

PAPER

[View Article Online](#)
[View Journal](#) | [View Issue](#)Cite this: *Mater. Adv.*, 2020,
1, 371

Poly(*N*-isopropyl acrylamide)–poly(ethylene glycol)–poly(*N*-isopropyl acrylamide) as a thermoreversible gelator for topical administration†

P. Haddow, W. J. McAuley, S. B. Kirton  and M. T. Cook *

Poly(*N*-isopropyl acrylamide)-*block*-poly(ethylene glycol)-*block*-poly(*N*-isopropyl acrylamide) is known to exhibit a thermally-induced solution-to-gel transition in water, which may be exploited for biomedical applications. This “thermoreversible gelator” has great potential for application in topical administration to the surfaces of the body such as the skin, eye, and vagina, but this has not yet been evaluated. This study evaluates PNIPAM₉₈–PEG₁₂₂–PNIPAM₉₈ as a thermoreversible gelator for vaginal administration, for the first time evaluating the effect of polymer concentration on gelation, studying rheological parameters relevant to topicals, measuring dissolution rates, stability and the phenomenon of mucoadhesion. Two drugs relevant to vaginal administration, progesterone and tenofovir disoproxil fumarate are investigated for use in the thermoreversible gelators, studying both hydrophobic and hydrophilic drug solubilisation and release. Throughout the study, comparison is made with poloxamer 407, the most commonly studied thermoreversible gelator. PNIPAM₉₈–PEG₁₂₂–PNIPAM₉₈ exhibits several advantages for topical administration, having low viscosity at room temperature to allow easy application, then exhibiting a gelation just below body temperature to form a viscous gel which is resistant to dissolution and relatively mucoadhesive. Drug release is highly dependent on temperature, with elevation to body temperature resulting in a dramatic retardation of progesterone release, which may be used in future medicines to provide sustained delivery of hydrophobic xenobiotics.

Received 5th March 2020,
Accepted 6th May 2020

DOI: 10.1039/d0ma00080a

rsc.li/materials-advances

Introduction

“Smart” materials respond to external stimuli by alteration of one or more of their physical properties. One class of smart material are thermoreversible gelators. These materials undergo a solution-to-gel transition when warmed above a critical temperature (T_{gel}), exhibiting the reverse gel-to-solution phase change upon cooling.¹ Thermoreversible gelators with a T_{gel} occurring between room temperature (*ca.* 25 °C) and body temperature (37 °C internally) may pass through an applicator or syringe, then form a viscous gel on contact with the body.² This *in situ* gelation may be used for injectables, where the gel may act as a drug depot or scaffold for therapeutic cells,³ and for topical administration to either skin or mucosal membranes.⁴ Thermoreversible gelators

are particularly attractive for vaginal application, as the material may pass through an applicator before forming a viscous gel upon contact with the body which enhances retention of bioactives to prolong local effect or absorption⁵ at a site where retention is often poor.

The most widely studied polymer which exhibits thermoreversible gelation is poloxamer 407 (also called Pluronic F127). Poloxamer 407 is an ABA triblock copolymer with the structure poly(ethylene glycol)₁₀₁-*block*-poly(propylene glycol)₆₅-*block*-poly(ethylene glycol)₁₀₁.⁶ Heating concentrated (> 15% w/v) poloxamer 407 solutions results in a hierarchical process of desolvation of the poly(propylene glycol) block, which drives micellization and in turn the formation of a face-centred cubic mesophase gel.⁷ This property makes poloxamer 407 an attractive excipient for topical use, where the body's heat may trigger gelation.⁸ However, poloxamer 407 has several drawbacks including: a low T_{gel} of *ca.* 25 °C and below,⁹ rapid dissolution, and low mucoadhesion.⁶ Thus, there is a need for novel thermoreversible gelators with modulated properties.

Poly(*N*-isopropyl acrylamide)-*block*-poly(ethylene glycol)-*block*-poly(*N*-isopropyl acrylamide) (PNIPAM–PEG–PNIPAM) exhibits thermoreversible gelation in aqueous solution.¹⁰ PNIPAM exhibits

Centre for Research in Topical Drug Delivery and Toxicology, Department of Clinical and Pharmaceutical Sciences, School of Life and Medical Sciences, University of Hertfordshire, UK. E-mail: m.cook5@herts.ac.uk

† Electronic supplementary information (ESI) available: Methods: HPLC methods and validation; calculation of critical micelle concentrations, solubilising powers, and micelle:water partition coefficients. Results: calibration curves, DOSY NMR, additional rheograms, DLS data for poloxamer 407, and mathematical fits to drug release profiles. See DOI: 10.1039/d0ma00080a

a lower critical solution temperature (LCST), resulting in a coil-to-globule transition above a critical temperature, typically 32 °C.¹¹ In the PNIPAM-PEG-PNIPAM copolymers, this transition leads to the formation of micelles, believed to have flower-like or fractal-like structures depending on molecular weight.^{12,13} At high concentrations (above *ca.* 20% w/v), PNIPAM-PEG-PNIPAM forms physical gels, hypothesized to be a result of entanglement of PNIPAM blocks,¹⁴ though this phenomenon has been studied in only a limited number of publications. It has previously been demonstrated that 20% (w/v) solutions of a PNIPAM (2.3 kDa)-PEG (4.6 kDa)-PNIPAM (2.3 kDa) copolymer undergo transition to a gel state when warmed above *ca.* 26 °C.¹⁴ Teodorescu *et al.*¹⁰ studied molecular weight effects on PNIPAM-PEG-PNIPAM gelation at concentrations of 20% (w/v) and below, varying PEG molecular weight between 1 and 6 kDa and PNIPAM between 5 and 30 kDa. It was found that only 4 and 6 kDa PEG with 20 or 30 kDa PNIPAM blocks resulted in gel formation, as determined by rheometry, with T_{gel} s of 38–43 °C. De Graaf *et al.* have demonstrated PNIPAM (16 kDa)-PEG (6 kDa)-PNIPAM (16 kDa) and PNIPAM (32 kDa)-PEG (6 kDa)-PNIPAM (32 kDa) exhibited gelation temperatures of 35 and 33 °C in phosphate buffered saline, allowing them to form a gel when injected into mice.¹⁵ This gel released paclitaxel in a controlled fashion and did not cause acute systemic toxicity. PNIPAM-PEG-PNIPAM is a potential thermoreversible gelator for topical drug delivery. However, understanding of its behaviour in drug delivery is limited and there are no studies evaluating its performance for topical application. Additionally, the solubilisation of drugs in concentrated PNIPAM-PEG-PNIPAM is limited,¹⁵ and there are no studies of saturation solubilisation and release in these systems. Furthermore, a critical evaluation of this novel material against poloxamer 407 has not been conducted. In order for novel excipients to translate into biomedical applications, clear advantages of these materials over the current approaches must be demonstrated. While gels are a traditional dosage form for topical application, there is a drive to generate novel materials with added functionality,¹⁶ and translation must be considered whilst driving this innovation.

In this research we study thermoreversible gelation in PNIPAM-PEG-PNIPAM with a view to topical drug delivery. The effect of concentration on T_{gel} and gel strength is studied, along with a critical evaluation of rheological performance against poloxamer 407. The stability of these materials is also reported for the first time. The materials are then compared side-by-side for the preparation of vaginal formulations, as an exemplar mucosal route, revealing a fundamental understanding of the behaviour of hydrophilic and hydrophobic drugs in these thermoreversible gelators.

Experimental

Materials

N-Isopropyl acrylamide (NIPAM) (97%), copper(I) bromide (CuBr) (98%), tetrahydrofuran (THF) (99%) and triethylamine (99.5%) were purchased from Sigma-Aldrich (UK). Tris[2-(dimethylamino)ethyl]amine (Me6TREN) (99%) and 2-bromoisobutyl

bromide (BiBB) (97%) were purchased from Alfa Aesar (UK). Isopropyl alcohol (99%) and dichloromethane (99%) were purchased from Fisher Scientific (UK). Bipyridine (99%), dimethylaminopyridine (DMAP) (97%) and PEG 10 kDa were purchased from Aldrich (UK). Aluminium oxide, neutral, Brockmann I was purchased from Acros Organics (UK). Dialysis tubing with a molecular weight cut off (MWCO) of 3500 Da was purchased from Medicell Membranes Ltd (UK) and soaked in deionised H₂O before use. GPC EasiVial poly(methyl methacrylate) mixed standards and a poly(methyl methacrylate) single standard (72 kDa) were purchased from Agilent (UK). Deionised H₂O was used in all experiments. All reagents were used as supplied, unless otherwise indicated.

Vaginal fluid simulant (VFS) was prepared by dissolution of NaCl (3.510 g, 0.060 mol), KOH (1.400 g, 0.025 mol), Ca(OH)₂ (0.222 g, 0.003 mol), bovine serum albumin (0.018 g), lactic acid (2.000 g, 0.022 mol), acetic acid (1.000 g, 0.011 mol), glycerol (0.160 g, 0.002 mol), urea (0.400 g, 0.007 mol) and glucose (5.000 g, 0.028 mol) in 1 L of deionised water. The solution was then adjusted to pH 4 with 1 M hydrochloric acid. This formula follows that recommended by Owen and Katz.¹⁷

Porcine vaginal tissue was purchased from WetLab (UK) and supplied frozen. The porcine vaginal mucosa was removed by incision with a scalpel. Once the tissue was removed it was cut into 1 cm² squares which were attached to the texture analyse probe using adhesive pads.

