

Figure 1: Chemical structure of A) cholesteryl-PAA; B) Fmoc-PAA; C) Dansyl- PAA; D) propofol; E) griseofulvin; F) prednisolone

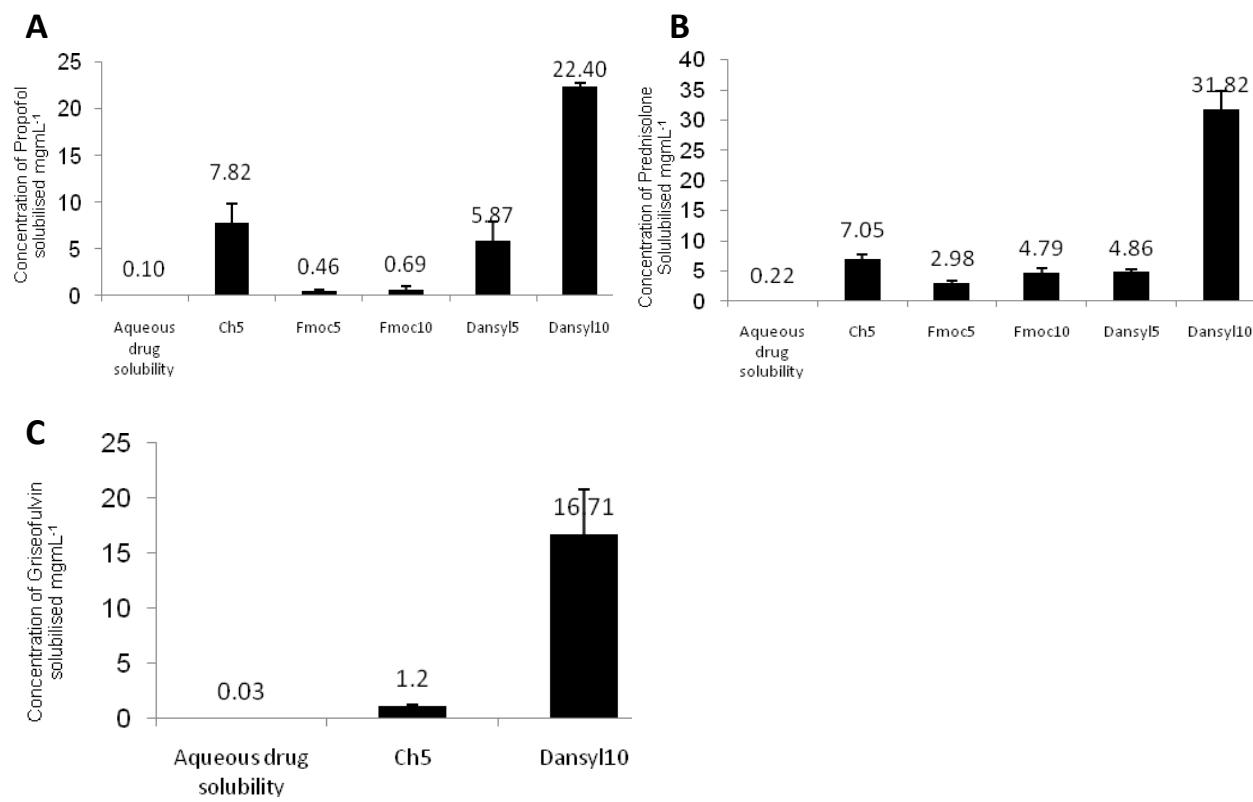


Figure 2. Maximum drug concentration solubilised by PAA amphiphiles. A) propofol, B) prednisolone and C) griseofulvin.

