

HaCaT-ras II-4 | 300495

General Information

Description	<p>HaCaT-ras II-4 is a cell line derived from HaCaT cells. It is a clone of HaCaT-ras II-4, which was established from a Balb/c-nu/nu mouse. The cell line is characterized by its high growth rate and its ability to form colonies in soft agar. It is a clone of HaCaT-ras II-4, which was established from a Balb/c-nu/nu mouse. The cell line is characterized by its high growth rate and its ability to form colonies in soft agar.</p>
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Organism	Human
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Tissue	Epithelial
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Synonyms	HaCaT-ras clone II-4, HaCaT II-4, II-4
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Characteristics

Age	62 years
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Gender	Male
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Ethnicity	White
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Cell type	Epithelial
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Growth properties	Adherent
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References and Safety

Citation	HaCaT-ras II-4 (Cytion 300495)
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Biosafety level	1
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NCBI_TaxID	9606
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CellSaurusAccession	CVCL_3868
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Product sheet

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GMO Status GMO-S1: (HaCaT-ras II-4) c-Ha-Ras

Protein expression P53 (+), CEA (+),

Tumorigenic Balb/c-nu/nu.

Karyotype ()

Culture Medium DMEM, w: 4.5 g/L, w: 4 mM L-, w: 3.7 g/L NaHCO₃, w: 1.0 mM (Cytion 820300a)

Supplements 10% FBS

Dissociation Reagent 1:1 EDTA (0.05%) (0.1%) PBS Ca₂+ /EDTA.

Subculturing

- 1.
2. PBS (T25, 5-10 T75,)
3. EDTA 0.05% - 1-2 T25 2.5 T75
4. 37 10
5. /EDTA (0.05%, 0.025% EDTA),
6. 1-2
7. FBS
- 8.

Seeding density 1×10^4 /

Fluid renewal

Freeze medium (FBS) + 10% DMSO

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Thawing and Culturing Cells

1. Thaw the vial rapidly in a water bath at 37°C. Remove the vial and centrifuge at 300 x g for 3 minutes. Discard the supernatant and resuspend the cells in 10 ml of complete DMEM medium.
2. Seed the cells into a 75 cm² flask containing 30 ml of complete DMEM medium. Incubate at 37°C with 5% CO₂ until cells reach 70-80% confluency.
3. Harvest the cells by trypsinization and seed into a 75 cm² flask containing 30 ml of complete DMEM medium. Incubate at 37°C with 5% CO₂ until cells reach 70-80% confluency.
4. Harvest the cells by trypsinization and seed into a 75 cm² flask containing 30 ml of complete DMEM medium. Incubate at 37°C with 5% CO₂ until cells reach 70-80% confluency.
5. Harvest the cells by trypsinization and seed into a 75 cm² flask containing 30 ml of complete DMEM medium. Incubate at 37°C with 5% CO₂ until cells reach 70-80% confluency.
6. Harvest the cells by trypsinization and seed into a 75 cm² flask containing 30 ml of complete DMEM medium. Incubate at 37°C with 5% CO₂ until cells reach 70-80% confluency.
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8. Harvest the cells by trypsinization and seed into a 75 cm² flask containing 30 ml of complete DMEM medium. Incubate at 37°C with 5% CO₂ until cells reach 70-80% confluency.

Incubation Atmosphere 37°C, 5% CO₂, humidified

Flask Coating None

Freezing Procedure Harvest cells by trypsinization and seed into a 75 cm² flask containing 30 ml of complete DMEM medium. Incubate at 37°C with 5% CO₂ until cells reach 70-80% confluency.

Shipping Conditions Harvest cells by trypsinization and seed into a 75 cm² flask containing 30 ml of complete DMEM medium. Incubate at 37°C with 5% CO₂ until cells reach 70-80% confluency.

Storage Conditions Harvest cells by trypsinization and seed into a 75 cm² flask containing 30 ml of complete DMEM medium. Incubate at 37°C with 5% CO₂ until cells reach 70-80% confluency.

Genotype / Phenotype / HLA

Sterility The cells are free of mycoplasma contamination. PCR screening for mycoplasma contamination is negative.