Synthesis of PEG macroinitiator

A 10 kDa PEG macroinitiator was synthesised using reported methods.¹⁸ DMAP (1.17 g, 9.6 mmol) in anhydrous dichloromethane (8 mL) was mixed with triethylamine (0.89 mL, 6.4 mmol) and cooled to 0 °C. BiBB (1.97 mL, 16.0 mmol) in anhydrous dichloromethane (8 mL) was then added to the DMAP and triethylamine solution. A solution of PEG (10 kDa, 16.0 g, 1.6 mmol) in dichloromethane (160 mL) was subsequently added dropwise over 1 h. When the addition of PEG was complete, the reaction was allowed to rise to room temperature and the reaction allowed to proceed with stirring for 18 h. The solution was filtered, and approximately half of the solvent in the filtrate removed *in vacuo*. The PEG initiator was then precipitated in cold diethyl ether (480 mL) and filtered. The crude PEG macroinitiator in the retentate was then recrystallised from absolute ethanol (300 mL) overnight. The resulting solid was filtered and washed with cold diethyl ether and dried *in vacuo* (yield = 86.5%).

Synthesis of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ by atom-transfer radical polymerisation (ATRP)

NIPAM was polymerised from the PEG macroinitiator by ATRP following established procedures.¹² Firstly, the CuBr catalyst (28.7 mg, 200 µmol) was sealed in a round bottom flask and degassed for 30 min by nitrogen purging. Separately, the Me6TREN ligand (53.5 µL, 200 µmol), NIPAM monomer (2.0 g, 17.7 mmol) and PEG 10 kDa macroinitiator (1.0 g, 100 µmol) were dissolved in deionised water (10 mL), within a sealed round bottomed flask. The solution was then degassed by nitrogen



Mater. Adv., 2020, 1, 371–386 | 373

by centrifugation (10 min at 14 500 rpm) and the clear supernatant was analysed by HPLC. Each experiment was performed in triplicate.

Release of progesterone and tenofovir disoproxil fumarate from thermoreversible gelators

The release of progesterone and tenofovir disoproxil fumarate from the thermoreversible gelators (20% w/v poloxamer 407 or 50% w/v PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈) was investigated using Franz diffusion cells equipped with a cellulose membrane (MWCO 3.5 kDa) at both 25 and 37 °C. The experiments used Franz cells with an average phosphate buffer saline (PBS) receiver fluid volume of 12.3 mL and an average bore area of 174 mm². The release of 50 µg mL⁻¹ progesterone or 200 µg mL⁻¹ tenofovir disoproxil fumarate was investigated, where this concentration did not violate sink conditions. A 20% w/v ethanol solution in PBS containing 50 µg mL⁻¹ progesterone and a 200 µg mL⁻¹ tenofovir disoproxil fumarate solution in PBS were used as controls. The Franz cells were then placed into a water bath for 30 min to allow the receiver fluid to reach the temperature of the surrounding water (25 or 37 °C). All Franz cells were dosed with 200 µL of sample and the release of drug measured at regular intervals by sampling receiver fluid (500 or 200 µL for the progesterone and tenofovir disoproxil fumarate respectively). Receiver fluid was replaced with an equal volume of pre-warmed PBS. The samples were analysed by HPLC (method and calibration (Fig. S1) in ESI†). The experiment was repeated 4 times.

Assessing the adhesion and mucoadhesion of thermoreversible gelators

The adhesion and mucoadhesion of the thermoreversible gelators (20% w/v poloxamer 407 or 50% w/v PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈) was assessed using a TAX.T Plus Texture Analyser (Stable Micro Systems, UK) with a poly(propylene) probe with a surface area of 1.25 cm². This probe was used to determine adhesion values, whereas mucoadhesion was evaluated using porcine vaginal mucosa tissue (WetLab, UK). The thermoreversible gelator was placed into a water bath at 25 °C, the probe was lowered until contact between the probe or vaginal mucosa tissue was made. The temperature of the water bath was then increased to 37 °C and held at this temperature for 2 min to ensure gel formation, and mimic *in situ* gelation. Once a gel had formed, the probe was withdrawn at a rate of 10.0 mm s⁻¹ until complete detachment was observed, as recommended in prior studies.¹⁹ The maximum force of detachment and the area under the force-displacement curve were determined using Texture Exponent 32 software (Stable Micro Systems, UK) and designated the “force of adhesion” and “work of adhesion”, respectively. All adhesion testing was performed 6 times. The mucoadhesion testing was performed using three different porcine vaginal tissues where three samples of vaginal tissue were taken from each vagina. Data was expressed as a mean of triplicate experiments ($N = 3$), where N was the mean value of the three measurements taken from a single vagina.

Stability of thermoreversible gelators

The stability of thermoreversible gelators (20% w/v poloxamer 407 or 50% w/v PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈) was assessed over 12 weeks. The polymer solutions were prepared in HPLC vials sealed with parafilm and placed in the refrigerator (4 °C) or in ovens set at 25 and 40 °C representing room temperature and accelerated storage conditions, respectively. At weekly intervals, the samples were lyophilised and the molecular weight determined by GPC, as described previously. Sufficient samples were prepared so that each weekly measurement was taken on a separate sample. The experiment was performed in triplicate.

Statistical analysis

All statistical analysis was conducted on Prism5 (GraphPad, USA). Data was analysed by one-way analysis of variance (ANOVA) with Tukey *post hoc* analysis unless otherwise stated. Data presented as means $\pm$ standard deviation.

Results and discussion

PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ was synthesised in a two-step process by covalent attachment of an ATRP initiator to PEG₁₂₂ using a simple S_N2 reaction with DMAP-activated BiBB followed by ATRP polymerisation from the macroinitiator using a Cu(I)Br/ME₆TREN catalyst/ligand complex (Fig. 1(i)).^{12,18} GPC confirmed that the resultant copolymer had a monomodal molecular weight distribution with a polydispersity index of 1.88 (Fig. 1(iii)), in line with previous publications.¹² The degree of polymerisation was calculated by ¹H NMR (Fig. 1(iv)), finding that the polymer had the structure PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈. DOSY NMR (Fig. S2, ESI†) unambiguously confirmed attachment of PNIPAM to PEG by demonstrating that the two blocks have the same diffusion coefficient.

All blocks of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ are water-soluble at room temperature, but the copolymer is expected to transition to an amphiphilic state when warmed due to the desolvation of PNIPAM. The comparator pharmaceutical excipient, poloxamer 407, is shown in Fig. 1(ii). This ABA copolymer exhibits water solubility at temperatures below *ca.* 20 °C but transitions to an amphiphilic state when warmed due to the desolvation of the central poly(propylene oxide) block. Desolvation of the central block leads to a hierarchical process of micelle formation (hydrodynamic radius *ca.* 10 nm) and subsequent packing into a face-centred cubic lattice to form a transparent gel mesophase.^{7,20}

The thermoresponsive gelation process exhibited by PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ was studied by rheometry with variation of polymer concentration between 20–50% (w/v) (Fig. 2(i)). The copolymer exhibited dramatic increases in G' and G'' above a critical temperature (T_{thick}), at *ca.* 30–35 °C believed to be associated with the desolvation of PNIPAM.²¹ All rheograms exhibited a gelation temperature (T_{gel}) above *ca.* 35 °C, at which point the absolute value of G' surpassed that of G'' , indicative of a transition from a viscous fluid to an elastic gel state.



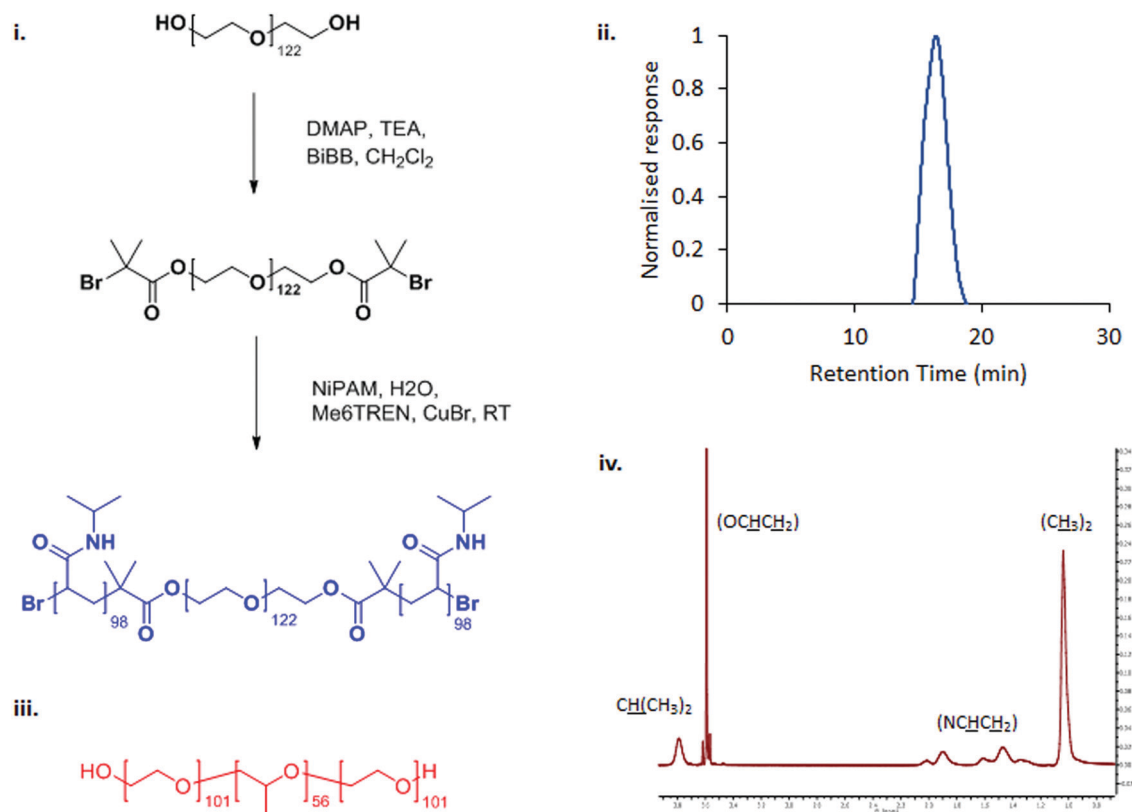


Fig. 1 The synthetic route to PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (i) and chemical structure of poloxamer 407 (iii). PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ was characterised by GPC (ii) and ¹H NMR (iv).