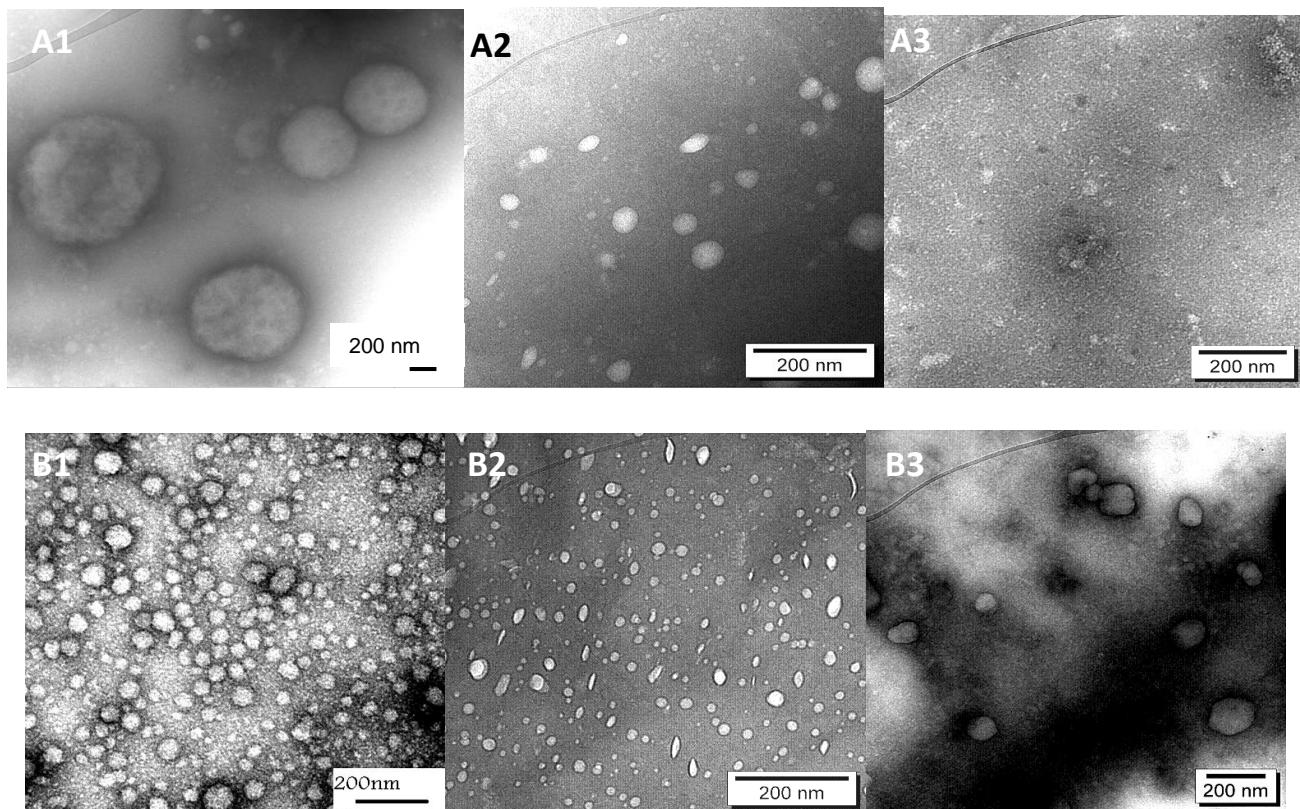


Figure 3. Negative-stained TEM of A) Ch₅ formulations with 1) propofol, 2) prednisolone and 3) griseofulvin. B) Dansyl₁₀, 1) propofol, 2) prednisolone and 3) griseofulvin. All the formulations consisted of 6 mgmL⁻¹ polymer and 10:1 initial drug: polymer mass ratio. Bar=200nm

