

# **Certificate of Analysis**

# Strain C57BL/6 Mouse Mesenchymal Stem Cells With RFP

Catalog No. MUBMX-01201 Lot Number: 120329J01

Cryopreservation Date: 2012-03-29 Passage Number: 9

### Viability

Cells are assayed for viability post-thaw using vital staining assay with trypan blue. Specification: Cells should exhibit  $\ge 80\%$  viability.

## Sterility

Bacterial and Fungal Contamination: Samples are inoculated and cultured in blood agar plate, thioglycolate broth, tryptocase soy broth and sabouraud dextrose agar. Specification: No growth must be observed.

Mycoplasma: Samples are tested for mycoplasma contamination using a PCR-based assay and direct culture.

Specification: Results must be negative.

Endotoxin: Samples are tested for endotoxin contamination with LAL test. Specification: Results must show a concentration of  $\leq 25$ EU/ml.

# Purity

Cells are assayed for purity using flow cytometric analysis of cell surface antigen expression after cryopreservation. Cells are immunofluorescently stained with fluorochrome-conjugated antibodies specific to cell surface antigens CD29, CD34, CD44, CD117 and Sca 1. Specification: Cells must show  $\geq$  70% positivity for expression of cell surface antigens CD29, CD34, CD44 and Sca-1. Cells must show  $\leq$  5% positivity for expression of cell surface antigens CD117.

### **Proliferation Ability**

Cells are characterized by their ability to proliferate in culture with an attached well-spread morphology for  $\geq$  5 passages, and  $\leq$  5% cells exhibit spontaneous differentiation in each passage.

### **RFP Expression**

Expression of constitutive RFP is assayed by visual inspection of RFP fluorescence signal. Specification: The results must indicate  $\geq$ 80% of cells are visually inspected for RFP fluorescence signal during extensive subcultivation.



# **Differentiation Ability**

Cells are assayed after cryopreservation for their ability of tri-lineage differentiation. Cells must be able to differentiate to osteocytes, adipocytes and chondrocytes when cultured in the appropriate differentiation media.

#### **Results:**

All specifications have been met.

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