These values of T_{thick} and T_{gel} are attractive for biomedical applications where the polymer solutions will have low viscosity at room temperatures, but thicken when warmed by the body. Poloxamer 407 exhibited a markedly different thermoresponsive gelation to PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈. Low viscosity gels ($G' \text{ ca. } 100 \text{ Pa}$) were formed at 15% (w/v), with T_{gel} at 45 °C. Viscous gels ($G' \text{ ca. } 10 \text{ kPa}$) were formed at 20 and 25% (w/v) with $T_{\text{gel}} < 25 \text{ °C}$. At 30% (w/v) the materials were gels over the whole temperature range studied.

T_{gel} , T_{thick} and the maximum absolute value of G' reached (G'_{max}) were extracted from the rheograms shown in Fig. 2 and Fig. S3 (ESI†) and are presented in Fig. 3. PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (Fig. 3(i)) exhibited a monotonic, near-linear, decrease in T_{gel} from 44 to 36 °C across the concentration range studied. A linear fit ($R^2 = 0.96$) indicates that the rate of T_{gel} depression is $-0.25 \text{ °C g}^{-1} \text{ dL}$, allowing tight control of T_{gel} over this range. T_{thick} decreased from 36 to 29 °C between 20 and 50% (w/v) concentration, allowing thickening to occur upon warming by the body. An inverse proportionality between T_{gel} and concentration has previously been observed in PNIPAM-poly(*N,N*-dimethylacrylamide)-PNIPAM solutions, and was attributed to the depressed LCST observed in PNIPAM solutions at high concentrations.²² G'_{max} increased with concentration from 1.5 kPa at 20% (w/v) to a maximum of 12.7 kPa at 45% (w/v). An increase in gel strength with concentration was also observed by Kirkland *et al.*²² when studying

PNIPAM-poly(*N,N*-dimethylacrylamide)-PNIPAM. The authors rationalise this phenomenon using the theory of Semenov *et al.*,²³ that telechelic polymers with associating end-groups form flower-like micelles bridged by polymer chains resulting in elasticity. Kirkland *et al.* suggest that a greater number of polymer chains results in additional bridges formed between micellar domains, increasing viscosity.²² It is also conceivable that a greater number of micelles increases the degree of overlap between these aggregates, in a manner analogous to that observed in poloxamer 407.²⁴ Values of T_{gel} for PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ were typically *ca.* 5 °C greater than T_{thick} , whereas poloxamer 407 gave a sharper transition at 20 and 25% (w/v) with T_{gel} within 1 °C of T_{thick} .

Poloxamer 407 exhibited a far greater dependence of T_{gel} on concentration than PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (Fig. 3(ii)), with T_{gel} decreasing from 45 to 21 °C as the concentration was increased from 15 to 25% (w/v). At 30% (w/v) no T_{gel} was determined, with the rheograms exhibiting $G' > G''$ at all temperatures. These findings are in line with established phase behaviour of pluronics.²⁰ Of the concentrations studied, only 17.5 and 20% (w/v) poloxamer 407 samples (T_{gel} of 30 and 5 °C, respectively) exhibited $T_{\text{thick}}/T_{\text{gel}}$ at a temperature suitable for *in situ* thickening upon contact with the body (*i.e.* 25 °C < $T_{\text{thick}} < 37 \text{ °C}$), albeit with a T_{gel} which may be reached at real room temperatures in warmer climates, *e.g.* WHO climatic zones III and IV (30 °C).²⁵ The T_{thick} of 15% poloxamer 407 was at body



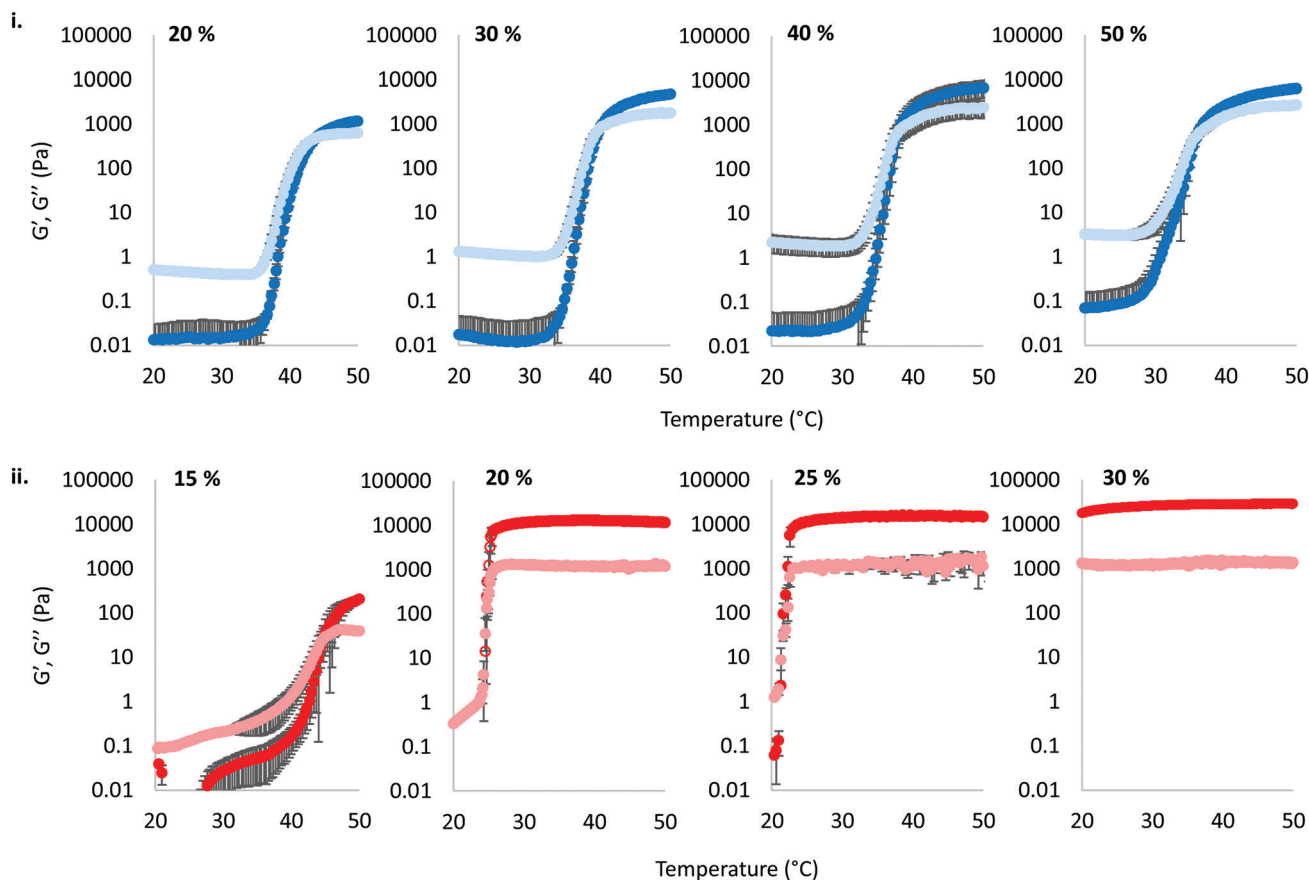


Fig. 2 Temperature ramp rheograms of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (i) and poloxamer 407 (ii) with variation of concentration (% w/v) at a fixed shear stress (1 Pa) and frequency (1 Hz). G' is presented as the dark colour, whilst the light colour corresponds to G'' . Data presented as mean $\pm$ standard deviation, $n = 3$.

temperature (37 $^{\circ}\text{C}$). Poloxamer 407's G'_{max} was proportional to concentration, and had a value of 6.5 kPa at 17.5% (w/v) poloxamer 407 and 12.7 kPa at 20% (w/v). Due to the larger gel strength of the 20% (w/v) formulation and its prevalence in the literature,⁶ this concentration was selected for comparison with PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ in future experiments. It is known that poloxamer 407 forms a gel *via* the packing of micelles into a cubic structure and increasing concentration leads to an increased volume fraction of micelles, and thus a greater degree of overlap between the micelles.²⁴ The increase in G'_{max} with concentration observed for the poloxamer samples is attributed to this greater degree of overlap and thus internal friction. The gels formed by 50% (w/v) PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ were translucent, whilst the 20% (w/v) poloxamer 407 gels were clear (Fig. 3(iii)). This indicates that the poloxamer micelles are sufficiently small to avoid scattering of visible light, but that the PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ aggregates are larger and the Tyndall effect is observed.

