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## Introduction

Peptides and proteins occupy a central role as the functional biomolecules that drive most processes in living systems. Following ribosomal synthesis, additional molecular complexity is added to peptides and proteins through post-translational modifications (PTMs) that serve to expand and diversify the functionality possible from the 20 proteinogenic amino acids.<sup>1,2</sup> These PTMs range from the alkylations and acylations that drive gene expression, to the addition of large glycan and lipid chains – many of which underpin crucial protein–protein interactions and therefore endow additional functionality to their cognate proteins.<sup>2–4</sup> Furthermore, a large number of therapeutics known as ‘biologics’,<sup>5–9</sup> including antibody–drug conjugates (ADCs), rely on the functionalisation of peptides and proteins with distinct entities.<sup>10</sup> The importance of protein modifications in biology and the pharmaceutical/biotechnological industry has led to the search for new methods that can be used for efficient access to modified peptides and proteins in homogeneous form (either with PTMs or unnatural ‘designer’ modifications, *e.g.* PEGylation)<sup>11</sup> from fully unprotected peptides and proteins.

To date, a number of so called ‘bioconjugation’ methods have been developed that enable site-selective functionalisation

of peptides and proteins.<sup>12</sup> Necessitating high degrees of chemo- and regio-selectivity in order to generate homogeneous protein conjugates, the development of such methods has focussed on two overarching approaches: (1) the incorporation of non-proteinogenic amino acids or engineered peptide sequences which take advantage of unique chemical (bio-orthogonal)<sup>13</sup> or chemoenzymatic reactivity,<sup>14</sup> and (2) the utilisation of selective chemical reactions at proteinogenic amino acids.<sup>15</sup> While both approaches have been successfully executed for the synthesis of peptide and protein conjugates, targeting native amino acids for late-stage modification provides a more operationally simple approach. However, this remains a challenging endeavour given that the reaction must be selective in the presence of a plethora of functionality that decorate the side chains of proteins, and reaction conditions must be mild and biocompatible (*i.e.* aqueous solvent, pH 6–8, temperatures  $\leq 37\text{ }^{\circ}\text{C}$ ).

Cysteine (Cys) represents an attractive amino acid for the development of functionalisation chemistry due to the unique nucleophilicity of its thiol side chain. This provides the potential for chemoselectivity,<sup>16</sup> while the relatively low natural abundance of Cys in proteins (*ca.* 1.8%)<sup>17</sup> provides an avenue for enhanced regioselectivity.<sup>18</sup> Maleimide reagents are frequently employed to covalently modify Cys residues both in academic and industrial settings. Notably this chemistry is employed for the functionalisation of monoclonal antibodies with cytotoxic payloads (*e.g.* brentuximab vedotin, also known as Adcetris®).<sup>10</sup> However, the resulting conjugates can suffer from poor stability due to retro-Michael reactivity following conjugation.<sup>19</sup>

This drawback has prompted the exploration of alternative chemistry that could be used to generate stable Cys conjugates chemoselectively. While several elegant Cys conjugation methods have been reported recently, including a number of

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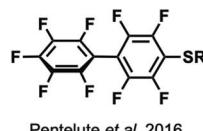
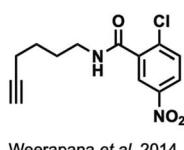
<sup>c</sup>School of Life and Environmental Sciences, The University of Sydney, Sydney, NSW 2006, Australia

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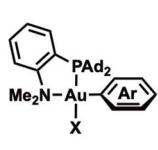
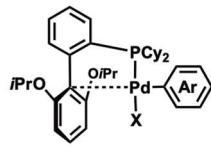
† Electronic supplementary information (ESI) available. See DOI: 10.1039/d1sc03127a



