Migration of mesenchymal stem cells to tumor xenograft models and in vitro drug delivery

by doxorubicin

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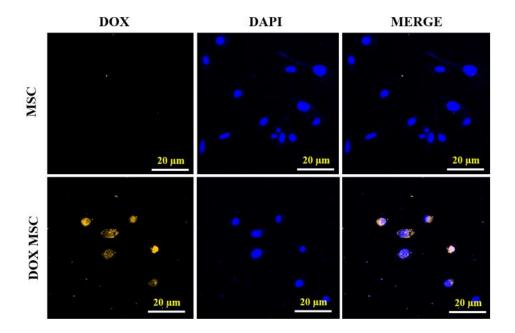
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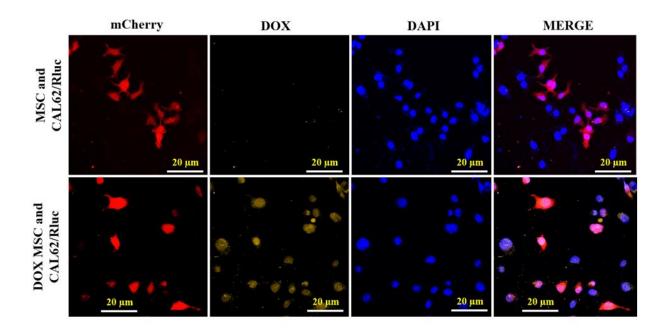
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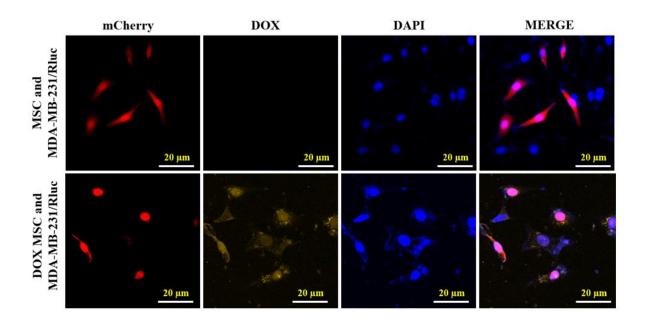
Supplementary Files



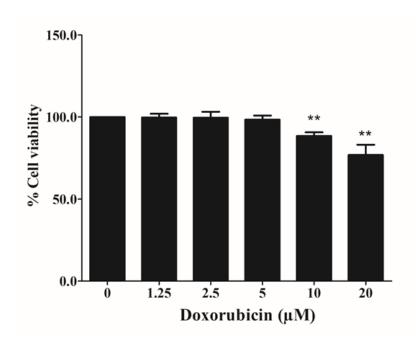
Suppl. Fig 1. DOX fluorescence in MSCs. DOX-pretreated MSCs were plated for 24 h, and the presence of DOX fluorescence was then visualized by confocal microscopy (20× magnification).



Suppl. Fig 2. Confocal microscopic image of Doxorubicin fluorescence in MSCs and MDA-MB-231/Rluc cells. DOX-pretreated MSCs and MDA-MB-231/Rluc cells were co-cultured for 24 h, and the DOX fluorescence in both cells was then visualized. MDA-MB-231/Rluc cells expressed mCherry, but the MSCs did not (20× magnification).



Suppl. Fig 3. Confocal microscopic image of doxorubicin fluorescence in MSCs and CAL62/Rluc cells. DOX-pretreated MSCs and CAL62/Rluc cells were co-cultured for 24 h, and the DOX fluorescence in both cells was visualized. MDA-MB-231/Rluc cells expressed mCherry, but the MSCs did not (20× magnification).



Suppl. Fig 4. Effect of DOX on MSC viability by CCK8 assay for 12 h. Data from three independent results were expressed as the means \pm standard deviation (SD), and a p-value < 0.05 was considered significant, according to the Student's t-test.