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A step-wise synthetic approach is necessary to access γ -conjugates of folate: folate-conjugated prodigiosenes†

Carlotta Figliola,‡ Estelle Marchal,‡ Brandon R. Groves and Alison Thompson *

Despite the vast literature that describes reacting folic acid with a pharmacophore, this route is ineffective in providing the correct regioisomer of the resulting conjugate. We herein present a step-wise route to the preparation of nine folate conjugates of the tripyrrolic prodigiosene skeleton. The strict requirement for step-wise construction of the folate core is demonstrated, so as to achieve conjugation at only the desired γ -carboxylic acid and thus maintain the α -carboxylic site for folate receptor (FR α) recognition. Linkages *via* ethylenediamine, polyethylene glycol and glutathione are demonstrated.

Introduction

Prodigiosin is a tripyrrolic, red pigmented natural product, produced by Gram positive and Gram negative bacteria, including certain strains of *Serratia marcescens* (Fig. 1).^{1,2} Prodigiosin is well known to exhibit immunosuppressive, antimicrobial and anticancer properties, yet modern uses extend from sunscreen³ to antibacterial dyes for silk⁴ and cotton.⁵ Known mechanisms of bioactivity include H⁺/Cl⁻ exchange, oxidative DNA cleavage through Cu(II) chelation and signal-transduction interference.⁶⁻¹⁵ Synthetic mimics of prodigiosin, named prodigiosenes,¹⁶ with modifications on the A-, B- or C-ring¹⁷⁻²⁴ have been shown to maintain the biological activity of the parent compound yet with 100-fold improvements in selectivity towards malignant cells and with 100-fold reduction of systemic toxicity in mice compared to the natural product (R = CO₂*i*Pr, R¹ = Me, Fig. 1).²⁵ Furthermore, the study of these synthetic analogues has enriched the knowledge of structure-activity relationships (SAR) for a better understanding of the biological activity of prodigiosin.

Folic acid (vitamin B9, Fig. 2) is considered²⁶ a promising biomarker for triple-negative breast cancer (TNBC).²⁷ More than 50% of TNBCs overexpress folate receptor alpha (FR α), as do a limited number of other cancers²⁸ (e.g. ovarian cancers)^{29,30} along with those tissues where TNBC is likely to result in metastasis, such as brain and lung.³¹ Importantly, FR α is present on only a few types of healthy cells (activated macrophages and proximal tubules of kidneys),³² and, crucially, the uptake of the vitamin folic acid (as a folate salt) is not mediated

by FR α .³³⁻³⁵ Consequently, cognisant that folate-conjugated drugs do not generally enter normal cells,³⁶ folic acid has been employed in the synthesis of multiple drug-delivery systems for chemotherapeutic treatment and imaging.

Given the potency of prodigiosenes and the targeting ability of folate, we herein report the synthesis of the first series of folate-prodigiosene conjugates. In doing so, we emphasize the ineffectiveness of the much-utilised direct synthetic approach to couple unmodified folic acid with the desired pharmacophore. Our observations and comments contrast with the vast literature that apparently describes the folate-targeted bioactivity of unpurified and uncharacterised material that results from such direct coupling. We categorically demonstrate that this direct approach, despite its wide use by others, gives rise to mixtures that include both regioisomers featuring the pharmacophore coupled to both carboxyl sites of folic acid. Instead, we demonstrate the requirement for step-wise construction of the folate core, so as to achieve conjugation at only the desired γ -carboxylic acid and thus maintenance of the α -carboxylic site for FR α recognition.³⁷

Results and discussion

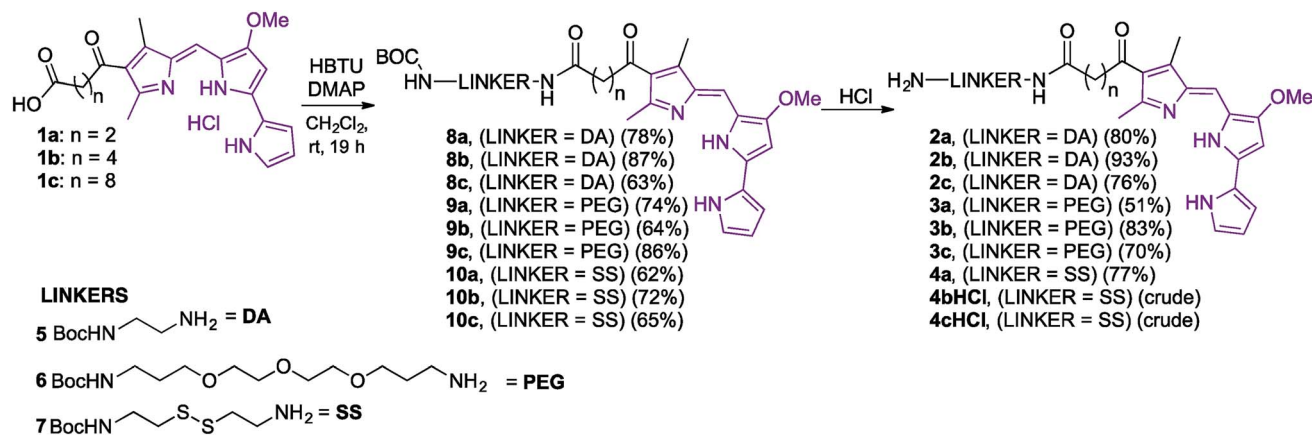
Our folate-prodigiosene conjugates (Fig. 3) feature prodigiosene **1** as the pharmacophore,³⁸ with varied aliphatic spacer length ($n = 2, 4, 8$) leading to the folate moiety. The conjugated carbonyl group on the β -position of the C-ring has been shown to facilitate purification of prodigiosene intermediates²¹ and so it, and the adjacent β -methyl substituent, were incorporated into the synthetic strategy. Linkers were chosen to increase either the lipophilicity³⁹ (ethylenediamine, DA) or the water-solubility and biocompatibility^{40,41} (polyethylene glycol, PEG) of the conjugates, or to enable intracellular redox-mediated cleavage by glutathione (disulfide, SS, Fig. 3).⁴²⁻⁴⁴

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‡ Equal contributions.





Scheme 1 Synthesis of linker-prodigiosene adducts 2–4.

coupling of folic acid with the pharmacophore. Following this approach, many literature reports appear to describe the direct activation of folic acid, followed by addition of the desired amine to afford conjugated folates.^{48–61} However, we met Strategy A with considerable scepticism as the presence of two carboxylic acid moieties within folic acid renders two regioisomeric coupling products clearly tangible. Given the nature of those two carboxylic acid groups, we were unable to find any rationale by which to convince ourselves that direct reaction would result in production of the required γ -conjugate. Furthermore, as conjugates to folic acid are often soluble only in DMSO the separation of the two regioisomers represents a considerable challenge. Adding to our skepticism, we found the literature regarding this direct synthetic approach to be incomplete and unreliable, due to either vagueness of purification techniques or lack of characterization such as to give assurance of purity. Indeed, upon close inspection it appears that there are many published reports that simply use the uncharacterised crude precipitate, obtained following the mixing of folic acid, coupling agent and pharmacophore, in studies relating to targeting using folate. Cognisant that coupling with DCC *etc.* involves ureas and other synthetic intermediates, and that these often persist in crude product mixtures, we remained unconvinced that Strategy A would be fruitful as we required a procedure that gave unequivocal access to the desired γ -conjugate of folic acid bearing prodigiosenes.

Retrosynthetic Strategies B and C (Fig. 4) involve the step-wise construction of suitably protected and activated folic acid from pteric acid and *L*-glutamic acid, allowing for the selective conjugation of pharmacophore at the desired γ -carboxyl site. Strategy B requires that the γ -activated, α -protected glutamic acid moiety be coupled with the pharmacophore prior to reaction with the pterate portion of what will become folate.^{62–64} Strategy C requires that the γ -activated, α -protected glutamic acid moiety first be coupled with activated pterate portion to create γ -activated, α -protected folate prior to reaction with the pharmacophore.⁶⁵ Solid-state resin approaches^{66,67} recognise the virtues of a step-wise approach yet, compared to the lure of Strategy A, both step-wise strategies obviously demand long

synthetic sequences and tedious purification processes and have not enjoyed widespread adoption. Indeed, the vast majority of published routes involving folate conjugates steer away from these step-wise approaches.⁶⁸

Given the evident popularity of the direct approach offered by Strategy A, we wanted to unequivocally assess the regioselective outcome of reacting folic acid with a coupling agent and an alcohol. We thus treated folic acid with DCC and NHS, and analysed the crude product mixture. Based on the phenyl group signals in the ¹H NMR spectrum (see ESI[†]), the crude NHS-adduct was determined to contain α - and γ -adducts in an approx. 25 : 75 α : γ ratio. A solution of the adduct mixture in DMSO was treated with the linker-prodigiosene conjugate **2a** in an attempt to form the desired amide linkage. Attempts to isolate pure material (**12a**) from the precipitate thus formed were completely unsuccessful (Scheme 2). Rather, the crude mixture was poorly soluble and decomposition occurred on chromatographic media (alumina and silica). Treatment of the NHS-adduct mixture with **2b** and **2c** also resulted in bulk precipitation, and again separation and purification of the highly insoluble material was not possible. Still intrigued by the apparent widespread success of Strategy A, we treated folic acid with DCC, NHS and a simple amine to thus employ another route by which to gain insight regarding the regioselectivity of activation.⁵⁷ A solution of the 25 : 75 α : γ folate–NHS adduct in DMSO was treated with octyl amine, as a model amine, and the resulting precipitate analyzed using ¹H NMR spectroscopy. Both α - and γ -regioisomers were present, along with the NHS-activated folic acid, and purification proved unsuccessful given the lack of solubility of the material. In short, direct activation following Strategy A was found to be wholly unsatisfactory in the quest towards isolated and pure γ -appended conjugates of folic acid.

With firm evidence that the direct reaction of folic acid with coupling agent and alcohol/amine does not regioselectively provide the desired γ -regioisomer, we turned our attention to strategies that used sequential protection/activation. We first turned to the applicability of retrosynthetic Strategy B, involving step-wise construction of the folate moiety so as to ensure



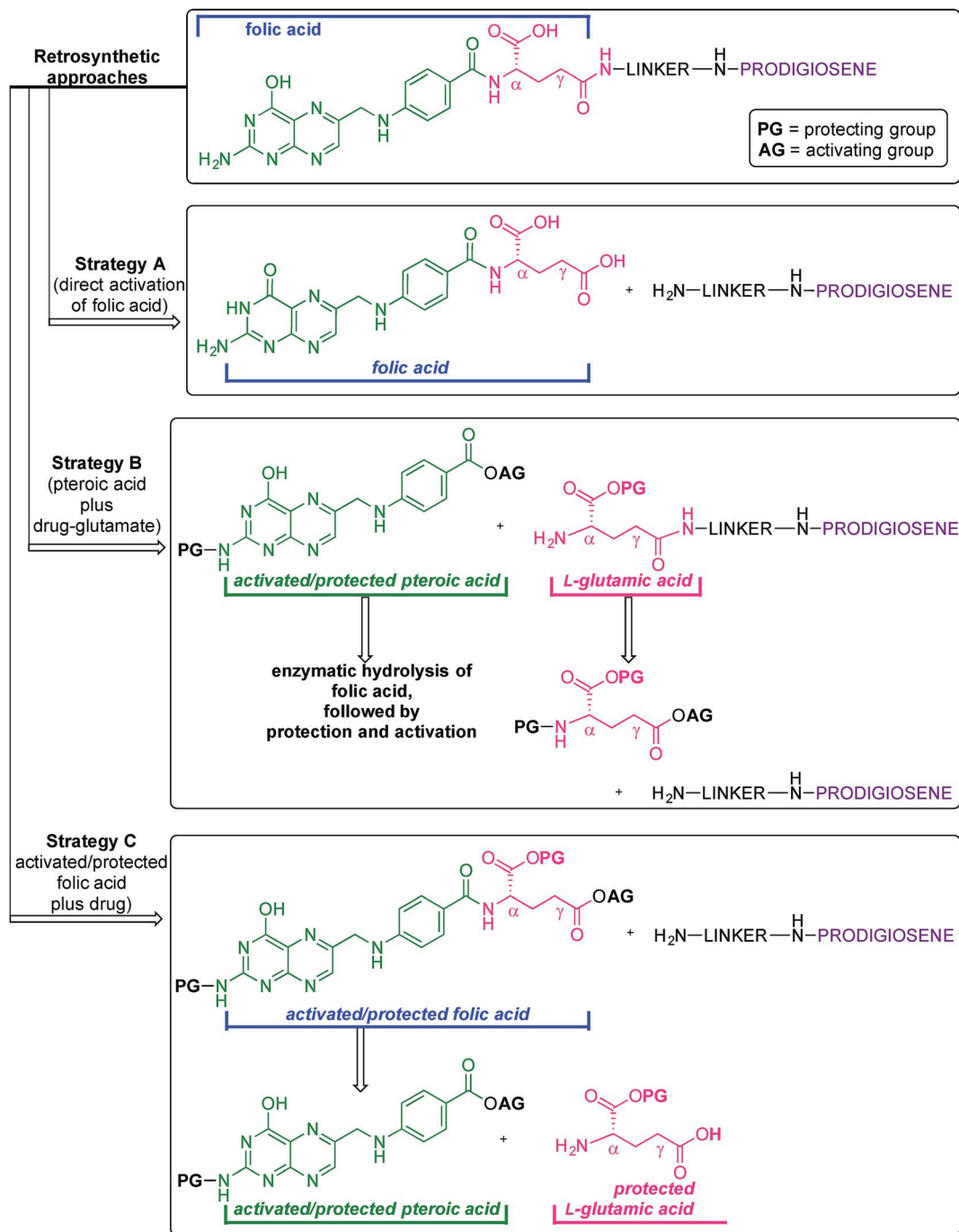
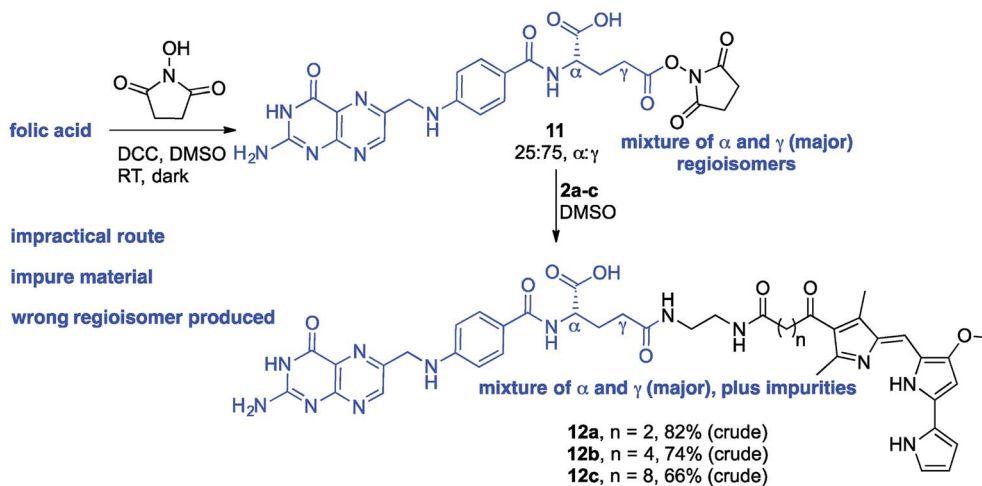


Fig. 4 Retrosynthetic approaches for the synthesis of folate-prodigiosene conjugates.