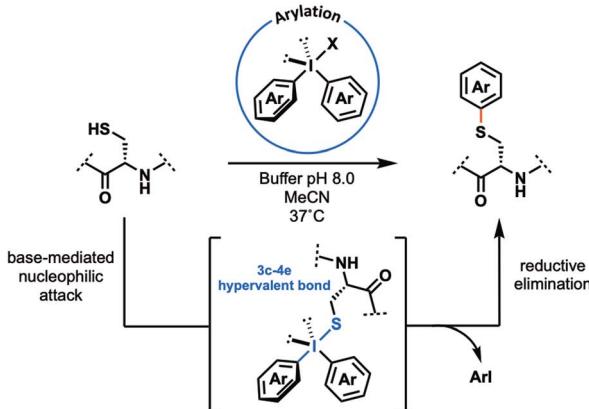
## (a) SnAr transformations



## (b) Oxidative addition complexes



## (c) This work: diaryliodonium salt mediated arylation at cysteine

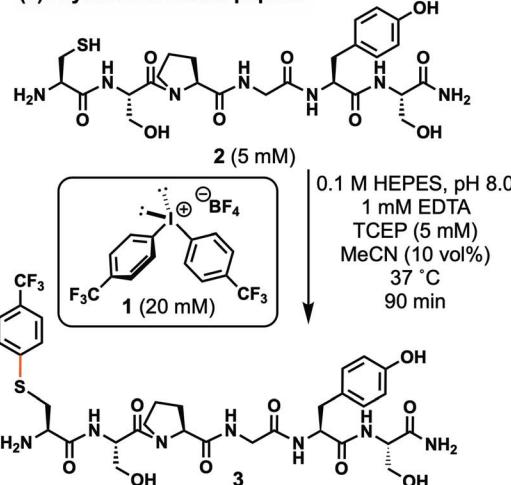


**Scheme 1** Previously reported (a) nucleophilic aromatic substitution (SnAr) and (b) oxidative addition of transition metal complexes for Cys arylation on peptides and proteins; R = functional cargo; X = halide or pseudohalide; (c) proposed use of diaryliodonium salts for direct arylation of Cys to generate thioether-linked peptide and protein conjugates described in this work.

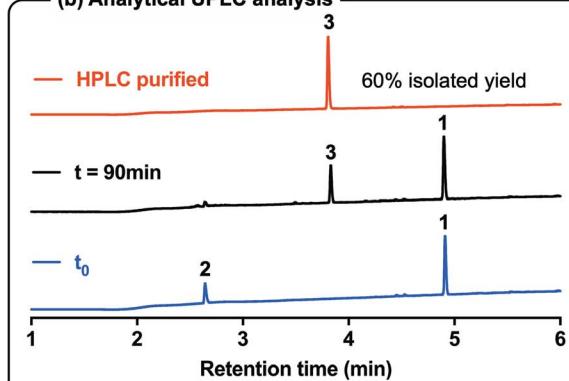
novel Cys arylation methodologies (Scheme 1a and b),<sup>20</sup> these approaches still suffer from modest chemoselectivity,<sup>21</sup> the need for highly specific peptide sequences,<sup>22</sup> or potential problems with biocompatibility owing to the use of transition metal reagents.<sup>23,24</sup>

Hypervalent iodine chemistry has recently emerged as an attractive means to modify peptides and proteins.<sup>25–30</sup> A report from Kirkwood *et al.* in 1947 that diphenyliodonium chloride reacts with Cys in boiling water to produce phenylcysteine<sup>31</sup> provided an encouraging proof of principle that this class of reagents could be used for direct functionalisation of peptides and proteins through the thiol side chain of Cys residues. Furthermore the air- and moisture-stability of hypervalent iodine reagents,<sup>32</sup> coupled with their simple access from cheap and readily available aryl iodide and aryl boronate precursors<sup>33</sup> led us to propose their application in the development of a platform for the chemoselective arylation of peptides and proteins at Cys (Scheme 1c). The proposed mechanism of arylation involves initial nucleophilic attack of the electrophilic iodine(III) centre by the thiolate of Cys to afford a 3-centre 4-electron hypervalent bond, followed by reductive elimination to

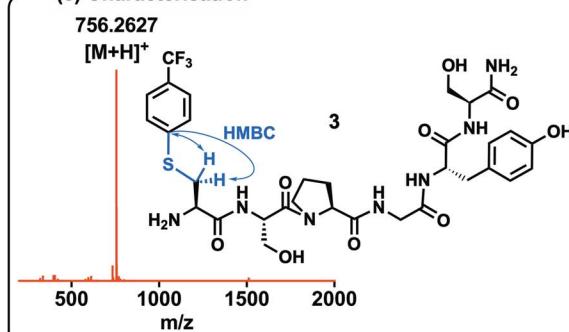
## (a) Arylation of model peptide



## (b) Analytical UPLC analysis



## (c) Characterisation



**Scheme 2** (a) Arylation of model peptide 2 at Cys with 1 to generate aryl conjugate 3; (b) monitoring of arylation reaction by analytical UPLC; (c) HR-ESI-MS of conjugate 3 and confirmation of Cys chemoselectivity through HMBC.

release an aryl iodide and generate the target thioether linkage (Scheme 1c).<sup>34</sup> Herein, we report the development of diaryliodonium salts as reagents for the late-stage modification of peptides and proteins at Cys. We demonstrate the power of the methodology *via* the chemoselective functionalisation of the mucin 1 variable number tandem repeat (MUC1 VNTR), an affibody to the epidermal growth factor receptor Ac-Cys-Z<sub>EGFR:1907</sub> (ZEGFR),<sup>35</sup> and the protein histone 2A (H2A).<sup>36</sup>