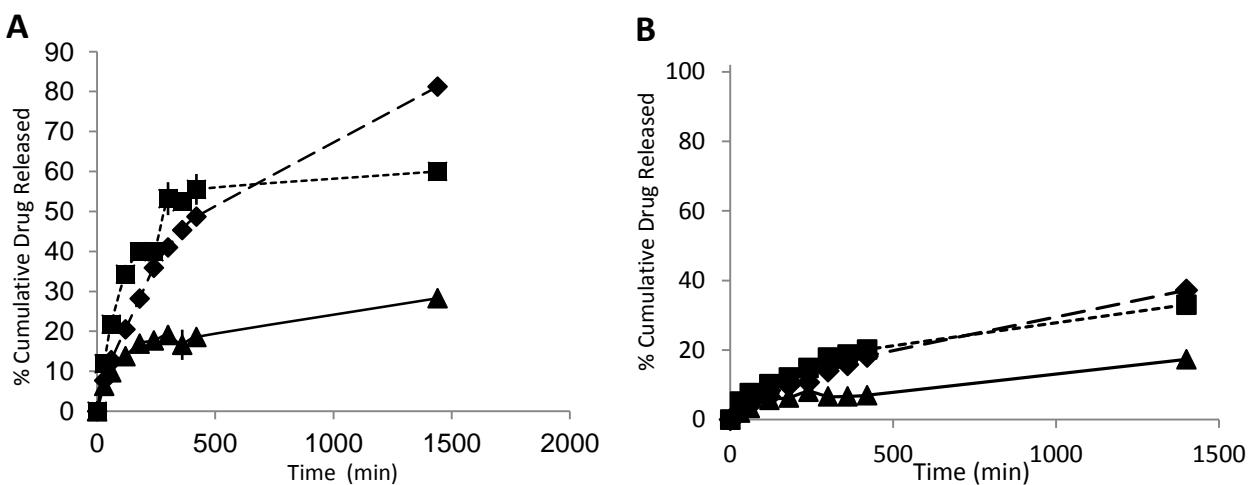


Figure 4. *In vitro* drug release of hydrophobic drugs from A) Ch₅ and B) Dansyl₁₀ formulations carried out in sink conditions. ♦ propofol ; ■ prednisolone ▲ griseofulvin. Data presented as n=3, ave ± s.d.

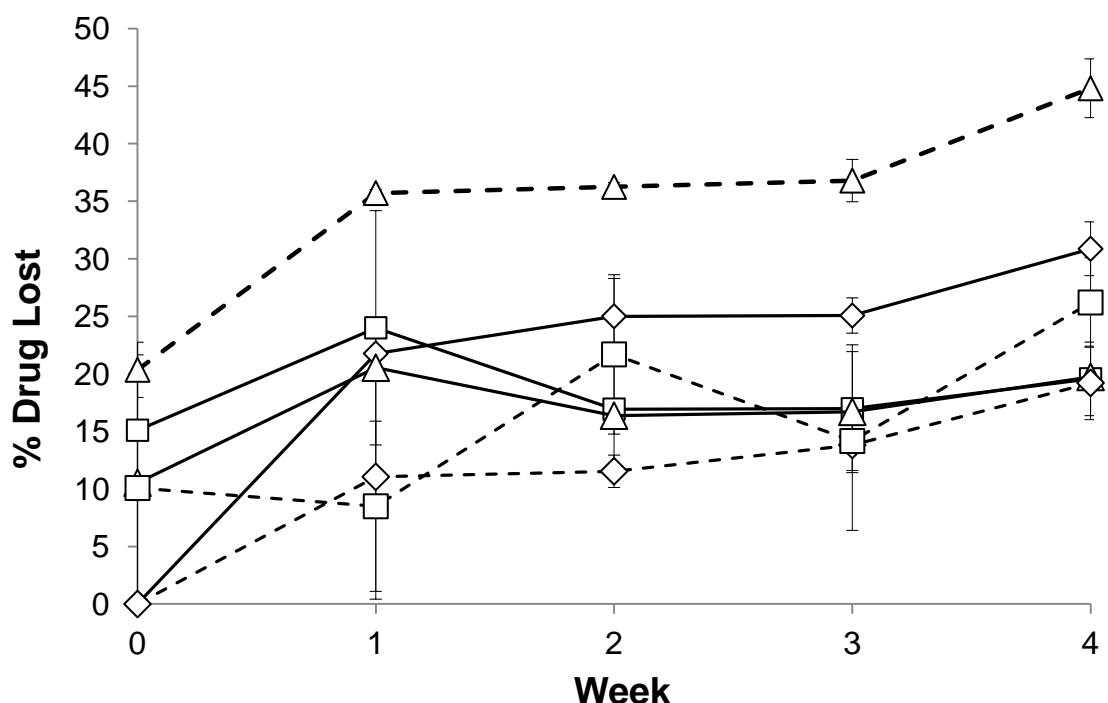


Figure 5. Percentage drug lost from Ch₅ (solid line) and Dansyl₁₀ (dashed line) formulations over 4 weeks stored in 55 % humidity, at room temperature and protected from light. Propofol stored in solution, prednisolone and griseofulvin formulations stored as freeze dried ‘cakes’. ◊ propofol ; □ prednisolone; △ griseofulvin. Data presented as n=3, ave ± s.d.

