

Supplementary Information

Highly enhanced leukemia therapy and oral bioavailability from a novel amphiphilic prodrug of cytarabine

Jing Liu^{a, 1}, Jing Liu^{ab, 1}, Dajuan Zhao^a, Naxin Ma^a and Yuxia Luan^{a*}

^aSchool of Pharmaceutical Science, Shandong University, 44 West Wenhua Road, Jinan, Shandong Province, 250012, P. R. China. Fax: (86) 531-88382548; Tel: (86) 531-88382007; E-mail: yuxialuan@sdu.edu.cn

^bChia Tai Tian Qing Pharmaceutical Group Co., Ltd., Lianyungang, Jiangsu Province, 222000, P. R. China

¹These authors contributed equally to this work.

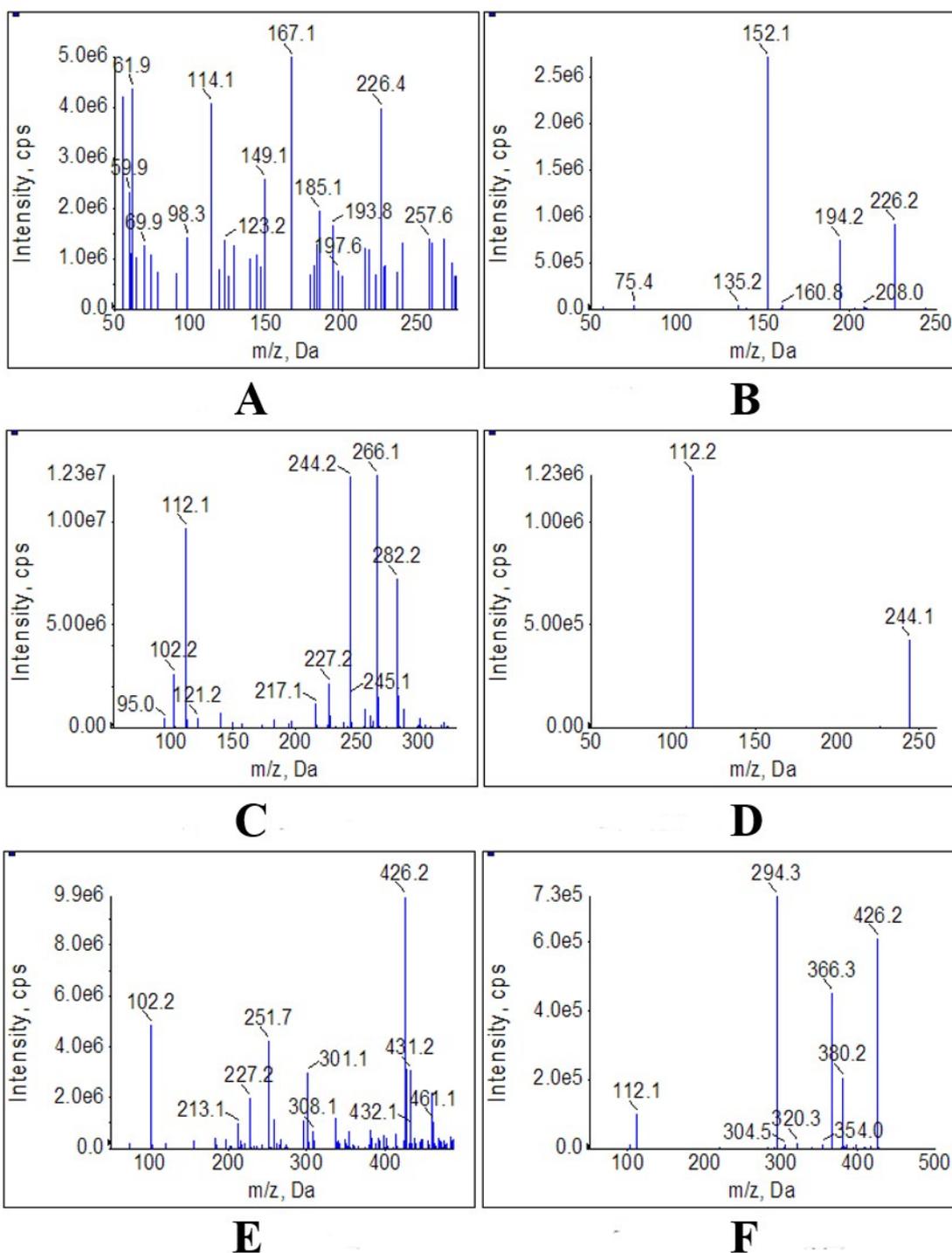


Fig. S1 The first order mass chromatogram (left) and second stage of mass chromatogram (right) of internal standard preparation acyclovir (A, B), cytarabine (C, D) and LA-Ara (E, F).