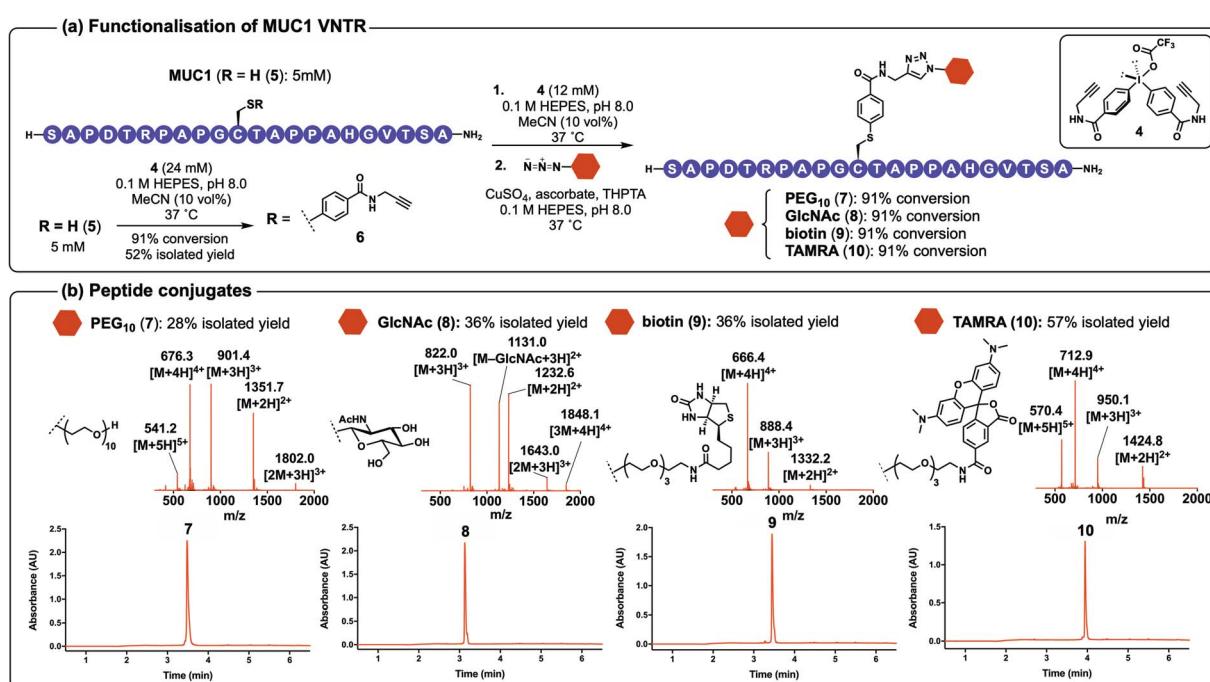
## Results and discussion

The development of the direct Cys arylation chemistry began with the synthesis of a suitable diaryliodonium salt (**1**) that could be used to optimise the proposed chemistry on a model peptide system (Scheme 2). To this end, we chose to adopt a one-pot procedure reported by Olofsson *et al.*<sup>37</sup> that uses readily available aryl iodide and aryl boronic acid precursors to synthesise the bis(4-trifluoromethylphenyl)iodonium tetrafluoroborate salt **1** (Scheme 2, see ESI† for synthetic details). With the bis(4-trifluoromethylphenyl)iodonium reagent **1** in hand we next investigated conditions for the proposed direct arylation chemistry on an unprotected model peptide bearing a Cys residue, namely H-CSPGYS-NH<sub>2</sub> (**2**) that was prepared by Fmoc-strategy solid-phase peptide synthesis (SPPS) (Scheme 2a, see ESI† for synthetic details). Several reaction conditions and buffers were systematically screened on this model system (see ESI†). Optimal reaction conditions that emerged involved the use of aqueous 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid (HEPES) buffer at pH 8.0, tris(2-carboxyethyl)phosphine (TCEP) to prevent cystine formation, with the addition of 10 vol% acetonitrile (MeCN) as co-solvent, which was necessary to dissolve the diaryliodonium salt **1** (Scheme 2a). Importantly, the reactions did not need to be conducted under an inert atmosphere and proceeded to 90% conversion in 90 min when performed open to air [as judged by analytical ultra-performance liquid-chromatography (UPLC) analysis] (Scheme 2b). The target arylation product **3** was isolated in 60% yield following high-performance liquid-chromatography (HPLC) purification and was characterised by nuclear

