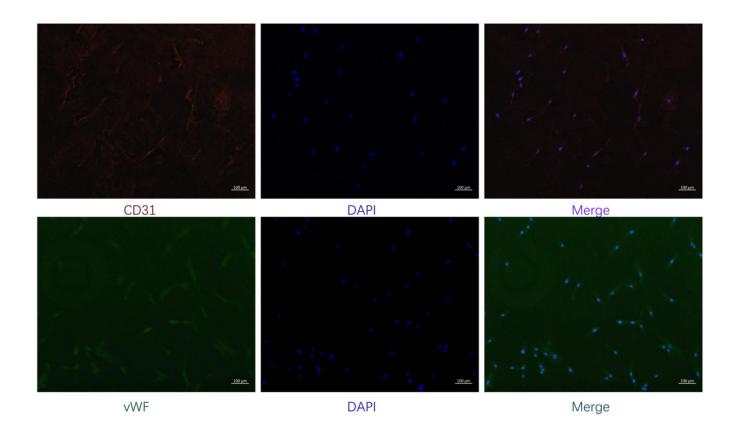
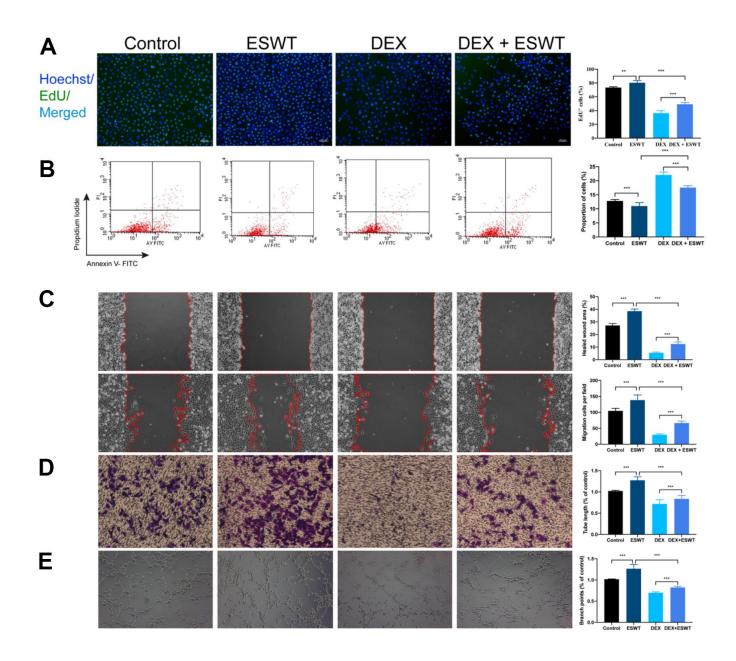
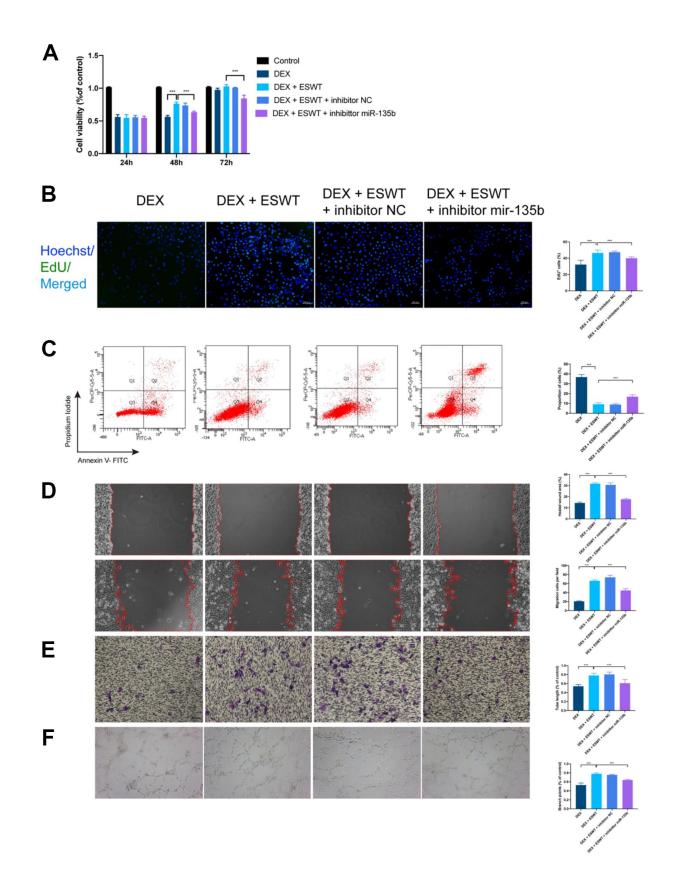
SUPPLEMENTARY FIGURES



