This article can be cited before page numbers have been issued, to do this please use: H. Yu, J. V. Rowley, D. Green and P. Thornton, Mater. Adv., 2020, DOI: 10.1039/D0MA00280A.

This is an Accepted Manuscript, which has been through the Royal Society of Chemistry peer review process and has been accepted for publication.

Accepted Manuscripts are published online shortly after acceptance, before technical editing, formatting and proof reading. Using this free service, authors can make their results available to the community, in citable form, before we publish the edited article. We will replace this Accepted Manuscript with the edited and formatted Advance Article as soon as it is available.

You can find more information about Accepted Manuscripts in the Information for Authors.

Please note that technical editing may introduce minor changes to the text and/or graphics, which may alter content. The journal’s standard Terms & Conditions and the Ethical guidelines still apply. In no event shall the Royal Society of Chemistry be held responsible for any errors or omissions in this Accepted Manuscript or any consequences arising from the use of any information it contains.
Fucose-Modified Thermoresponsive Poly(2-Hydroxypropyl Methacrylate) Nanoparticles for Controlled Doxorubicin Release from an Injectable Depot

Huayang Yu, Jason V. Rowley, David C. Green and Paul D. Thornton

The modification of poly(2-hydroxypropyl methacrylate) with a single fucose group per polymer chain enabled macromolecular self-assembly, and the formation of thermoresponsive nanoparticles. In addition, poly(2-hydroxypropyl methacrylate) formed a self-healing material that may act as an injectable vehicle and depot for poly(2-hydroxypropyl methacrylate) nanoparticle delivery and localisation at a particular site. In combination, the injectable depot permits the controlled release of doxorubicin from the fucose presenting nanoparticles that it contains following injection. Such thermoresponsive materials are highly promising candidates for the treatment of diseases that may be remedied by exploiting targeted fucose-cell binding, such as pancreatic cancer.

Poly(2-hydroxypropyl methacrylate) (PHPMA) is non-toxic and biocompatible. PHPMA-containing block copolymers can be readily formed by reversible addition-fragmentation chain-transfer (RAFT) aqueous dispersion polymerisation. Such polymers have been reported to self-assemble into a plethora of arrangements dependent on their macromolecular configuration, and the environment in which they are maintained. For instance, poly(glycidyl methacrylate)-b-PHPMA may be manipulated to form spheres, worms or vesicles dependent on the composition of the block copolymer, and the temperature of the aqueous solution in which the polymer is dispersed. PHPMA, and PHPMA-based materials have been applied to the thermally-triggered release of drug molecules, and controlled release applications.

Implantable polymer-based materials for drug and cell delivery have attracted significant recent attention. Such materials may facilitate the immobilisation of drug-loaded nanoparticles at the target site, ensuring extensive drug release at diseased areas, and minimal drug release to healthy tissue. This enhances the efficacy of the treatment, reducing the side-effects that are caused by drug interaction with non-target, healthy cells. An implantable depot that enables the controlled and prolonged release of fucose-presenting nanoparticles for long-term pancreatic cancer treatment is a noteworthy long-term goal.

We report that the simple modification of PHPMA polymer chains with a single fucose unit enables the polymer to self-assemble, forming nanoparticles in aqueous solution. Such nanoparticles are stable at room temperature but undergo a morphological change upon heating to physiological temperature, which may be exploited to actuate therapeutic payload release. Unmodified PHPMA can form an injectable depot that undergoes a sol-gel transition at a temperature slightly above physiological temperature. Consequently, the depot, which is self-healing and can withhold guest molecules itself, may act as an injectable host for the controlled release of doxorubicin (Dox) from fucose-capped PHPMA nanoparticles at a localised site.