Overall, the rheological behaviour of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ and poloxamer 407 with temperature are distinct. Where 20–50% (w/v) PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ solutions would increase in viscosity when warmed from room to body temperature, this was observed only in 17.5 and 20% (w/v) poloxamer 407 solutions. The lower concentration dependence

of T_{gel} seen for PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ is attractive where dilution effects have been observed to affect poloxamer 407's gelation *in vivo*.²⁶ Achieving *in situ* thickening at high polymer concentrations also allows for viscous gels to be formed with > 40% w/v PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈, achieving G'_{max} values of 11–13 kPa, whilst retaining a $T_{\text{thick}}/T_{\text{gel}}$ close to body temperature.

PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (50% (w/v)) was further explored as a smart material for drug delivery. Its T_{gel} of 36 $^{\circ}\text{C}$ is highly attractive for administration onto or into the body at sites where the temperature is 37 $^{\circ}\text{C}$. In particular, the material may have application in vaginal drug delivery where the local temperature is expected to be 37 $^{\circ}\text{C}$. Rheology was used to simulate the topical application process and determine the time required for the gel phase to form (Fig. 4(i)). PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (50% (w/v)) was oscillated at 1 Pa shear stress and a frequency of 1 Hz while temperature was varied. The sample was set constant to room temperature (25 $^{\circ}\text{C}$) for 60 s before holding at body temperature (37 $^{\circ}\text{C}$) for 240 s. The transition from 25 to 37 $^{\circ}\text{C}$ led to immediate thickening of the sample, which formed a gel after 87 ± 5 s and plateaued at a G' of ca. 700 Pa. This relatively long gelation time may have clinical implications, where a patient may be asked to remain still until gel formation has occurred. The same



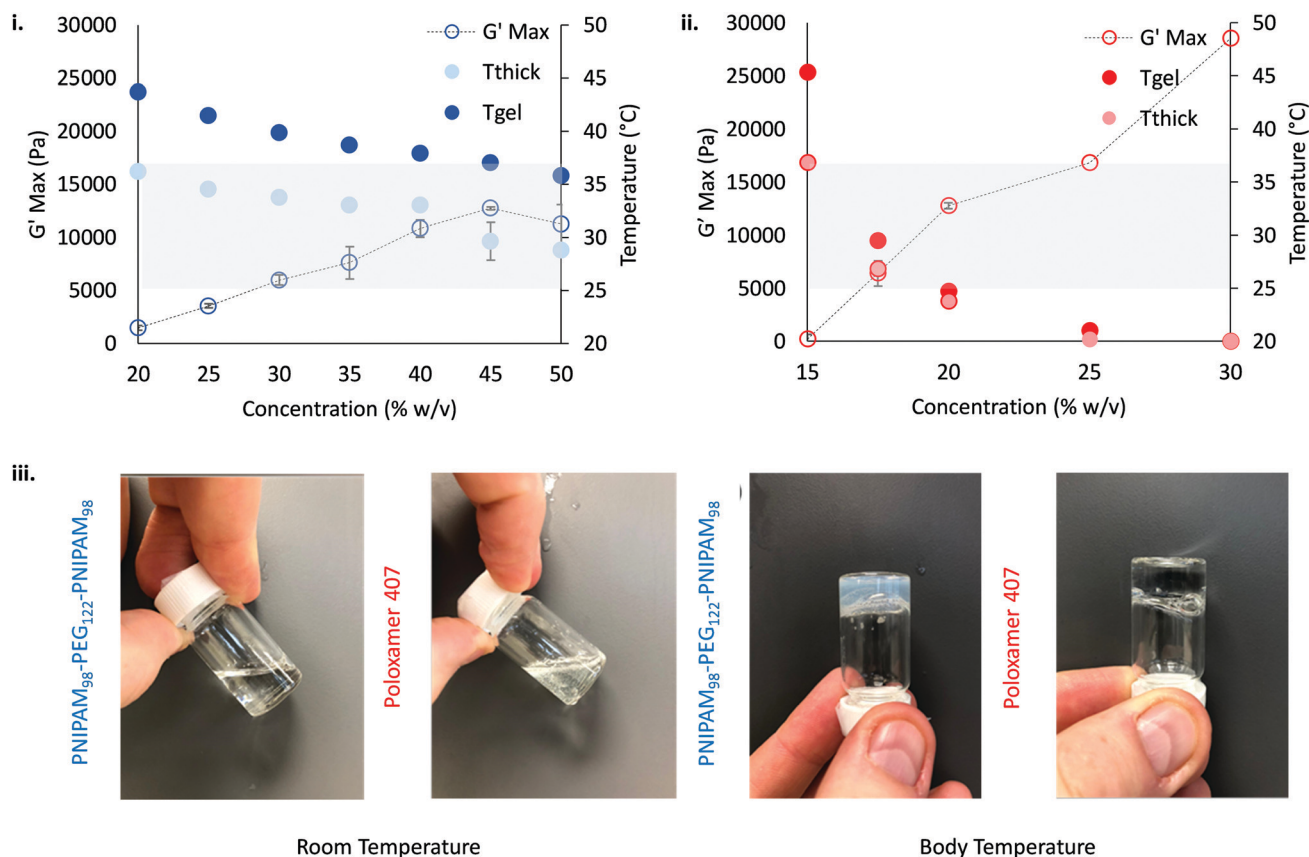


Fig. 3 T_{thick} (light circle), T_{gel} (dark circle), and G'_{max} (hollow circle with dashes) as a function of concentration for PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (i, blue) and poloxamer 407 (ii, red). The temperature range which would allow for *in situ* thickening of polymer solutions is overlaid in grey. Please note that T_{thick} and T_{gel} are omitted for 30% (w/v) poloxamer 407 as they were not present in the rheogram. Data presented as mean $\pm$ standard deviation, $n = 3$. Images of the thermoresponsive gelators are included (iii) at room temperature and body temperature, 37 °C.

experimental procedure for poloxamer 407 (20% (w/v)) (Fig. S4, ESI†) gave a time to gel of 27 ± 5 s, with a plateau at *ca.* 5.5 kPa indicating a more rapid and rigid gel formation for this sample. The PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ gel formed at 37 °C was subjected to an oscillatory stress sweep which demonstrated thinning above *ca.* 200 Pa with a gel yield observed at 247 ± 72 Pa, which was not significantly different ($p > 0.05$ by *t*-test) than poloxamer 407 (20% (w/v)) which exhibited a yield at 256 ± 58 Pa (Fig. S5, ESI†). The yield strength is an indicator of the force required to make the sample undergo viscous flow, and may be taken as the point where G' becomes lower than G'' during an oscillatory stress sweep in the gel phase.²⁷ An oscillatory frequency sweep of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (50% (w/v)) at 37 °C confirms that the structure is a rheological gel, with $G' > G''$ at all frequencies measured (Fig. 4(iii)). Finally, the PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (50% (w/v)) was cycled between 25 and 37 °C at 1 Pa and 1 Hz (Fig. 4(iv)), which indicated that the thermoresponsive gelation process was reversible and repeatable.

The effect of physiological fluids on the phase transition of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ and poloxamer 407 was then evaluated by preparing polymer solutions in VFS (Fig. 5).¹⁷ The rheological temperature ramp for PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (50% (w/v)) was altered considerably in VFS relative to

deionised water. T_{gel} shifted to 32.9 ± 0.2 °C in VFS relative to 35.8 ± 0.2 °C in deionised water. This depression of T_{gel} was also seen in poloxamer 407 which formed a gel at 22.3 ± 0.4 °C in VFS compared to 24.7 ± 0.2 °C in deionised water. The depression of T_{gel} seen in PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ is attributed to salting-out of the PNIPAM block. Chaotropes have been shown to reduce LCSTs through an increased surface tension between PNIPAM hydrophobic domains (*i.e.* isopropyl groups and the carbon backbone) and their hydration layer.²⁸ T_{gel} depression in poloxamer 407 is also attributed to salting-out, where salts are known to entropically promote the micellization process.²⁹ VFS did not affect G'_{max} of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (50% (w/v)), which was 11.7 ± 1.5 and 11.2 ± 1.8 kPa in VFS and deionised water, respectively ($p < 0.05$ by *t*-test). However, the depression in T_{gel} did result in a greater value of G' at 37 °C for PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (50% (w/v)) in VFS compared in deionised water, giving values of 4.8 ± 2.8 kPa and 1.1 ± 0.8 kPa, respectively. G' of poloxamer 407 (20% (w/v)) at 37 °C was unaffected by the presence of VFS, with both VFS and deionised water leading to values of G' of *ca.* 13 kPa at this temperature.

A significant limitation of poloxamer 407 as a thermoresponsive gelator is its rapid dissolution in physiological fluids.⁶ 20% (w/v) solutions of poloxamer 407 dissolved in VFS within





Fig. 4 (i) Determination of gelation time for PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (50% (w/v)). (ii) Oscillatory stress sweep of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (50% (w/v)), and (iii) oscillatory frequency sweep. (iv) Thermal cycling of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (50% (w/v)) to demonstrate reversibility of the gelation process. G' and G'' are shown as dark and light blue markers, respectively. Data presented as mean $\pm$ standard deviation, $n = 3$.

