

Label-Free Density-Based Detection of Adipocytes of Bone Marrow Origin Using Magnetic Levitation

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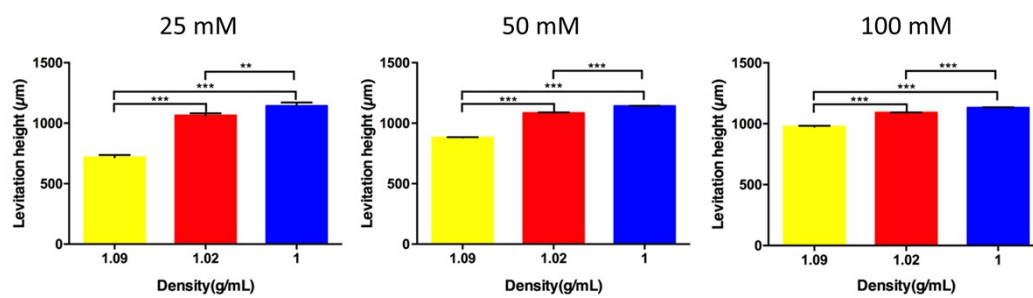
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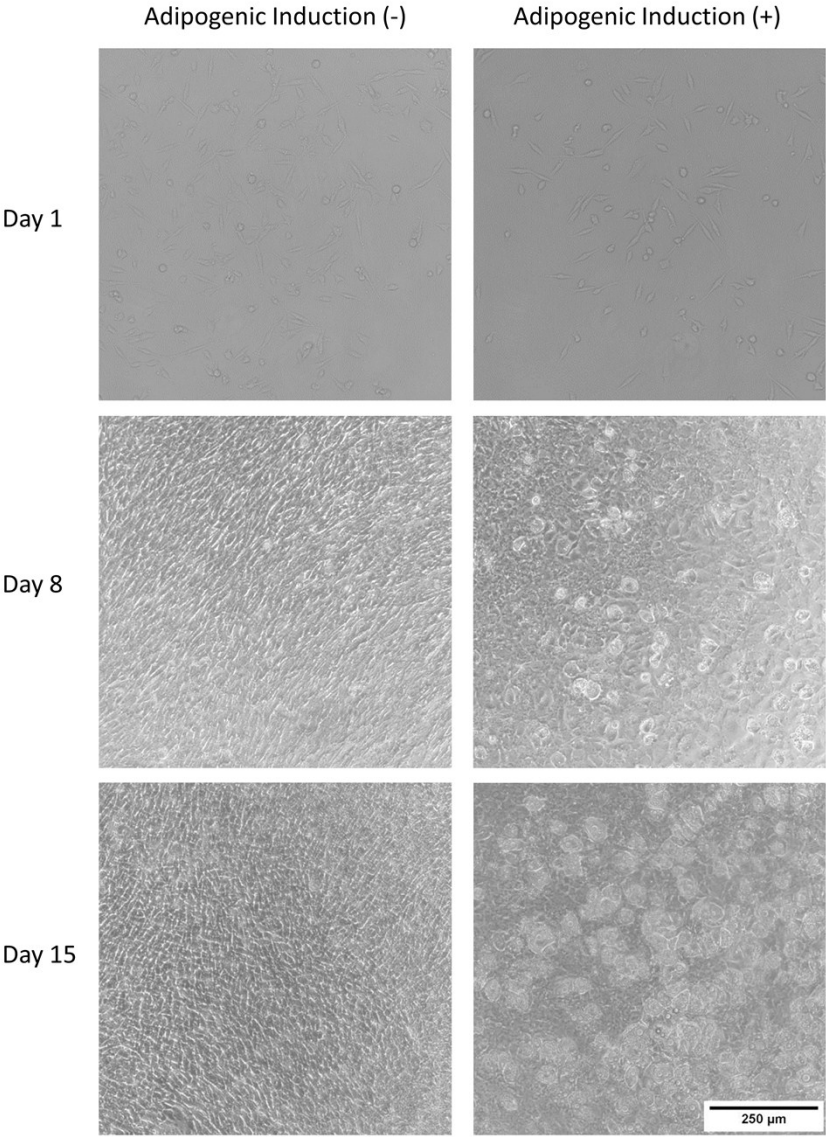
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Figure captions