Introduction

Effective cancer treatment in the absence of extremely intrusive side-effects remains a key challenge in contemporary medicine. Providing a remedy that destroys cancer cells, whilst leaving healthy cells relatively unaffected must be realised for this challenge to be overcome. Encapsulating a toxic anti-cancer drug within a shielding polymeric carrier protects the drug from interacting with cells, thus preventing the damage of non-cancerous cells, and the aggressive side-effects that are associated with this occurring during chemotherapy. Extensive drug loading into polymeric nanoparticles, and a mechanism for the programmed release of the drug from the nanoparticles, are essential features of effective treatments. Additionally, next-generation drug delivery vehicles must present cell-binding groups that enable extensive therapeutic interaction with target cells only.

Fucose plays a key role in mammalian development, immunity and cancer metastasis. Increased expression of fucosyltransferases is associated with pancreatic cancer, whereby enzymes accelerate malignant transformation through the fucosylation of sialylated precursors, and pancreatic cancer cells display enhanced fucose uptake. Pancreatic cancer is an extremely aggressive cancer with a distinctive long survival rate; it is estimated that the 5-year survival rate is 5% owing to its aggressiveness, and a lack of effective therapies. Currently, patients that undergo treatment with the chemotherapy drug Gemcitabine (Gemzar®) have a median survival time of 5.7 months. A plausible explanation for this abysmal inefficiency is the extremely ineffective delivery of anticancer drugs to the tumour site. There is enormous demand for the creation of fucose-presenting nanoparticles to be deployed as increasingly advanced drug delivery systems for the treatment of pancreatic cancer.
Experimental

Materials and methods

L(-)-Fucose 97 % was purchased from ACROS Organics. Chloroform (HPLC grade), dichloromethane (DCM) (HPLC grade) and acetone were obtained from WVR Chemicals. HPMA, Phosphate buffered saline tablet, N,N-dimethylformamide, 2,2'-azobis(2-methyl-propionamidine) dihydrochloride (AAPH) granular 97 %, 4-cyano-4-((phenylcarbonothioylthio)pentanoic acid N-succinimidyl ester (SCPDB), dialysis tubing benzoylated 2000 Da, deuterium oxide 99.9 atom D% and acetic acid 99-100% were purchased from Sigma-Aldrich. Triethylamine and tetrahydrofuran (THF, analytical grade) were obtained from Thermo Scientific. 4-Fucose was dissolved in 10.0 mL of deionised water. Sodium meta-periodate was obtained from Thermo Scientific. 4-Fucose was dissolved in 10.0 mL of deionised water. 

SCPD Conjugation to Amine-Bearing fucose

0.118 g of fucose-amine and 0.213 g of SCPDB RAFT agent were dissolved in 5.0 mL of deionised water and 20.0 mL of acetone, respectively. The SCPDB RAFT agent solution was added dropwise into the fucose-amine solution with stirring followed by the addition of 3.0 mL of acetic acid. The mixture was stirred at 60 °C overnight. Following this, the solution underwent rotary evaporation at 50 °C to remove acetone and freeze-dried for 48 hours. The fucose-RAFT agent was washed multiple times with DCM, being recovered by centrifugation for 30 minutes at 4500 rpm, before finally being dried in a vacuum oven at room temperature overnight.

HPMA Polymerisation from the fucose-RAFT agent

0.052 g of fucose-RAFT agent was dissolved in 3.0 mL and 2.0 mL of acetone and deionised water. Then, 0.091 g of AAPH was dissolved in the mixture (mole ratio of fucose-RAFT agent: AAPH = 1:3). 1.608 g of HPMA was added dropwise into the reaction which was stirred at 55 °C overnight (mole ratio of fucose-RAFT agent: HPMA = 1:100). The product underwent rotary evaporation at 50 °C for 20 minutes and the recovered product was freeze-dried overnight. Finally, fucose-HPMA was washed multiple times with deionised water, being recovered by centrifugation for 30 minutes at 4500 rpm, before finally being freeze-dried for 48 hours.