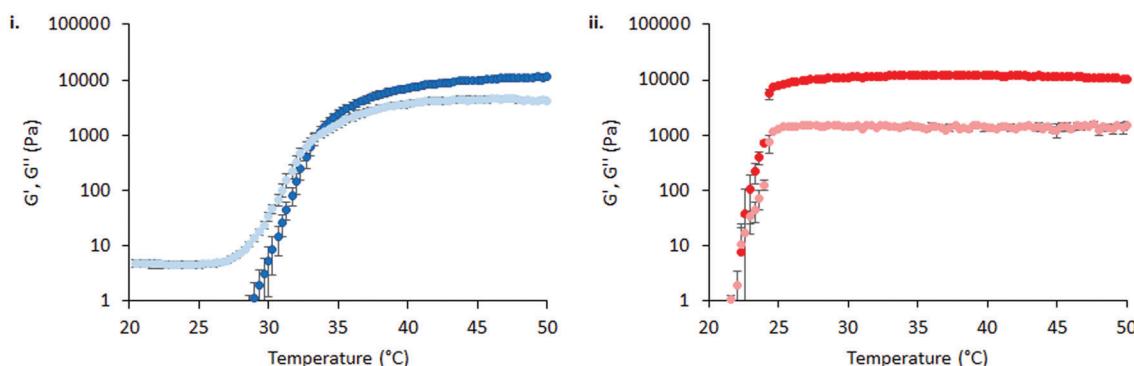


Fig. 5 Temperature ramp rheograms of 50% (w/v) PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (i) and 20% (w/v) poloxamer 407 (ii) in VFS at a fixed shear stress (1 Pa) and frequency (1 Hz). G' and G'' are shown as dark and light markers, respectively. Data presented as mean $\pm$ standard deviation, $n = 3$.

60 min (Fig. 6). PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (50% (w/v)) had a significantly ($p < 0.05$) greater resistance to dissolution, requiring 230 min for dissolution to occur. A control of 50% (w/v) poloxamer 407 dissolved after 140 min, which indicated that this difference is not solely explained by concentration. It is hypothesised that where poloxamer 407 gels are composed of non-interacting polymeric micelles, the liberation of micelles into the dissolution medium occurs rapidly. PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ micelles are thought to be bridged by polymeric unimers which reduce the favourability of micelle liberation into the dissolution medium.^{14,23} These experiments were conducted in a large excess of dissolution media, and the dissolution process in the real, smaller, volumes of

physiological fluid present *in vivo* is likely to occur over a longer period of time.

The mucoadhesion of poloxamer 407 is weak,³⁰ which limits its residence time on mucosal membranes. Mucoadhesion is mediated by several different factors, which are not mutually exclusive. Briefly, entanglements may occur between macromolecules in the dosage form and mucin glycoproteins coating the mucosa, which are supported by non-covalent interactions, enhancing adhesion.³¹ The viscosity of a gel will enhance retention, whilst the movement of moisture from the mucosa to the dosage form will either improve or decrease mucoadhesion depending on levels of hydration.³² Several other theories exist.³³ The most common method to determine mucoadhesion is by



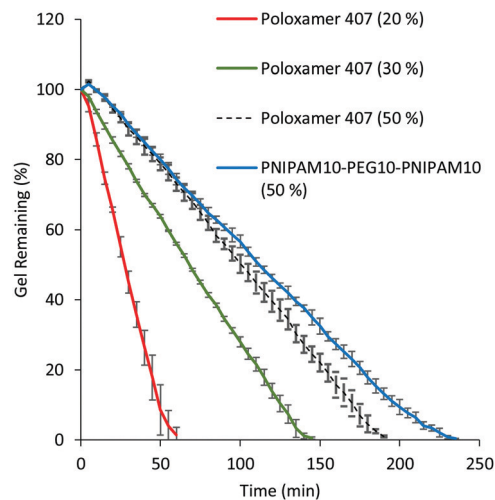


Fig. 6 Dissolution of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ 50% (w/v) (blue) and 20 (red), 30 (green) and 50% (w/v) (black) poloxamer 407 in VFS. Data presented as mean $\pm$ standard deviation ($n = 3$).

measuring the force-displacement curve during removal of the dosage form from a membrane, where the peak force is termed the “force of adhesion” and the total area under the curve is the “work of adhesion”.³⁴ These adhesion processes were studied for 50% (w/v) solutions of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ as well as 20, 30, and 50% (w/v) poloxamer 407. Only 20% (w/v) solutions of poloxamer 407 exhibit T_{gel} at a relevant temperature (25 °C), but 30 and 50% (w/v) solutions were explored to account for differences in concentration between the two thermoreversible gelators and understand whether adhesion processes are affected by chemical structure or concentration. Firstly, the adhesion of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ and poloxamer 407 to a poly(propylene) probe was assessed (Fig. 7). In this control experiment, van der Waals forces are believed to be the

major contributor to adhesion. The data demonstrates that the adhesion of poloxamer 407 increases with concentration, and that 50% (w/v) solutions of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ and poloxamer 407 had equivalent ($p > 0.05$) adhesion to the probe. The mucoadhesion of 20% (w/v) poloxamer 407 and 50% (w/v) PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ is shown in Fig. 7. PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (50% (w/v)) had greater force and work of adhesion than 20% (w/v) poloxamer 407 ($p < 0.01$) which is desirable for mucosal drug delivery. A control of 50% (w/v) poloxamer 407 was equivalent to the PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (50% (w/v)) which demonstrates that this enhanced adhesion is likely to be related to concentration, rather than enhanced specific intermolecular interactions between polymer and mucosa. However, concentrations of poloxamer 407 above 20% (w/v) did not exhibit a T_{gel} in the range required and thus are not appropriate for *in situ* gelation with topical administration. Overall, PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ allows the formation of *in situ* gelators ($25 < T_{gel} < 37$ °C) at 50% (w/v), which imparts a greater level of mucoadhesion than seen for 20% (w/v) poloxamer 407. 50% (w/v) poloxamer 407 had equivalent mucoadhesion to 50% (w/v) PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ but it is not a thermoreversible gelator, existing in a gel phase at all temperatures studied.

The thermoresponsive polymers were then evaluated for their ability to solubilise two drugs relevant to intravaginal drug delivery: progesterone and tenofovir disoproxil fumarate. Progesterone is administered intravaginally to treat low fertility, premenstrual syndrome, and puerperal depression, as well as support *in vivo* fertilisation and reduce the risk of pre-term delivery in pregnancy.^{35–38} Intravaginal tenofovir disoproxil fumarate has been shown to reduce transmission of HIV in a macaque model.³⁹ In addition to their relevance to vaginal drug delivery, progesterone and tenofovir disoproxil fumarate were selected as they have large differences in their water solubilities.



Fig. 7 Adhesion (i) and mucoadhesion (ii) of 50% (w/v) PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ and poloxamer 407 onto a poly(propylene) probe or porcine vaginal tissue, respectively.



Progesterone was determined to have a water solubility of $9.8 \pm 0.1 \mu\text{g mL}^{-1}$ at 25°C , whilst tenofovir disoproxil fumarate's solubility was approximately 1000-fold greater at $9.5 \pm 1.0 \text{ mg mL}^{-1}$. The solubility of the two drugs in the presence of the polymers at a range of concentrations between $5 \mu\text{g mL}^{-1}$ and 10 mg mL^{-1} was determined at 25 and 37°C (Fig. 8). It was found that for both polymers the solubility of progesterone (Fig. 8(i)) was greatly increased above a critical concentration, believed to be the critical micelle concentration (CMC). The CMC could then be extracted from the data, giving values of 0.26 and 1.88 mg mL^{-1} at 25°C for PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ and poloxamer 407 respectively, which rose to 0.68 and 1.95 mg mL^{-1} at 37°C . The CMC of poloxamer 407 is on the same order of magnitude as that reported previously (*ca.* 3 mg mL^{-1}) for the pure polymer,⁴⁰ and the deviation is attributed to the presence of progesterone, as reported for other drugs.⁴¹ Tenofovir disoproxil

fumarate (Fig. 8(ii)) solubility was not enhanced by increasing concentration of either polymer, even above the CMCs. The data indicates that the formation of micelles occurs for both polymers above a CMC at 25 and 37°C , at which point the micelles provide a locus for solubilisation for the relatively hydrophobic progesterone, but tenofovir disoproxil fumarate is excluded from the micelle. The solubilities of both drugs are increased at 37°C , above and below the critical micelle concentration, with a more pronounced effect seen in tenofovir disoproxil fumarate which is attributed to its enhanced water solubility at this temperature.