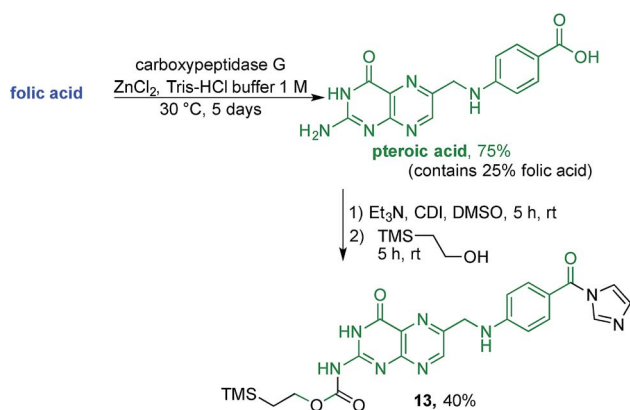
conjugation at the desired γ -carboxyl site. As shown in Fig. 4, the critical step entails coupling of suitably protected (the base portion) and activated (the acid portion) pteric acid with a derivative of L-glutamic acid that has been pre-linked to the prodigiosene pharmacophore by way of the γ -position. This key coupling would result in the folate-prodigiosene conjugate as a single isomer, courtesy of the previous regiospecific activation of L-glutamic acid.^{62–65,68,69} To embark upon Strategy B, pteric acid was prepared following literature procedure through hydrolysis of folic acid using carboxypeptidase G in the

presence of ZnCl_2 at 30 °C.⁶⁵ Carboxypeptidase G was purchased from Sigma and used as is (7 mg; 20 units, since 1 unit = 1 $\mu\text{mol min}^{-1}$ at optimal conditions of 30 °C and pH 7.3). The enzyme operates optimally at pH = 7.3. However, in our hands the reported 0.1 M Tris-HCl buffer was insufficient to maintain stable pH after the addition of folic acid. Therefore, the initial buffer concentration was increased to 1 M and the pH subsequently stabilised in the range of 7.2–7.3 *via* the addition of solid buffer. After 5 days, ^1H NMR spectroscopic analysis of the crude product mixture revealed 75% conversion of folic acid to





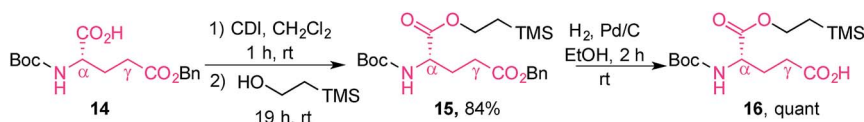
Scheme 2 Activation of folic acid with NHS and conjugation to linker-prodigiosenes 2a–c.



Scheme 3 Preparation of the activated Teoc-protected pteric acid 13.

ptericoic acid. The carboxylic acid of the crude ptericoic acid was activated using CDI and the primary amine then protected with a trimethylsilylethoxycarbonyl (Teoc) group, to give 13 after purification using column chromatography on silica (Scheme 3). This purification also allowed the removal of folic acid remaining from the initial step.

The L-glutamic acid moiety of folic acid was prepared from its commercially available Boc-L-glutamic acid 5-benzyl ester derivative (14), which was first esterified at the α -position using CDI and 2-(trimethylsilyloxy)ethanol, in order to obtain 15. Subsequent hydrogenolysis of the benzyl ester at the γ -position gave the protected amino acid 16, ready for coupling with prodigiosene (Scheme 4).⁶⁵

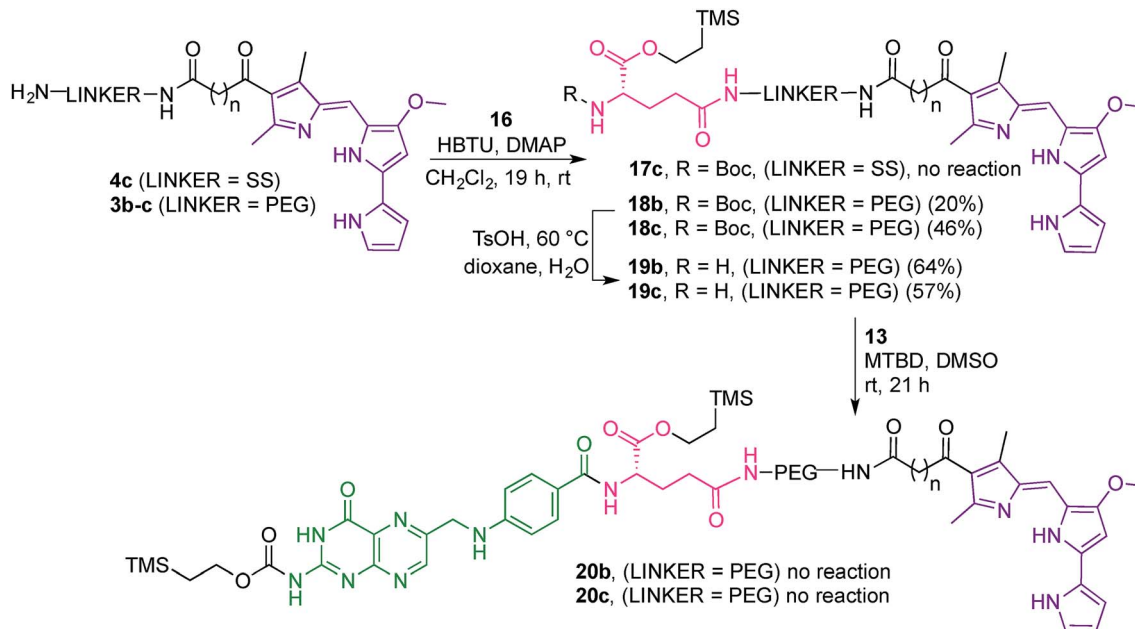


Scheme 4 Preparation of the protected glutamic acid derivative 16.

Attempts to couple 16 with the dithio linker-prodigiosene conjugate 4c·HCl in the presence of HBTU and DMAP were unsuccessful and only starting material was recovered (Scheme 5). However, coupling between 3b–c and the protected amino acid 16 successfully afforded substrates 18b–c, thus crucially connecting the prodigiosene pharmacophore with the γ -position of L-glutamic acid, by way of a PEG linker. Boc-deprotection in the presence of TsOH in a mixture of dioxane/water gave the amines 19b–c. Strategy B (Fig. 4) then requires coupling of the drug-appended glutamic acid moiety with the activated ptericoic acid, but attempts to couple 19b–c to 13 resulted in only partial recovery of the substrate prodigiosenes 19b–c (Scheme 5).

In turning our attention to Strategy C (Fig. 4), the L-glutamic acid derivative 16 was first deprotected at the amino position, and then coupled to the activated ptericoic acid to successfully afford 22. This γ -carboxylic acid was then reacted with NHS to give 23 which is activated in the desired position of what will become folate (Scheme 6).⁶⁵ The amines 2–4a–c were reacted with a solution of 23, to provide conjugates 24–26a–c. Subsequently, the Teoc protecting group was removed with TBAF, and the resulting tetrabutylammonium salt precipitated *via* the addition of water. Dissolution of the salt in DMSO, followed by reaction with NaOAc, afforded eight sodium folate–prodigiosene conjugates 27–28a–c and 29a–b. However, addition of TBAF to 26c caused the cleavage of the S–S bond and the corresponding folate–prodigiosene conjugate 29c could not be isolated (Scheme 6). Clearly the step-wise approach to building folate acid, such that only the γ -carboxyl site is available for coupling, requires more synthetic effort than the direct approach. However, in contrast to Strategy A, the step-wise





Scheme 5 Attempts to synthesise Teoc-protected folic acid-prodigiosene conjugates 20.

approaches of Strategies B and C tolerate substantiation of purity and characterization at each synthetic step and should thus be the approaches adopted for the preparation of regioisomerically pure desired conjugates of folate.

Conclusions

In summary, the first series of folate-prodigiosene conjugates have been synthesised, and full characterisation acquired. In order to increase the lipophilicity, the water-solubility and the ease of cleavage, three linkers were chosen to link the folate and the prodigiosene moieties (DA, PEG and SS respectively). Cognisant that the α -carboxylate of folate is involved in the interaction with $\text{FR}\alpha$, three synthetic approaches were investigated for the preparation of the targeted γ -adduct. The direct and most commonly used literature method involves coupling the unprotected folic acid, activated *in situ*, to the amino-counterpart. Despite the widespread use of this direct approach, often presented along with unsubstantiated claims as to purity and constitution of the product(s) thus obtained, we determined it to be wholly unsatisfactory as it gives mixtures of α - and γ -regioisomers, along with various ureas and other synthetic intermediates presenting as impurities, which are challenging to separate. We evaluated this shortcoming at the activation stage, using simply NHS, as well as through reaction with octyl amine and the prodigiosene pharmacophore. In order to access the required γ -adduct, the hydrolysis of folic acid into its two core components (pterioic acid and L-glutamic acid) was necessary. The subsequent step-wise synthesis of folic acid, with the α -carboxylic acid protected, allowed synthesis of only the desired regioisomer thus, for the first time, connecting folic acid with the prodigiosene pharmacophore. Next steps will

involve evaluation of the cytotoxicity of the conjugates, alongside optimization of drug release and internalisation.

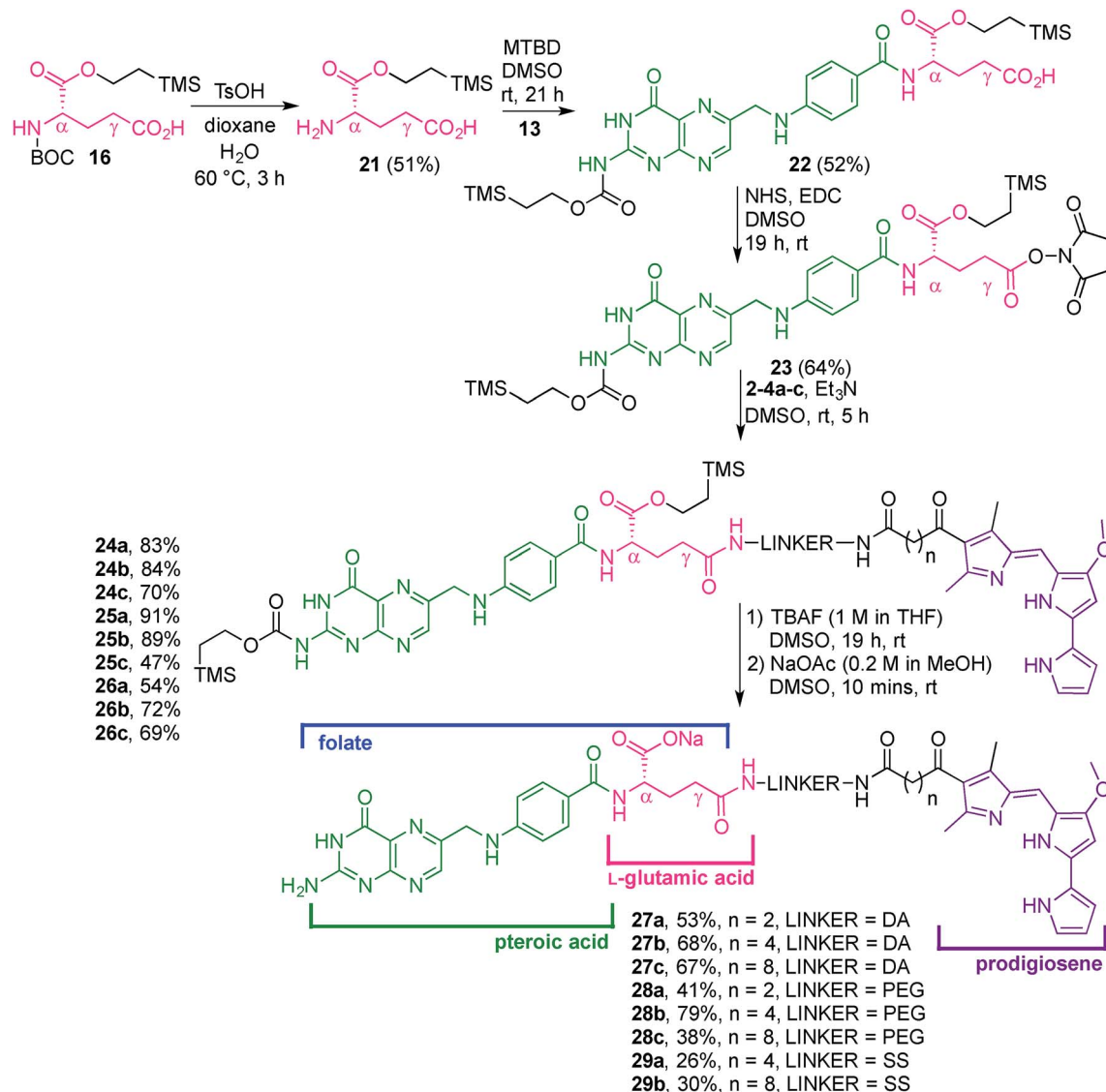
Experimental section

All chemicals were purchased and used as received unless otherwise indicated. Moisture sensitive reactions were performed in oven-dried glassware and under a positive pressure of nitrogen. Air- and moisture-sensitive compounds were introduced *via* syringe or cannula through a rubber septum. Flash chromatography was performed using either Silicycle ultra-pure silica (230–400 mm) or 150 mesh Brockmann III activated neutral/basic alumina oxide, as indicated. The NMR spectra were recorded using a 500 MHz spectrometer instrument using CDCl_3 , MeOD or DMSO as solvents and are reported in part per million (ppm). Internal solvents were referenced at 7.26 ppm for ^1H and at 77.16 ppm for ^{13}C when using CDCl_3 , at 3.31 ppm for ^1H and at 49.00 ppm for ^{13}C when using MeOD and at 2.50 ppm for ^1H and at 39.5 ppm for ^{13}C when using DMSO- d_6 . Coupling constants (J) are given in Hertz (Hz). Mass spectra were obtained using TOF and LCQ Duo ion trap instruments operating in $\text{ESI}^{+/-}$ mode, as indicated. Compounds **1a-c**,²¹ **5-7**,^{39,40,45} **11**,⁵⁷ pterioic acid,⁶⁵ **13**,⁶⁵ **15-16**,⁶⁵ and **21-23** (ref. 65) were prepared following literature procedures.

General procedure 1 (GP1)

Amines **5-7** (2 eq.), HBTU (1.2 eq.) and DMAP (1.2 eq.) were added to a solution of prodigiosene **1** (1 eq.) in anhydrous CH_2Cl_2 (0.01 M) and the reaction mixture was stirred at room temperature for 16 h under N_2 atmosphere. The solvent was removed under reduced pressure and the crude mixture was partitioned between EtOAc and NaHCO_3 (aq. sat solution). The aqueous layer was extracted with EtOAc ($\times 3$). The combined





Scheme 6 Synthesis of folate–prodigiosene conjugates 27–28a–c and 29a–b.

organic layers were washed with water ($\times 5$), brine, dried over Na_2SO_4 and concentrated under reduced pressure.