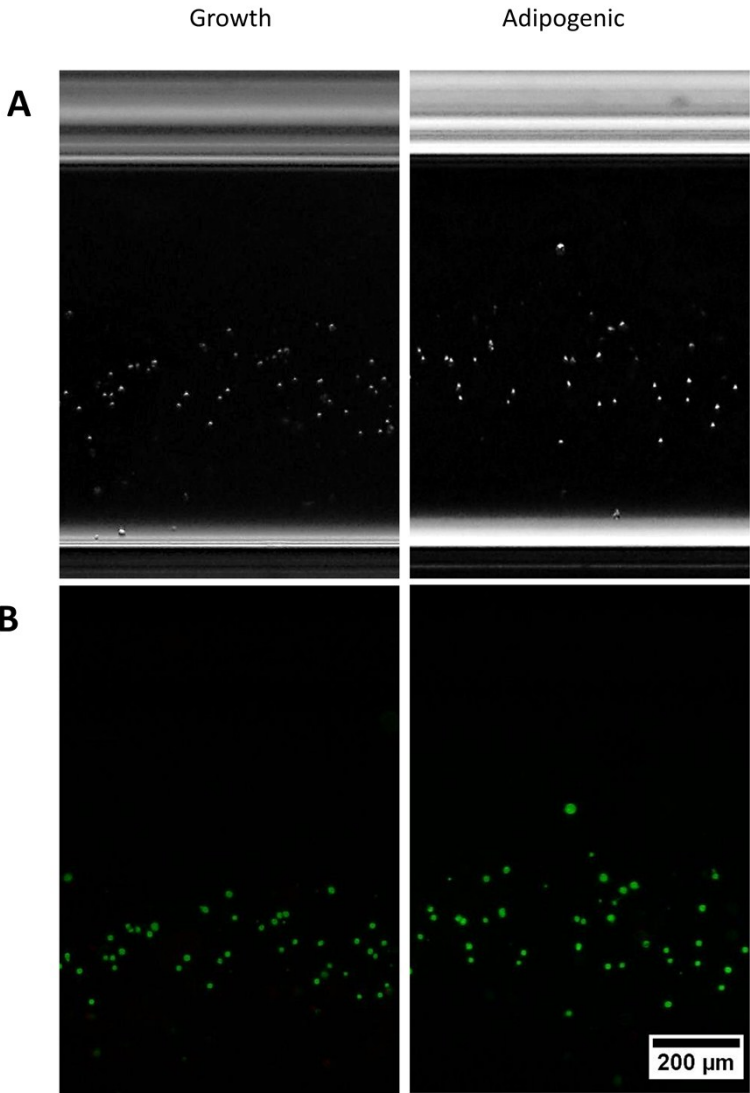
Supplementary Figure S1. Levitation heights of beads with different densities (1, 1.02 and 1.09 g/mL) at 25, 50 and 100 mM Gd^{3+} containing medium. Data are presented as mean of replicates with error bars (\pm SD) and one-way ANOVA with Tukey's multiple comparisons test was performed for statistical analysis. Statistical significance was defined as *: $p < 0.05$; **: $p < 0.01$; ***: $p < 0.001$.