Dynamic light scattering revealed a pronounced influence of progesterone on the micellization of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈. 10 mg mL^{-1} solutions of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ exhibited no detectable particulates at 25°C but when warmed to 37°C transitioned to a 138 nm nanoparticle, which shrunk and remained at a constant hydrodynamic diameter of



Fig. 8 Solubilisation of progesterone (i) and tenofovir disoproxil fumarate (ii) by PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (blue) and poloxamer 407 (red) at 25 and 37°C (light and dark colours, respectively). Inset: Schematic representation of the micellization process. (iii) The scattering counts of a 10 mg mL^{-1} PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ solution saturated with progesterone as a function of temperature. (iv) Hydrodynamic diameter of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ nanoparticles saturated with progesterone at 25 (light blue), 37 (dark blue), and 40 (black) $^\circ\text{C}$, and PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ without progesterone at 40°C (black dashed).



ca. 90 nm at temperatures at 40 °C and above (Fig. S6, ESI†). In the presence of progesterone, nanoparticle aggregates were detected at 25 °C. Furthermore, at 38 °C and above, a pronounced transition was observed (Fig. 8(iii)). The hydrodynamic diameter of the nanoparticles in the presence of progesterone at 25, 37, and 40 °C were 15 ± 1 , 15 ± 1 , and 73 ± 0 nm, respectively (Fig. 8(iv)), demonstrating the change in nanostructure associated with the transition at 38 °C. It is hypothesised that the presence of progesterone drives the formation of micellar nanoparticles below the LCST-type transition of PNIPAM, which, when triggered by a rise in temperature, leads to the aggregation or rearrangement of these micelles. The size of these particles is close to those found without progesterone at 40 °C (88 ± 1 nm) (Fig. 8(iv)). Prior study of PNIPAM-PEG-PNIPAM reports PNIPAM (16 kDa)-PEG (6 kDa)-PNIPAM (16 kDa) formed flower-like micelles with a hydrodynamic diameter of 62 nm,¹³ whereas lower molecular weight PNIPAM (4 kDa)-PEG (6 kDa)-PNIPAM (4 kDa) formed fractal-like structures with hydrodynamic diameters of up to 400 nm.¹² Due to the closer molecular weight of the constituent unimers, and similar hydrodynamic diameters, it is believed that PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ forms flower-like micelles in a similar manner to PNIPAM (16 kDa)-PEG (6 kDa)-PNIPAM (16 kDa).

The solubilising power and micelle:water partition coefficients of the polymer/drug mixtures were calculated from the progesterone solubilisation data in Fig. 8(i) and are included in Table 1.⁴² Briefly, the solubilising power reflects the unit mass increase in progesterone solubilised per unit mass of polymer, whereas the micelle:water partition coefficient reflects the concentration of drug within the micelle relative to that free in solution. PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ had a solubilising power approximately 1.5-fold greater than poloxamer 407 at both temperatures, and a micelle:water partition coefficient around twice as large. These differences were demonstrated to be statistically significant by two-way ANOVA with Bonferroni *post hoc* testing ($p < 0.0001$). It is known that the greater the molecular weight of the relatively hydrophobic poly(propylene oxide) domains, the greater the solubilising power of poloxamers.⁴³ The microphase separated NIPAM domains (DP: 196) in PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ have a greater degree of polymerisation (and molecular weight) than the poly(propylene oxide) in poloxamer 407 (DP: 65) which may lead to a greater solubilising power. These considerations are balanced against the free energy of the solubilisation process, which will differ between the two polymers.⁴³

Temperature increased the mean value of solubilising power for both polymers, however this increase was only statistically

significant ($p < 0.05$) for PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈. As no structural transition was observed between 25 and 37 °C by dynamic light scattering, it is hypothesised that this small increase is the result of entropic effects favouring polymer-progesterone interactions at the elevated temperature. Poloxamer exhibited a structural transition at 26 °C (Fig. S7, ESI†), increasing in hydrodynamic diameter from ca. 9 to 22 nm, however this did not affect the solubilisation of progesterone.

The saturation solubilities of progesterone and tenofovir disoproxil fumarate were then evaluated in the thermoreversible gels at high concentrations (Fig. 9). Strikingly, progesterone solubility is dramatically enhanced at 37 °C, relative to 25 °C, in the polymer solutions where the effect is small in water alone. This has important implications for drug delivery across a membrane, where liberation of drug from a dosage form is dictated by its thermodynamic activity within that base.⁴⁴ The greater the degree of saturation, the greater the driving force for liberation. If progesterone is loaded into the solutions at room temperature then applied to the body and warmed then the thermodynamic activity is expected to decrease, thus liberation from the dosage form is expected to be slowed at this higher temperature. Despite the differences in solubilising power and concentration of the two polymers, progesterone saturation solubility in both polymer mixtures at 37 °C was equivalent, at 505.0 ± 7.8 and $504.8 \pm 30.2 \mu\text{g mL}^{-1}$ for poloxamer 407 and PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈, respectively, equivalent to a ca. 45-fold increase relative to the saturation solubility in water ($11.3 \mu\text{g mL}^{-1}$). The molarities of the two solutions are approximately 16.5 and 18.2 mM for poloxamer 407 and PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ respectively, and the greater solubilising power of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ is expected to lead to a much larger increase in progesterone solubility in this material. However, nanostructures present at the low concentrations studied for solubilising power calculations ($< 10 \text{ mg mL}^{-1}$) may not be equivalent to those found at 50% (w/v) in the PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈. It is known that poloxamer exists as spherical micelles of approximately 10 nm diameter across a wide range of concentrations, however the nanostructure of the PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ materials at 50% (w/v) has never been studied and block copolymers may exist in different phases dependent on concentration.²⁴ It is further suggested that the high concentration of the PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ has driven the micellar transition observed in Fig. 8(iii) at 38 °C to a temperature lower than 37 °C, in line with the depression in T_{gel} seen at high concentrations by rheometry. The solubilising power of poloxamer 407 (Table 1) would suggest that progesterone solubility would be unaffected by temperature, however behaviour at this high concentration (20% w/v) may differ from that in

Table 1 Solubilising power and micelle:water partition coefficients calculated from data presented in Fig. 8

Polymer	Solubilising power ($\times 10^{-3}$)		Micelle:water partition coefficient	
	25 °C	37 °C	25 °C	37 °C
Poloxamer 407	1.53 ± 0.07	1.65 ± 0.22	1.30 ± 0.06	1.21 ± 0.16
PNIPAM ₉₈ -PEG ₁₂₂ -PNIPAM ₉₈	2.45 ± 0.22	2.86 ± 0.09	2.37 ± 0.08	2.54 ± 0.08



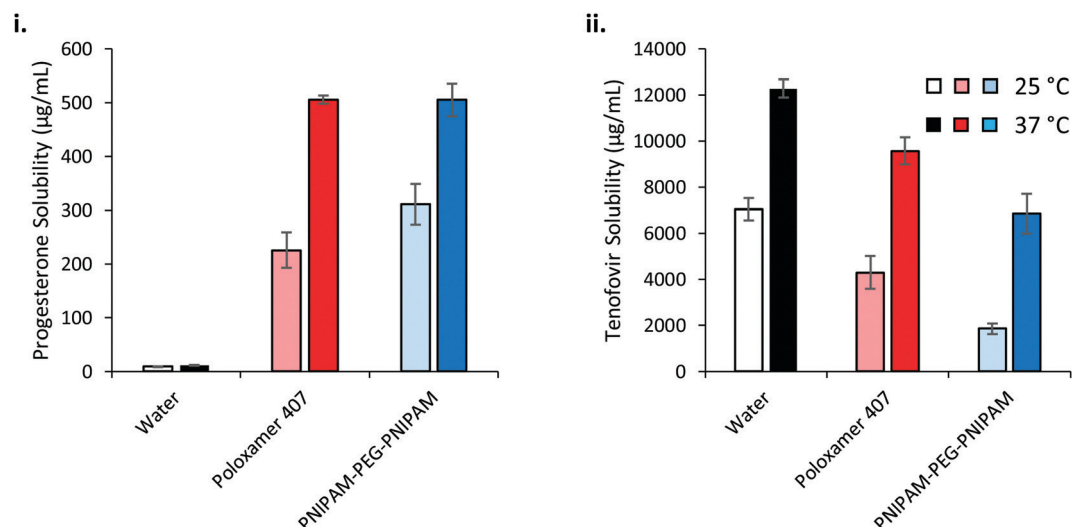


Fig. 9 Saturation solubility of progesterone (i) and tenofovir disoproxil fumarate (ii) in 20% (w/v) poloxamer 407 (red) and 50% (w/v) PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (blue) at 25 and 37 °C (light and dark colours, respectively).

dilute solution, as has been reported for micelle:water partition coefficients.⁴⁵

Tenofovir disoproxil fumarate is excluded from the micelles formed by either polymer, and as such the increase in solubility observed in these polymers at elevated temperatures is attributed solely to the improvement in solubility caused by heat alone (Fig. 9(ii)). The reduction in solubility seen in the polymer samples is attributed to the decreased volume fraction of water in these mixtures. Adjusting for the phase volume of water accounts for this depression.