magnetic resonance (NMR) spectroscopy and high-resolution electrospray ionisation mass-spectrometry (HR-ESI-MS) (Scheme 2c, see ESI† for details). Importantly, the chemoselectivity for the side chain of Cys was confirmed by a heteronuclear multiple bond correlation (HMBC) experiment revealing a correlation between the aromatic *ipso*-carbon and the methylene protons of Cys (Scheme 2c, see ESI†). Following the success of this initial model system, we explored the effect of aryl substituents on reactivity with Cys (see ESI† for details). Consistent with the previously established reactivity reported for this class of reagents,<sup>34,38</sup> we observed a decrease in the rate of arylation and overall efficiency of the reaction when moving from an electron withdrawing aryl system to more electron rich systems (e.g. *p*-methoxyphenyl, see ESI†).<sup>39</sup>

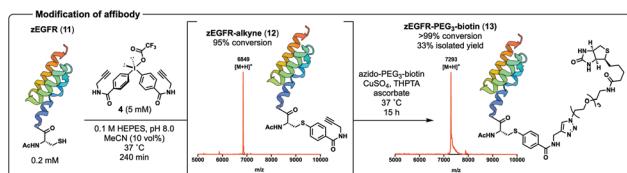
Having demonstrated the direct arylation of a simple model system, we next sought to synthesise a diaryliodonium salt equipped with an alkyne handle that would enable late-stage Cu(i)-catalysed azide–alkyne cycloaddition (CuAAC)<sup>40,41</sup> chemistry on peptide and protein substrates, taking advantage of the variety of azides that can be obtained commercially or through simple synthetic procedures. Attempts to prepare the tetrafluoroborate analogue of compound **4** (Scheme 3a) *via* the same procedure used to prepare **1** provided the product in extremely low yield and purity. Therefore, compound **4** was synthesised *via* an alternative route with a modified version of a protocol reported by DiMagno *et al.* (see ESI† for synthetic details).<sup>42</sup>

With compound **4** in hand we aimed to apply our methodology to a larger peptide substrate containing 11 of the 20 proteinogenic amino acids, namely, MUC1 VNTR (**5**).<sup>43</sup> The peptide was synthesised by Fmoc-strategy SPPS with a Cys



**Scheme 3** (a) Arylation of peptide MUC1 VNTR **5** with diaryliodonium salt **4** to afford **6** and subsequent functionalisation by CuAAC in one-pot to furnish **7**, **8**, **9**, and **10**; (b) characterisation of MUC1 VNTR conjugates **7** (PEG<sub>10</sub>), **8** (GlcNAc), **9** (biotin), and **10** (TAMRA) by ESI-MS and analytical UPLC.





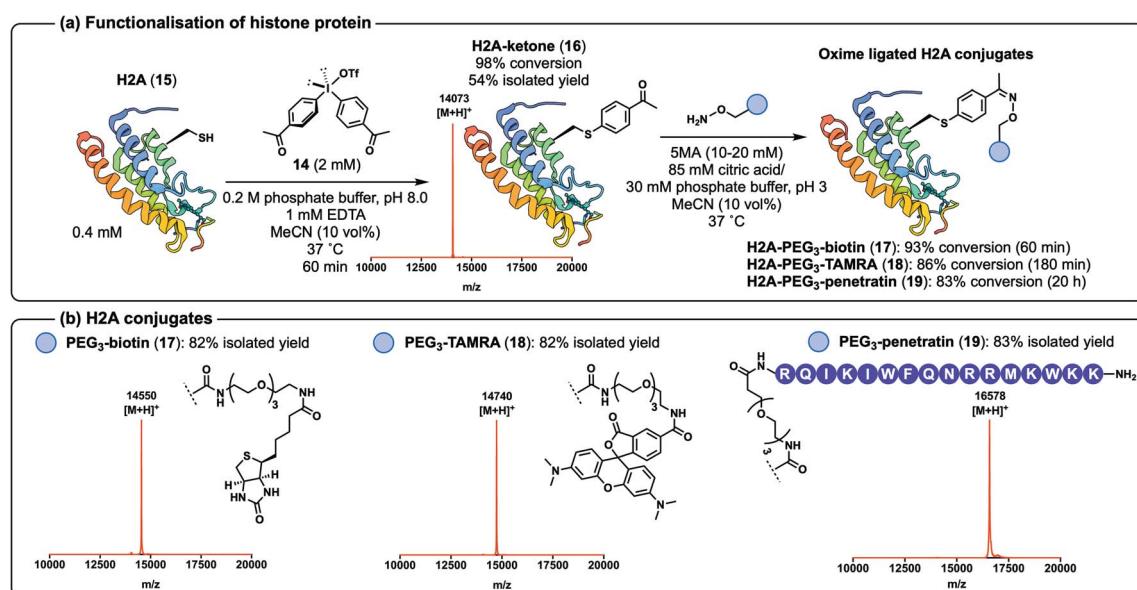
**Scheme 4** Arylation of zEGFR affibody **11** with **4** to afford the intermediate **12**, which was converted to biotin conjugate **13** in one-pot using the CuAAC reaction. Structure of zEGFR from PDB (ID: 2KZI).

