

## Human Primary Dermal Fibroblasts

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

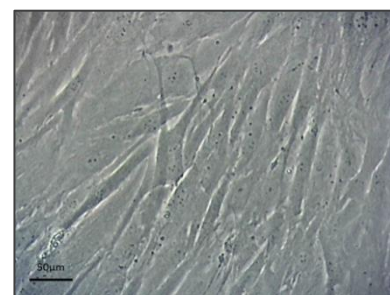
The primary cell isolate was prepared from human tissue obtained with full ethical permission. Skin tissue was dissected to obtain explants of dermis which were seeded into tissue culture treated dishes. Cell outgrowth from explanted tissue was promoted using customised medium optimised for primary human dermal fibroblast cells. A fibroblast population was isolated using FSP-1 immuno-magnetic selection. Propagated cells were banked and cryopreserved under liquid nitrogen. The cell population was analysed by fluorescence-activated flow cytometry.

### DONOR TISSUE FEATURES

- Female donor, Caucasian, 59 years old
- Isolated from abdominal skin
- Additional donor history available on request

### CELL CHARACTERISTICS

Batch number:	14-1906
Vial content:	0.5x10 <sup>6</sup> cells
Appearance:	Monolayer of spindle-shaped cells
Seeding density:	5,000 cells/cm <sup>2</sup>
Culture medium:	AvantiCell medium (DF-HNM-01) recommended
Recovery from frozen:	94%
Population doubling:	2-3 days
Mycoplasma test:	Negative (by real-time PCR)
Virus tests:	Negative for HIV1, HIV2, HBV, HCV (by serology screen)
Other tests:	Negative for yeast, fungus, bacteria



Cell morphology. Cells in culture were photographed using a phase contrast microscope. (Bar 50µm)

### FLOW CYTOMETRY CELL ANALYSIS

Cell Marker	Target Description	Population Positive*
Ecto-5' nucleotidase	Interstitial fibroblast marker	99.10%
α-smooth muscle actin (α-SMA)	Myofibroblast marker	90.75%

\*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