The liberation of drugs from the thermoreversible gelators across cellulose membrane was then explored at loading concentrations below the saturation limit (Fig. 10), maintaining sink conditions throughout the experiments. Control solutions of each drug were included to ensure that permeation across cellulose membrane was not rate-limiting, and each of these experiments give a characteristic curve associated with donor depletion in finite dosing experiments.⁴⁶ Tenofovir disoproxil fumarate release from the two polymer solutions was equivalent at 25 °C. However, drug release was significantly ($p < 0.05$) enhanced at 37 °C for poloxamer 407 but remained constant in the case of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈. For example, at 8 h the cumulative release at 25 °C was $58.3 \pm 1.5\%$ and $61.0 \pm 3.8\%$ for PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ and poloxamer 407, respectively, but $53.8 \pm 4.3\%$ and $86.6 \pm 5.3\%$ for the same samples at 37 °C. As drug release is occurring from a planar dosage form, the Higuchi model was applied to the data. This simple model is based on a linear fit to the fractional drug release with the square-root of time.⁴⁷ The model gave $R^2 > 0.98$ in all cases, supporting the principle that the gels act as a matrix controlled release system from which Fickian diffusion occurs (Fig. S8, ESI†).⁴⁷ Heat will enhance diffusion out of the matrix when release is purely diffusion controlled as the Stokes-Einstein equation predicts that the diffusion coefficient of a molecule scales linearly with temperature, when viscosity is constant.⁴⁸ The control, a solution of tenofovir disoproxil

fumarate exhibits this phenomenon. In the poloxamer 407 samples, which are a gel at both temperatures tested, an enhanced release from the gels is attributed to the increased diffusion coefficient, despite the increased viscosity of the system. For PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈, a phase transition occurs between the two temperatures tested. The identical release rate at 37 °C relative to 25 °C is attributed to the formation of this gel phase, which provides a more tortuous path for liberation to occur where tenofovir is known to be excluded from the micelles, counteracting the increased diffusion coefficient.⁴⁷

Progesterone release from PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ and poloxamer 407 samples was dramatically different than that exhibited by tenofovir disoproxil fumarate. Progesterone release from PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ at 25 °C followed Higuchi kinetics (Fig. S8, ESI†), and 100% of the drug was released after 32 h. Increasing the temperature to 37 °C retarded the release of progesterone significantly, with 100% drug liberated only after 144 h. This release profile did not fit the Higuchi model, but exhibited a good fit ($R^2: 0.98$) to the Korsmeyer-Peppas power law, albeit with an extreme exponent ($n = 2.42$) designating non-Fickian super case II transport kinetics.⁴⁹ This retardation of release from PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ at 37 °C is attributed to the formation of the gel phase, which provides microphase separated domains in which drug solubilisation may occur, and from which liberation becomes disfavoured, prolonging the release until 144 h. Additionally, polymer entanglements and an increased tortuosity within the samples may contribute to this effect. Progesterone release from poloxamer 407 was equivalent at 25 and 37 °C, where the material is in the gel phase at both temperatures. Both cases gave Higuchi diffusion-controlled release in the first 60% of the profile ($R^2 > 0.98$), which is consistent with prior studies of drug delivery across a membrane from poloxamer gels.⁸ This behaviour is substantially different than the PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ system which exhibited clear temperature-dependence on drug release, and a switch from Higuchi kinetics to a non-Fickian





Fig. 10 Release of $50 \mu\text{g mL}^{-1}$ progesterone and $200 \mu\text{g mL}^{-1}$ tenofovir disoproxil fumarate from 50% (w/v) PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (blue) and 20% (w/v) poloxamer 407 (red). A control formulation of the drugs dissolved in a 1:1 water: ethanol mixture (open circles) demonstrates that liberation from the polymer gel is rate-limiting. Please note that for the tenofovir release experiments, the next time point determined for the gels (24 h) contained 100% drug liberation.

release mechanisms. Higuchi kinetics following Fickian diffusion may be rationalised below the T_{gel} where release of drug is controlled by factors such as partitioning and the viscosity of the medium. Super case II kinetics have been reported for chemically cross-linked PNIPAM hydrogels,⁵⁰ where Fickian diffusion is believed to be combined with polymer relaxation and possible swelling effects. This swelling/relaxation behaviour may combine with nanostructural rearrangement to give rise to the lengthy lag exhibited in the progesterone release studies from PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ at 37 °C, though this was not explicitly studied. It has been demonstrated by DLS that progesterone drives the formation of aggregates even at low temperatures, and the slow depletion of progesterone from the gel may also lead to structural rearrangement. Pure PNIPAM hydrogels also exhibit the long lag time seen for PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ at 37 °C.⁵⁰ The release of progesterone from PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ was significantly slower than poloxamer 407 at 37 °C. For example, at 72 h, $75.5 \pm 1.6\%$ of the entrapped progesterone was delivered from the 20% (w/v) poloxamer 407 sample, where only $25.3 \pm 6.7\%$ of the drug had been liberated from PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈

($p < 0.0001$ by t -test). This gives the PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ an advantage over poloxamer 407 when the sustained release of hydrophobic drugs is required. The total $10 \mu\text{g}$ progesterone dose was delivered from the $200 \mu\text{L}$ PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ formulation over 6 days, requiring a 5 mL dose to achieve the approximately $250 \mu\text{g}$ delivered from the standard Crinone gel over the same period.⁵¹

The impact of progesterone and tenofovir disoproxil fumarate ($50 \mu\text{g mL}^{-1}$) on the rheology of the thermoreversible gelators was assessed by temperature ramp (Fig. S9, ESI†). In the presence of progesterone, the gelation temperature of poloxamer 407 (20%) remained at 25 °C. Whereas the gelation temperature of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (50%) was slightly reduced from 35.8 ± 0.4 to 34.2 °C. It is known drug-polymer interactions may decrease the LCST of PNIPAM, which may explain this slight reduction in T_{gel} .⁵² The inclusion of progesterone resulted in a decrease in gel strength for poloxamer 407 (20%) from 12.7 ± 0.5 to 10.2 kPa while G'_{max} of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ did not vary, having values of 11.2 ± 1.8 and 10.7 kPa in undoped and progesterone-doped solutions, respectively. Tenofovir disoproxil fumarate did not alter the





Fig. 11 Stability of poloxamer 407 (i) and PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (ii) at 4, 25 and 40 °C. Variation in the number-average molecular weight is shown on the left. GPC traces are shown on the right, with week 0 shown in black, and week 12 at 4, 25 and 40 °C shown in green, blue and red, respectively.

gelation temperatures of poloxamer 407 (20%) and PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (50%), having values of 24 and 35.4 °C, respectively. Tenofovir disoproxil fumarate reduced $G'_{\max}$ to 9.7 kPa in the poloxamer 407 samples, but the value measured for PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ (10.2 kPa) was within the error of the undoped samples. Overall, the PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ samples retained their ability to act as thermo-reversible gelators after the addition of both model drugs.

The stability of P407 and PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ was assessed in aqueous solution at 4, 25, or 40 °C, reflecting refrigerated storage, storage at room temperature, and an accelerated storage condition, respectively (Fig. 11). Accelerated storage at 40 °C aimed to predict longer-term storage at room temperature. GPC analysis demonstrated that both polymers exhibit small reductions in molecular weight over 12 weeks, with losses accelerated in the PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ sample at elevated temperatures. However, at 4 and 25 °C the reduction in number-average molecular weight of PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ was not statistically significant ($p < 0.05$). GPC traces (Fig. 11(i)) of poloxamer at weeks 0 and 12 are near-identical, with the accelerated storage condition (red) exhibiting a small increase in a shoulder at low molecular weight.

PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ traces exhibit a clear shift to a lower molecular weight under accelerated storage conditions (Fig. 11(ii)). Under all conditions the trace for PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ traces remained monomodal with no shoulder. Thus, hydrolysis was not observed at the ester linkages present between individual polymer blocks, but amide hydrolysis is possible. Poloxamer 407 does not possess hydrolytically unstable ester or amide linkages, and its molecular weight remained constant throughout the study. This is the first report on the long-term stability of a PNIPAM-PEG copolymer in water. Future studies on the stability of the materials should expand this experiment in-line with ICH standards.⁵³ Additionally, it is not known whether the degradation exhibited in the accelerated storage condition will be seen at 25 °C, where typically small extrapolations are made from this data only when the degradation routes have been established.⁵³

Conclusions

This study investigates the use of thermoreversible PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ gelators as smart materials for topical



administration for the first time, with a critical comparison to poloxamer 407. The low dependence of T_{gel} on concentration allows PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ to exhibit *in situ* gelation at concentrations up to 50% (w/v), where T_{gel} may be finely tuned to temperatures just below 37 °C. Thickening occurred only at temperatures above 28 °C, ensuring that increases in viscosity do not occur at room temperature. This gives the system advantages over poloxamer 407 where values of T_{gel} are highly concentration dependant. Additionally, of the poloxamer 407 concentrations examined, T_{gel} typically occurred near or below room temperature (25 °C), making the materials unattractive in warmer climates (e.g. WHO climatic zones II (25 °C), III (30 °C) and IV (30 °C)).²⁵ PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ gels exhibit enhanced mucoadhesion and reduced dissolution compared to poloxamer 407, giving PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ significant advantages for mucosal drug delivery, where these two attributes are linked to retention at the site of administration. PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ was also found to be stable in solution at room temperature over 12 weeks.

The solubilisation and release of two drugs relevant to vaginal drug delivery was investigated. Progesterone was selected as a hydrophobic drug, and tenofovir disoproxil fumarate was used as a relatively hydrophilic species. PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ was able to greatly enhance the solubility of progesterone due to inclusion of the drug into polymer micelles, with a greater solubilising power than poloxamer 407, whereas tenofovir disoproxil fumarate was excluded from the micelles in both cases. The release of these two drugs was then evaluated using a Franz diffusion cell system at 25 and 37 °C. Release of progesterone from the poloxamer gels was identical at both temperatures, whereas tenofovir disoproxil was liberated more rapidly at the elevated temperature. PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈, however, gave an unusual temperature-responsive retardation of release of both drugs at elevated temperatures, controlling progesterone release over 6 days. This effect may be a means of controlling delivery of other xenobiotics or retaining drug on body surfaces for local effect. Overall, PNIPAM₉₈-PEG₁₂₂-PNIPAM₉₈ exhibits unique properties as a thermoreversible gelator for drug delivery with advantages over poloxamer 407 and could act as an advanced material for future topical therapies.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The Centre for Research in Topical Drug Delivery and Toxicology at the University of Hertfordshire funded the studentship of Peter Haddow. The Royal Society are thanked for partial funding of the research (RG160139).