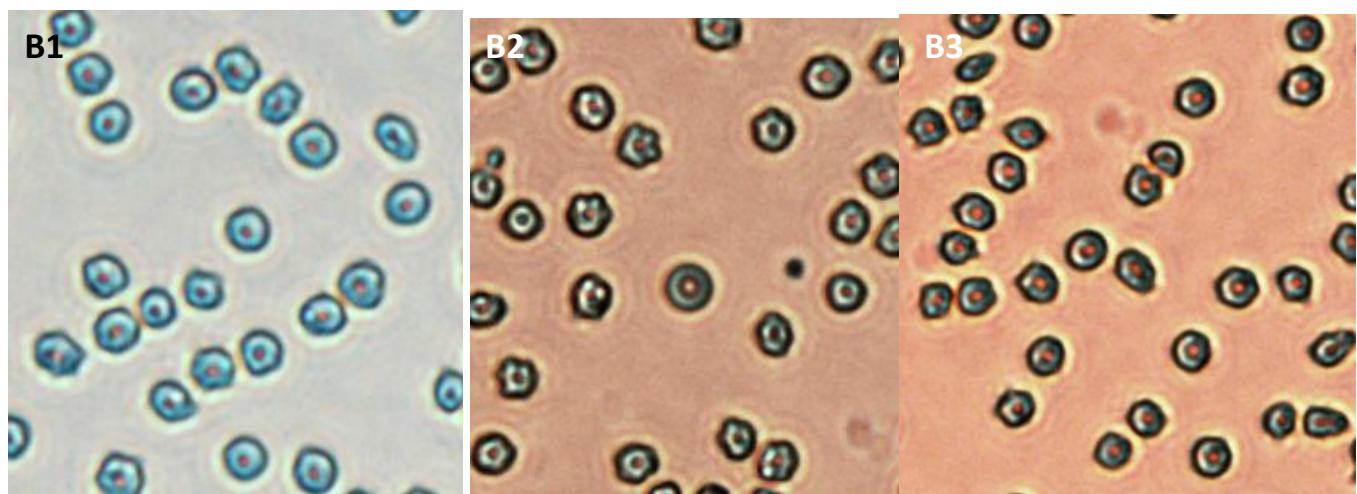
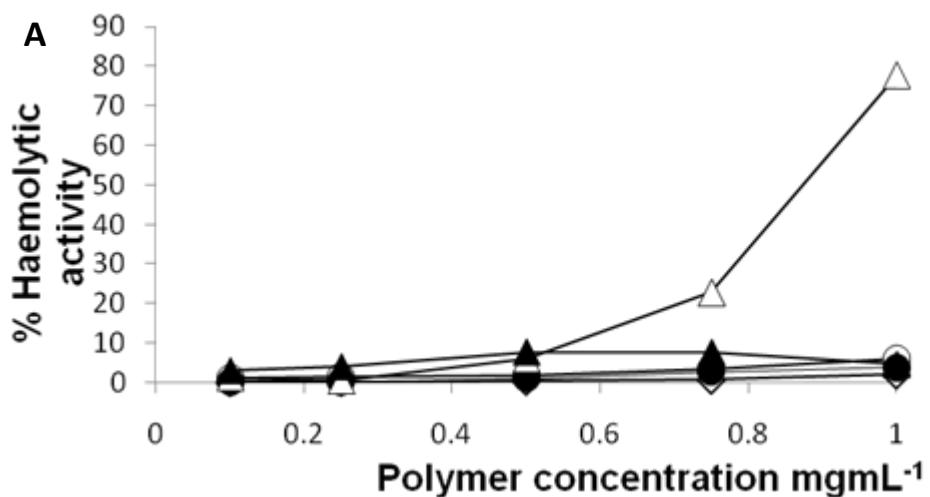