HPMA synthesis: HPMA polymerisation from 4-Cyano-4-((phenylcarbonothioylthio)pentanoic acid

0.020 g of 4-Cyano-4-((phenylcarbonothioylthio)pentanoic acid RAFT agent was dissolved in 3.0 mL and 2.0 mL of acetone and deionised water, respectively. Then, 0.059 g of AAPH was dissolved into the mixture (mole ratio of RAFT agent: AAPH = 1:3). 1.025 g of HPMA was added dropwise into the reaction and stirred at 55 °C overnight (mole ratio of fucose-RAFT agent: HPMA = 1:100). The product was rotary evaporated at 50 °C for 20 minutes and freeze dried for 48 hours. Finally, PHPMA was washed with deionised water, centrifugation for 30 minutes at 4500 rpm and freeze dried for 48 hours.

PHPMA gel synthesis: HPMA Polymerisation from 4-Cyano-4-((phenylcarbonothioylthio)pentanoic acid

As a representative procedure to produce PHPMA, 0.020 g of 4-Cyano-4-((phenylcarbonothioylthio)pentanoic acid RAFT agent was dissolved in 3.0 mL and 2.0 mL of acetone and deionised water, respectively. Then, 0.059 g of AAPH was dissolved into the mixture (mole ratio of RAFT agent: AAPH = 1:3). 2.050 g of HPMA was added dropwise into the reaction, which was stirred at 55 °C overnight (mole ratio of fucose-RAFT agent: HPMA = 1:200).
Results and discussion

The overall route to fucose-capped PHPMA nanoparticles that are able to withhold, and gradually release Dox is presented in Fig. 1, and detailed reaction schemes provided in Schemes S1-S2. Fucose was successfully oxidised by sodium meta-periodate, as confirmed by the proton corresponding to the aldehyde group being present in the $^1$H NMR spectrum (Fig. S1). Fucose-amine was produced by sequential reaction with ethylene diamine and sodium cyanoborohydride, and the structure confirmed by $^{13}$C NMR (Fig. S3). The peaks present in the region 40-55 ppm are assigned to the conjugated diamine. RAFT agent conjugation to fucose-amine, to enable RAFT polymerisation to occur from the fucose-amine initiator, was confirmed by the peaks in the aromatic region of Fig. S5, which remained after thorough washing with DCM. Different chain lengths (20, 40, 60, 80 and 100 repeat units) of fucose-PHPMA were synthesised to ascertain the optimal polymer for stable nanoparticle formation. The average molecular weight values of fucose-PHPMA with 20, 40, 60, 80 and 100 repeat units were assessed by $^1$H NMR spectroscopy using the proton environments of the RAFT agent aromatic group and peaks f, g, h, e, d and i to confirm the desired polymers were produced (Figs. S6-S10). APC was used to further evaluate polymer molecular weight (Table S1).

The products from each step of the overall reaction (Scheme S1) were analysed by FTIR spectroscopy. The aldehyde group of opened fucose (Fig. S17 black) confirms fucose oxidation by sodium meta-periodate. Peaks corresponding to C=S, C-S and aromatic groups can be found in the FTIR spectra of the fucose-RAFT agent, and fucose-PHPMA, confirming successful synthesis of activated fucose and the desired final product, respectively (Fig. S17 and Fig. S18). Additionally, a strong and broad peak corresponding to the alcohol group is present in the fucose-PHPMA FTIR spectrum further confirming PHPMA polymerisation from the fucose-RAFT agent (Fig. S17). This route to product, in which the RAFT agent is covalently conjugated to fucose, ensures that each PHPMA chain possesses a desired terminal fucose unit.

Nanoparticles were produced from the various fucose-PHPMA batches by coacervation. Nanoparticle size increased with polymer chain length when Dox was absent from the nanoparticles, and also when Dox was loaded into the nanoparticles (Tables 1 and 2). At room temperature, the particle sizes of blank and Dox-loaded nanoparticles were comparable both in terms of dimensions and PDI values. In all instances, the PDI values were less than 0.3, confirming a stable dispersion formed. Conversely, PHPMA of the same number of repeat units, but lacking fucose, did not form stable nanoparticles.

