Supporting information

Doxorubicin resistant cancer cells activate myeloid-derived suppressor cells by releasing PGE₂

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Gene	Forward primer	Reverse primer
name		
NOS2	CCAAGCCCTCACCTACTTCC	CTCTGAGGGCTGACACAAGG
Arg-1	CTCCAAGCCAAAGTCCTTAGAG	AGGAGCTGTCATTAGGGACATC
GAPDH	AGGTCATCCCAGAGCTGAACG	ACCCTGTTGCTGTAGCCGTAT
β-Actin	ACGGCCAGGTCATCACTATTC	AGGAAGGCTGGAAAAGAGCC
TNF-α	TCTATGGCCCAGACCCTCAC	GACGGCAGAGAGGAGGTTGA
COX2	TGAGTACCGCAAACGCTTCT	CTCCCCAAAGATAGCATCTGG
MMP9	TGAGCTGGACAGCCAGACACTAAA	TCGCGGCAAGTCTTCAGAGTAGTT

Supplementary Table S1. Primers used for Real-time PCR

Supplementary Figures

Supplementary Figure S1



Supplementary Figure S1. Bone marrow (BM) cells were cultured with PBS or conditioned medium from 4T1 or Doxorubicin-resistant 4T1 cells (4T1/DOX) for 4 days in the presence of GM-CSF and IL-6, the frequency of M-MDSCs and G-MDSCs in the BM cells was analyzed by flowcytometry.

Supplementary Figure S2



Supplementary Figure S2. MicroRNA expression profiles in MDSCs treated by PBS, Supernatant from 4T1 or 4T1/DOX.

Supplementary Figure S3



Supplementary Figure S3. BM cells were cultured with 4T1 cells and 4T1/DOX cells-derived conditioned medium; in the presence of GM-CSF and IL-6 for 24 h, and the expression of miR-10a in M-MDSCs or G-MDSCs was measured by qRT-PCR. The data represent Mean \pm SD, (n=5). ** *p*< 0.01, ****p*<0.0001.

Supplementary Figure S4



Supplementary Figure S4. The expression of miR-10a in MDSCs sorted from tumor xenograft of 4T1 tumor bearing mice receiving repeated injections of doxorubicin (5mg/kg) or PBS was analyzed by qRT-PCR. The data represent Mean \pm SD, (n=5). ** p < 0.01.