

## **MELANOCYTES**

ATCC® Normal Human Primary Epidermal Melanocytes from Neonatal Foreskin, when grown in Dermal Cell Basal Medium supplemented with Melanocyte Growth Kit components, provides an ideal cell system for propagation in low-serum (less than 1.0% FBS) conditions in the absence of cholera toxin and phorbol 12-myristate 13-acetate (PMA).

Each lot of ATCC® Normal Human Primary Epidermal Melanocytes is:

- Cryopreserved in the second passage to ensure the highest viability and plating efficiency.
- Performance tested together with ATCC® Primary Cell Solutions™ media, kit supplements and reagents to guarantee optimum reliability.
- Thoroughly tested for sample purity as part of the ATCC commitment to quality.

Applications for use might include research related to melanoma; dermal response to UV radiation; psoriasis and other skin diseases; skin trauma (e.g., wound repair, scars, burns); and cosmetic research (e.g., skin lightening compounds, skin protecting compounds).

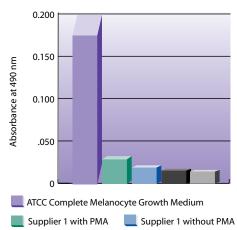
#### OPTIMIZED GROWTH MEDIUM MAKES A DIFFERENCE

Primary melanocytes are effectively supported by the complete ATCC® Primary Cell Solutions™ dermal cell system consisting of Dermal Cell Basal Medium supplemented with the Melanocyte Growth Kit. This unique formulation — without cholera toxin or PMA —is designed to produce cultures with:

- Functional expression of relevant biomarkers;
- Normal morphology; and
- Superior growth rates.

Use of this complete system eliminates the need for additional components such as feeder layers, extracellular matrix proteins or other substrates to enhance attachment and proliferation.

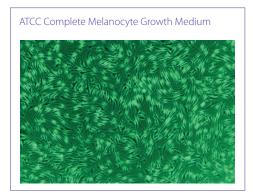
### Melanocytes Grown in Different Brands of Low Serum or Serum-Free Media: Levels of L-DOPA Oxidase Activity



ATCC Primary Cell Solutions melanocytes were taken from liquid nitrogen and cultures initiated. The cells were grown for 4 days and then seeded in triplicate into a 24-well plate at 2,500 cells/cm<sup>2</sup> and grown for 6 days in different brands of low serum or serum-free media plus or minus PMA. The medium was not changed during the incubation period. L-DOPA Oxidase activity was measured by adding 500 µl of assay buffer, 300 µl MBTH reagent, and 200 µl of 5 mM L-DOPA to each well, incubating for 30 min at 37°C, and then measuring absorbance at 490 nm using a Wallac VICTOR2™ MultiLabel Counter. The higher the absorbance at 490 nm, the greater the level of tyrosine activity. a functional marker for melanocytes.

Supplier 1 with PMA Supplier 3 without PMA, without FBS

### Comparison of Melanocyte Morphology and Cell Density in Different Medias\*







<sup>\*</sup> Day 6 of the L-DOPA Oxidase functional biomarker experiment.

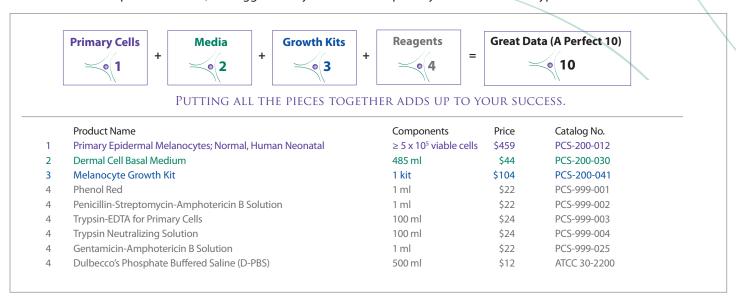
Growth Rate Comparison: Primary Melanocytes Cultured in Different Brands of Medium (21.2 Days)\*

Medium	Number of Doublings	Average Doubling Time (hrs)	Number of Passages
ATCC Complete Melanocyte Growth Medium	15	33.5	\ 4
Supplier 1 Medium without PMA	7	73.6	\3
Supplier 2 Medium without PMA	5.5	88.9	3

<sup>\*</sup>This experiment was conducted while various lots of ATCC® Primary Cell Solutions™ melanocytes were undergoing QC testing. When the QC-specification for population doublings was achieved (15) the experiment was concluded at the end of 21.2 days of testing.

#### **ORDERING INFORMATION**

To achieve the best possible results, we suggest that you order a complete system for each cell type:



Additional cells/cell types will be added in the coming months. Visit us online at **www.atcc.org** to bookmark the primary cell page for easy reference.

# SUPERIOR QUALITY. EXPERT SUPPORT. RELIABLE RESULTS.

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