(Z)-tert-Butyl (2-(4-(2-((4-methoxy-1*H*,1'*H*-[2,2'-bipyrryl]-5-yl)methylene)-3,5-dimethyl-2*H*-pyrrol-4-yl)-4-oxobutanamido)ethyl)carbamate (8a). Compound **8a** was obtained according to GP1. The crude mixture was purified by column chromatography using A_2O_3 type III basic (CH_2Cl_2 : MeOH 100 : 0, 99 : 1), followed by A_2O_3 type III neutral (CH_2Cl_2 : MeOH, 100 : 0, 99 : 1) to afford **8a** (0.098 g, 78%) as dark red glass. ^1H NMR (CDCl_3 , 500 MHz) 1.41 (s, 9H), 2.21 (s, 3H), 2.38 (s, 3H), 2.47 (t, $J = 6.1$ Hz, 2H), 3.02 (t, $J = 6.3$ Hz, 2H), 3.24 (br s, 2H), 3.31 (br s, 2H), 3.96 (s, 3H), 5.14 (br s, 1H), 6.04 (s, 1H), 6.20–6.21 (m, 1H), 6.35 (br s, 1H), 6.72 (d, $J = 3.4$ Hz, 1H), 6.76 (s, 1H), 6.89 (s, 1H). ^{13}C NMR (CDCl_3 , 125 MHz) 12.6, 14.6, 28.5, 29.8, 30.7, 38.3, 40.4, 58.7, 79.5, 96.1, 110.8, 111.9, 113.9, 122.7, 123.3, 126.4, 128.2, 129.8, 141.0, 142.8, 156.7, 160.9, 169.0, 173.7, 196.2.

HRMS (TOF) (m/z): $[\text{M} + \text{H}]^+$ calcd for $\text{C}_{27}\text{H}_{36}\text{N}_5\text{O}_5$, 510.2711; found, 510.2719.

(Z)-tert-Butyl (2-(6-(2-((4-methoxy-1*H*,1'*H*-[2,2'-bipyrryl]-5-yl)methylene)-3,5-dimethyl-2*H*-pyrrol-4-yl)-6-oxohexanamido)ethyl)carbamate (8b). Compound **8b** was obtained according to GP1. The crude mixture was purified by column chromatography using A_2O_3 type III basic (CH_2Cl_2 : MeOH 100 : 0, 99 : 1), precipitated with CH_2Cl_2 : hexanes (1 : 9), filtered and washed with hexanes to afford **8b** (0.100 g, 87%) as orange glass. ^1H NMR (CD_3OD , 500 MHz) 1.42 (s, 9H), 1.68–1.70 (m, 4H), 2.22–2.26 (m, 2H), 2.39 (s, 3H), 2.66 (s, 3H), 2.81 (t, $J = 6.6$ Hz, 2H), 3.15 (t, $J = 6.0$ Hz, 2H), 3.25 (t, $J = 6.1$ Hz, 2H), 3.92 (s, 3H), 6.08 (s, 1H), 6.26–6.27 (m, 1H), 6.76 (s, 2H), 7.02 (s, 1H). ^{13}C NMR (CD_3OD , 125 MHz) 12.3, 15.4, 25.1, 26.7, 28.8, 30.2, 37.0, 40.5, 41.0, 43.1, 59.0, 96.7, 111.2, 111.3, 114.1, 123.6, 123.7, 128.3, 128.8, 129.8, 142.6, 144.3, 162.1, 169.6, 176.3, 200.0 (1 carbon



atom unaccounted for). HRMS (TOF) (m/z): $[M + H]^+$ calcd for $C_{29}H_{40}N_5O_5$, 538.3024; found, 538.3012.

(Z)-tert-Butyl (2-(10-((4-methoxy-1H,1'H-[2,2'-bipyrryl]-5-yl)methylene)-3,5-dimethyl-2H-pyrrol-4-yl)-10-oxodecanamido)ethyl)carbamate (8c). Compound **8c** was obtained according to GP1. The crude mixture was purified by column chromatography using Al_2O_3 type III neutral (EtOAc 100%) followed by Al_2O_3 type III neutral (CH_2Cl_2 : MeOH 99 : 1) and Al_2O_3 type III basic (CH_2Cl_2 : MeOH 99.5 : 0.5, 99 : 1, 98.5 : 0.5) to afford **8c** as a red glass (0.116 g, 63%). 1H NMR ($CDCl_3$, 300 MHz) 1.26 (s, 8H), 1.42 (s, 9H), 1.58–1.60 (m, 4H), 2.10–2.13 (m, 5H), 2.38 (s, 3H), 2.62 (t, $J = 7.2$ Hz, 2H), 3.24 (br s, 2H), 3.31 (br s, 2H), 3.97 (s, 3H), 5.05 (br s, 1H), 6.06 (s, 1H), 6.18 (s, 1H), 6.25 (br s, 1H), 6.71–6.72 (s, 2H), 6.93 (s, 1H). ^{13}C NMR ($CDCl_3$, 100 MHz) 12.4, 14.0, 24.3, 25.8, 28.5, 29.2, 29.3, 29.4, 36.8, 40.5, 40.7, 42.8, 58.7, 79.7, 96.1, 110.7, 112.1, 113.8, 123.4, 123.6, 126.2, 128.3, 129.7, 132.3, 140.5, 142.2, 157.1, 161.1, 169.1, 174.0, 198.3. HRMS (TOF) (m/z): $[M + Na]^+$ calcd for $C_{33}H_{47}N_5NaO_5$, 616.3436; found, 616.3490.

(Z)-tert-Butyl (18-(2-((4-methoxy-1H,1'H-[2,2'-bipyrryl]-5-yl)methylene)-3,5-dimethyl-2H-pyrrol-4-yl)-15,18-dioxo-4,7,10-trioxa-14-azaoctadecyl)carbamate (9a). Compound **9a** was obtained according to GP1. The crude mixture was purified by column chromatography using Al_2O_3 type III basic (EtOAc : hexanes 50 : 50, 70 : 30 EtOAc 100 : 0, CH_2Cl_2 : MeOH 100 : 0, 99 : 1) to afford **9a** as an orange glass (0.124 g, 74%). 1H NMR ($CDCl_3$, 500 MHz) 1.42 (s, 9H), 1.73–1.79 (m, 4H), 2.25 (br s, 3H), 2.38 (s, 3H), 2.49 (t, $J = 6.5$ Hz, 2H), 3.02 (t, $J = 6.5$ Hz, 2H), 3.19–3.20 (m, 2H), 3.34 (q, $J = 6.0$ Hz, 2H), 3.51–3.59 (m, 8H), 3.62–3.64 (m, 4H), 3.96 (s, 3H), 5.02 (br s, 1H), 6.02 (s, 1H), 6.22 (br s, 1H), 6.42 (br s, 1H), 6.72 (br s, 1H), 6.78 (br s, 1H), 6.88 (s, 1H). ^{13}C NMR ($CDCl_3$, 125 MHz) 12.5, 14.7, 28.6, 29.1, 29.8, 30.5, 37.8, 38.2, 38.6, 58.6, 69.6, 69.9, 70.3 ($\times 2$), 70.6, 79.0, 95.9, 110.8, 111.9, 113.8, 122.8, 123.4, 126.4, 128.2, 129.9, 140.7, 142.6, 156.2, 160.6, 168.8, 172.8, 196.0 (1 carbon atom unaccounted for). HRMS (TOF) (m/z): $[M + Na]^+$ calcd for $C_{35}H_{51}N_5Na_1O_8$, 692.3630; found, 692.3627.

(Z)-tert-Butyl (20-(2-((4-methoxy-1H,1'H-[2,2'-bipyrryl]-5-yl)methylene)-3,5-dimethyl-2H-pyrrol-4-yl)-15,20-dioxo-4,7,10-trioxa-14-azaicosyl)carbamate (9b). Compound **9b** was obtained according to GP1. The crude mixture was purified by column chromatography using Al_2O_3 type III neutral (hexanes : EtOAc 50 : 50, 40 : 60, 30 : 70, 20 : 80, 10 : 90, 0 : 100, then CH_2Cl_2 : MeOH 99 : 1), followed by twice Al_2O_3 type III basic (CH_2Cl_2 : MeOH, 100 : 0, 99 : 1) to afford **9b** (0.104 g, 64%) as dark red solid. 1H NMR ($CDCl_3$, 500 MHz) 1.42 (s, 9H), 1.63–1.66 (m, 4H), 1.70–1.80 (m, 4H), 2.14–2.19 (m, 5H), 2.38 (s, 3H), 2.65–2.69 (m, 2H), 3.19 (d, $J = 5.5$ Hz, 2H), 3.34 (dd, $J = 12.0$ and 6.0 Hz, 2H), 3.48–3.62 (m, 12H), 3.97 (s, 3H), 5.02 (m, 1H), 6.04 (s, 1H), 6.19–6.21 (m, 1H), 6.39 (br s, 1H), 6.72 (d, $J = 3.6$ Hz, 1H), 6.76 (s, 1H), 6.91 (s, 1H). ^{13}C NMR ($CDCl_3$, 125 MHz) 12.3, 13.9, 23.7, 25.4, 28.5, 29.1, 29.7, 36.6, 37.8, 38.5, 42.3, 58.6, 70.0, 70.2 ($\times 2$), 70.6, 78.9, 96.1, 110.6, 111.9, 113.9, 123.3, 123.5, 126.1, 128.1, 129.7, 140.3, 142.3, 156.2, 161.0, 169.1, 172.9, 197.5 (2 carbon atoms unaccounted for). HRMS (TOF) (m/z): $[M + H]^+$ calcd for $C_{37}H_{55}N_5NaO_5$, 698.4123; found, 698.4102.

(Z)-tert-Butyl (24-(2-((4-methoxy-1H,1'H-[2,2'-bipyrryl]-5-yl)methylene)-3,5-dimethyl-2H-pyrrol-4-yl)-15,24-dioxo-4,7,10-trioxa-14-azatetracosyl)carbamate (9c). Compound **9c** was obtained according to GP1. The crude mixture was purified by column chromatography using Al_2O_3 type III basic (EtOAc : hexanes 60 : 40, 80 : 20, EtOAc : MeOH 100 : 0, 99 : 1) followed by Al_2O_3 type III neutral (EtOAc : hexanes 60 : 40, 80 : 20, EtOAc : MeOH 100 : 0, 99 : 1, 98 : 2) and Al_2O_3 type III neutral (EtOAc 100%) to afford **9c** as a red glass (0.200 g, 86%). 1H NMR ($CDCl_3$, 500 MHz) 1.28 (br s, 8H), 1.43 (s, 9H), 1.56–1.63 (m, 4H), 1.75 (quint., $J = 6.0$ Hz, 4H), 2.12 (t, $J = 7.5$ Hz, 2H), 2.20 (s, 3H), 2.39 (s, 3H), 2.63 (t, $J = 7.5$ Hz, 2H), 3.19–3.23 (m, 2H), 3.34 (q, $J = 6.0$ Hz, 2H), 3.49–3.64 (m, 12H), 3.97 (s, 3H), 4.97 (br s, 1H), 6.04 (s, 1H), 6.19–6.21 (m, 2H), 6.72 (d, $J = 3.6$ Hz, 1H), 6.75 (br s, 1H), 6.91 (s, 1H). ^{13}C NMR ($CDCl_3$, 125 MHz) 12.4, 14.1, 24.3, 25.9, 28.6, 29.1, 29.3, 29.4, 29.5, 29.8, 36.9, 38.0, 38.6, 42.8, 58.6, 69.7, 70.3 ($\times 2$), 70.6, 70.7, 77.4, 77.6, 79.0, 96.1, 110.7, 112.1, 113.8, 123.3, 126.2, 128.3, 129.6, 140.5, 142.1, 156.2, 160.9, 169.0, 173.2, 198.2 (1 carbon atom unaccounted for). HRMS (TOF) (m/z): $[M + Na]^+$ calcd for $C_{41}H_{63}N_5Na_1O_8$, 776.4569; found, 776.4558.

(Z)-tert-Butyl (2-((2-(4-(2-((4-methoxy-1H,1'H-[2,2'-bipyrryl]-5-yl)methylene)-3,5-dimethyl-2H-pyrrol-4-yl)-4-oxobutanamido)ethyl)disulfanyl)ethyl)carbamate (10a). Compound **10a** was obtained according to GP1. The crude mixture was purified by column chromatography using Al_2O_3 type III neutral (EtOAc : hexanes 20 : 80, 30 : 70, 40 : 60, 50 : 50, 60 : 40, 70 : 50, 100 : 0, EtOAc : MeOH 99 : 1) to afford **10a** as a red glass (0.052 g, 72%). 1H NMR ($CDCl_3$, 300 MHz) 1.43 (s, 9H), 2.28 (s, 3H), 2.38 (s, 3H), 2.55 (t, $J = 6.4$ Hz, 2H), 2.75–2.82 (m, 4H), 3.04 (t, $J = 6.4$ Hz, 2H), 3.41–3.45 (m, 2H), 3.55 (q, $J = 6.4$ Hz, 2H), 3.97 (s, 3H), 5.05 (br s, 1H), 6.03 (s, 1H), 6.23 (dd, $J = 3.6$, 2.4 Hz, 1H), 6.54 (t, $J = 5.1$ Hz, 1H), 6.23 (dd, $J = 3.6$, 2.4 Hz, 1H), 6.52–6.56 (m, 1H), 6.73 (dd, $J = 3.6$, 1.2 Hz, 1H), 6.81 (br s, 1H), 6.88 (s, 1H). ^{13}C NMR ($CDCl_3$, 125 MHz) 12.6, 14.5, 28.5, 30.5, 38.2, 38.4, 39.6, 58.6, 77.4, 79.7, 96.0, 110.8, 111.9, 113.9, 122.7, 123.5, 126.3, 128.1, 130.0, 140.5, 142.8, 156.0, 160.7, 168.9, 173.2, 195.9 (1 carbon atom unaccounted for). HRMS (TOF) (m/z): $[M + H]^+$ calcd for $C_{29}H_{40}N_5O_5S_2$, 602.2465; found, 602.2456.

(Z)-tert-Butyl (2-((2-(6-(2-((4-methoxy-1H,1'H-[2,2'-bipyrryl]-5-yl)methylene)-3,5-dimethyl-2H-pyrrol-4-yl)-6-oxohexanamido)ethyl)disulfanyl)ethyl)carbamate (10b). Compound **10b** was obtained according to GP1. The crude mixture was purified by column chromatography using Al_2O_3 type III neutral (EtOAc : hexanes 20 : 80, 30 : 70, 40 : 60, 50 : 50, 60 : 40, 70 : 30, 100 : 0 then EtOAc : MeOH 99/1) to afford **10b** as a red-brown glass (0.090 g, 75%). 1H NMR ($CDCl_3$, 300 MHz) 1.43 (s, 9H), 1.66–1.68 (m, 4H), 2.21–2.26 (m, 5H), 2.39 (s, 3H), 2.69 (t, $J = 6.3$ Hz, 2H), 2.75 (t, $J = 6.6$ Hz, 2H), 2.82 (t, $J = 6.3$ Hz, 2H), 3.37–3.43 (m, 2H), 3.55 (q, $J = 6.1$ Hz, 2H), 3.98 (s, 3H), 5.02 (br s, 1H), 6.05 (s, 1H), 6.21 (dd, $J = 3.6$, 2.4 Hz, 1H), 6.59 (br s, 1H), 6.73 (dd, $J = 3.6$, 1.2 Hz, 1H), 6.77 (br s, 1H), 6.92 (s, 1H). ^{13}C NMR ($CDCl_3$, 125 MHz) 12.4, 14.0, 23.7, 25.3, 28.5, 36.4, 38.0, 38.3, 38.4, 39.6, 42.3, 58.7, 79.7, 96.2, 110.7, 112.0, 114.0, 123.5, 126.2, 128.2, 129.7, 140.5, 142.3, 156.0, 161.1, 169.1, 173.3, 197.6 (1



carbon atom unaccounted for). HRMS (TOF) (m/z): $[M + H]^+$ calcd for $C_{31}H_{44}N_5O_5S_2$, 630.2778; found, 630.2770.

(Z)-tert-Butyl (2-((2-(10-(2-((4-methoxy-1H,1'H-[2,2'-bipyrrol]-5-yl)methylene)-3,5-dimethyl-2H-pyrrol-4-yl)-10-oxodecanamido)ethyl)disulfanyl)ethyl)carbamate (10c).