References

- 1 M. A. Ward and T. K. Georgiou, *Polymers*, 2011, **3**, 1215–1242.
- 2 K. Al Khateb, E. K. Ozhmukhametova, M. N. Mussin, S. K. Seilkhanov, T. K. Rakhypbekov, W. M. Lau and V. V. Khutoryanskiy, *Int. J. Pharm.*, 2016, **502**, 70–79.
- 3 L. Yu and J. Ding, *Chem. Soc. Rev.*, 2008, **37**, 1473–1481.
- 4 E. Giuliano, D. Paolino, M. Fresta and D. Cosco, *Pharmaceutics*, 2018, **10**, E159.
- 5 M. T. Cook and M. B. Brown, *J. Controlled Release*, 2018, **270**, 145–157.
- 6 M. A. Abou-Shamat, J. Calvo-Castro, J. L. Stair and M. T. Cook, *Macromol. Chem. Phys.*, 2019, 1900173.
- 7 C. Wu, T. Liu, B. Chu, D. K. Schneider and V. Graziano, *Macromolecules*, 1997, **30**, 4574–4583.
- 8 G. Dumortier, J. L. Grossiord, F. Agnely and J. C. Chaumeil, *Pharm. Res.*, 2006, **23**, 2709–2728.
- 9 A. Fakhari, M. Corcoran and A. Schwarz, *Heliyon*, 2017, **3**, e00390.
- 10 M. Teodorescu, I. Negru, P. O. Stanescu, C. Drăghici, A. Lungu and A. Sârbu, *React. Funct. Polym.*, 2010, **70**, 790–797.
- 11 M. Heskins and J. E. Guillet, *J. Macromol. Sci., Part A: Pure Appl. Chem.*, 1968, **2**, 1441–1455.
- 12 S. K. Filippov, A. Bogomolova, L. Kabarov, N. Velychkivska, L. Starovoytova, Z. Cernochova, S. E. Rogers, W. M. Lau, V. V. Khutoryanskiy and M. T. Cook, *Langmuir*, 2016, **32**, 5314–5323.
- 13 A. J. De Graaf, K. W. M. Boere, J. Kemmink, R. G. Fokkink, C. F. Van Nostrum, D. T. S. Rijkers, J. Van Der Gucht, H. Wienk, M. Baldus, E. Mastrobattista, T. Vermonden and W. E. Hennink, *Langmuir*, 2011, **27**, 9843–9848.
- 14 H. H. Lin and Y. L. Cheng, *Macromolecules*, 2001, **34**, 3710–3715.
- 15 A. J. De Graaf, I. I. Azevedo Próspero Dos Santos, E. H. E. Pieters, D. T. S. Rijkers, C. F. Van Nostrum, T. Vermonden, R. J. Kok, W. E. Hennink and E. Mastrobattista, *J. Controlled Release*, 2012, **162**, 582–590.
- 16 D. Limón, K. Talló Domínguez, M. L. Garduño-Ramírez, B. Andrade, A. C. Calpena and L. Pérez-García, *Colloids Surf., B*, 2019, **181**, 657–670.
- 17 D. H. Owen and D. F. Katz, *Contraception*, 1999, **59**, 91–95.
- 18 M. Garcia, M. P. Beecham, K. Kempe, D. M. Haddleton, A. Khan and A. Marsh, *Eur. Polym. J.*, 2015, **66**, 444–451.
- 19 J. Bassi da Silva, S. B. de S. Ferreira, A. V. Reis, M. T. Cook and M. L. Bruschi, *Polymers*, 2018, **10**, 1–19.
- 20 M. Malmsten and B. Lindman, *Macromolecules*, 1992, **25**, 5440–5445.
- 21 B. Jeong, S. W. Kim and Y. H. Bae, *Adv. Drug Delivery Rev.*, 2012, **64**, 154–162.
- 22 S. E. Kirkland, R. M. Hensarling, S. D. McConaughy, Y. Guo, W. L. Jarrett and C. L. McCormick, *Biomacromolecules*, 2008, **9**, 481–486.
- 23 A. N. Semenov, J. F. Joanny and A. R. Khokhlov, *Macromolecules*, 1995, **28**, 1066–1075.
- 24 R. K. Prud'homme, G. Wu and D. K. Schneider, *Langmuir*, 1996, **12**, 4651–4659.



- 25 W. Grimm, *Drug Dev. Ind. Pharm.*, 1993, **19**, 2795–2830.
- 26 K. Edsman, J. Carlfors and R. Petersson, *Eur. J. Pharm. Sci.*, 1998, **6**, 105–112.
- 27 M. Castro, D. W. Giles, C. W. Macosko and T. Moaddel, *J. Rheol.*, 2010, **54**, 81–94.
- 28 Y. Zhang, S. Furyk, L. B. Sagle, Y. Cho, D. E. Bergbreiter and P. S. Cremer, *J. Phys. Chem. C*, 2007, **111**, 8916–8924.
- 29 D. R. Perinelli, M. Cespi, S. Pucciarelli, L. Casettari, G. F. Palmieri and G. Bonacucina, *Colloids Surf., A*, 2013, **436**, 123–129.
- 30 G. Niu, F. Du, L. Song, H. Zhang, J. Yang, H. Cao, Y. Zheng, Z. Yang, G. Wang, H. Yang and S. Zhu, *J. Controlled Release*, 2009, **138**, 49–56.
- 31 N. A. Peppas and Y. Huang, *Adv. Drug Delivery Rev.*, 2004, **56**, 1675–1687.
- 32 M. T. Cook and V. V. Khutoryanskiy, *Int. J. Pharm.*, 2015, **495**, 991–998.
- 33 J. D. Smart, *Adv. Drug Delivery Rev.*, 2005, **57**, 1556–1568.
- 34 V. V. Khutoryanskiy, *Macromol. Biosci.*, 2011, **11**, 748–764.
- 35 SPC Crinone 8% Progesterone Vaginal Gel, <https://www.medicines.org.uk/emc/product/1283/smpe>, accessed 31 October 2019.
- 36 SPC Cyclogest 200 mg, <https://www.medicines.org.uk/emc/product/5568/smpe>, accessed 31 October 2019.
- 37 SPC Lutigest 100 mg vaginal tablets, <https://www.medicines.org.uk/emc/product/3635/smpe>, accessed 31 October 2019.
- 38 SPC Utrogestan Vaginal 200 mg Capsules, <https://www.medicines.org.uk/emc/product/3244/smpe>, accessed 31 October 2019.
- 39 J. M. Smith, R. Rastogi, R. S. Teller, P. Srinivasan, P. M. M. Mesquita, U. Nagaraja, J. M. McNicholl, R. M. Hendry, C. T. Dinh, A. Martin, B. C. Herold and P. F. Kiser, *Proc. Natl. Acad. Sci. U. S. A.*, 2013, **110**, 16145–16150.
- 40 P. Alexandridis and T. Alan Hatton, *Colloids Surf., A*, 1995, **96**, 1–46.
- 41 A. M. Bodratti and P. Alexandridis, *J. Funct. Biomater.*, 2018, **9**, 11.
- 42 A. R. Tehrani-Bagha and K. Holmberg, *Materials*, 2013, **6**, 580–608.
- 43 A. M. Bodratti and P. Alexandridis, *J. Funct. Biomater.*, 2018, **9**, E11.
- 44 F. P. Schwarb, G. Imanidis, E. W. Smith, J. M. Haigh and C. Surber, *Pharm. Res.*, 1999, **16**, 909–915.
- 45 P. Singla, S. Chabba and R. K. Mahajan, *Colloids Surf., A*, 2016, **504**, 479–488.
- 46 D. Selzer, M. M. A. Abdel-Mottaleb, T. Hahn, U. F. Schaefer and D. Neumann, *Adv. Drug Delivery Rev.*, 2013, **65**, 278–294.
- 47 S. Dash, P. N. Murthy, L. Nath and P. Chowdhury, *Acta Pol. Pharm. – Drug Res.*, 2010, **67**, 217–223.
- 48 F. Akomeah, T. Nazir, G. P. Martin and M. B. Brown, *Eur. J. Pharm. Sci.*, 2004, **21**, 337–345.
- 49 M. L. Bruschi, *Strategies to Modify the Drug Release from Pharmaceutical Systems*, 2015.
- 50 Y. Danyuo, C. J. Ani, A. A. Salifu, J. D. Obayemi, S. Dozie-Nwachukwu, V. O. Obanawu, U. M. Akpan, O. S. Odusanya, M. Abade-Abugre, F. McBagonluri and W. O. Soboyejo, *Sci. Rep.*, 2019, **9**, 1–14.
- 51 M. Campana-Seoane, A. Peleteiro, R. Laguna and F. J. Otero-Espinar, *Int. J. Pharm.*, 2014, **477**, 495–505.
- 52 D. C. Coughlan and O. I. Corrigan, *Int. J. Pharm.*, 2006, **313**, 163–174.
- 53 ICH, Q1A(R2) Stability Testing of New Drug Substances and Products, 2003.

