# HaCaT-ras II-4

300495

## GENERAL INFORMATION

**Description**

[illegible] HaCaT-ras II-4 [illegible] HaCaT [illegible] c-Ha-ras (EJ) onogene[illegible] G418 [illegible] Boukamp [illegible] 1990

[illegible] HaCaT-ras II-4 [illegible] Balb/c-nu/nu [illegible]

[illegible] HaCaT-ras II-4 [illegible]

[illegible]

[illegible] HaCaT-ras II-4 [illegible]

[illegible] HaCaT-ras II-4 [illegible] c-Ha-ras (EJ) [illegible] HaCaT-ras II-4 [illegible]

| | |
|---|---|
| **Tissue** | [illegible] |

**Synonyms**

- HaCaT-ras clone II-4
- HaCaT II-4
- II-4

## CLASSIFICATION

| | |
|---|---|
| **Organism** | Human |
| **Sex** | Male |

| | |
|---|---|
| **Growth properties** | Adherent |
| **GMO status** | GMO-S1 |

**GMO status details**

GMO-S1: This human keratinocyte line (HaCaT-ras II-4) contains a plasmid encoding c-Ha-Ras oncogene sequences introduced by transfection, enabling transformed growth behavior. The construct is integrated into HaCaT-derived keratinocytes. This classification applies only within Germany and may differ elsewhere.

| | |
|---|---|
| **Biosafety level** | 1 |

## CHARACTERISTICS

| | |
|---|---|
| **Cell type** | □□□□□□□□□□□□□ |
| **Tumourigenic** | □□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□ Balb/c-nu/nu |

**Karyotype**

□□□□□□□□□□□ (□□□□□□□□□□□□□□□□□□□)

## IDENTIFIERS

**Citation**

HaCaT-ras II-4 (□□□□□□□□□□□□□□□□□□□□ Cytion 300495)

| | |
|---|---|
| **NCBI TaxID** | 9606 |
| **Cellosaurus** | [CVCL_3868] |

## EXPRESSION AND SUSCEPTIBILITY

**Protein expression**

P53 (+), CEA (+),

## HANDLING AND CULTURE

**Culture medium**

DMEM, w: 4.5 □□□□/□□□□ □□□□□□, w: 4 □□□□□□□□□□□ □□□-□□□□□□□□□, w: 3.7 □□□□/□□□□ NaHCO3, w: 1.0 □□□□□□□□ □□□□ □□□□□□□□□□□□□□□□ (□□□□□□□□□□□□□ Cytion 820300a)

| | |
|---|---|
| **Incubation atmosphere** | 37°C, 5%CO2, □□□□□□□□□□□□□ |

**Supplements**

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| | |
|---|---|
| **Dissociation reagent** | □□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□ 1:1 □□□ EDTA (□□□□□□ 0.05%) □□□□□□□□ □□□ (□□□□□□: 0.1%) □□□□□□□□□□□□□□□□□□□□□□□□□□ □□□□□□□ PBS □□□□□□□□□ Ca2+ □□□ Mg2+ □□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□/EDTA □□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□ □□□□□□□□□□□□□□□□□□□□□□□□□ TrypLETM Express (Life Technologies) □□□□□□□□□□□□/EDTA □□□ □□□ □□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□ |

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### Subculturing

[illegible]: [illegible]
[illegible]: [illegible] PBS ([illegible]) 3-5 [illegible]. [illegible] T25 [illegible] 5-10 [illegible]. [illegible] T75 [illegible]
[illegible] EDTA: [illegible] EDTA 0.05% [illegible] - [illegible] 1-2 [illegible] T25 [illegible] 2.5 [illegible] T75
[illegible]: [illegible] 37 [illegible] 10 [illegible]
[illegible]/EDTA: [illegible]/EDTA [illegible] ([illegible] 0.05%, EDTA 0.025%) [illegible]—[illegible] 1 [illegible]. [illegible] T25 [illegible] 2.5 [illegible]. [illegible] T75
[illegible]: [illegible] 1-2 [illegible]
[illegible]: [illegible] FBS [illegible]
[illegible]: [illegible]

| | |
|---|---|
| **Seeding density** | 1 x 10^4 [illegible]/[illegible].^² |
| **Fluid renewal** | 2 [illegible] |

### 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

[illegible] PCR [illegible]
[illegible]

| | |
|---|---|
| **Freeze medium** | [illegible] ([illegible] FBS) + 10% DMSO [illegible] CM-1 ([illegible] Cytion 800100) [illegible] |

### Storage conditions

[illegible] −150 [illegible] −196 °C [illegible] −80 °C [illegible]

### Shipping conditions

[illegible] −78 °C [illegible]

## DONOR

| | |
|---|---|
| **Donor age** | 62 □□ |
| **Ethnicity** | □□□□□□□□□□ |

## LEGAL AND SAFETY

### Safety precautions

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### Warranty

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|---|---|
| **Third-party agreements apply** | Yes |