Compound **10c** was obtained according to GP1. The crude mixture was purified by column chromatography using Al_2O_3 type III basic (EtOAc : hexanes 50 : 50, 70 : 30 EtOAc 100%) followed by Al_2O_3 type III neutral (EtOAc : hexanes 50 : 50, 60/40) to afford **10c** as a red glass (0.138 g, 65%). 1H NMR ($CDCl_3$, 300 MHz) 1.28 (s, 8H), 1.43 (s, 9H), 1.58–1.62 (m, 4H), 2.14 (s, 3H), 2.18 (t, $J = 7.4$ Hz, 2H), 2.39 (s, 3H), 2.62 (t, $J = 7.4$ Hz, 2H), 2.75 (t, $J = 6.7$ Hz, 2H), 2.81 (t, $J = 5.8$ Hz, 2H), 3.39–3.43 (m, 2H), 3.54 (q, $J = 6.1$ Hz, 2H), 3.98 (s, 3H), 5.06 (br s, 1H), 6.06 (s, 1H), 6.18 (t, $J = 3.0$ Hz, 1H), 6.41 (br s, 1H), 6.72 (d, $J = 3.0$ Hz, 2H), 6.93 (s, 1H). ^{13}C NMR ($CDCl_3$, 100 MHz) 12.4, 14.0, 24.3, 25.8, 28.5, 29.3, 29.5, 36.7, 37.9, 38.2, 38.6, 39.7, 42.8, 58.7, 79.7, 96.1, 110.7, 112.1, 113.9, 123.4, 123.6, 126.2, 128.2, 129.8, 140.3, 142.2, 156.1, 161.0, 169.1, 173.7, 198.2 (2 carbon atoms unaccounted for). HRMS (TOF) (m/z): $[M + H]^+$ calcd for $C_{35}H_{52}N_5O_5S_2$, 686.3404; found, 686.3386.

(Z)-N-(2-Aminoethyl)-4-(2-((4-methoxy-1H,1'H-[2,2'-bipyrrol]-5-yl)methylene)-3,5-dimethyl-2H-pyrrol-4-yl)-4-oxobutanamide (2a).

A solution of **8a** (0.038 g, 0.070 mmol) in MeOH (1 mL) was treated with HCl (1.2 mL, 2 M solution in Et_2O , 2.40 mmol) and the reaction mixture was stirred at room temperature for 18 h. The solvent was removed under reduced pressure and the crude mixture was dissolved in MeOH (1.5 mL). NaOH (10% aq. solution) was added to the solution until a colour change occurred (dark red to orange). The solvent was removed under reduced pressure and the precipitate was filtered and washed with H_2O to afford **2a** (0.023 g, 80%) as orange solid. Mp 119 °C. 1H NMR ($DMSO-d_6$, 500 MHz) 2.36 (s, 3H), 2.41 (t, $J = 6.7$ Hz, 2H), 2.56 (t, $J = 6.3$ Hz, 2H), 2.69 (s, 3H), 2.99 (t, $J = 6.8$ Hz, 2H), 3.05 (dd, $J = 12.1, 6.2$ Hz, 2H), 3.88 (s, 3H), 6.19 (s, 1H), 6.25 (br s, 1H), 6.70 (s, 1H), 6.79 (br d, 1H, $J = 3.4$ Hz), 7.14 (br s, 1H), 7.78–7.79 (m, 1H), 11.81 (s, 1H). ^{13}C NMR ($DMSO-d_6$, 125 MHz) 11.8, 15.4, 29.4, 37.3, 41.4, 42.4, 58.5, 96.3, 110.1 (2C), 113.1, 122.1, 122.9, 126.3, 127.5, 128.2, 140.8, 142.1, 159.4, 167.3, 171.6, 195.3 (1 carbon atom unaccounted for). HRMS (TOF) (m/z): $[M + H]^+$ calcd for $C_{22}H_{28}N_5O_3$, 410.2187; found, 410.2177.

(Z)-N-(2-Aminoethyl)-6-(2-((4-methoxy-1H,1'H-[2,2'-bipyrrol]-5-yl)methylene)-3,5-dimethyl-2H-pyrrol-4-yl)-6-oxohexanamide (2b).

A solution of **8b** (0.100 g, 0.186 mmol) in a mixture of MeOH : $CHCl_3$ (1 : 1, 4 mL) was treated with HCl (3.3 mL, 2 M solution in Et_2O , 6.60 mmol) and the reaction mixture was stirred at room temperature for 18 h. The solvent was removed under reduced pressure and the crude mixture was dissolved in MeOH (3.8 mL). NaOH (10% aq. solution) was added to the solution until a colour change occurred (dark red to orange). The solvent was removed under reduced pressure and the precipitate was filtered and washed with H_2O to afford **2b** (0.076 g, 93%) as orange solid. 1H NMR (CD_3OD , 500 MHz) 1.68–1.72 (m, 4H), 2.24–2.27 (m, 2H), 2.40 (s, 3H), 2.66 (s, 3H), 2.79–2.83 (m, 2H), 3.28–3.32 (m, 2H), 3.92 (s, 3H), 6.08 (s, 1H), 6.25–6.27 (m, 1H), 6.76–6.78 (m, 2H), 7.02–7.03 (m, 1H). ^{13}C NMR (CD_3OD , 125 MHz) 12.3, 15.4, 25.1, 26.6, 37.0, 41.5, 41.7,

43.1, 59.00, 96.7, 111.2, 111.3, 114.1, 123.5, 123.7, 128.2, 128.9, 129.8, 142.7, 144.2, 162.0, 169.6, 176.6, 199.9. HRMS (TOF) (m/z): $[M + H]^+$ calcd for $C_{24}H_{32}N_5O_3$, 438.2500; found, 438.2488.

(Z)-N-(2-Aminoethyl)-10-(2-((4-methoxy-1H,1'H-[2,2'-bipyrrol]-5-yl)methylene)-3,5-dimethyl-2H-pyrrol-4-yl)-10-oxodecanamide (2c). A solution of **8c** (0.073 g, 0.12 mmol) in a mixture of $CHCl_3$: MeOH (1 : 1, 6 mL) was treated with HCl 12 N (0.024 mL, 0.29 mmol) and the reaction mixture was stirred at room temperature for 7 h. The solvent was removed under vacuum and NaOH (10% aq. solution) was added drop-wise under stirring until the red solution turned orange-brown. The solvent was removed under vacuum and the precipitate was washed with water to give a dark red solid (0.066 g, 76%). Mp 88 °C. 1H NMR ($DMSO-d_6$, 500 MHz) 1.22–1.32 (m, 8H), 1.47–1.49 (m, 2H), 1.56–1.59 (m, 2H), 2.03–2.06 (m, 2H), 2.34 (s, 3H), 2.67 (s, 3H), 2.72–2.74 (m, 2H), 2.99–3.03 (m, 2H), 3.25–3.35 (m, 2H), 3.88 (s, 3H), 6.19 (s, 1H), 6.25 (s, 1H), 6.70 (s, 1H), 6.79 (s, 1H), 7.14 (s, 1H), 7.68 (br s, 1H), 11.81 (br s, 1H). ^{13}C NMR ($DMSO-d_6$, 125 MHz) 11.7, 15.3, 23.7, 25.3, 28.6, 28.7, 28.8, 28.9, 35.4, 41.4, 41.8, 42.2, 58.4, 96.2, 110.1 (2C), 113.0, 122.4, 122.9, 126.3, 127.4, 128.2, 140.6, 142.1, 159.3, 167.3, 172.1, 196.6. HRMS (TOF) (m/z): $[M + H]^+$ calcd for $C_{28}H_{40}N_5O_3$, 494.3126; found, 494.3129.

(Z)-N-(3-(2-(2-(3-Aminopropoxy)ethoxy)ethoxy)propyl)-4-(2-((4-methoxy-1H,1'H-[2,2'-bipyrrol]-5-yl)methylene)-3,5-

dimethyl-2H-pyrrol-4-yl)-4-oxobutanamide (3a). A solution of **9a** (0.125 g, 0.18 mmol) in a mixture of MeOH/ $CHCl_3$ (1 : 1, 4 mL) was treated with HCl 12 N (0.031 mL, 0.37 mmol) at room temperature. After 30 min additional HCl 12 N (0.030 mL, 0.36 mmol) was added and the reaction mixture stirred for an additional 3 hours. The solvent was removed under reduced pressure and the crude mixture was dissolved in MeOH (2 mL). NaOH (10% aq. solution) was added drop-wise under stirring until the red solution turned orange-brown. The solvent was removed under vacuum and the crude solid was purified using flash chromatography on Al_2O_3 basic type III (EtOAc 100%, CH_2Cl_2 : MeOH 98 : 2, 95 : 5, 90 : 10 then MeOH 100%) to afford **3a** as a dark red glass (0.105 g, 51%). 1H NMR ($CDCl_3$, 500 MHz) 1.68–1.73 (m, 4H), 2.20 (s, 3H), 2.34 (s, 3H), 2.48 (t, $J = 6.5$ Hz, 2H), 2.84 (t, $J = 6.5$ Hz, 2H), 2.97 (t, $J = 6.5$ Hz, 2H), 3.28 (q, $J = 6.5$ Hz, 2H), 3.47–3.60 (m, 12H), 3.95 (s, 3H), 6.03 (s, 1H), 6.18–6.19 (s, 1H), 6.68–6.71 (m, 2H), 6.76 (s, 1H), 6.86 (s, 1H). ^{13}C NMR ($CDCl_3$, 125 MHz) 12.5, 14.5, 29.3, 30.4, 30.7, 37.4, 38.0, 40.1, 58.6, 69.3, 69.9, 70.0 ($\times 2$), 70.2, 70.5, 96.0, 110.7, 111.7, 113.9, 122.7, 123.5, 126.4, 128.1, 129.8, 140.6, 142.7, 160.7, 168.9, 173.2, 196.1. HRMS (TOF) (m/z): $[M + H]^+$ calcd for $C_{30}H_{44}N_5O_6$, 570.3286; found, 570.3295.

N-(3-(2-(2-(3-Aminopropoxy)ethoxy)ethoxy)propyl)-6-(2-((4-

methoxy-1H,1'H-[2,2'-bipyrrol]-5-yl)methylene)-3,5-dimethyl-2H-pyrrol-4-yl)-6-oxohexanamide (3b). A solution of **9b** (0.104 g, 0.150 mmol) in a mixture of MeOH : $CHCl_3$ (1 : 1, 3.4 mL) was treated with HCl 12 N (0.55 mL, 6.60 mmol) and the reaction mixture was stirred at room temperature for 25 h. The solvent was removed under reduced pressure and the crude mixture was dissolved in MeOH (2 mL). NaOH (6 M aq. solution) was added to the solution until a colour change occurred (dark red to orange). The solvent was removed under reduced pressure



and the crude mixture was purified using flash chromatography on Al₂O₃ basic type III (CH₂Cl₂ : MeOH, 9 : 1) to afford **3b** (0.074 g, 83%) as a dark red glass. ¹H NMR (CDCl₃, 500 MHz) 1.63 (br s, 4H), 1.66–1.71 (m, 2H), 1.72–1.77 (m, 2H), 2.15 (s, 3H), 2.17 (br s, 2H), 2.37 (s, 3H), 2.65 (br s, 2H), 2.76 (t, 2H, *J* = 6.6 Hz), 3.30–3.34 (m, 2H), 3.52 (t, 4H, *J* = 5.8 Hz), 3.56 (ddd, 12H, *J* = 24.1, 10.1, 8.0 Hz), 3.96 (s, 3H), 6.04 (s, 1H), 6.18–6.19 (m, 1H), 6.63 (br s, 1H), 6.71 (d, 1H, *J* = 3.5 Hz), 6.74 (br s, 1H), 6.91 (s, 1H). ¹³C NMR (CDCl₃, 125 MHz) 12.4, 14.0, 23.8, 25.5, 29.1, 29.8, 31.0, 33.3, 36.7, 37.7, 39.7, 42.4, 58.6, 69.6, 69.9, 70.2, 70.6, 96.2, 110.7, 112.0, 113.9, 123.4 (×2), 126.2, 128.2, 129.7, 140.5, 142.3, 161.1, 169.1, 173.0, 197.6. HRMS (TOF) (*m/z*): [M + H]⁺ calcd for C₃₂H₄₈N₅O₆, 598.3599; found, 598.3577.

N-(3-(2-(2-(3-aminopropoxy)ethoxy)ethoxy)propyl)-10-(2-((4-methoxy-1*H*,1'*H*-[2,2'-bipyrrol]-5-yl)methylene)-3,5-dimethyl-2*H*-pyrrol-4-yl)-10-oxodecanamide (**3c**). A solution of **9c** (0.113 g, 0.150 mmol) in a mixture of MeOH : CHCl₃ (1 : 1, 3.2 mL) was treated with HCl 12 N (0.65 mL of 12 M aqueous solution, 7.80 mmol) and the reaction mixture was stirred at room temperature for 5 h. The solvent was removed under reduced pressure and the crude mixture was dissolved in MeOH (2 mL). NaOH (2 M aq. solution) was added to the solution until a colour change occurred (dark red to orange). The solvent was removed under reduced pressure and the crude mixture was purified by flash chromatography using Al₂O₃ basic type III (CH₂Cl₂ : MeOH, 9 : 1) to afford **3c** (0.068 g, 70%) as a dark red glass. ¹H NMR (CDCl₃, 500 MHz) 1.24–1.27 (m, 8H), 1.56–1.60 (m, 4H), 1.63 (br s, 4H), 1.65–1.79 (m, 4H), 2.11 (d, 2H, *J* = 7.8 Hz), 2.15 (s, 3H), 2.38 (s, 3H), 2.62 (t, 2H, *J* = 7.4 Hz), 2.77 (t, 2H, *J* = 6.7 Hz), 3.33 (dd, 2H, *J* = 12.2, 6.0 Hz), 3.51–3.63 (m, 12H), 3.97 (s, 3H), 6.05 (s, 1H), 6.17–6.19 (m, 1H), 6.39–6.43 (br m, 1H), 6.70–6.73 (m, 2H), 6.92 (s, 1H). ¹³C NMR (CDCl₃, 125 MHz) 12.4, 14.1, 24.3, 25.9, 29.1, 29.3, 29.4, 29.5, 33.4, 36.9, 37.8, 39.8, 58.6, 69.6 (2C), 70.1, 70.3 (2C), 70.7, 96.1, 110.7, 112.1, 113.8, 123.3, 123.6, 126.2, 128.2, 129.7, 140.4, 142.2, 160.9, 169.1, 173.2, 198.1 (2 carbons unaccounted for). HRMS (TOF) (*m/z*): [M + H]⁺ calcd for C₃₆H₅₆N₅O₆, 654.4225; found, 654.4203.

