

Figure S1. Apoptosis of MCF-7 cells induced by cisplatin (Ci-Apoptotic-MCF-7) and exosomes from macrophages cocultured with Ci-Apoptotic-MCF-7 cells (Ci-Co-exo) increases proliferation, migration and invasion ability of MCF-7 cells

(A) Flow cytometry analysis of Annexin V-FITC/PI co-stained untreated (left panel) and apoptotic (right panel) MCF-7 cells induced by cisplatin. (B) Quantification of the flow cytometry results. (C) Proliferation of MCF-7 cells in control (CON), MCF-7^{Ci-Mac-exo} and MCF-7 Ci-Co-exo groups was measured over 72 h by MTS assay. Representative images (upper panel) and quantification (lower panel) of migration (D) and invasion (E) assays of CON, MCF-7^{Ci-Mac-exo} and MCF-7 Ci-Co-exo cells. Each bar represents the average number of cells from 5 fields. Results are typical of three independent experiments. Data represent means \pm S.E. ($\bar{x}\pm$ s) (n=3). * p<0.05, **

p<0.01 and *** p<0.001 indicate statistical significance in comparisons between the MCF-7^{Ci-Mac-exo} or MCF-7 ^{Ci-Co-exo} groups with the CON group. ### p<0.001 indicates statistical significance in comparisons between the MCF-7 ^{Ci-Co-exo} group with the MCF-7 ^{Ci-Mac-exo} group.

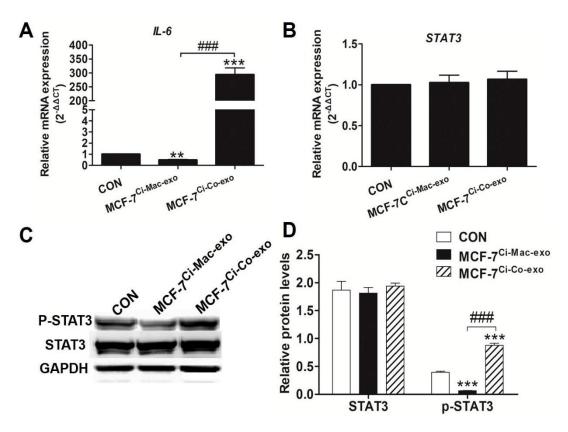


Figure S2. IL-6/ STAT3 signaling pathway is activated after MCF-7 cells are cocultured with Ci-Co-exo.

IL-6 (A) and STAT3 (B) mRNA levels were assessed using RT-qPCR assay in CON, MCF- $7^{\text{Ci-Mac-exo}}$ and MCF- $7^{\text{Ci-Co-exo}}$ cells. GAPDH was used as the internal standard. (C) Western blotting for STAT3 and p-STAT3, with GAPDH used as the loading control. (D) Bar charts illustrate the relative protein abundance of STAT3 and p-STAT3 compared to GAPDH in CON, MCF- $7^{\text{Ci-Mac-exo}}$ and MCF- $7^{\text{Ci-Co-exo}}$ cells based on densitometry of Western blotting. Results are typical of three independent experiments. Data represent means \pm S.E. ($\bar{x}\pm$ s) (n=3). ** p<0.01 and *** p<0.001 indicate statistical significance in comparisons of the MCF- $7^{\text{Ci-Mac-exo}}$ and MCF- $7^{\text{Ci-Co-exo}}$ groups to the CON group. ### p<0.001 indicates statistical significance in comparisons of the MCF- $7^{\text{Ci-Mac-exo}}$ group.

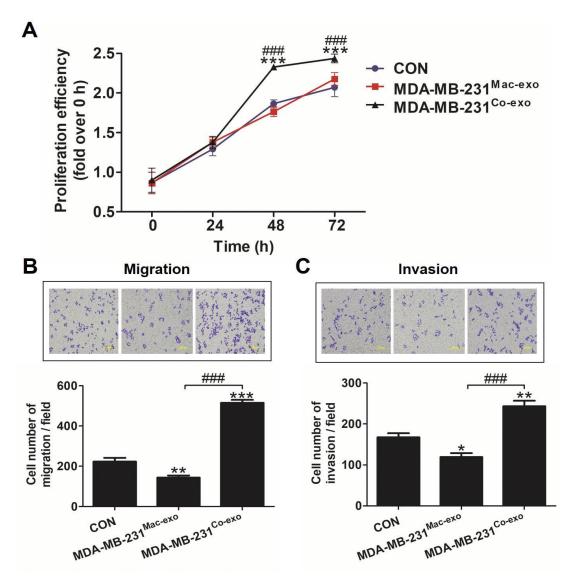


Figure S3. Co-exo increases proliferation ability and promotes cellular migration and invasion of MDA-MB-231 cells

(A) Proliferation of MDA-MB-231 cells in control (CON), MDA-MB-231^{Mac-exo} and MDA-MB-231^{Co-exo} groups was measured over 72 h by MTS assay. Results are typical of three independent experiments. Data represent means \pm S.E. ($\bar{x}\pm$ s) (n=3). Representative images (upper panel) and quantification (lower panel) of migration (B) and invasion (C) assays of CON, MDA-MB-231^{Mac-exo} and MDA-MB-231^{Co-exo} cells. Each bar represents the average number of cells from 5 fields. * p<0.05, ** p<0.01 and **** p<0.001 indicate statistical significance in comparisons of the MDA-MB-231^{Mac-exo} or MDA-MB-231^{Co-exo} groups with the CON group. ### p<0.001 indicates statistical significance in comparisons of the MDA-MB-231^{Mac-exo} group.

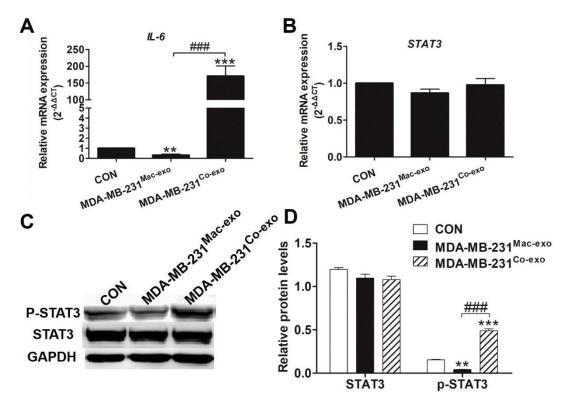


Figure S4. IL-6/ STAT3 signaling pathway is activated after MDA-MB-231 cells are co-cultured with Co-exo.

IL-6 (A) and STAT3 (B) mRNA levels were assessed using RT-qPCR assay in CON, MDA-MB-231^{Mac-exo} and MDA-MB-231^{Co-exo} cells. GAPDH was used as the internal standard. (C) Western blotting for STAT3 and p-STAT3, with GAPDH used as the loading control. (D) Bar charts illustrate the relative protein abundance of STAT3 and p-STAT3 compared to GAPDH in CON, MDA-MB-231^{Mac-exo} and MDA-MB-231^{Co-exo} cells based on densitometry of Western blotting. Results are typical of three independent experiments. Data represent means \pm S.E. ($\bar{x}\pm$ s) (n=3). ** p<0.01 and *** p<0.001 indicate statistical significance in comparisons of the MDA-MB-231^{Mac-exo} or MDA-MB-231^{Co-exo} groups with the CON group. ### p<0.001 indicates statistical significance in comparisons of the MDA-MB-231^{Mac-exo} group.