Figure 6. Effect of PAA and its amphiphilic PAA polymers on bovine red blood cells. **A)** % Haemolysis of $\diamond\text{PAA}$; $\triangle\text{Dansyl}_5$; $\blacktriangle\text{Dansyl}_{10}$; $\circ\text{Fmoc}_5$; $\bullet\text{Fmoc}_{10}$. Data presented as $n=3$, ave \pm s.d. **B)** Morphology of red blood cells upon 4h incubation with 1) PBS control; 2) Dansyl₅; 3) Dansyl₁₀ (1 mg mL^{-1}) (100x magnification)

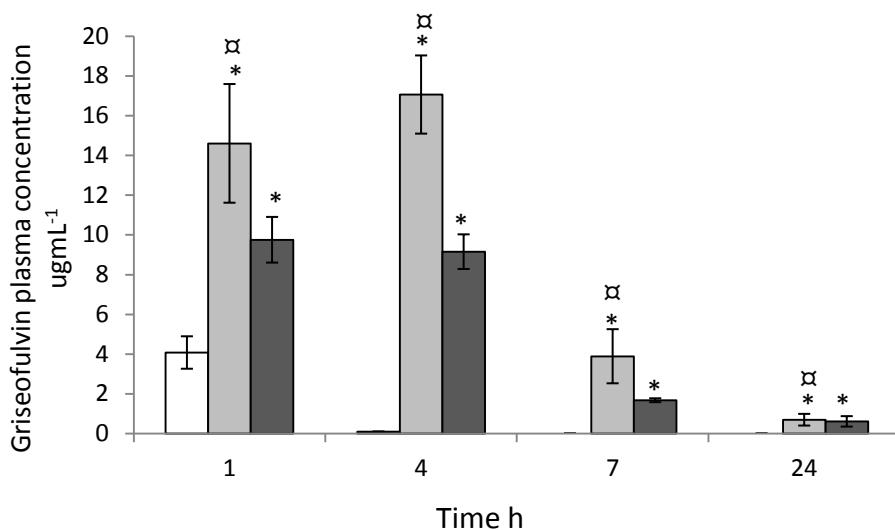


Figure 7. Mean plasma griseofulvin concentration ($\mu\text{g mL}^{-1}$) following administration of griseofulvin by oral gavage in rats. □ Griseofulvin in water; ■ Ch5, griseofulvin and ■ Dansyl₁₀, griseofulvin. Data presented as n=4, ave \pm s.d. * p<0.0001 polymer formulations vs. griseofulvin in water, ☺ p < 0.001 Dansyl₁₀, vs. Ch₅.