(Z)-**N**-(2-(2-(2-aminoethyl)disulfanyl)ethyl)-4-(2-((4-methoxy-1*H*,1'*H*-[2,2'-bipyrrol]-5-yl)methylene)-3,5-dimethyl-2*H*-pyrrol-4-yl)-4-oxobutanamide (**4a**). A solution of **10a** (0.052 g, 0.077 mmol) in a mixture of MeOH : CHCl₃ (1 : 1, 4 mL) was treated with HCl 12 N (0.024, 0.077 mmol). The reaction was stirred overnight at room temperature and additional HCl 12 N (0.024, 0.077 mmol) was added. The solution was concentrated under reduced pressure and the crude mixture dissolved in MeOH (3 mL). NaOH (10% aq. solution) was added dropwise under stirring until the red solution turned orange-brown. The solvent was removed under vacuum and the crude solid was purified using flash chromatography on Al₂O₃ basic type III (EtOAc 100%, CH₂Cl₂ : MeOH, 95 : 5) to afford **4a** as a dark red glass (0.033 g, 77%) ¹H NMR (CDCl₃, 300 MHz) 2.25 (s, 3H), 2.38 (s, 3H), 2.53 (t, *J* = 6.3 Hz), 2.73–2.80 (m, 4H), 2.97–3.05 (m, 4H), 3.56 (q, *J* = 6.8 Hz, 2H), 3.97 (s, 3H), 6.03 (s, 1H), 6.22 (dd, *J* = 3.8, 2.7 Hz, 1H), 6.38 (t, *J* = 5.7 Hz, 1H), 6.73 (dd, *J* = 3.8, 1.2 Hz, 1H), 6.79 (s, 1H), 6.89 (s, 1H). ¹³C NMR (CDCl₃, 125 MHz) 12.6, 14.8, 30.6, 38.0, 38.2, 38.5, 40.8, 42.7, 58.7, 96.0, 110.9, 111.9, 113.8, 122.7, 123.2, 126.5, 128.2, 129.8, 142.6, 160.7, 168.9,

173.1, 196.0 (1 carbon atom unaccounted for). HRMS (TOF) (*m/z*): [M + H]⁺ calcd for C₂₄H₃₂N₅O₃S₂, 502.1941; found, 502.1933. HRMS (TOF) (*m/z*): [M + H]⁺ calcd for C₂₄H₃₂N₅O₃S₂, 502.1933; found, 502.1941.

(S,Z)-2-(Trimethylsilyl)ethyl 2-((*tert*-butoxycarbonyl)amino)-25-(2-((4-methoxy-1*H*,1'*H*-[2,2'-bipyrrol]-5-yl)methylene)-3,5-dimethyl-2*H*-pyrrol-4-yl)-20,25-dioxo-9,12,15-trioxa-5,19-diazapentacosan-1-oate (**18b**). Compound **3b** was obtained following GP1. The crude mixture was purified using Al₂O₃ type III neutral (CH₂Cl₂ 100%, CH₂Cl₂/MeOH 99/1, 98/2) followed by Al₂O₃ type III basic (CH₂Cl₂ 100%, CH₂Cl₂/MeOH 99/1) to afford **18b** as a red glass (0.020 g, 20%). ¹H NMR (CDCl₃, 300 MHz) 0.02 (s, 9H), 0.95–1.01 (m, 2H), 1.42 (s, 9H), 1.65–1.67 (m, 4H), 1.73–1.81 (m, 4H), 1.87–1.95 (m, 1H), 2.08–2.13 (m, 1H), 2.16–2.20 (m, 3H), 2.26 (s, 3H), 2.39 (s, 3H), 2.67–2.69 (m, 2H), 3.31–3.37 (m, 4H), 3.51–3.64 (m, 12H), 3.96 (s, 3H), 4.14–4.20 (m, 3H), 5.44 (d, 1H, *J* = 8.1 Hz), 6.02 (s, 1H), 6.22 (dd, 1H, *J* = 3.6, 2.7 Hz), 6.45 (br s, 1H), 6.59 (br s, 1H), 6.72 (dd, 1H, *J* = 3.6, 1.2 Hz), 6.79 (br s, 1H), 6.89 (s, 1H). ¹³C NMR (CDCl₃, 125 MHz), −1.4 (3C), 12.5, 14.7, 17.5, 23.9, 25.5, 28.5 (3C), 28.8, 29.2, 29.8, 32.8, 36.7, 37.8, 37.9, 42.4, 53.5, 58.6, 63.9, 70.0 (×2), 70.2, 70.6 (4C), 80.0, 96.0, 110.8, 111.9, 113.6, 123.1, 123.3, 126.6, 128.4, 129.3, 141.2, 141.9, 156.0, 160.8, 168.9, 172.1, 172.6, 173.0, 197.6. HRMS (TOF) (*m/z*): [M + Na]⁺ calcd for C₄₇H₇₄N₆Na₁O₁₁Si₁, 949.5077; found, 949.5086.

(S,Z)-2-(Trimethylsilyl)ethyl 2-((*tert*-butoxycarbonyl)amino)-29-(2-((4-methoxy-1*H*,1'*H*-[2,2'-bipyrrol]-5-yl)methylene)-3,5-dimethyl-2*H*-pyrrol-4-yl)-20,29-dioxo-9,12,15-trioxa-5,19-diazanonacosan-1-oate (**18c**). Compound **3c** was obtained following GP1. The crude mixture was purified using Al₂O₃ type III basic (EtOAc 100%, EtOAc/MeOH 99/1, CH₂Cl₂/MeOH 98/2), followed by Al₂O₃ type III basic (CH₂Cl₂/MeOH 99/1) and Al₂O₃ type III neutral (CH₂Cl₂/MeOH 99/1, 98/2) to afford **18c** as a red glass (0.130 g, 46%). ¹H NMR (CDCl₃, 300 MHz) 0.03 (s, 9H), 0.96–1.02 (m, 2H), 1.28 (s, 8H), 1.43 (s, 9H), 1.59–1.62 (m, 4H), 1.76 (quint., 4H, *J* = 6.0 Hz), 1.87–1.97 (m, 1H), 2.12 (t, 3H, *J* = 7.5 Hz), 2.21–2.26 (m, 5H), 2.39 (s, 3H), 2.64 (t, 2H, *J* = 7.3 Hz), 3.33 (q, 4H, *J* = 6.0 Hz), 3.52–3.64 (m, 12H), 3.96 (s, 3H), 4.16–4.21 (m, 3H), 5.43 (d, 1H, *J* = 8.1 Hz), 6.03 (s, 1H), 6.21–6.23 (m, 2H), 6.56 (br s, 1H), 6.71–6.73 (m, 1H), 6.79 (br s, 1H), 6.91 (s, 1H). ¹³C NMR (CDCl₃, 100 MHz), −1.4 (3C), 12.4, 14.3, 17.5, 24.3, 25.9, 28.4 (3C), 28.8, 29.1, 29.2, 29.3, 29.4, 29.5, 32.8, 36.9, 37.7, 37.9, 42.8, 53.5, 58.6, 63.9, 69.9, 70.1, 70.5 (4C), 77.4, 77.6, 80.0, 96.0, 110.8, 112.1, 113.8, 123.3, 123.5, 126.3, 128.2, 129.7, 142.1, 155.9, 168.9, 172.1, 172.6, 173.4, 198.2. HRMS (TOF) (*m/z*): [M + Na]⁺ calcd for C₅₁H₈₂N₆Na₁O₁₁Si₁, 1005.5703; found, 1005.5713.

(S,Z)-2-(Trimethylsilyl)ethyl 2-amino-25-(2-((4-methoxy-1*H*,1'*H*-[2,2'-bipyrrol]-5-yl)methylene)-3,5-dimethyl-2*H*-pyrrol-4-yl)-20,25-dioxo-9,12,15-trioxa-5,19-diazapentacosan-1-oate (**19b**). Prodigiosene **18b** (0.020 g, 0.021 mmol) was dissolved in a mixture of dioxane : water (1 : 1, 1 mL) and PTSA (0.016 g, 0.052 mmol) was added. The reaction was stirred at 60 °C for 3 hours and additional PTSA (0.016 g, 0.052 mmol) was added. After 16 h the solvent was removed under reduced pressure the crude solid was purified using Al₂O₃ type III basic (CH₂Cl₂/MeOH 99/1) to afford **19b** as an orange glass (0.011 g, 64%). ¹H



NMR (CDCl₃, 300 MHz) 0.04 (s, 9H), 0.96–1.02 (m, 2H), 1.65–1.68 (m, 4H), 1.73–1.82 (m, 5H), 2.04–2.13 (m, 1H), 2.15–2.29 (m, 2H), 2.32–2.40 (m, 2H), 2.68 (s, 3H), 2.72 (s, 3H), 3.30–3.43 (m, 2H), 3.50–3.64 (m, 5H), 3.96 (s, 3H), 4.15–4.21 (m, 2H), 6.01 (s, 1H), 6.23–6.25 (m, 1H), 6.45 (t, 1H, *J* = 5.1 Hz), 6.51 (t, 1H, *J* = 4.8 Hz), 6.72 (d, 1H, *J* = 3.6 Hz), 6.84 (br s, 1H), 6.89 (s, 1H). ¹³C NMR (CDCl₃, 100 MHz), –1.4 (3C), 12.5, 14.9, 17.6, 23.9, 25.5, 29.2, 29.3, 30.5, 33.0, 36.7, 37.8, 42.4, 54.2, 58.6, 63.5, 70.0, 70.1, 70.2, 70.6 (4C), 95.9, 110.8, 111.9, 113.5, 123.0, 123.2, 126.7, 128.4, 129.2, 141.4, 141.7, 160.6, 168.8, 172.5, 173.0, 175.9, 197.6.0. HRMS (TOF) (*m/z*): [M + Na]⁺ calcd for C₄₂H₆₆N₆Na₁O₉Si₁, 849.4553; found, 849.4557.

(*S,Z*)-2-(Trimethylsilyl)ethyl 2-amino-29-(2-((4-methoxy-1*H*,1'*H*-[2,2'-bipyrrrol]-5-yl)methylene)-3,5-dimethyl-2*H*-pyrrol-4-yl)-20,29-dioxo-9,12,15-trioxa-5,19-diazanonacosan-1-oate (19c). Prodigiosene **18c** (0.130 g, 0.13 mmol) was dissolved in a mixture of dioxane : water (1 : 1, 6 mL) and PTSA (0.063 g, 0.32 mmol) was added. The reaction was stirred at 60 °C for 3 hours and additional PTSA (0.063 mg, 0.32 mmol) was added. After 16 h the reaction mixture was quenched by addition of NaHCO₃ solid until the dark red solution turned dark brown. The solvent were removed under vacuum and the crude solid was purified using Al₂O₃ type III basic (CH₂Cl₂/MeOH 98/2, 95/5) to afford **19c** as an orange glass (0.066 g, 57%). ¹H NMR (CDCl₃, 300 MHz) 0.03 (s, 9H), 0.96–1.01 (m, 2H), 1.24–1.27 (m, 11H), 1.58–1.60 (m, 4H), 1.74 (quint., 4H, *J* = 6.1 Hz), 2.09–2.14 (m, 3H), 2.19 (s, 3H), 2.29 (t, 2H, *J* = 7.3 Hz), 2.38 (s, 3H), 2.63 (t, 2H, *J* = 7.3 Hz), 3.32 (q, 4H, *J* = 6.1 Hz), 3.38–3.43 (m, 1H), 3.50–3.64 (m, 13H), 3.96 (s, 3H), 4.15–4.21 (m, 2H), 6.04 (s, 1H), 6.19 (dd, 1H, *J* = 3.6, 2.7 Hz), 6.24 (br s, 1H), 6.49 (br s, 1H), 6.71 (dd, 1H, *J* = 3.6, 1.1 Hz), 6.74–6.75 (m, 1H), 6.91 (s, 1H). ¹³C NMR (CDCl₃, 125 MHz), –1.4 (3C), 12.4, 14.3, 17.6, 24.3, 25.9, 29.2, 29.3, 29.4, 29.5, 29.8, 30.5, 32.9, 36.9, 37.8 (×2), 42.8, 54.1, 58.6, 63.4, 70.0 (×2), 70.2, 70.6 (4C), 96.0, 110.7, 112.1, 113.7, 123.2, 123.5, 126.3, 128.3, 129.5, 140.8, 142.0, 160.8, 179.0, 172.4, 173.3, 175.9, 198.2. HRMS (TOF) (*m/z*): [M + Na]⁺ calcd for C₄₆H₇₄N₆Na₁O₉Si₁, 905.5179; found, 905.5145.

General procedure 2 (GP2)

A solution of linker-prodigiosene conjugate (1 eq.) and **23** (1 eq.) in anhydrous DMSO (0.05 M) was treated with NEt₃ (2 eq.). The resulting reaction mixture was stirred at room temperature for 5 h under N₂ atmosphere. Water was added to the reaction mixture until a precipitate formed, which was isolated *via* microfiltration, then washed 3 times with water and dried under vacuum.

Teoc-FA-DA-prod(C2) (24a). Compound **24a** was obtained according to GP2 as an orange solid (0.096 g, 83%). Mp 148–150 °C. ¹H NMR (DMSO-*d*₆, 500 MHz) 0.02 (s, 9H), 0.07 (s, 9H), 0.93–0.96 (m, 2H), 1.02–1.06 (m, 2H), 1.37 (s, 6H), 1.92–2.27 (m, 4H), 2.36 (s, 3H), 2.68 (s, 3H), 2.73 (s, 1H), 2.98–3.00 (m, 4H), 3.88 (s, 3H), 4.10–4.16 (m, 2H), 4.25–4.30 (m, 2H), 4.35 (br s, 1H), 4.57 (br s, 2H), 6.17 (s, 1H), 6.25 (s, 1H), 6.67 (br d, 2H, *J* = 7.0 Hz), 6.79 (br s, 1H), 6.83 (br s, 1H), 7.11 (br s, 1H), 7.66 (br d, 3H, *J* = 7.2 Hz), 8.16 (br s, 1H), 8.78 (s, 1H), 11.70 (br s, 1H). ¹³C NMR (DMSO-*d*₆, 125 MHz), –1.5 (6C), 11.9, 15.4, 16.8, 17.1,

26.4, 28.2, 29.5, 31.8, 37.3 (2C), 46.0, 52.3, 58.4, 62.4, 64.1, 77.6, 96.3, 110.1, 111.2 (2C), 113.1, 121.3, 122.1, 122.9, 126.4, 127.5, 128.2, 129.0 (2C), 129.8, 140.9, 142.2, 148.9, 150.7, 151.4, 154.9, 155.3, 155.6, 159.5, 160.9, 166.4, 167.4, 171.6, 171.8, 172.3, 195.3. HRMS (TOF) (*m/z*): [M + H]⁺ calcd for C₅₂H₆₉N₁₂NaO₁₀Si₂, 1077.4793; found, 1077.4748.

Teoc-FA-DA-prod(C4) (24b). Compound **24b** was obtained according to GP2 as orange solid (0.096 g, 84%). Mp 138–140 °C. ¹H NMR (DMSO-*d*₆, 500 MHz) 0.00 (s, 9H), 0.05 (s, 9H), 0.86–0.87 (m, 3H), 0.91–0.94 (m, 2H), 1.02–1.05 (m, 3H), 1.24–1.28 (m, 1H), 1.36 (br s, 3H), 1.54 (br s, 4H), 1.91–2.07 (m, 4H), 2.18 (br s, 2H), 2.34 (s, 3H), 2.67 (s, 3H), 2.72 (br s, 2H), 3.06 (m, 4H), 3.87 (s, 3H), 4.09–4.12 (m, 2H), 4.28–4.29 (m, 2H), 4.58 (br s, 3H), 6.18 (s, 1H), 6.24 (s, 1H), 6.65 (br d, 2H, *J* = 7.0 Hz), 6.68 (s, 1H), 6.78 (br s, 1H), 6.99 (br s, 1H), 7.13 (s, 1H), 7.65 (br d, 2H, *J* = 7.0 Hz), 7.77 (br s, 1H), 7.85 (br s, 1H), 8.28–8.31 (m, 1H), 8.81 (s, 1H), 11.55 (br s, 2H), 11.81 (br s, 1H). ¹³C NMR (DMSO-*d*₆, 125 MHz), –1.5 (6C), 11.8, 15.4, 16.9, 17.1, 23.4, 25.0, 26.4, 28.2, 31.8, 35.5, 38.3, 38.5, 41.6, 46.0, 52.3, 58.4, 62.4, 64.6, 79.2, 96.2, 110.1, 111.3, 113.0, 121.4, 122.3, 122.9, 126.3, 127.5, 128.2, 128.7, 129.1, 129.9, 140.8, 142.2, 149.1, 149.2, 150.7, 151.9, 154.5, 155.0, 159.4, 159.6, 166.4, 167.4, 171.6, 172.2, 172.3, 196.4. HRMS (TOF) (*m/z*): [M + H]⁺ calcd for C₅₄H₇₃N₁₂O₁₀Si₂, 1105.5106; found, 1105.5086.