![Fig. 1 Reaction outline for the creation of fucose-PHPMA nanoparticles that are capable of encapsulating, and releasing Dox at physiological temperature.](image-url)

Table 1. Nanoparticle size and PDI values for nanoparticles formed from fucose-PHPMA (DMF:H$_2$O, 1:1 volume ratio), maintained at room temperature. The nanoparticles were measured both when vacant and loaded with Dox.

| Polymer          | Size (nm) | PDI  |
|------------------|-----------|------|
| Fucose-PHPMA$_{20}$ | 93 ±5     | 0.170|
| Fucose-PHPMA$_{40}$ | 106 ±6    | 0.205|
| Fucose-PHPMA$_{60}$ | 121 ±1    | 0.169|
| Fucose-PHPMA$_{80}$ | 163 ±6    | 0.165|
| Fucose-PHPMA$_{100}$ | 217 ±3   | 0.151|
| Fucose-PHPMA$_{20}$(Dox)$^\dagger$ | 91 ±2    | 0.209|
| Fucose-PHPMA$_{40}$(Dox)$^\dagger$ | 108 ±4   | 0.218|
| Fucose-PHPMA$_{60}$(Dox)$^\dagger$ | 122 ±1   | 0.186|
| Fucose-PHPMA$_{80}$(Dox)$^\dagger$ | 165 ±2   | 0.217|
| Fucose-PHPMA$_{100}$(Dox)$^\dagger$ | 219 ±2   | 0.181|

$^\dagger$ Denotes Dox-loaded nanoparticles.
(Table S2), emphasising the necessity for fucose conjugation to PHPMA for stable nanoparticle assembly. Fucose-PHPMA$_{100}$ with and without Dox-loading were shown as spherical nanoparticles by SEM analysis (Fig. 2 (a) and (b), respectively).

To assess the thermoresponse of fucose-PHPMA nanoparticles, their dimensions were assessed by DLS at 37 °C. The particle size of all types of fucose-PHPMA nanoparticles decreased significantly at 37 °C, signifying a thermo-responsive behaviour (Table 2). PDI values of the particles at 37 °C were slightly higher than those at room temperature, however all samples had PDI values less than 0.3, confirming their stability against aggregation. Significantly smaller artefacts were present in particles with PDI values less than 0.3, confirming their stability against aggregation.

The nanoparticles were measured both when vacant and loaded with Dox. Since fucose-PHPMA$_{100}$ nanoparticles at room temperature PBS buffer solution and (b) at 37 °C in PBS buffer solution.

Table 2. Nanoparticle size and PDI values obtained for nanoparticles formed from fucose-PHPMA, maintained in aqueous solution at 37 °C. The nanoparticles were measured both when vacant and loaded with Dox.

| Polymer               | Size (nm) | PDI  |
|-----------------------|-----------|------|
| Fucose-PHPMA$_{10}$   | 12 ±0.2   | 0.203|
| Fucose-PHPMA$_{40}$   | 16 ±2.5   | 0.215|
| Fucose-PHPMA$_{60}$   | 19 ±1.2   | 0.217|
| Fucose-PHPMA$_{80}$   | 22 ±1.3   | 0.218|
| Fucose-PHPMA$_{100}$  | 32 ±3.3   | 0.232|
| Fucose-PHPMA$_{100}$ (Dox) | 11 ±1.6 | 0.238|
| Fucose-PHPMA$_{40}$ (Dox) | 16 ±0.8 | 0.251|
| Fucose-PHPMA$_{60}$ (Dox) | 19 ±1.3 | 0.256|
| Fucose-PHPMA$_{80}$ (Dox) | 22 ±1.5 | 0.233|
| Fucose-PHPMA$_{100}$ (Dox) | 32 ±1.0 | 0.201|

