ensures the stability and uniformity of expression among the cells in this line.

Organism

Human

Tissue

Cervix

Disease

Carcinoma

Synonyms

HeLa Kyoto EGFP- α -tubulin/H2B-mCherry, HeLa H2B-mRFP and mEGFP-alpha-tubulin

Characteristics

Age

30 years

Gender

Female

Ethnicity

African American

Morphology

Epithelial-like cells with mosaic stone shape

Growth properties

Monolayer, adherent

Regulatory Data

Citation

HK EGFP-alpha-tubulin/H2B-mCherry (Cytion catalog number 300670)

HK EGFP-alpha-tubulin/H2B-mCherry Cells | 300670

Biosafety level 1**NCBI_TaxID** 9606**CellosaurusAccession** CVCL_L802**Depositor** The Ellenberg Lab (EMBL)**GMO Status** GMO-S1: This HeLa Kyoto line contains EGFP- α -tubulin and H2B-mCherry constructs for simultaneous imaging of microtubules and chromatin. This classification applies only within Germany and may differ elsewhere.

Biomolecular Data

Protein expression EGFP-alpha-tubulin, H2B-mCherry: Location/Gene: 1..589 / Pcmv, 652..1029 H2B, 1042..1752 / mCherry, 2983..3777 / KanR/NeoR**Viruses** Negative for HIV, HBV and HCV.**Products** CMV Promotor, Histone H2B, Neomycin, Phosphotransferase

Handling

Culture Medium DMEM, w: 4.5 g/L Glucose, w: 4 mM L-Glutamine, w: 3.7 g/L NaHCO₃, w: 1.0 mM Sodium pyruvate (Cytion article number 820300a)**Supplements** Supplement the medium with 10% FBS**Dissociation Reagent** Accutase**Doubling time** 24 hours**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.**Seeding density** 1×10^4 cells/cm²

HK EGFP-alpha-tubulin/H2B-mCherry Cells | 300670

Fluid renewal 2 to 3 times per week

Post-Thaw Recovery After thawing, plate the cells at 5×10^4 cells/cm² and allow the cells to recover from the freezing process and to adhere for at least 24 hours.

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.

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.

HK EGFP-alpha-tubulin/H2B-mCherry Cells | 300670

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

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.