Teoc-FA-DA-prod(C4) (24c). Compound **24c** was obtained according to GP2 as a dark red solid (0.098 g, 70%). Mp 124–126 °C. ¹H NMR (CDCl₃, 500 MHz) 0.02 (s, 9H), 0.05 (s, 9H), 0.99–1.05 (m, 4H), 1.21–1.25 (m, 8H), 1.56 (s, 4H), 1.98 (s, 1H), 2.13–2.17 (m, 3H), 2.26–2.34 (m, 8H), 2.57–2.61 (m, 2H), 3.21 (br s, 1H), 3.33 (s, 1H), 3.43–3.50 (m, 2H), 3.94 (s, 3H), 4.20–4.29 (m, 4H), 4.58 (s, 2H), 4.65–4.68 (m, 1H), 5.56 (br s, 1H), 5.97 (s, 1H), 6.25 (br s, 1H), 6.59 (br s, 2H), 6.72 (br s, 1H), 6.84 (s, 2H), 6.96–6.97 (m, 2H), 7.12 (br s, 1H), 7.19 (br s, 1H), 7.63–7.64 (m, 2H), 8.74 (s, 1H). ¹³C NMR (CDCl₃, 125 MHz), –1.5 (6C), 11.8, 15.3, 16.8, 17.1, 23.8, 25.2, 26.3, 28.2, 28.8, 28.9, 31.7, 35.4, 37.8 (2C), 41.8, 46.1, 52.2, 58.4, 62.4, 63.6, 77.6, 96.2, 110.1, 111.2 (2C), 111.3, 113.0, 121.2, 121.3, 122.4, 122.9, 126.3, 127.4, 128.2, 129.0, 129.6 (2C), 140.6, 142.1, 148.5, 150.7 (×2), 152.2, 155.2, 155.6, 159.4, 162.0, 166.3, 167.3, 171.5, 172.3, 196.6. HRMS (TOF) (*m/z*): [(M + H)]⁺ calcd for C₅₈H₈₁N₁₂O₁₀Si₂, 1161.5732; found, 1161.5700.

Teoc-FA-PEG-prod(C2) (25a). Compound **25a** was obtained according to GP2 as an orange-brown solid (0.146 g, 91%). Mp 114–115 °C. ¹H NMR (CDCl₃, 500 MHz) 0.01 (s, 9H), 0.06 (s, 9H), 0.96–1.00 (m, 2H), 1.02–1.06 (m, 2H), 1.20 (t, *J* = 7.0 Hz, 1H), 1.71 (dt, *J* = 21.5, 6.0 Hz, 4H), 2.10–2.14 (m, 2H), 2.19–2.29 (m, 2H), 2.33 (s, 6H), 2.49 (t, *J* = 6.7 Hz, 2H), 2.97 (t, *J* = 6.7 Hz, 2H), 3.23–3.32 (m, 4H), 3.46–3.63 (m, 12H), 3.92 (s, 3H), 4.16–4.19 (m, 2H), 4.28 (t, *J* = 7.7 Hz, 2H), 4.57–4.64 (m, 3H), 5.38 (br s, 1H), 5.93 (s, 1H), 6.26 (s, 1H), 6.56 (d, *J* = 7.2 Hz, 2H), 6.71 (s, 2H), 6.78 (s, 2H), 6.96 (br s, 1H), 7.39 (d, *J* = 6.5 Hz, 1H), 7.67 (d, *J* = 7.2 Hz, 2H), 8.74 (br s, 1H). ¹³C NMR (CDCl₃, 125 MHz), –1.41 (6C), 12.5, 15.4, 17.5, 17.7, 27.7, 29.0, 29.2, 30.4 (2C), 32.9, 37.6, 37.8, 38.1 (2C), 46.7, 53.1, 58.6, 63.9, 65.6, 65.9 (2C), 69.6, 69.7, 70.0, 70.1, 70.5 (2C), 95.5, 110.8, 111.5, 112.1 (2C), 113.9, 122.5 (×2), 123.8, 126.5, 127.7, 129.2, 129.8 (2C), 142.6, 149.0, 150.4, 151.5, 154.0, 154.7, 167.5, 168.2, 172.7, 173.1, 196.0 (2



carbons unaccounted for). HRMS (TOF) (m/z): $[M + H]^+$ calcd for $C_{60}H_{85}N_{12}O_{13}Si_2$, 1237.5892; found, 1237.5890.

Teoc-FA-PEG-prod(C4) (25b). Compound **25b** was obtained according to GP2 as a dark red solid (0.101 g, 89%). Mp 128–130 °C. 1H NMR ($CDCl_3$, 500 MHz) 0.03 (s, 9H), 0.07 (s, 9H), 0.98–1.01 (m, 2H), 1.04–1.07 (m, 2H), 1.61 (br s, 4H), 1.70–1.72 (m, 2H), 1.75–1.78 (m, 2H), 2.16–2.22 (m, 4H), 2.30–2.39 (m, 8H), 2.61 (br s, 2H), 3.23–3.36 (m, 4H), 3.50–3.60 (m, 12H), 3.94 (s, 3H), 3.97–3.99 (br m, 1H), 4.18–4.22 (m, 2H), 4.27–4.31 (m, 2H), 4.57 (br s, 2H), 4.61–4.66 (m, 2H), 5.44 (br s, 1H), 5.95 (s, 1H), 6.27 (s, 1H), 6.58 (d, 2H, $J = 7.2$ Hz), 6.72–6.74 (m, 2H), 6.81 (s, 1H), 6.90 (br s, 1H), 6.97 (br s, 1H), 7.41 (d, 1H, $J = 7$ Hz), 7.68 (d, 2H, $J = 7.2$ Hz), 8.74 (br s, 1H). ^{13}C NMR ($CDCl_3$, 125 MHz), –1.5 (6C), 12.3, 15.1, 17.4, 17.6, 23.7, 25.4, 27.7, 28.5, 29.0, 29.2, 29.7, 32.8, 36.5, 37.5, 37.8, 42.3, 46.7, 53.0, 58.6, 63.8, 65.7, 69.7, 70.0, 70.1, 70.4 (2C), 95.6, 110.8, 111.5, 112.1 (2C), 114.0, 122.5, 123.0, 123.6, 126.4, 127.6, 127.9, 129.1, 129.8 (2C), 139.9, 142.2, 149.1, 150.3, 151.5, 154.1, 154.6, 159.4, 161.9, 167.5, 168.3, 172.6, 172.7, 173.2, 197.4. HRMS (TOF) (m/z): $[(M + H + Na)/2]^{2+}$ calcd for $C_{62}H_{89}N_{12}NaO_{13}Si_2$, 644.3049; found, 644.3030.

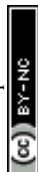
Teoc-FA-PEG-prod(C8) (25c). Compound **25c** was obtained according to GP2. The crude mixture was purified by column chromatography using Al_2O_3 type III neutral (CH_2Cl_2 : MeOH, 9 : 1) and **25c** was obtained as dark red glass (0.065 g, 47%). 1H NMR ($CDCl_3$, 500 MHz) 0.01 (s, 9H), 0.05 (s, 9H), 0.82–0.89 (m, 1H), 0.96–1.07 (m, 4H), 1.25 (br s, 8H), 1.57–1.58 (br m, 4H), 1.68–1.77 (m, 4H), 2.09–2.14 (m, 2H), 2.19–2.25 (m, 2H), 2.29–2.33 (m, 2H), 2.36 (s, 6H), 2.61 (t, 2H, $J = 7.1$ Hz), 3.27–3.34 (m, 4H), 3.47–3.58 (m, 12H), 3.94 (s, 3H), 4.17–4.22 (m, 2H), 4.26–4.31 (m, 2H), 4.62 (br s, 3H), 5.43 (br s, 1H), 5.98 (s, 1H), 6.24 (br s, 1H), 6.44–6.48 (m, 1H), 6.60 (d, 2H, $J = 8.3$ Hz), 6.72 (br s, 1H), 6.81–6.85 (m, 2H), 6.90 (br s, 1H), 7.40 (d, 1H, $J = 7.2$ Hz), 7.67 (d, 2H, $J = 8.3$ Hz), 8.77 (s, 1H). ^{13}C NMR ($CDCl_3$, 125 MHz), –1.4 (6C), 12.4, 15.2, 17.5, 17.7, 24.3, 25.9, 27.9, 29.0, 29.3 ($\times 2$), 29.4, 29.5 ($\times 2$), 29.8 (2C), 32.9, 36.9, 37.7, 38.0 (2C), 42.8, 46.9, 53.1, 58.7, 64.0, 66.1 (2C), 69.9, 70.0, 70.1, 70.2, 70.6, 95.7, 110.9, 111.9, 112.3 (2C), 114.0, 122.8, 123.5, 126.6, 127.9, 129.2, 130.1 (2C), 140.2, 142.2, 149.4, 150.3, 151.8, 154.0, 154.8, 159.6, 161.0, 167.3, 168.5, 172.6, 172.7, 173.5, 198.1. HRMS (TOF) (m/z): $[M + H]^+$ calcd for $C_{66}H_{97}N_{12}O_{13}Si_2$, 1321.6831; found, 1321.6807.

Teoc-FA-SS-prod(C2) (26a). Compound **26a** was obtained according to GP2 as an orange-brown solid (0.042 g, 54%). Mp 149–151 °C. 1H NMR ($CDCl_3$, 500 MHz) 0.00 (1 s, 9H), 0.04 (s, 9H), 0.95 (t, $J = 8.5$ Hz, 2H), 1.01 (t, $J = 8.5$ Hz, 2H), 1.25 (s, 1H), 2.16–2.20 (m, 2H), 2.26 (s, 6H), 2.33–2.39 (m, 2H), 2.45 (br s, 2H), 2.61 (br s, 2H), 2.71–2.72 (m, 2H), 2.92 (br s, 2H), 3.42–3.51 (m, 4H), 3.91 (s, 3H), 4.15 (t, $J = 8.5$ Hz, 2H), 4.24–4.32 (m, 3H), 4.55 (br s, 2H), 4.65–4.68 (m, 1H), 5.49 (br s, 1H), 5.92 (s, 1H), 6.23 (s, 1H), 6.53 (d, $J = 7.7$ Hz, 2H), 6.69 (s, 1H), 6.72 (s, 1H), 6.91 (s, 1H), 7.06 (br s, 1H), 7.36 (br s, 1H), 7.64 (d, $J = 7.7$ Hz, 2H), 8.71 (s, 1H). ^{13}C NMR ($CDCl_3$, 125 MHz), –1.4 (6C), 12.5, 15.6, 17.5, 17.7, 27.5, 28.5, 29.8, 30.3, 32.8 (2C), 38.0 (2C), 38.7, 38.8, 46.8, 53.1, 58.6, 64.0, 65.9, 95.4, 110.9, 111.3, 112.1, 112.2 (2C), 114.1, 122.4, 124.0, 126.5 (2C), 127.5, 129.2 (2C), 130.0, 142.9, 149.1, 150.4, 151.9, 154.1, 167.6, 168.1, 172.6, 173.2, 173.6, 196.0 (3 carbon atoms unaccounted for). HRMS (TOF) (m/z):

$[(M + H + Na)/2]^{2+}$ calcd for $C_{54}H_{73}N_{12}NaO_{10}S_2Si_2$, 596.2211; found, 596.2220.

Teoc-FA-SS-prod(C4) (26b). A solution of **4b** (0.128 g, 0.203 mmol) in a mixture of MeOH : $CHCl_3$ (1 : 1, 4 mL) was treated with HCl 12 N (1.2 mL, 14.4 mmol) and the reaction mixture was stirred at room temperature for 4 h. The solvent was removed under reduced pressure to obtain **4b** as dark purple solid (0.124 g), which was used in the next step without any further purification. HRMS (TOF) (m/z): $[M + H]^+$ calcd for $C_{26}H_{36}N_5O_3S_2$, 530.2254; found, 530.2264. Compound **26b** was obtained according to GP2. The crude mixture was purified by column chromatography using Al_2O_3 type III neutral (CH_2Cl_2 : MeOH, 9 : 1) and **26b** was obtained as an orange solid (0.176 g, 72%). Mp 135–137 °C. 1H NMR ($CDCl_3$, 500 MHz) 0.02 (s, 9H), 0.06 (s, 9H), 0.81–0.90 (m, 1H), 0.96–1.06 (m, 4H), 1.58 (br s, 4H), 2.09–2.19 (m, 4H), 2.26 (br m, 3H), 2.31 (s, 3H), 2.34–2.40 (m, 2H), 2.57 (br s, 2H), 2.69–2.78 (m, 4H), 3.40–3.53 (m, 4H), 3.93 (s, 3H), 4.16–4.32 (m, 4H), 4.58 (d, 2H, $J = 4.0$ Hz), 4.65–4.72 (m, 1H), 5.45–5.46 (m, 1H), 5.95 (s, 1H), 6.25–6.27 (m, 1H), 6.56 (d, 2H, $J = 8.6$ Hz), 6.72 (br s, 1H), 6.80 (s, 1H), 6.89–6.95 (m, 2H), 7.14 (t, 1H, $J = 5.6$ Hz), 7.32 (d, 1H, $J = 6.7$ Hz), 7.65 (d, 2H, $J = 8.6$ Hz), 8.75 (s, 1H). ^{13}C NMR ($CDCl_3$, 125 MHz), –1.4 (6C), 12.4, 15.4, 17.5, 17.7, 23.6, 25.3, 27.3, 29.8, 32.9, 36.4, 37.9, 38.1, 38.6, 38.8, 42.2, 46.8, 53.0, 58.7, 64.0, 66.0, 95.6, 111.0, 111.6, 112.2 (2C), 114.2, 122.5, 123.1, 124.0, 126.5, 127.5, 129.2 (2C), 129.9, 139.8, 142.3, 149.2, 149.9, 150.4, 151.8, 154.0, 154.7, 159.3, 161.5, 167.7, 168.3, 172.6, 173.2, 173.8, 197.5. HRMS (TOF) (m/z): $[(M + H + Na)/2]^+$ calcd for $C_{56}H_{77}N_{12}NaO_{10}S_2Si_2$, 610.2376; found, 610.2356.