± Denotes Dox-loaded nanoparticles.
specific temperature that lead to complete Dox release, with studies conducted that measured Dox release at various temperatures. To begin with, the Dox release study ran over a temperature range from 22 °C to 50 °C and the solution temperature was increased by 1 °C every 15 minutes (Fig. 4 (a)). Extensive Dox release occurred between 39 °C (19%) and 46 °C (47%) (Fig. 4 (a)). The temperature range of greatest Dox release was then narrowed to between 35 °C and 46 °C, by increasing the solution temperature by 1 °C every 40 minutes. Extensive Dox release (19.2 %, 0.041 mg) occurred between 41 °C and 42 °C (Fig. 4 (b)), demonstrating the potential of the reported materials to release Dox more extensively under moderate heating. Further Dox release studies were carried out to determine the time required for complete Dox release to occur at 41 °C. 22 % (0.048 mg) of Dox was released in the first hour (Fig. 4 (c)), with continuous release culminating in 51 % (0.126 mg), 74 % (0.183 mg) and 90 % (0.221 mg) Dox release after 24 hours, 48 hours and 72 hours, respectively (Fig. 4 (d)). Dox release from the fucose-PHPMA nanoparticles reached 100 % (0.245 mg) after 96 hours.

Next, a thermoresponsive PHPMA depot was created that may act as an injectable delivery vehicle, and host, for Dox-loaded fucose-

Fig. 4 (a) Dox release from fucose-PHPMA nanoparticles into solution increasing in temperature from 22 °C to 50 °C at a rate of 1 °C every 15 minutes. (b) Dox release from fucose-PHPMA nanoparticles into solution increasing in temperature from 35 °C to 46 °C at a rate of 1 °C every 40 minutes. (c) Dox release from fucose-PHPMA nanoparticles, monitored every five minutes at 41 °C. (d) Dox release from nanoparticles maintained at 41 °C over 7 days.

Fig. 5 The percentage of Dox released from fucose-PHPMA nanoparticles loaded in PHPMA depot maintained in PBS buffer solution at room temperature, and at 37 °C
Fig. 6 0.002 mg mL⁻¹ of rhodamine b was used to partially colour PHPMA<sub>200</sub>. a) Stretching PHPMA<sub>200</sub>; b-e) Demonstration of PHPMA<sub>200</sub> self-healing at 25 °C for one hour; f-i) Demonstration of PHPMA<sub>200</sub> self-healing at 25 °C for one hour in PBS buffer solution; j-m) Demonstration of PHPMA<sub>200</sub> self-healing at 37 °C for one hour; n-q) Demonstration of PHPMA<sub>200</sub> self-healing at 37 °C for one hour in PBS buffer solution.
PHPMA$_{200}$ nanoparticles that enables localised drug release (Fig. S23). PHPMA$_{80}$ and PHPMA$_{200}$ depots were synthesised by RAFT polymerisation (Scheme S2). PHPMA depot formation occurred upon polymer/DMSO solution being added to PBS buffer solution. The glass transition temperature of the depots was greater with increased polymer chain length (Fig. S24), and the water content of the two depots were 87.1 % (PHPMA$_{80}$) and 82.5 % (PHPMA$_{200}$), by mass (Table S3). The loss modulus of PHPMA$_{200}$ depot formed in PBS buffer solution was constant across the 0.1-100 frequency range indicating that the material was stable and inelastic (Fig. S26).

PHPMA$_{80}$ and PHPMA$_{200}$ depots were then created in the presence of Dox fucose-PHPMA$_{100}$ nanoparticles (Fig. S27). Due to the enhanced thermal and mechanical stability of the PHPMA$_{200}$ depot, Dox release from nanoparticles embedded within this material was studied at room temperature and at 37 °C (Fig. 5). Dox release increased up to 168 hours in solution maintained at 37 °C, after 43.9 % (0.011 mg) Dox release was recorded after 288 hours. Only a negligible amount of Dox release (2.3 %, 0.0006 mg) occurred at room temperature after 288 hours. The polymer nanoparticles are unable to penetrate the PHPMA depot, ensuring that they remain at the injection site following their administration.

