## Intracellular Delivery of DNA and Protein by a Novel Cell-Permeable Peptide Derived from DOT1L

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## Supplementary figures and figure legends

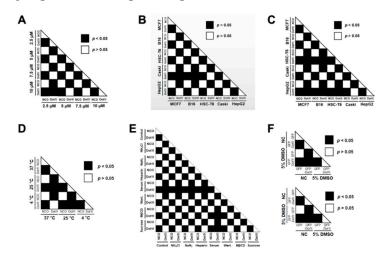


Figure S1 Statistical analysis.

- A. Corresponding to Figure 2B.
- B. Corresponding to Figure 2D of w/o DMSO pretreatment part.
- C. Corresponding to Figure 2D of 5% DMSO pretreatment part.
- D. Corresponding to Figure 3B.
- E. Corresponding to Figure 3D.
- F. Corresponding to Figure 7C.

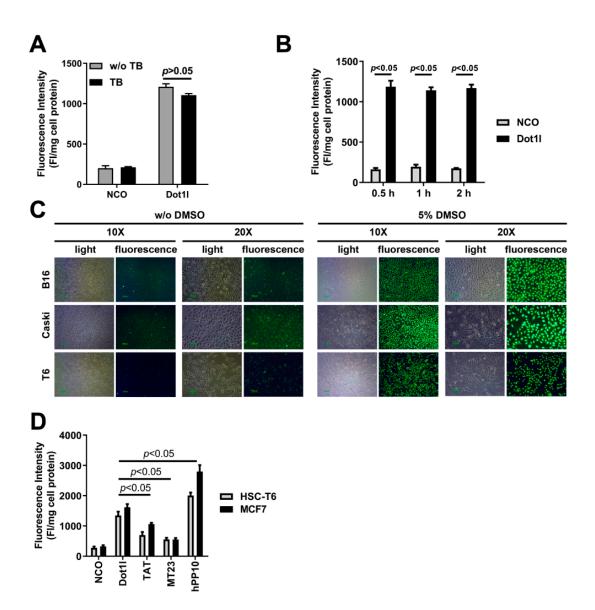


Figure S2 Penetration efficiency of Dot1l in different conditions.

- A. Quantification of Dot1l penetration with or without TB treatment.
- B. Effects of different incubation time to the Dot1I penetration.
- C. Penetration efficiency of Dot1I in different cell lines.
- D. Comparison of penetration efficiency from different CPPs in MCF7 and HSC-T6 cells.

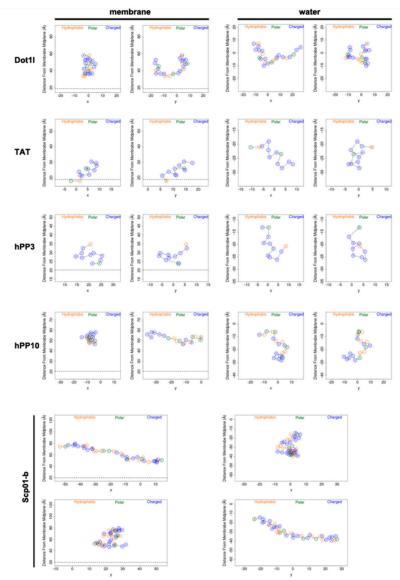


Figure S3 Prediction of different CPP-membrane interaction in aqueous and lipid bilayer environment.

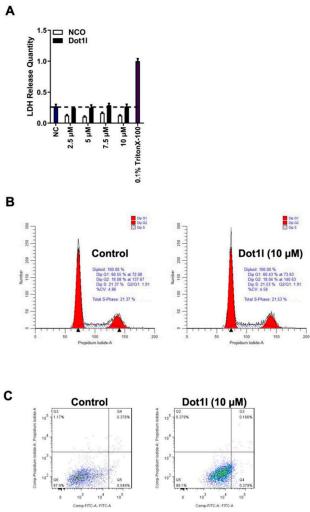


Figure S4 Dot1I safety examined by LDH release, cell cycle and cell apoptosis assay.

A. LDH release quantity assay in MCF7 cells treated with Dot1l at different concentrations,

B. Cell cycle examined by flow cytometry.

C. Cell apoptosis examined by flow cytometry with or without Dot1l incubation.

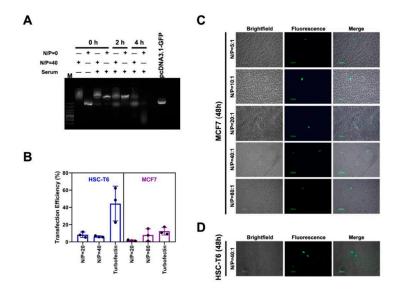


Figure S5 Stability of Dot1l/pDNA complex and Dot1l peptide mediated pDNA transfection.

A. Agarose gel examination of Dot1I peptide and pcDNA3.1-GFP complex at different N/P ratio in 50% serum solution.

B. Transfection efficiency quantification of Dot1l peptide/pDNA in different cell lines.

C. Dot1l peptide mediated pDNA transfection in MCF7 cells (48 hrs) examined by fluorescence microscope.

D. Dot1l peptide mediated pDNA transfection in HSC-T6 cells (48 hrs) examined by fluorescence microscope.