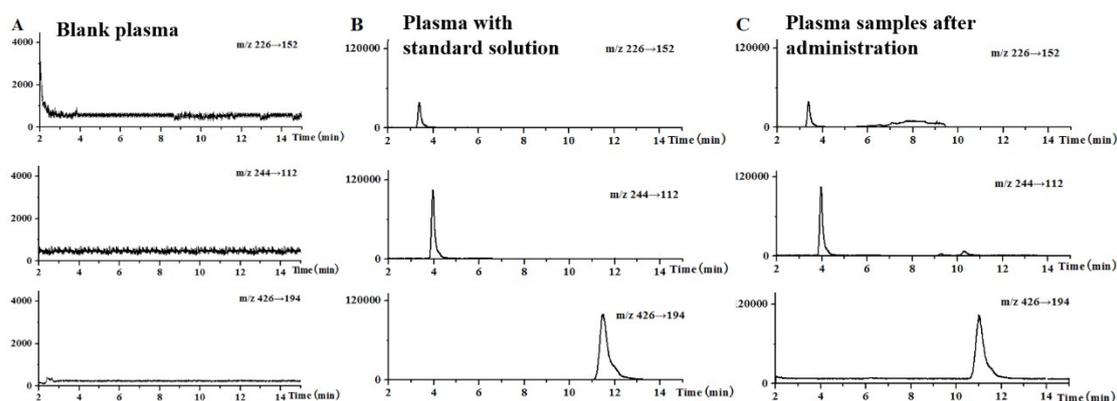


Fig. S2 Representative MRM chromatograms of acyclovir (m/z 226→152), cytarabine (m/z 244→112) and LA-Ara(m/z 426→294) in rat plasma samples.

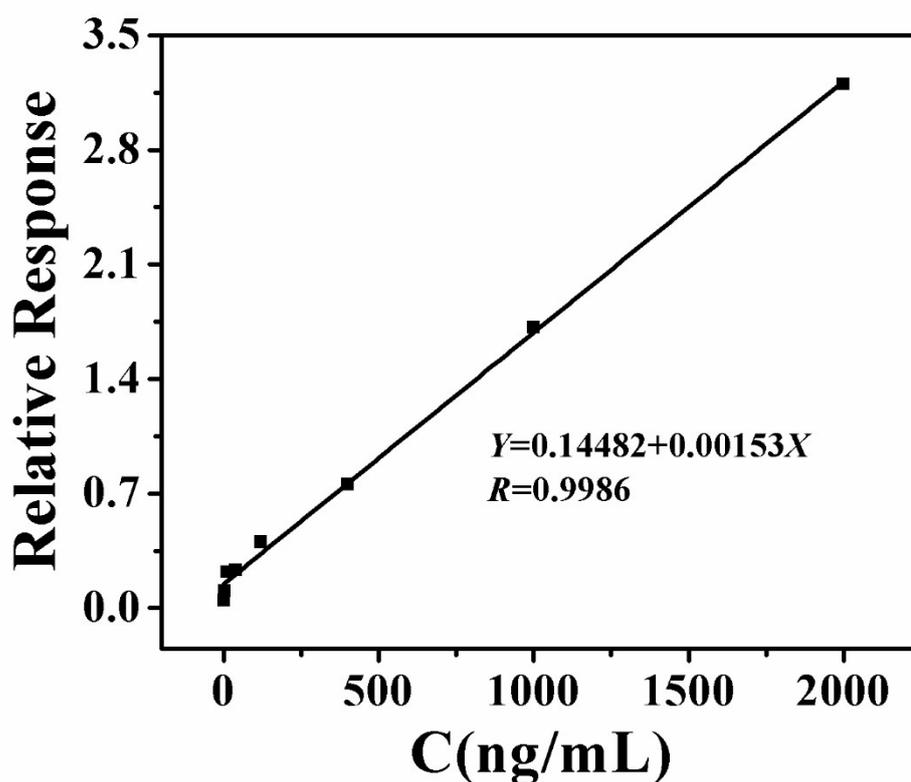


Fig. S3 Standard curve of Ara-C in the plasma of rats. Linear relationship is well within concentration of Ara-C range from 1.2 to 2000 ng·mL⁻¹.