Further studies revealed that the PHPMA$_{200}$ depot possesses self-healing properties. Two batches of the PHPMA$_{200}$ material were produced; one containing aqueous rhodamine b solution incorporated and the other containing water only, for illustrative purposes. The PHPMA$_{200}$ material can be stretched to 24.0 cm unevenly, which corresponds to 1200% elongation (Fig. 6a). Rhodamine b-containing material was placed in contact with blank PHPMA$_{200}$ material, in the presence and absence of PBS buffer solution at 25 °C and at 37 °C (Fig. 6c, g, k and o). Over 1 hour, the two independent materials connected to reveal the self-healing properties of the PHPMA$_{200}$ depot, indicating its potential suitability to be maintained for extended periods, and self-heal if damaged, in vivo (Fig. 6, e, i, m and q). The PHPMA$_{200}$ depot could also withhold rhodamine b in the absence of nanoparticles and be manipulated to form distinct and stable forms in PBS buffer solution. Rhodamine b-containing material could be formed upon injection onto a glass sheet as a viscous liquid (Fig. 7a). Upon the addition of PBS buffer solution, a depot formed that maintained both the loaded rhodamine b, and a defined structure. These materials could be rearranged and floated on the buffer solution, with the fusion of letters D and E occurring as further evidence of a self-healing material (Fig. 7c). The material dimensions and rhodamine b payload were both retained in PBS buffer solution, before the shapes were retrieved from solution and placed on a different glass sheet (Fig. 7d). The letters could be removed intact from solution and placed on a new glass sheet.

**Conclusions**

Thermally-responsive fucose-modified PHPMA nanoparticles were synthesised by RAFT polymerisation. A single fucose group per polymer chain was sufficient to instigate polymer self-assembly to yield nanoparticles in aqueous solution. Spherical fucose-PHPMA nanoparticles that contained or lacked Dox were very stable at room temperature and at 37 °C, as determined by DLS and SEM analysis. Dox loading into the nanoparticles was achieved by coacervation; negligible Dox release from the nanoparticles occurred at room temperature, whilst extensive release occurred upon the solution temperature increasing to 41 °C. An injectable PHPMA$_{200}$ depot was synthesised by RAFT polymerisation as a delivery vehicle for fucose-PHPMA$_{100}$ nanoparticles. This material possessed self-healing properties in PBS buffer solution and offers great promise to enable nanoparticle localisation at a target site for the prolonged release of anticancer therapeutics.

**Conflicts of interest**

There are no conflicts to declare.

