

Supplemental Information

**Targeting Exosomal EBV-LMP1 Transfer
and miR-203 Expression via the NF-κB Pathway:
The Therapeutic Role of Aspirin in NPC**

Lielian Zuo, Yan Xie, Jinyong Tang, Shuyu Xin, Lingzhi Liu, Siwei Zhang, Qijia Yan, Fanxiu Zhu, and Jianhong Lu

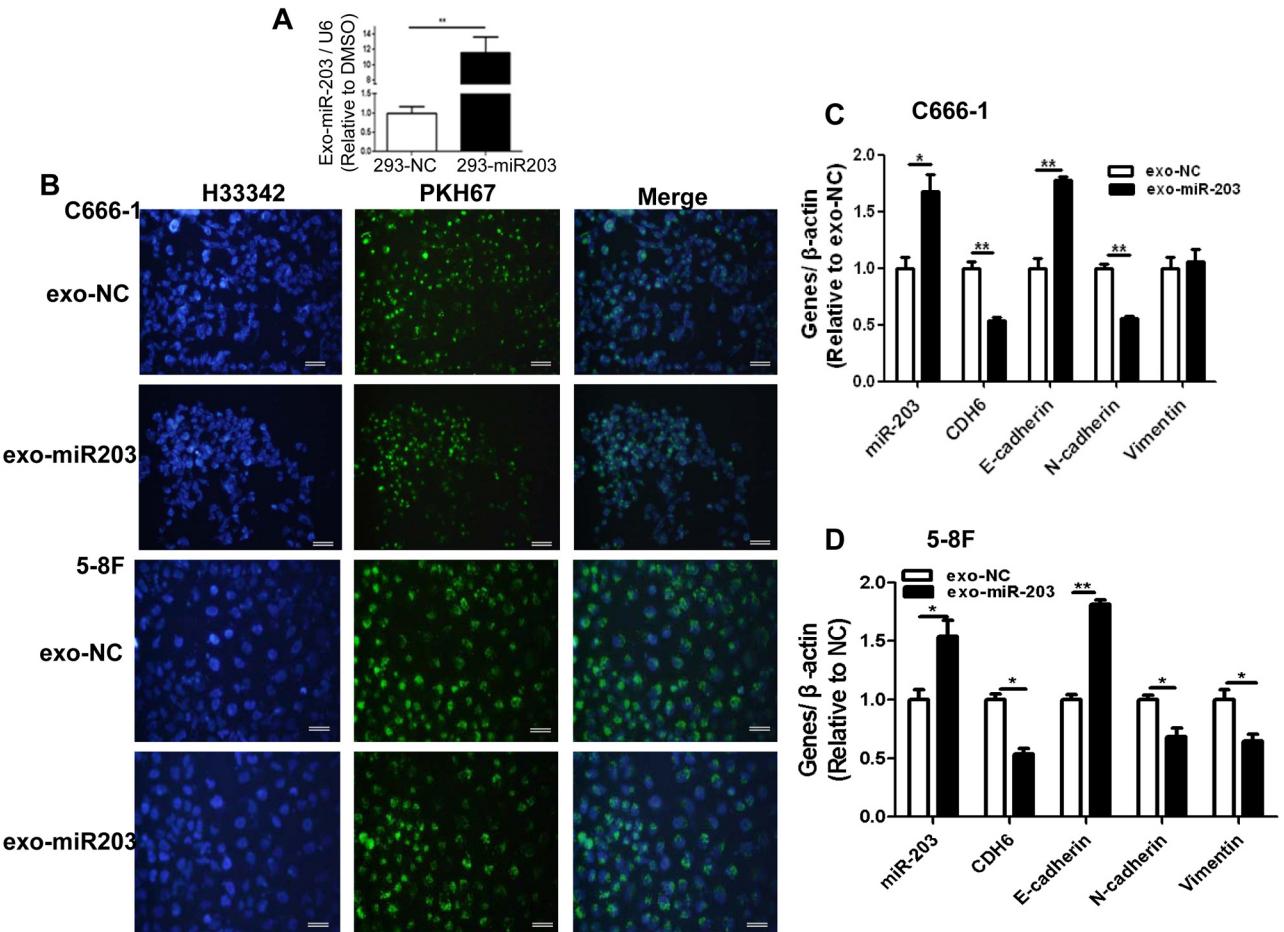


Figure S1. Exosomal miR-203 uptake inhibits EMT potential in recipient NPC cells. (A) The detection of exosome miR-203 by qPCR. The exosomes (Exo) were derived from 293-miR203 and 293-NC control cells respectively. (B) NPC C666-1 or 5-8F cells co-cultured with exo-miR203. Green indicates PKH67-labelled exosomes and blue indicates cell nuclei with H33342 staining. Scale bar, 50μm. (C) The detection of miR-203, CDH6, and the EMT markers in C666-1 cells by qPCR after co-culture. (D) The detection of miR-203, CDH6, and the EMT markers in 5-8F cells after co-culture. The results are the means ± SD ($n = 3$). * $p < 0.05$, ** $p < 0.01$.

Table S1.**Primers used for qPCR.**

Gene names	Primer names	Primer sequences
LMP1	LMP1-L	TGAACACCACCACGATGACT
	LMP1-R	GTGCGCCTAGGTTTGAGAG
CDH6	CDH6-L	TTCCTCAGGGTGCCGATATC
	CDH6-R	GTTGCCATCCTCTGTGCAT
E-cadherin	E-cadherin-L	TTCTGGAAGGAATGGAGGAGTC
	E-cadherin-R	ACCTGGAATTGGGCAAATGTG
N-cadherin	N-cadherin-L	GACAATGCCCTCAAGTGT
	N-cadherin-R	CCATTAAGCCGAGTGATGGT
Vimentin	Vimentin-L	AGATGGCCCTTGACATTGAG
	Vimentin-R	TGGAAGAGGCAGAGAAATCC
β -actin	β -actin-L	TCACCAACTGGGACGACATG
	β -actin-R	GTCACCGGAGTCCATCACGAT