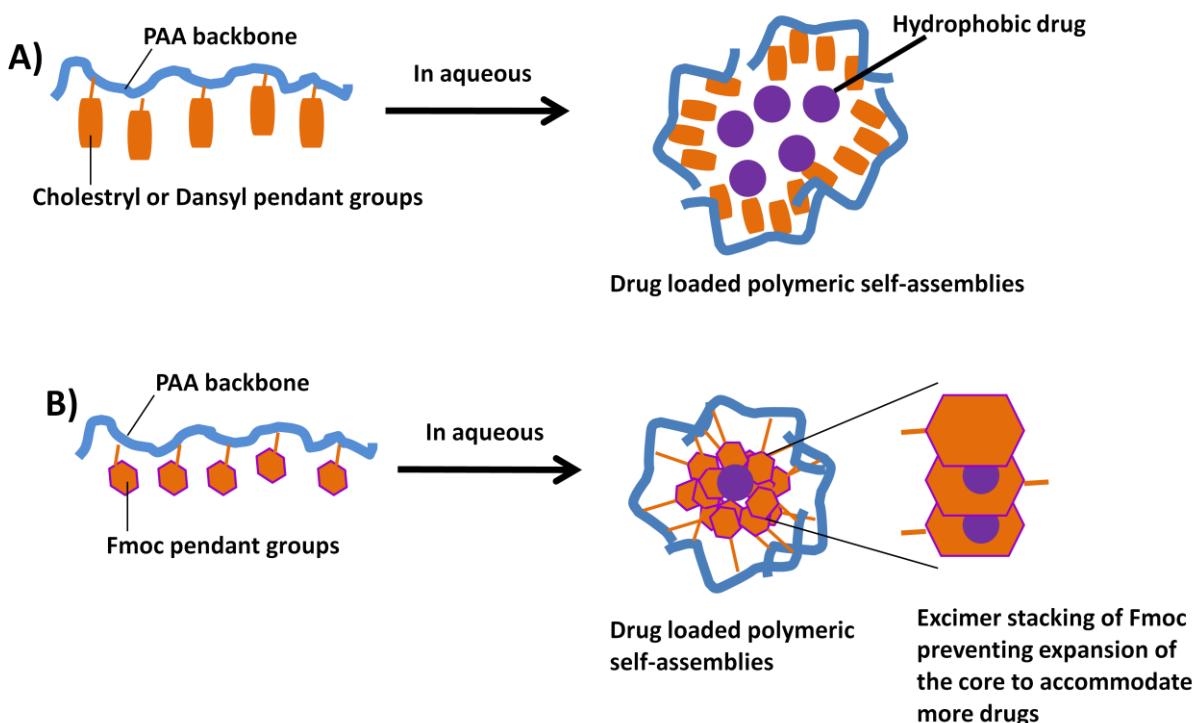


Figure 8. Proposed drug loaded polymeric self-assemblies structures in aqueous environment A) Ch and Dansyl-PAA and B) Fmoc-PAA

Legends to figures

Figure 1. Chemical structure of A) cholesteryl- PAA; B) Fmoc - PAA; C) Dansyl- PAA; D) propofol; E) griseofulvin; F) prednisolone

Figure 2. Maximum drug concentration solubilised by PAA amphiphilic. A) propofol, B) prednisolone and C) griseofulvin.

Figure 3. Negative-stained TEM of A) Ch₅ formulations with 1) propofol, 2) prednisolone and 3) griseofulvin. B) Dansyl₁₀, 1) propofol, 2) prednisolone and 3) griseofulvin. All the formulations consisted of 6 mgmL⁻¹ polymer and 10:1 initial drug: polymer mass ratio. Bar=200nm

Figure 4. *In vitro* drug release of hydrophobic drugs from A) Ch₅ and B) Dansyl₁₀ formulations carried out in sink conditions. ◆ propofol ; ■ prednisolone ▲ griseofulvin. Data presented as n=3, ave ± s.d.

Figure 5. Percentage drug lost from Ch₅ (solid line) and Dansyl₁₀ (dashed line) formulations over 4 weeks stored in 55 % humidity, at room temperature and protected from light. Propofol stored in solution, prednisolone and griseofulvin formulations stored as freeze dried ‘cakes’. ◇ propofol ; □ prednisolone; △ griseofulvin. Data presented as n=3, ave ± s.d.

Figure 6. Effect of PAA and its amphiphilic PAA polymers on bovine red blood cells. **A)** % Haemolysis of ◇PAA; △Dansyl₅; ▲Dansyl₁₀; ○Fmoc₅; ● Fmoc₁₀. Data presented as n=3, ave ± s.d. **B)** Morphology of red blood cells upon 4h incubation with 1) PBS control; 2) Dansyl₅; 3) Dansyl₁₀ (6 mgmL⁻¹) (100x magnification)

Figure 7. Mean plasma griseofulvin concentration (μgmL⁻¹) following administration of griseofulvin by oral gavage in rats. Griseofulvin in water: □ Ch₅, griseofulvin and □ Dansyl₁₀, griseofulvin. Data presented as n=4, ave± s.d. * p<0.0001 polymer formulations vs. griseofulvin in water, ‡ p < 0.001 Dansyl₁₀, vs. Ch₅

Figure 8. Proposed drug loaded polymeric self-assemblies structures in aqueous environment A) Ch and Dansyl-PAA and B) Fmoc-PAA