**Notes and references**

1. J. Jacob, J. T. Haponiuk, S. Thomas and S. Gopi, Mater. Today Chem., 2018, 9, 43-55.
2. X. R. Qiao, X. F. Wang, Y. Shang, Y. Li and S. Z. Chen, Cancer Commun., 2018, 38, 13.
3. F. Canfarotta, L. Lezina, A. Guerreiro, J. Czulak, A. Petukhov, A. Daks, K. Smolinska-Kempisty, A. Poma, S. Piletsky and N. A. Barlev, Nano Lett., 2018, 18, 4641-4646.
4. M. V. Novoselova, H. M. Loh, D. B. Trushina, A. Ketkar, T. O. Abakumova, T. S. Zatsepin, M. Kakran, A. M. Brzozowska, H. H. Lau, D. A. Gorin, M. N. Antipina and A. I. Brichkina, ACS Appl. Mater. Interfaces, 2020, 12, 5610-5623.
5. M. Miyazaki, E. Yuba, H. Hayashi, A. Harada and K. Kono, Bioconjugate Chem., 2018, 29, 44-55.
6. A. Bordat, T. Boissenot, J. Nicolas and N. Tsapis, Adv. Drug Deliv. Rev., 2019, 138, 167-192.
7. Y. Qu, B. Y. Chu, X. W. Wei, M. Y. Lei, D. R. Hu, R. Y. Zha, L. Zhong, M. Y. Wang, F. F. Wang and Z. Y. Qian, J. Control. Release, 2019, 296, 93-106.
8. D. J. Price, M. Khuphe, R. P. W. Davies, J. R. McLaughlan, N. Ingram and P. D. Thornton, *Chem. Commun.*, 2017, **53**, 8687-8690.
9. M. Khuphe, N. Ingram and P. D. Thornton, *Nanoscale*, 2018, **10**, 14201-14206.
10. B. S. McAvan, M. Khuphe and P. D. Thornton, *Eur. Polym. J.*, 2017, **87**, 468-477.
11. F. Araste, K. Abnous, M. Hashemi, S. M. Taghdisi, M. Ramezani and M. Alibolandi, *J. Control. Release*, 2018, **292**, 141-162.
12. W. Q. Cao, J. Zhou, A. Mann, Y. Wang and L. Zhu, *Biomacromolecules*, 2011, **12**, 2697-2707.
13. M. Khuphe, C. S. Mahon and P. D. Thornton, *Biomater. Sci.*, 2016, **4**, 1792-1801.
14. M. Schneider, E. Al-Shareef and R. S. Haltiwanger, *Glycobiology*, 2017, **27**, 601-618.
15. C. Lepage, R. Capacaccia, M. Hackl, V. Lemmens, E. Molina, D. Pierannunzio, M. Sant, A. Trama, J. Faire and E. W. Grp, *Eur. J. Cancer*, 2015, **51**, 2169-2178.
16. S. Boeck, M. Haas, S. Kruger and V. Heinemann, *In Vivo*, 2014, **28**, 1189-1192.
17. N. J. W. Penfold, J. R. Whatley and S. P. Armes, *Macromolecules*, 2019, **52**, 1653-1662.
18. J. Madsen, S. P. Armes, K. Bertal, S. MacNeil and A. L. Lewis, *Biomacromolecules*, 2009, **10**, 1875-1887.
19. S. Sugihara, S. P. Armes, A. Blanazs and A. L. Lewis, *Soft Matter*, 2011, **7**, 10787-10793.
20. N. J. Warren, M. J. Derry, O. O. Mykhaylyk, J. R. Lovett, L. P. D. Ratcliffe, V. Ladmiral, A. Blanazs, L. A. Fielding and S. P. Armes, *Macromolecules*, 2018, **51**, 8357-8371.
21. N. Orakdogen and B. Sanay, *J. Appl. Polym. Sci.*, 2018, **135**, 13.
22. H. Y. Yu, N. Ingram, J. V. Rowley, S. Parkinson, D. C. Green, N. J. Warren and P. D. Thornton, *J. Mat. Chem. B*, 2019, **7**, 7795-7795.
23. W. D. Wang, L. Xiang, L. Gong, W. H. Hu, W. J. Huang, Y. J. Chen, A. B. Asha, S. Srinivas, L. Y. Chen, R. Narain and H. B. Zeng, *Chem. Mat.*, 2019, **31**, 2366-2376.
24. M. H. Chen, L. L. Wang, J. J. Chung, Y. H. Kim, P. Atluri and J. A. Burdick, *ACS Biomater. Sci. Eng.*, 2017, **3**, 3146-3160.
25. A. R. Town, M. Giardiello, R. Gurjar, M. Siccardi, M. E. Briggs, R. Akhtar and T. O. McDonald, *Nanoscale*, 2017, **9**, 6302-6314.
26. B. Maiti, S. Samanta, S. Sarkar, S. Alam and T. Govindaraju, *ACS Applied Bio Materials*, 2020, **3**, 3544-3552.
27. L. P. Datta, S. Manchineella and T. Govindaraju, *Biomaterials*, 2020, **230**, 119633.
The modification of poly(2-hydroxypropyl methacrylate) with a single fucose group per polymer chain enabled macromolecular self-assembly, and the formation of thermoresponsive nanoparticles that could selectively release doxorubicin.