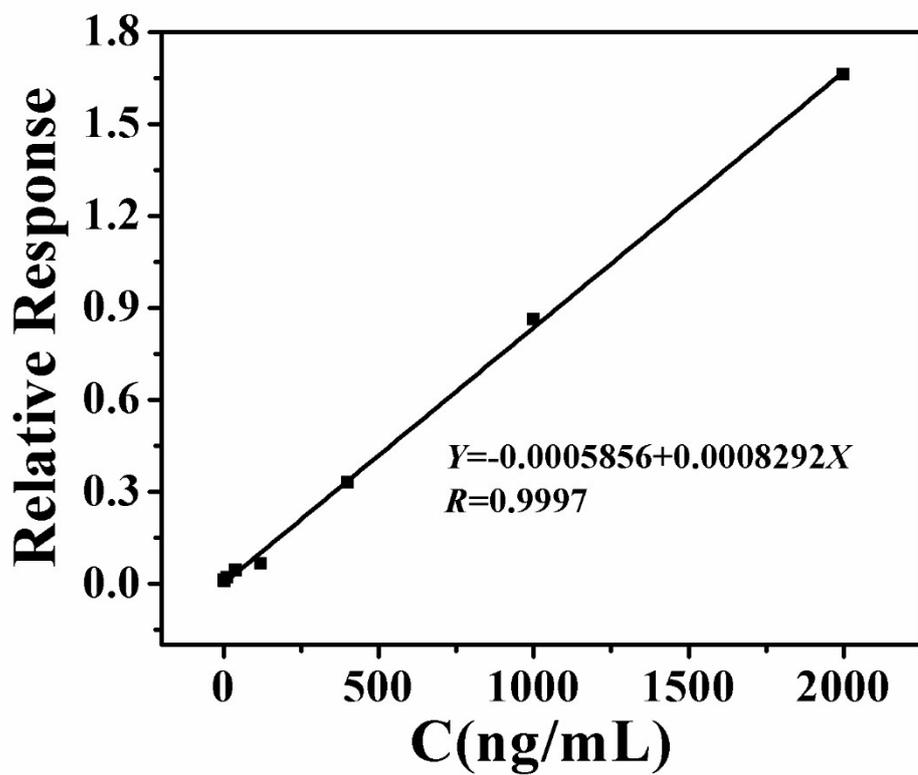


Fig. S4 Standard curve of LA-Ara in the plasma of rats. Linear relationship is well within concentration of LA-Ara range from 1.2 to 2000 ng·mL⁻¹.