Teoc-FA-SS-prod(C8) (26c). A solution of **4c** (0.150 g, 0.22 mmol) in a mixture of MeOH : $CHCl_3$ (1 : 1, 4 mL) was treated with HCl 12 N (0.036 mL, 0.44 mmol). After an additional 30 min HCl 12 N (0.030 mL, 0.36 mmol) was added and the reaction mixture stirred for a further 12 hours. The solvent was removed under reduced pressure to afford **4c** as a red glass (0.130 g), which was used in the next step without any further purification. HRMS (TOF) (m/z): $[M + H]^+$ calcd for $C_{30}H_{44}N_5O_3S_2$, 586.2880; found, 586.2856. Compound **26c** was obtained according to GP2 as a dark red glass (0.086 g, 69%). 1H NMR ($CDCl_3$, 500 MHz) 0.00 (s, 9H), 0.04 (s, 9H), 0.96–1.00 (m, 4H), 1.02–1.04 (m, 8H), 1.21–1.24 (m, 4H), 1.53–1.55 (m, 4H), 2.10–2.13 (m, 3H), 2.21 (app br s, 4H), 2.34–2.36 (m, 5H), 2.57–2.58 (m, 2H), 2.68–2.69 (m, 2H), 2.72–2.75 (m, 2H), 3.40–3.52 (m, 4H), 3.92 (m, 3H), 4.17–4.20 (m, 2H), 4.26 (t, $J = 8.2$ Hz, 2H), 4.58 (s, 2H), 4.64–4.68 (m, 1H), 5.59 (br s, 1H), 5.97 (s, 1H), 6.21 (s, 1H), 6.56 (d, $J = 7.5$ Hz, 2H), 6.70 (s, 2H), 6.83 (s, 2H), 7.25–7.26 (m, 2H), 7.35 (d, $J = 6.0$ Hz, 1H), 7.62 (d, $J = 7.5$ Hz, 2H), 8.74 (s, 1H). ^{13}C NMR ($CDCl_3$, 125 MHz), –1.4 (6C), 12.4, 14.9, 17.5, 17.7, 24.3, 25.8, 28.1, 29.1, 29.2, 29.4 ($\times 2$), 32.8, 36.6, 37.9, 38.0, 38.6, 42.8, 46.9, 52.9, 58.7, 64.1, 66.0, 95.7, 110.9, 111.8, 112.2 (2C), 114.0, 122.5, 123.4, 123.8, 126.4, 127.7, 129.2 (2C), 130.0, 139.9, 142.1, 149.3, 149.8, 150.4, 151.8, 154.1, 154.8, 159.8, 161.3, 167.6, 168.5, 168.6, 172.6, 173.2, 174.0, 198.1 (1 carbon atom unaccounted for). HRMS (TOF) (m/z): $[(M + H + Na)/2]^{2+}$ calcd for $C_{60}H_{85}N_{12}NaO_{10}S_2Si_2$, 638.2689; found, 638.2663.



General procedure 3 (GP3)

A solution of (1 eq.) Teoc-FA-LINKERS-prod. (1 eq.) in anhydrous DMSO (0.1 M) was treated with TBAF (1 M solution in THF, 2 eq.) and AcOH (1.25/0.1 mmol). The resulting reaction mixture was stirred at room temperature for 18 h under N₂ atmosphere. EtOAc was added to the reaction mixture until a precipitate formed, which was filtered using Milipore® and washed 3 times with water. The obtained dark red solid was dissolved in DMSO (0.1 M), treated with NaOH (0.2 M solution in MeOH) and stirred at room temperature for 10 min. Water was added and a precipitate was formed. The resulting sodium salt was isolated *via* microfiltration, then washed 3 times with water and dried under vacuum.

FA-DA-prod(C2) (27a). Compound 27a was obtained according to GP3 as dark red solid (0.040 g, 53%). Mp decomp. >230 °C. ¹H NMR (DMSO-d₆, 500 MHz, 80 °C) 1.39 (s, 3H), 1.96–2.22 (m, 7H), 2.68 (s, 3H), 3.00 (s, 4H), 3.90 (s, 3H), 4.32 (br s, 1H), 4.49 (s, 2H), 6.16 (br s, 1H), 6.26 (br s, 1H), 6.67 (br s, 4H), 6.78 (br s, 1H), 7.09 (br s, 1H), 7.64 (br s, 3H), 7.90 (br s, 1H), 8.65 (br s, 1H), 11.61 (br s, 1H). ¹³C NMR (DMSO-d₆, 125 MHz) 11.8, 15.4, 26.8, 28.2, 29.4, 32.0, 36.6, 37.3, 38.4 (2C), 45.9, 52.3, 58.4, 96.3, 110.2, 111.2 (2C), 113.1, 121.4, 122.1, 122.9, 127.6, 127.9, 128.2, 128.9 (2C), 140.9, 142.0, 148.5, 150.7, 153.9, 155.6, 156.3, 159.4, 161.2, 166.3, 167.4, 171.8, 174.0, 195.3 (1 carbon unaccounted for). HRMS (TOF) (*m/z*): [(M – H–Na)/2]^{2–} calcd for C₄₁H₄₂NaN₁₂O₈, 415.1630; found, 415.1648.

FA-DA-prod(C4) (27b). Compound 27b was obtained according to GP3 as dark red solid (0.036 g, 68%). Mp decomp. >200 °C. ¹H NMR (DMSO-d₆, 500 MHz, 60 °C) 1.54 (br s, 4H), 1.85–1.95 (m, 2H), 2.08–2.27 (m, 6H), 2.34 (s, 3H), 2.68 (s, 3H), 2.73 (br s, 2H), 3.06 (br s, 4H), 3.87 (s, 3H), 4.23–4.31 (m, 1H), 4.47 (d, 2H, *J* = 5.6 Hz), 6.19 (s, 1H), 6.24 (br s, 1H), 6.64 (d, 2H, *J* = 8.6 Hz), 6.69 (s, 1H), 6.78–6.79 (m, 1H), 6.87–6.91 (m, 2H), 6.78 (br s, 1H), 7.13 (br s, 1H), 7.65 (d, 2H, *J* = 8.4 Hz), 7.79–7.85 (m, 2H), 8.15 (d, 1H, *J* = 5.4 Hz), 8.63 (s, 1H), 11.81 (br s, 1H). ¹³C NMR (DMSO-d₆, 125 MHz) 11.8, 15.3, 23.4, 24.9, 26.6, 28.2, 32.0, 35.5, 38.3, 38.5, 41.6, 45.9, 52.1, 58.4, 96.3, 110.2, 111.2 (2C), 113.1, 121.4, 122.3, 122.9, 126.3, 127.6, 127.9, 128.2, 128.9 (2C), 140.8, 142.1, 148.5, 150.8, 153.8, 156.3, 159.3, 161.0, 161.1, 166.3, 167.4, 171.7, 172.2, 173.9, 196.3. HRMS (TOF) (*m/z*): [M – H–Na/2]^{2–} calcd for C₄₃H₄₆NaN₁₂O₈, 429.1786; found, 429.1796.

FA-DA-prod(C8) (27c). Compound 27c was obtained according to GP3 as a black solid (0.025 g, 67%). Mp decomp. >205 °C. ¹H NMR (DMSO-d₆, 500 MHz) 1.24–1.27 (m, 8H), 1.45–1.48 (m, 2H), 1.55–1.57 (m, 2H), 1.88–1.93 (m, 1H), 2.01–2.09 (m, 3H), 2.15–2.19 (m, 2H), 2.34 (s, 3H), 2.67 (s, 3H), 2.72 (t, 2H, *J* = 7.0 Hz), 3.00–3.10 (m, 5H), 3.30 (br s, 4H), 3.87 (s, 3H), 4.28–4.30 (m, 1H), 4.48 (d, 2H, *J* = 5.5 Hz), 6.19 (s, 1H), 6.24 (s, 1H), 6.64 (d, 2H, *J* = 8.2 Hz), 6.69 (s, 1H), 6.79 (s, 1H), 6.89–6.91 (m, 2H), 7.14 (s, 1H), 7.65 (d, 2H, *J* = 8.2 Hz), 7.74 (s, 1H), 7.84 (s, 1H), 8.17 (d, 1H, *J* = 7.5 Hz), 8.64 (s, 1H), 11.43 (s, 1H), 11.81 (s, 1H). ¹³C NMR (DMSO-d₆, 125 MHz) 11.7, 15.3, 23.7, 25.1, 27.0, 28.2, 28.7 (×2), 28.9, 32.0, 35.4, 41.8, 45.9, 52.3, 58.4, 77.5, 96.2, 110.1, 111.2 (2C), 113.0, 121.5, 122.4, 122.9, 126.3, 127.5, 127.9, 128.2, 128.8 (2C), 140.7, 142.0, 148.5, 150.7, 154.0, 155.6, 156.2, 159.3, 161.2, 166.1, 167.3, 171.8, 172.3, 173.0, 174.2, 196.6 (1 carbon

unaccounted for). HRMS (TOF) (*m/z*): [(M + H–Na)]⁺ calcd for C₄₇H₅₅NaN₁₂O₈, 915.4271; found, 915.4232.

FA-PEG-prod(C4) (28a). Compound 28a was obtained according to GP3 as a black solid (0.010 g, 41%). Mp 185 °C. ¹H NMR (DMSO-d₆, 500 MHz) 1.57–1.63 (m, 4H), 1.86–1.93 (m, 1H), 2.01–2.07 (m, 1H), 2.17 (t, 2H, *J* = 7.5 Hz), 2.35 (s, 3H), 2.40 (t, 2H, *J* = 6.7 Hz), 2.69 (s, 3H), 2.97 (t, 2H, *J* = 6.7 Hz), 3.04–3.10 (m, 4H), 3.34–3.50 (m, 12H), 3.87 (s, 3H), 4.24–4.28 (m, 1H), 4.48 (d, 2H, *J* = 5.5 Hz), 6.19 (s, 1H), 6.24 (s, 1H), 6.64 (d, 2H, *J* = 8.7 Hz), 6.70 (s, 1H), 6.79 (s, 1H), 6.88–6.90 (m, 2H), 7.13 (s, 1H), 7.64 (d, 2H, *J* = 8.7 Hz), 7.78–7.81 (m, 2H), 8.12 (d, 1H, *J* = 7.0 Hz), 8.64 (s, 1H), 11.82 (br s, 1H). ¹³C NMR (DMSO-d₆, 125 MHz) 11.8, 15.4, 26.6, 29.3, 29.4 (2C), 32.0, 35.8, 37.3 (2C), 45.9, 52.3, 58.4, 68.0 (2C), 68.1, 69.5, 69.7 (4C), 96.3, 110.1, 111.2 (2C), 113.1, 121.4, 122.1, 122.9, 126.3, 127.6, 127.9, 128.2, 128.9 (2C), 140.8, 142.1, 148.5, 150.7, 153.8, 159.4, 166.2, 167.4, 171.5, 173.9, 195.3 (3 carbon atoms unaccounted for). HRMS (TOF) (*m/z*): [(M – H–Na)/2]^{2–} calcd for C₄₉H₅₈NaN₁₂O₁₁, 495.2179; found, 495.2177.

FA-PEG-prod(C4) (28b). Compound 28b was obtained according to GP3 as dark red solid (0.065 g, 79%). Mp decomp. >170 °C. ¹H NMR (DMSO-d₆, 500 MHz) 1.55–1.64 (m, 8H), 1.89–1.93 (m, 1H), 2.02–2.10 (m, 3H), 2.15–2.19 (m, 2H), 2.35 (s, 3H), 2.68 (s, 3H), 2.73 (br s, 2H), 3.02–3.10 (m, 4H), 3.36 (dd, 4H, *J* = 12.6, 6.3 Hz), 3.45–3.48 (m, 8H), 3.87 (s, 3H), 4.23–4.30 (m, 1H), 4.48 (d, 2H, *J* = 5.7 Hz), 6.19 (s, 1H), 6.24 (br s, 1H), 6.64 (d, 2H, *J* = 8.6 Hz), 6.70 (s, 1H), 6.80 (br s, 1H), 6.88–6.91 (m, 2H), 7.13 (s, 1H), 7.65 (d, 2H, *J* = 8.6 Hz), 7.73–7.80 (m, 2H), 8.14 (d, 1H, *J* = 7.4 Hz), 8.64 (s, 1H), 11.81 (br s, 1H). ¹³C NMR (DMSO-d₆, 125 MHz) 11.8, 15.33, 23.5, 25.1, 26.7, 29.3, 29.4 (2C), 32.0, 35.5, 35.7, 35.8 (2C), 41.2, 41.6, 45.9, 58.4, 68.0 (2C), 68.1, 69.5, 69.7 (4C), 96.3, 110.2, 111.2 (2C), 113.1, 121.4, 122.3, 122.9, 126.3, 128.2, 128.9 (2C), 140.8, 142.1, 148.5, 148.6, 150.7, 153.8, 159.4, 161.0, 166.2, 167.4, 171.5, 172.0, 173.9, 196.5. HRMS (TOF) (*m/z*): [M–Na][–] calcd for C₅₁H₆₃NaN₁₂O₁₁, 1019.4745; found, 1019.4718.

FA-PEG-prod(C8) (28c). Compound 28c was obtained according to GP3 as dark red solid (0.021 g, 38%). Mp decomp. >150 °C. ¹H NMR (DMSO-d₆, 500 MHz) 1.25–1.26 (br m, 8H), 1.44–1.50 (m, 2H), 1.53–1.64 (m, 4H), 2.03 (t, 2H, *J* = 7.1 Hz), 2.14–2.28 (m, 4H), 2.34 (s, 3H), 2.67 (s, 3H), 2.72 (t, 2H, *J* = 7.1 Hz), 3.06 (dd, 4H, *J* = 12.9, 6.8 Hz), 3.33–3.39 (m, 4H), 3.43–3.49 (m, 8H), 3.87 (s, 3H), 4.22–4.29 (m, 1H), 4.48 (d, 2H, *J* = 5.7 Hz), 6.19 (s, 1H), 6.64 (d, 2H, *J* = 8.3 Hz), 6.69 (s, 1H), 6.78–6.80 (m, 1H), 6.85–6.92 (m, 3H), 7.13 (br s, 1H), 7.64 (d, 2H, *J* = 8.3 Hz), 7.69–7.72 (m, 1H), 7.77–7.81 (m, 1H), 8.14 (d, 1H, *J* = 7.0 Hz), 8.64 (s, 1H), 11.43 (br s, 1H), 11.81 (br s, 1H). ¹³C NMR (DMSO-d₆, 125 MHz) 11.8, 15.3, 23.8, 25.3, 26.7, 28.6, 28.7, 28.8, 29.0, 29.3, 29.4, 32.0, 35.4, 35.7, 35.8, 41.8, 45.9, 52.3, 58.4, 68.0, 68.1, 69.5 (×2), 69.7 (4C), 96.3, 110.2, 111.2 (2C), 113.1, 121.4, 122.4, 122.9, 126.3, 127.5, 127.9, 128.2, 128.9 (2C), 140.7, 142.1, 148.5, 148.6, 150.7, 153.8, 153.9, 159.4, 166.2, 167.4, 171.5, 172.0, 173.8, 197.7. HRMS (TOF) (*m/z*): [(M – H–Na)/2][–] calcd for C₅₅H₇₀NaN₁₂O₁₁, 537.2649; found, 537.2664.

FA-SS-prod(C4) (29a). Compound 29a was obtained according to GP3 as a dark red solid (0.009 g, 26%). ¹H NMR (DMSO-d₆, 300 MHz, 50 °C) 1.86–1.96 (m, 1H), 2.03–2.17 (m, 1H), 2.19–



2.23 (m, 2H), 2.35 (s, 3H), 2.42 (t, 2H, $J = 6.6$ Hz), 2.68 (s, 3H), 2.72–2.96 (m, 4H), 3.00 (t, 1H, $J = 7.0$ Hz), 3.30–3.36 (m, 5H), 3.89 (s, 3H), 4.29 (br s, 1H), 4.49 (d, 1H, $J = 4.5$ Hz), 6.20 (s, 1H), 6.25 (s, 1H), 6.62–6.70 (m, 3H), 6.89–6.93 (m, 3H), 7.04–7.07 (m, 1H), 7.13 (s, 1H), 7.65 (d, 2H, $J = 7.5$ Hz), 8.06 (t, 1H, $J = 5.7$ Hz), 8.15–8.17 (m, 1H), 8.64 (s, 1H), 11.82 (br s, 1H). ^{13}C NMR (DMSO- d_6 , 125 MHz) 11.8, 15.4, 26.5, 28.2, 29.3, 31.9, 37.2 (2C), 37.3, 37.9, 38.1 (2C), 39.0, 45.9, 52.1, 58.4, 96.3, 110.1 (2C), 111.1, 113.1, 121.3, 122.1, 122.9, 126.3, 127.6, 127.9, 128.1, 128.9 (2C), 140.9, 142.0, 148.6, 150.7, 153.8, 159.3, 166.2, 167.3, 171.7 ($\times 2$), 173.8, 195.1 (1 carbon atom unaccounted for). HRMS (TOF) (m/z): $[(M - H - Na)/2]^{2-}$ calcd for $\text{C}_{43}\text{H}_{46}\text{NaN}_{12}\text{O}_8\text{S}_2$, 461.1507; found, 461.1530.

