Supporting Information

Biological Assessment of Triazine Dendrimers as Candidate Platforms for Nanomedicine: Toxicological Profiles, Solution Behavior, Biodistribution, and Drug Release and Efficacy in a PEGylated, Paclitaxel Construct

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Figure S1. The average volume distribution plots for dendrimer **1** is water (blue), 10mM NaCl (red), PBS (green), and saline (black) at 2mg/mL.

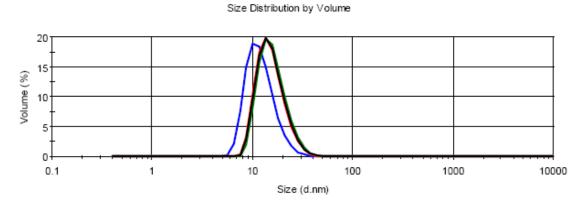


Figure S2. Summary of the hydrodynamic size of **1** in different solvents.

Solvent	η, cP	Z-Avg, nm	PdI	Int-Peak, nm	%Int	Vol-Peak, nm	%Vol
Water	0.8872	15.6 (0.3)	0.173 (0.009)	16.9 (0.3)	97.2 (0.5)	12.4 (0.4)	100 (0)
10 mM NaCl	0.8894	18.5 (0.4)	0.098 (0.006)	20.5 (0.4)	100 (0)	15.6 (0.4)	100 (0)
PBS	0.8941	19.4 (0.5)	0.105 (0.007)	21.6 (0.6)	100 (0)	16.3 (0.3)	100 (0)
Saline	0.9025	18.9 (0.3)	0.101 (0.010)	21.0 (0.4)	100 (0)	15.8 (0.4)	100 (0)

Figure S3. The average zeta potential for 1 in 10 mM NaCL measured on a Malvern Zetasizer Nano ZS instrument at 25° C with an applied voltage of 120 V.

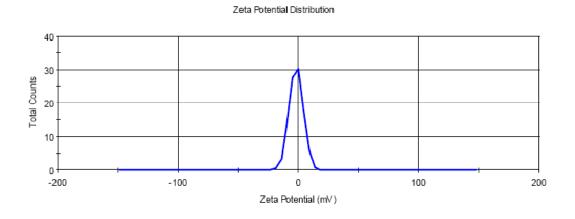


Figure S4. LS174T Morphology. Human colon carcinoma cells were treated with 1 and Taxol at 300 uM paclitaxel equivalent for 24h. Morphology was assessed by phase contrast light microscopy at 200x magnification.

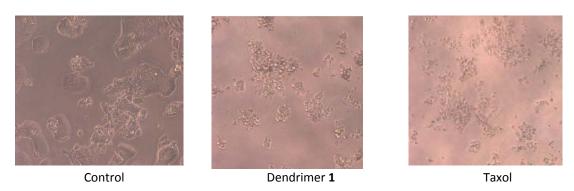


Figure S5. The effect of **1** and Taxol on LS174T caspase activity. Cells were treated for 24h (A) and 48h (B) with $3x10^{-7}$ to 300 μ M paclitaxel equivalent concentration of **1** (blue) and Taxol (purple) or control media. The positive control was 10mM acetaminophen. The negative control was media. Data are represented as %negative media control caspase 3 activity. Bars correspond to the mean + SE of 3 individual samples. *P < 0.05, significantly different from negative control by ANOVA, with post hoc testing by Dunnett's t test.

