

## Human Primary Glioblastoma Cancer Cells

A primary cell isolate with application in cell-based screening and life science research

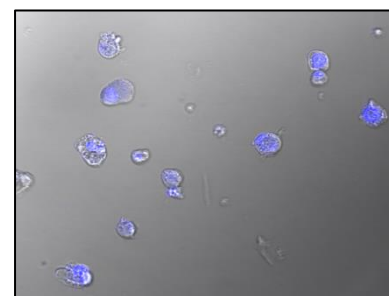
The primary cell isolate was prepared from human tissue obtained with full ethical permission. Tissue was dissociated by enzymatic digestion and cells were harvested and washed by conventional filtration and centrifugation. Cells were grown in culture using medium optimised for primary glioblastoma cancer cells, and were banked and cryopreserved under liquid nitrogen. Cell population analysis was performed by fluorescence flow cytometry.

### DONOR TISSUE FEATURES

- Male donor, 75 years
- Glioblastoma multiforme, tumour stage IV
- Additional donor history available on request

### CELL CHARACTERISTICS

Batch number:	11-1309
Vial content:	0.5x10 <sup>6</sup> cells
Appearance:	Dependent upon culture conditions
Seeding density:	13,500 cells/cm <sup>2</sup> (adherent culture), 2x10 <sup>5</sup> cells/ml (suspension culture)
Culture medium:	AvantiCell medium (GM-HCM-01) recommended
Recovery from frozen:	> 90%
Mycoplasma test:	Negative (by luminescent-based mycoplasma assay)
Virus tests:	HIV1, HIV2, HAV, HBV, HCV, HTLV1/2 (Negative by real time PCR screen)
Other tests:	Yeast, fungus (Negative)



Cell morphology. DAPI stained neurospheres in suspension culture fixed in situ (Magnification: x40).

### FLOW CYTOMETRY CELL ANALYSIS

Cell Marker	Target Description	Population Positive*
CD33	Cancer stem cell marker	Negative
CD56	Neuronal progenitor cell marker	75%
A2B5	Glial progenitor cell marker	25%

\*Percentage of cells with fluorescence greater than the isotype control background

### USES AND RESTRICTIONS

- Store at -150°C. Once thawed do not re-freeze
- For research use ONLY — not suitable for *in vitro* diagnostic use or human or animal treatment
- Potential biohazard — handle with care

## Leaders in Cell Culture