FA-SS-prod(C4) (29b). Compound **29b** was obtained according to GP3 as dark red solid (0.047 g, 30%). Mp decomp. >210 °C. ^1H NMR (DMSO- d_6 , 500 MHz) 1.55 (br s, 4H), 1.84–1.96 (m, 2H), 2.02–2.12 (m, 2H), 2.16–2.22 (m, 2H), 2.35 (s, 3H), 2.68 (s, 3H), 2.73–2.78 (m, 6H), 3.29–3.32 (m, 4H), 3.87 (s, 3H), 4.23–4.32 (m, 1H), 4.48 (d, 2H, $J = 5.5$ Hz), 6.19 (s, 1H), 6.24 (br s, 1H), 6.66 (d, 2H, $J = 8.6$ Hz), 6.80 (br s, 1H), 6.88–6.93 (m, 2H), 6.99 (br s, 1H), 7.13 (br s, 1H), 7.64 (dd, 2H, $J = 8.6, 2.8$ Hz), 7.96–8.04 (m, 2H), 8.09–8.16 (m, 1H), 8.64 (s, 1H), 11.81 (br s, 1H). ^{13}C NMR (DMSO- d_6 , 125 MHz) 11.8, 15.3, 23.4, 25.0, 26.6, 32.0, 35.4, 37.2 ($\times 2$), 37.3, 37.9, 38.0, 41.6, 45.9, 52.3, 58.4, 96.3, 110.2, 111.2 (2C), 113.1, 121.4, 122.4, 122.9, 126.3, 127.9, 128.2, 128.9 (2C), 140.8, 142.1, 148.5, 150.7, 153.8, 156.3, 159.3, 161.1, 166.0, 166.2, 166.3, 167.3, 171.8, 172.3, 173.9, 196.5. HRMS (TOF) (m/z): $[M - Na]^-$ calcd for $\text{C}_{45}\text{H}_{51}\text{NaN}_{12}\text{O}_8\text{S}_2$, 951.3400; found, 951.3369.

Conflicts of interest

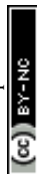
There is no conflict to declare.

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References

- 1 A. Fürstner, *Angew. Chem., Int. Ed.*, 2003, **42**, 3582–3603.
- 2 N. R. Williamson, P. C. Fineran, F. J. Leeper and G. P. Salmond, *Nat. Rev. Microbiol.*, 2006, **4**, 887–899.
- 3 R. K. Suryawanshi, C. D. Patil, H. P. Borase, C. P. Narkhede, A. Stevenson, J. E. Hallsworth and S. V. Patil, *Int. J. Cosmet. Sci.*, 2015, **37**, 98–107.
- 4 Y. Kim and J. Choi, *Fibers Polym.*, 2015, **16**, 1981–1987.
- 5 Y. Ren, J. Gong, R. Fu, Z. Li, Q. Li, J. Zhang, Z. Yu and X. Cheng, *Dyes Pigm.*, 2017, **138**, 147–153.
- 6 A. Fürstner and E. J. Grabowski, *ChemBioChem*, 2001, **2**, 706–709.
- 7 A. Fürstner, K. Reinecke, H. Prinz and H. Waldmann, *ChemBioChem*, 2004, **5**, 1575–1579.
- 8 M. S. Melvin, J. T. Tomlinson, G. Park, C. S. Day, G. R. Saluta, G. L. Kucera and R. A. Manderville, *Chem. Res. Toxicol.*, 2002, **15**, 734–741.
- 9 M. S. Melvin, J. T. Tomlinson, G. R. Saluta, G. L. Kucera, N. Lindquist and R. A. Manderville, *J. Am. Chem. Soc.*, 2000, **122**, 6333–6334.
- 10 B. Montaner, W. Castillo-Avila, M. Martinell, R. Oellinger, J. Aymami, E. Giralt and R. Perez-Tomas, *Toxicol. Sci.*, 2005, **85**, 870–879.
- 11 S. Ohkuma, T. Sato, M. Okamoto, H. Matsuya, K. Arai, T. Kataoka, K. Nagai and H. H. Wasserman, *Biochem. J.*, 1998, **334**, 731–741.
- 12 G. Park, J. T. Tomlinson, M. S. Melvin, M. W. Wright, C. S. Day and R. A. Manderville, *Org. Lett.*, 2003, **5**, 113–116.
- 13 R. I. Sáez Díaz, S. M. Bennett and A. Thompson, *ChemMedChem*, 2009, **4**, 742–745.
- 14 T. Sato, H. Konno, Y. Tanaka, T. Kataoka, K. Nagai, H. H. Wasserman and S. Ohkuma, *J. Biol. Chem.*, 1998, **273**, 21455–21462.
- 15 J. L. Seganish and J. T. Davis, *Chem. Commun.*, 2005, 5781–5783.
- 16 W. R. Hearn, M. K. Elson, R. H. Williams and J. Medina-Castro, *J. Org. Chem.*, 1970, **35**, 142–146.
- 17 E. Marchal, S. Rastogi, A. Thompson and J. T. Davis, *Org. Biomol. Chem.*, 2014, **12**, 7515–7522.
- 18 E. Marchal, D. A. Smithen, I. M. Uddin, A. W. Robertson, D. L. Jakeman, V. Mollard, C. D. Goodman, K. S. MacDougall, S. A. McFarland, G. I. McFadden and A. Thompson, *Org. Biomol. Chem.*, 2014, **12**, 4132–4142.
- 19 E. Marchal, M. I. Uddin, D. A. Smithen, L. A. Hawco, M. Lanteigne, D. P. Overy, R. G. Kerr and A. Thompson, *RSC Adv.*, 2013, **3**, 22967–22971.
- 20 S. Rastogi, E. Marchal, I. Uddin, B. Groves, J. Colpitts, S. A. McFarland, J. T. Davis and A. Thompson, *Org. Biomol. Chem.*, 2013, **11**, 3834–3845.
- 21 J. Regourd, A. Al-Sheikh Ali and A. Thompson, *J. Med. Chem.*, 2007, **50**, 1528–1536.
- 22 R. I. Sáez Díaz, J. Regourd, P. V. Santacroce, J. T. Davis, D. L. Jakeman and A. Thompson, *Chem. Commun.*, 2007, 2701–2703.
- 23 D. A. Smithen, A. M. Forrester, D. P. Corkery, G. Dellaire, J. Colpitts, S. A. McFarland, J. N. Berman and A. Thompson, *Org. Biomol. Chem.*, 2013, **11**, 62–68.
- 24 M. I. Uddin, S. Thirumalairajan, S. M. Crawford, T. S. Cameron and A. Thompson, *Synlett*, 2010, 2561–2564.
- 25 C. Ferrario and G. Batist, *Expert Opin. Drug Discovery*, 2014, **9**, 647–668.
- 26 A. Cheung, H. J. Bax, D. H. Josephs, K. M. Ilieva, G. Pellizzari, J. Opzoomer, J. Bloomfield, M. Fittall, A. Grigoriadis, M. Figini, S. Canevari, J. F. Spicer, A. N. Tutt and S. N. Karagiannis, *Oncotarget*, 2016, **7**, 52553–52574.



- 27 R. Paulmurugan, R. Bhethanabotla, K. Mishra, R. Devulapally, K. Foygel, T. Sekar, J. Ananta, T. Massoud and A. Joy, *Mol. Cancer Ther.*, 2016, **15**, 221–231.
- 28 Y. Lu and P. S. Low, *Adv. Drug Delivery Rev.*, 2002, **54**, 675–693.
- 29 J. Sudimack and R. J. Lee, *Adv. Drug Delivery Rev.*, 2000, **41**, 147–162.
- 30 F. Zagouri, M. A. Dimopoulos, E. Bournakis and C. A. Papadimitriou, *Eur. J. Gynaecol. Oncol.*, 2010, **31**, 268–277.
- 31 Y.-S. Yi, *Immune Netw.*, 2016, **16**, 337–343.
- 32 W. Xia, A. R. Hilgenbrink, E. L. Matteson, M. B. Lockwood, J. X. Cheng and P. S. Low, *Blood*, 2009, **113**, 438–446.
- 33 L. H. Matherly, Z. Hou and Y. Deng, *Cancer Metastasis Rev.*, 2007, **26**, 111–128.
- 34 J. M. Scott and D. G. Weir, *J. Cardiovasc. Risk*, 1998, **5**, 223–227.
- 35 R. Zhao, S. H. Min, Y. Wang, E. Campanella, P. S. Low and I. D. Goldman, *J. Biol. Chem.*, 2009, **284**, 4267–4274.
- 36 P. S. Low and S. A. SuKularatne, *Curr. Opin. Chem. Biol.*, 2009, **13**, 256–262.
- 37 S. Wang and P. S. Low, *J. Controlled Release*, 1998, **53**, 39–48.
- 38 C. L. A. Hawco, E. Marchal, M. I. Uddin, A. E. G. Baker, D. P. Corkery, G. Dellaire and A. Thompson, *Bioorg. Med. Chem.*, 2013, **21**, 5995–6002.
- 39 G. Huang, D. Pemp, P. Stadtmüller, M. Nimczick, J. Heilmann and M. Decker, *Bioorg. Med. Chem. Lett.*, 2014, **24**, 4209–4214.
- 40 S. Dhar, Z. Liu, J. Thomale, H. Dai and S. J. Lippard, *J. Am. Chem. Soc.*, 2008, **130**, 11467–11476.
- 41 S. Zalipsky, *Adv. Drug Delivery Rev.*, 1995, **16**, 157–182.
- 42 M. P. Grillo, *Expert Opin. Drug Metab. Toxicol.*, 2015, **11**, 1281–1302.
- 43 H. Joyce, A. McCann, M. Clynes and A. Larkin, *Expert Opin. Drug Metab. Toxicol.*, 2015, **11**, 795–809.
- 44 A. Sullivan, A. Gibson, B. K. Park and D. J. Naisbitt, *Expert Opin. Drug Metab. Toxicol.*, 2015, **11**, 357–368.
- 45 R. S. Shirazi, K. K. Ewert, C. Leal, R. N. Majzoub, N. F. Bouxsein and C. R. Safinya, *Biochim. Biophys. Acta, Biomembr.*, 2011, **1808**, 2156–2166.
- 46 I. R. Vlahov and C. P. Leamon, *Bioconjugate Chem.*, 2012, **23**, 1357–1369.
- 47 C. Chen, J. Ke, X. E. Zhou, W. Yi, J. S. Brunzelle, J. Li, E.-L. Yong, H. E. Xu and K. Melcher, *Nature*, 2013, **500**, 486–489.
- 48 M. P. Carrasco, E. A. Enyedy, N. I. Krupenko, S. A. Krupenko, E. Nuti, T. Tuccinardi, S. Santamaria, A. Rossello, A. Martinelli and M. A. Santos, *Med. Chem.*, 2011, **7**, 265–274.
- 49 S. K. Choi, T. Thomas, M.-H. Li, A. Kotlyar, A. Desai and J. R. Baker Jr, *Chem. Commun.*, 2010, **46**, 2632–2634.
- 50 A. Galbiati, C. Tabolacci, B. M. D. Rocca, P. Mattioli, S. Beninati, G. Paradossi and A. Desideri, *Bioconjugate Chem.*, 2011, **22**, 1066–1072.
- 51 L. Jiang, Z.-M. Gao, L. Ye, A.-Y. Zhang and Z.-G. Feng, *Polymer*, 2013, **54**, 5188–5198.
- 52 H. Jing, Z. Guo, W. Guo, W. Yang, P. Xu and X. Zhang, *Bioorg. Med. Chem. Lett.*, 2012, **22**, 3418–3424.
- 53 C. Lainé, E. Mornet, L. Lemiègre, T. Montier, S. Cammas-Marion, C. Neveu, N. Carmoy, P. Lehn and T. Benvegna, *Chemistry*, 2008, **14**, 8330–8340.
- 54 P. Li, Y. Wang, F. Zeng, L. Chen, Z. Peng and L. X. Kong, *Carbohydr. Res.*, 2011, **346**, 801–806.
- 55 M. A. Santos, E. A. Enyedy, E. Nuti, A. Rossello, N. I. Krupenko and S. A. Krupenko, *Bioorg. Med. Chem.*, 2007, **15**, 1266–1274.
- 56 R. K. Singh, D. Rai, D. Yadav, A. Bhargava, J. Balzarini and E. De Clercq, *Eur. J. Med. Chem.*, 2010, **45**, 1078–1086.
- 57 A. F. Trindade, R. F. M. Frade, E. M. S. Macoas, C. Graca, C. A. B. Rodrigues, J. M. G. Martinho and C. A. M. Afonso, *Org. Biomol. Chem.*, 2014, **12**, 3181–3190.
- 58 P. M. Valencia, M. H. Hanewich-Hollatz, W. Gao, F. Karim, R. Langer, R. Karnik and O. C. Farokhzad, *Biomaterials*, 2011, **32**, 6226–6233.
- 59 Y. Yang, Y. M. Zhang, Y. Chen, D. Zhao, J. T. Chen and Y. Liu, *Chemistry*, 2012, **18**, 4208–4215.
- 60 H. Zhang, Z. Cai, Y. Sun, F. Yu, Y. Chen and B. Sun, *J. Biomed. Mater. Res., Part A*, 2012, **100**, 2441–2449.
- 61 A. Bettio, M. Honer, C. Müller, M. Brühlmeier, U. Müller, R. Schibli, V. Groehn, A. P. Schubiger and S. M. Ametamey, *J. Nucl. Med.*, 2006, **47**, 1153–1160.
- 62 I. R. Vlahov, H. K. Santhapuram, P. J. Kleindl, S. J. Howard, K. Stanford and C. P. Leamon, *Bioorg. Med. Chem. Lett.*, 2006, **16**, 5093–5096.
- 63 W. A. Henne, S. A. Kularatne, J. Hakenjos, J. D. Carron and K. L. Henne, *Bioorg. Med. Chem. Lett.*, 2013, **23**, 5810–5813.
- 64 T. Suzuki, S. Hisakawa, Y. Itoh, N. Suzuki, K. Takahashi, M. Kawahata, K. Yamaguchi, H. Nakagawa and N. Miyata, *Bioorg. Med. Chem. Lett.*, 2007, **17**, 4208–4212.
- 65 M. Nomura, S. Shuto and A. Matsuda, *J. Org. Chem.*, 2000, **65**, 5016–5021.
- 66 C. P. Leamon, M. A. Parker, I. R. Vlahov, L. C. Xu, J. A. Reddy, M. Vetzal and N. Douglas, *Bioconjugate Chem.*, 2002, **13**, 1200–1210.
- 67 J. D. Seitz, J. G. Vineberg, E. Herlihy, B. Park, E. Melief and I. Ojima, *Bioorg. Med. Chem.*, 2015, **23**, 2187–2194.
- 68 E. Vlashi, L. E. Kelderhouse, J. E. Sturgis and P. S. Low, *ACS Nano*, 2013, **7**, 8573–8582.
- 69 J. Luo, M. D. Smith, D. A. Lantrip, S. Wang and P. L. Fuchs, *J. Am. Chem. Soc.*, 1997, **119**, 10004–10013.