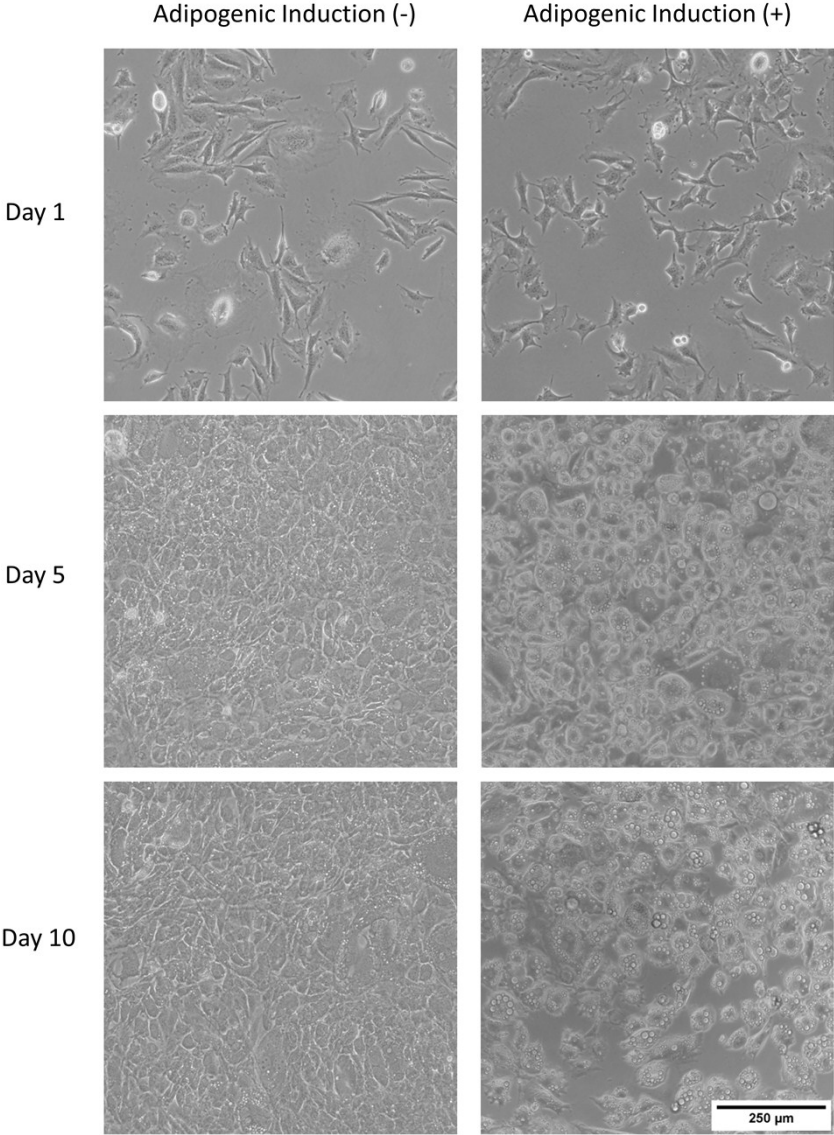
Supplementary Figure S2. Cell culture images of D1 ORL UVA cells that were cultured in standard growth medium and adipogenic differentiation medium over 15 days. Scale bar: 250 μm .



