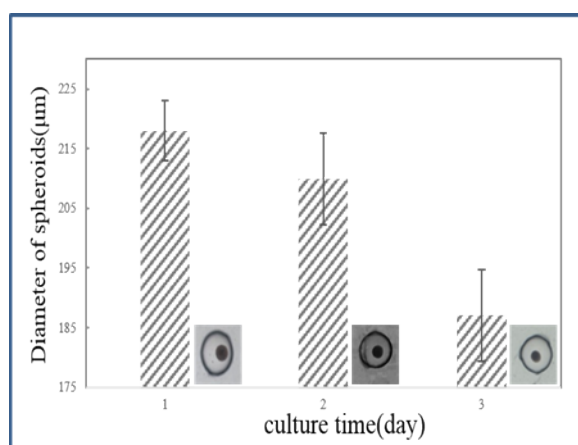
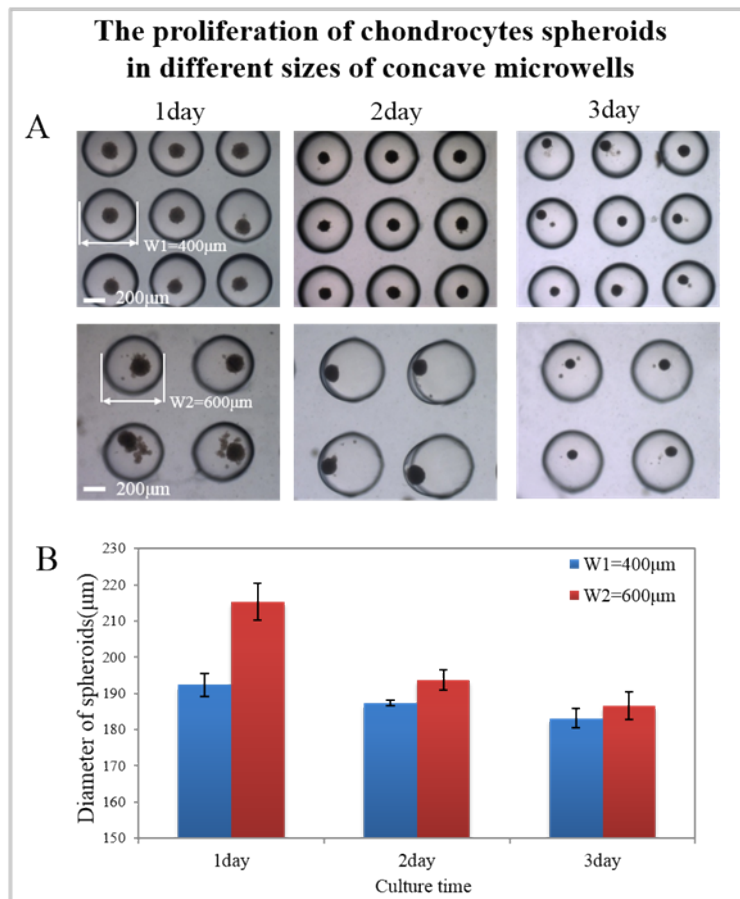


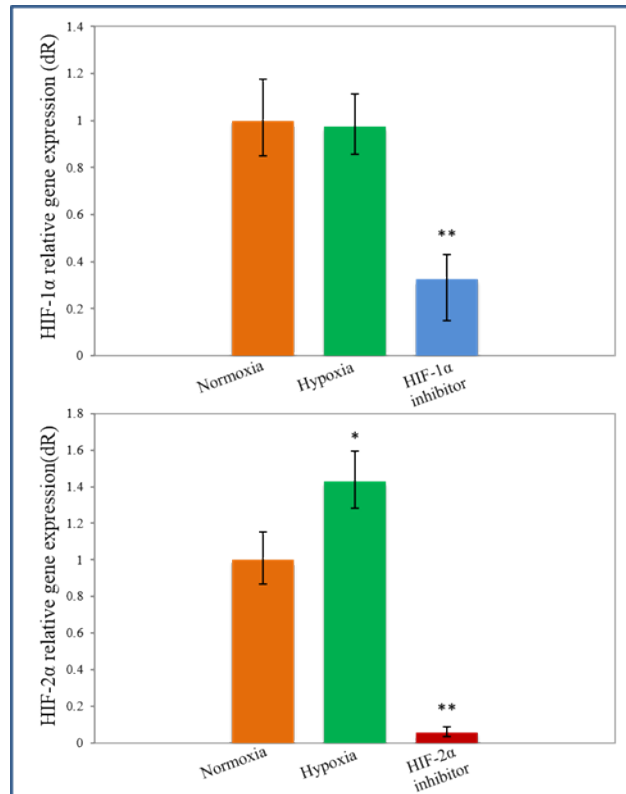
Supplemental figure 1. The viability test of chondrocytes cultured for up to 7 days using dead/live staining. Green color represented for live cells stained by Calcein AM and Red for dead cells stained by Eth-D.



Supplemental figure 2. The proliferation testing of chondrocytes cultured in concave microwells for 3 days. Error bars represent the standard deviations (mean \pm SD) obtained from six independent experiments.



Supplemental figure 3. The proliferation testing of chondrocytes cultured in concave microwells with different sizes over 3 days. A. Bright images of cellular spheroids cultured in microwells with diameter at 400µm, and 600µm, respectively. B. The averaged diameter of chondrocytes spheroids cultured in both sizes of microwells (mean \pm SD, n=10).



Supplemental figure 4. The effects of HIFs inhibitors on the mRNA gene expressions of HIF-1 α and HIF-2 α by quantitative RT-PCR assay. Chondrocyte spheroids were cultured for 3 days after treated with HIF-1 α and HIF-2 α inhibitors at the concentration of 30 μ M, respectively. Error bars were standard deviations, the data was shown as mean \pm SD (n=3), Statistic significance was calculated by Student's unpaired t-test; **, p<0.01 vs normoxic group *, p<0.05 vs normoxic group.