

Protocol for passaging of ASC/TERT1

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Evercyte Ord. No.:	CHT-001-0005
Designation:	ASC/TERT1, human mesenchymal stromal cells
Growth medium:	EGM TM -2 Endothelial Cell Growth Medium-2 (Lonza, Cat# CC-3162) supplemented with FBS and G418: <u>Final components:</u> EBM TM -2 basal medium (Lonza, Cat# CC-3156) Components of EGM TM -2 SingleQuots TM (Lonza, Cat# CC-4176: Hydrocortisone, hFGF, VEGF, R3-IGF-1, Ascorbic acid, hEGF, Heparin) 4 % FBS (PAN Biotech, Cat# P30-3031) 200 µg/ml G418 (InvivoGen, Cat# ant-gn5, 100 mg/ml, ready-to-use) <ul style="list-style-type: none">- take one bottle of EBM-2 basal medium (500 ml)- add 20 ml of FBS (ready-to-use)- add 200 µl of Hydrocortisone (EGMTM-2 SingleQuotsTM)- add 2 ml of hFGF (EGMTM-2 SingleQuotsTM)- add 500 µl VEGF (EGMTM-2 SingleQuotsTM)- add 500 µl of R3-IGF-1 (EGMTM-2 SingleQuotsTM)- add 500 µl of hEGF (EGMTM-2 SingleQuotsTM)- add 500 µl of Heparin (EGMTM-2 SingleQuotsTM)- add 500 µl of Ascorbic Acid (EGMTM-2 SingleQuotsTM)- add 1 ml of G418 stock solution - mix properly - store at 4°C for a maximum of 4 weeks- temper the medium to room temperature (not 37°C) before use
Additional reagents:	0.05% Trypsin-EDTA (Gibco, Cat# 25300-054, ready-to-use, stored at 4°C) Defined Trypsin Inhibitor (Gibco, Cat# R007100, ready-to-use, stored at 4°C) PBS (Sigma-Aldrich, Cat# D8537, ready-to-use, stored at RT)
Passaging of cells:	<ul style="list-style-type: none">- remove and discard the culture medium- wash the cells twice with PBS (each 160 µl/cm²), remove PBS completely- add Trypsin-EDTA solution (20 µl/cm²), make sure that all cells have been in contact with this solution- incubate the culture flask at 37°C for approximately 2-3 min- observe cell detachment under an inverted microscope

- as soon as all cells are detached (if necessary, agitate the cells by gently hitting the flask), add Defined Trypsin Inhibitor (20 $\mu\text{l}/\text{cm}^2$)
- resuspend the cells in growth medium (about 160 $\mu\text{l}/\text{cm}^2$) and aspirate the cells by pipetting
- centrifuge at 170 g for 5 min
- discard the supernatant, resuspend the cell pellet in the remaining droplet and add growth medium
- transfer appropriate aliquots of the cell suspension to culture vessels supplemented with growth medium (final volume of 240 $\mu\text{l}/\text{cm}^2$)
- a split ratio of 1:3 to 1:4 twice a week is recommended (after having reached about 80 % confluence)
- cultivate cells at 37°C in a humidified atmosphere with 5% CO₂

Related products: WJ-MSC/TERT273, human mesenchymal stromal cells (Evercyte, Cat# CHT-021-0273)