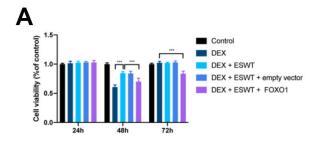
Supplementary Figure 1. Identification of BMECs. The representative CD31-positive and vWF-positive cells were identified as BMECs.

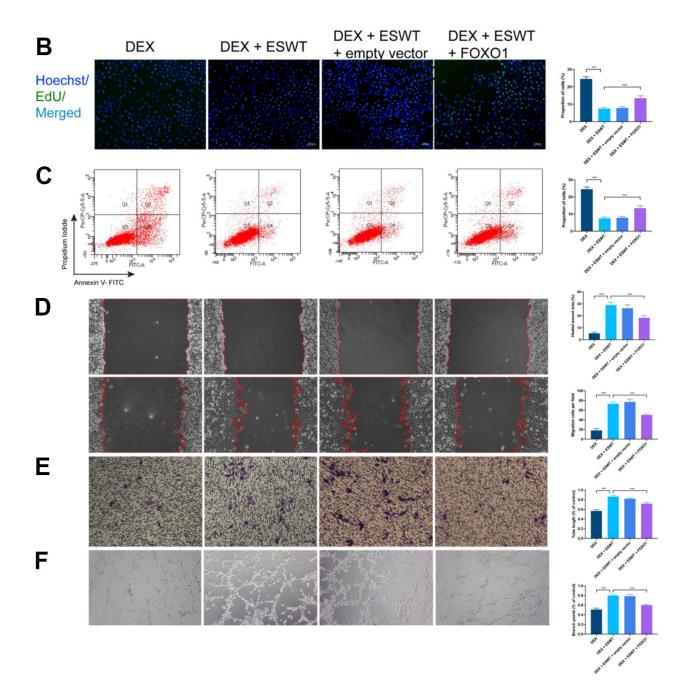


Supplementary Figure 2. Effect of ESWT on HUVECs treated with GCs. ECs were subjected to ESWT with 0.05 mJ/mm², 1000 shots followed by DEX with 180 μ M. (**A**) cell proliferation confirmed by EdU assay; (**B**) apoptosis rate of assessed through Annexin V-FITC/PI; (**C**) migration ability evaluated by wound healing assay; (**D**) migration ability evaluated by Transwell assay; (**E**) angiogenesis ability evaluated by tube formation assay. n=3 ^{**}P < .01, ^{***}P < .001.



Supplementary Figure 3. Effect of miR-135b on HUVECs treated with GCs. After transfection of inhibitor mir-135b, ECs were subjected to ESWT with 0.05 mJ/mm², 1000 shots followed by DEX with 180 μ M. (A) cell viability examined by CCK-8 analysis; (B) cell proliferation confirmed by EdU assay; (C) apoptosis rate of assessed through Annexin V-FITC/PI; (D) migration ability evaluated by wound healing assay; (E) migration ability evaluated by Transwell assay; (F) angiogenesis ability evaluated by tube formation assay. n=3 ^{**}P < .01, ^{***}P < .001.





Supplementary Figure 4. Effect of FOXO1 on HUVECs treated with GCs. After overexpression of FOXO1, ECs were subjected to ESWT with 0.05 mJ/mm², 1000 shots followed by DEX with 180 μ M. (A) cell viability examined by CCK-8 analysis; (B) cell proliferation confirmed by EdU assay; (C) apoptosis rate of assessed through Annexin V-FITC/PI; (D) migration ability evaluated by wound healing assay; (E) migration ability evaluated by Transwell assay; (F) angiogenesis ability evaluated by tube formation assay. n=3 ^{**}P < .01, ^{***}P < .001.