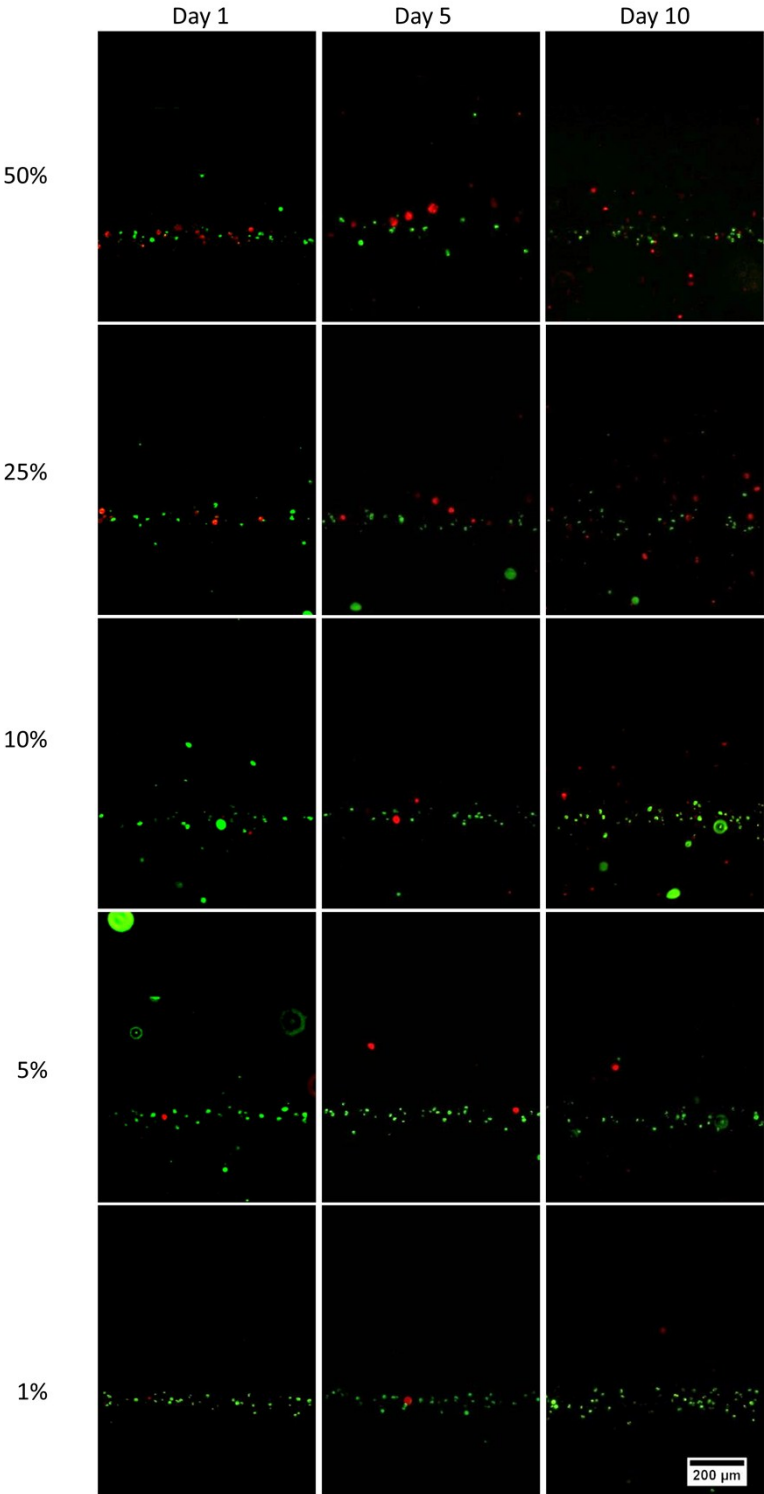
Supplementary Figure S3. Bright-field (A) and fluorescent (B) microscopy images of D1 ORL UVA cells cultured in growth or adipogenic differentiation medium at the 22nd day, in magnetic levitation device (25 mM Gd³⁺). Cell viability was visualized by live-dead (live: green, dead: red) staining. Dead cells were not detected in the system. Scale bar: 200 μ m.



Supplementary Figure S4. Cell culture images of 7F2 cells that were cultured in standard growth medium and adipogenic medium for 10 days. Scale bar: 250 μm .



Supplementary Figure S5. Fluorescent images of the levitated heterogeneous populations consisted of quiescent D1 ORL UVA cells (green) and adipogenic differentiated 7F2 cells (red) at different ratios (50%, 25%, 10%, 5% and 1%) after 1, 5 and 10 day culture. Scale bar: 200 μm .



Supplementary Figure S6. Fluorescent images of the levitated heterogeneous population consist of undifferentiated 7F2 (red) cells and D1 ORL UVA (green) cells mixed at 50% ratio after 1, 5 and 10 day culture. Scale bar: 200 μm .

