

Product sheet

HROG04 | 300932

Description [REDACTED] PD Dr. Michael Linnebacher [REDACTED]

Organism [REDACTED]

Tissue [REDACTED], R, [REDACTED]

Disease [REDACTED] ([REDACTED] IV)

Age 53 [REDACTED]

Gender [REDACTED]

Ethnicity [REDACTED]

Morphology [REDACTED]

Growth properties [REDACTED]

Citation HROG04 ([REDACTED] Cytion 300932)

Biosafety level 1

NCBI_TaxID 9606

CellSaurusAccession CVCL_4U39

Antigen expression HLA-A02 +, MHC class I -MHC class II -, Beta-microglobulin +, HLA-E +, HLA-G -, MIC A -, MIC-B -, ICAM-1 +, GFAP +, nestin +, vimentin +, S-100 +, GBM +, BTSC +

Mutational profile PTENW274L, 9p212.3(CDKN2A) [REDACTED]

Product sheet

HROG04 | 300932

Culture Medium DMEM:Ham's F12 (1:1), w: 3.1 g/L, w: 2.5 g/L L-Ascorbic acid, w: 15 mg/L HEPES, w: 0.5 g/L

Supplements FBS 10%

Dissociation Reagent Trypsin

Subculturing PBS, T25

Seeding density 1×10^4 cells/cm²

Fluid renewal 3-5 days

Freeze medium 50% basal medium + 40% FBS + 10% DMSO, CM-1

Thawing and Culturing Cells

1. Thaw cryovial in 37°C water bath
2. Transfer cells to 15 ml centrifuge tube
3. Centrifuge at 300 x g for 3 min
4. Remove supernatant and resuspend in 10 ml fresh medium
5. Seed cells into T25 flask
6. Incubate at 37°C, 5% CO₂
7. Monitor cell growth and confluency
8. Pass cells when 70-80% confluent

Incubation Atmosphere 37°C, 5% CO₂

