

Supporting Online Materials for

Role of Arsenic (+3 oxidation state) Methyltransferase in Arsenic Mediated APL

Treatment: An in vitro Investigation

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Table S1.

Gene	Primer Sequence (5'-3')	T _A (°C)	Cycle
AS3MT	FP: GACGTGCAGACCTACTACGG RP: ACCCAGACACTCACCCATA	55	35
MRP-1	FP: ATGTCACGTGGAATACCAGC RP: GAAGACTGAACTCCCTTCCT	57	35
MRP-2	FP: ACAGAGGCTGGTGGCAACC RP: ACCATTACCTTGTCACTGTCCATGA	59	35
β-actin	FP: GCTGTCACCTTCACCGTTCC RP: CTCCATCCTGGCCTCGCTGT	61	28

T_A: Annealing Temperature

Supplementary Figure Legends and Figures

Figure S1. Comparison of PML-RAR α fusion protein degradation in 293T cells with or without co-expression of AS3MT

Flag-PML-RAR α was co-transfected with vector or GFP-AS3MT in 293T cells. 24h after transfection, cells were treated with 2 μ M iAs^{III} for 12h. Then, degradation of PML-RAR α fusion protein was detected by western blot.

Figure S2. Protein Expression of MRP Isoforms in Different Cell Lines

Cellular localization of MRP1 (A) and MRP2 proteins (B) in HeLa, HepG2 and HEK293T cells were imaged with a laser scanning confocal microscope as described in materials and methods. The green fluorescence indicates MRP1 and 2, while the blue fluorescence indicates cell nucleus. Gene (mRNA) expressions of MRP1 and 2 were determined by RT-PCR (C) and MRP protein expressions in HeLa, HepG2 and HEK293T were determined by western blot (D). Moreover, protein expression of AS3MT in HepG2 cells was determined by western blot with anti-GFP antibody 24h after transfection of GFP-hAs3MT gene into HepG2 cells (E). Intracellular localization and expression of AS3MT was observed by a laser scanning confocal microscope (F).

Figure S1

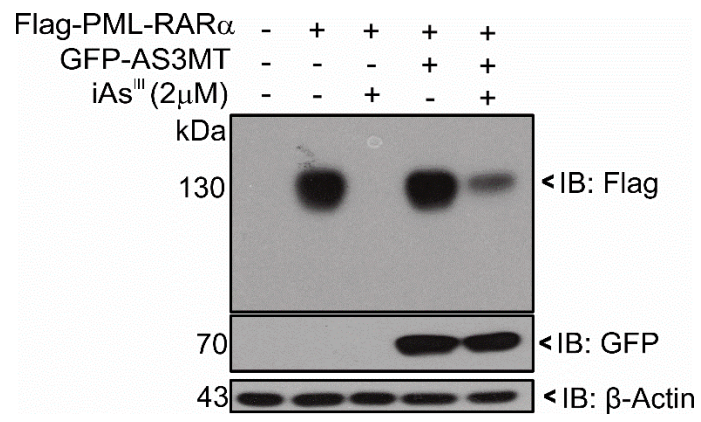


Figure S2