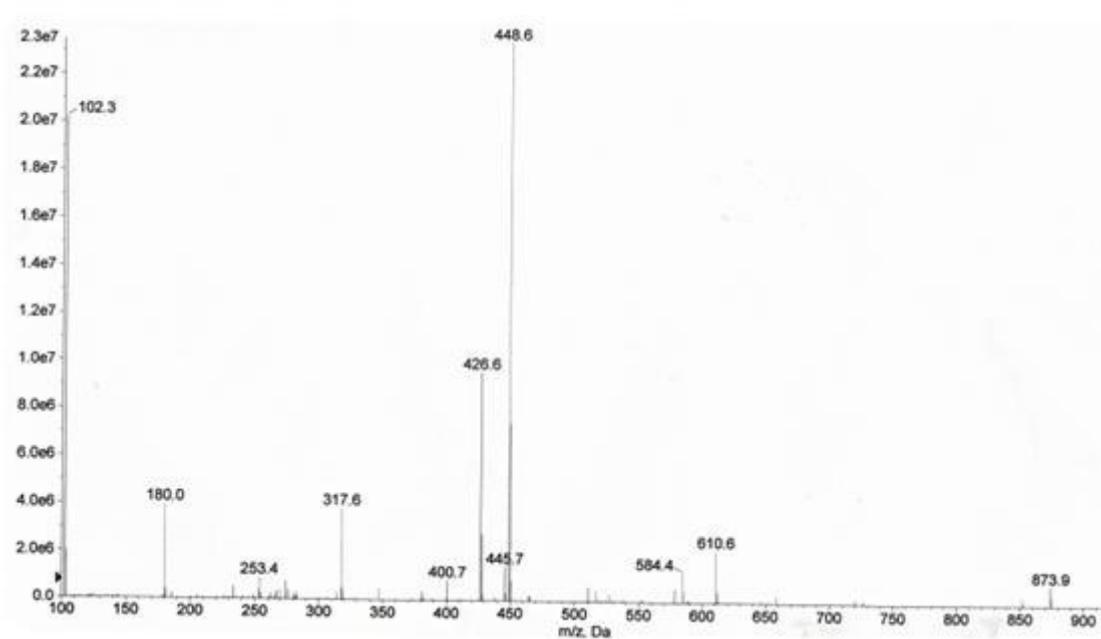


Fig. S7 Mass spectra of prodrug LA-Ara

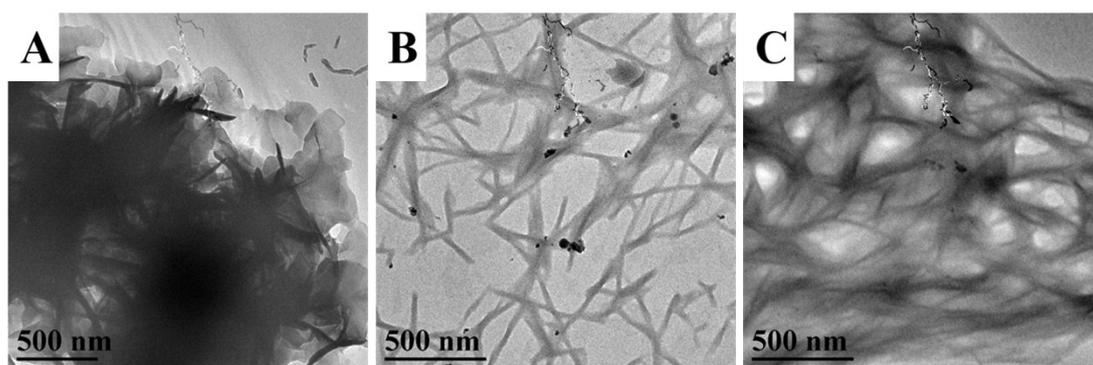


Fig. S8 TEM images of LA-Ara nanofibers after incubation for 6h at pH 1.2 (A), 4.5 (B) and 6.8 (C)

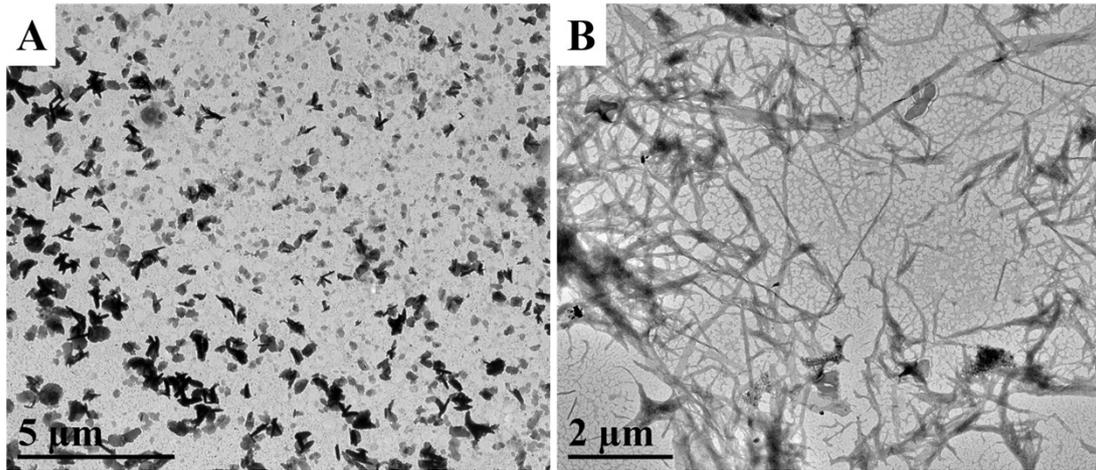


Fig. S9 TEM images of LA-Ara nanofibers after incubation at artificial gastric juice for 2 h (A) and at artificial intestinal fluid for 4 h (B).