Supporting Information

Slow, efficient and safe nanoplatform of tailored ZnS QD-Mycophenolic acid conjugates for *in vitro* drug delivery against Dengue virus 2 genome replication

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Fig. S1. Figure shows the UV-Vis absorption spectra of (a) Pure ZnS, (b) Pure MPA & (c) ZnS QD-MPA and Tauc plot of (d) Pure ZnS, (e) Pure MPA & (f) ZnS QD-MPA.



Fig. S2. Figure shows the Photo-luminescence spectra of (a) Pure ZnS (b) Pure MPA and (c) ZnS QD-MPA.



Fig. S3. CHNS elemental analysis. Chromatogram of ZnS QD MPA by CHNS analyser.



Fig. S4. Generation and Validation of DV2-replicon cells (a) Schematic representation of Invitro Transcription of DENV2 and generation of DV2-replicon cells. (b) Morphology of DV2-replicon cells. (c) ELISA test for NS1 protein secretion from DV2-replicon soup. (d) Luciferase assay for active replication of DENV2 RNA. (e) Immunostaining of DENV2 NS3 protein.



Fig. S5. Brightfield images of DV2-replicon cells: (a) The cytopathic effect of the cells after treatment of the compounds at various concentrations panel (i) only ZnS QD, (ii) MPA, (iii) ZnS QD-MPA. The MPA showed the highest CPE at ~5 μ M whereas MPA ZnS QD showed CPE ~2000 μ M. There was no CPE observed in the cells treated with ZnS QD only up to 3000 μ M



Fig. S6. Lethal Doses 50 (LD₅₀) of (a) Celgosivir (LD₅₀=77.69 \pm 1.579 μ M). (b) ZnS QD-Celgosivir (LD₅₀=186.7 \pm 1.099 μ M). (c) Curcumin (LD_{50}=27.86 \pm 1.197 μ M). (d) ZnS QD-Curcumin (LD_{50}=95.43 \pm 1.1494 μ M).



Fig. S7. Effective Concentration 50 (EC₅₀) of (a) Celgosivir (EC₅₀ = $6.231 \pm 1.412 \mu$ M). (b) ZnS QD-Celgosivir (EC₅₀ = $23.16 \pm 1.440 \mu$ M). (c) Curcumin (EC₅₀ = $2.193 \pm 1.068 \mu$ M). (d) ZnS QD-Curcumin (EC₅₀ = $8.268 \pm 1.105 \mu$ M).

(a)

(b)



Fig. S8. BCA Standard curve for protein estimation.



Fig. S9. Standard curve of MPA release in different pH buffers (a) Phosphate buffer, pH 5 (b) Phosphate buffer ,pH 7.2.