residue installed internally (in place of a serine) to eliminate steric bias at position 11 (Scheme 3a, see ESI† for synthetic details). Pleasingly, the arylation of **5** with diaryliodonium salt **4** under the optimised conditions proceeded to 91% conversion within 120 min (as indicated by UPLC-MS analysis), and the product MUC1-alkyne (**6**) peptide conjugate could be isolated in 52% yield following HPLC purification (see ESI† for synthetic details). Importantly, the thioether linkage in **6** was stable for 24 h in plasma, and phosphate buffered saline containing 100 mM glutathione, dithiothreitol or TCEP (see ESI† for details). Diaryliodonium reagent **4** was itself stable to storage in solid form at room temperature for 18 months.

Alkyne-containing peptide **6** was subsequently reacted with azides bearing a PEG<sub>10</sub>, *N*-acetyl glucosamine (GlcNAc), biotin or fluorescent TAMRA functionality under CuAAC conditions<sup>40,41</sup> to afford peptide conjugates **7–10** in good yields over the two-step process (see ESI† for details). However, arylation of peptide **5** with **4** and CuAAC could also be performed in a one-pot manner without any purification of the MUC1-alkyne intermediate **6** (Scheme 3a). This one-pot arylation-CuAAC chemistry on **5** was used to generate four peptide conjugates bearing PEG<sub>10</sub> (**7**), GlcNAc (**8**), biotin (**9**) and TAMRA (**10**) functionalities. Each of the reactions proceeded smoothly to

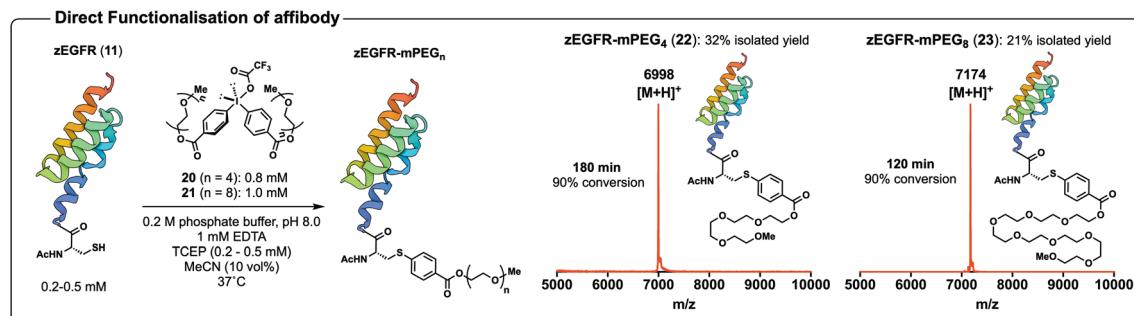
generate the conjugates with 91% conversion over the two steps (as judged by UPLC-MS analysis). These conjugates were then purified by HPLC to afford **7–10** in 28–57% isolated yield (Scheme 3b). We note that the modest yields in some cases were a result of poor recoveries from HPLC rather than inefficient reactions. The peptide conjugates were characterised by electrospray ionisation mass-spectrometry (ESI-MS) and analytical UPLC.

Having demonstrated the efficiency of the arylation chemistry on the MUC1 VNTR peptide, we next envisaged applying our novel protein conjugation platform to a more challenging system, namely the affibody zEGFR (**11**) equipped with an N-terminal Cys residue (Scheme 4, see ESI† for synthetic details). Pleasingly, arylation with diaryliodonium salt **4** proceeded to 95% conversion in 240 min (as judged by UPLC-MS analysis). In this case TCEP was necessary to prevent the formation of unreactive cystine during the reaction. The protein conjugate **12** was isolated in 45% yield following HPLC purification and characterised by ESI-MS, analytical UPLC, and matrix-assisted laser desorption ionisation-time of flight mass-spectrometry (MALDI-ToF MS) (Scheme 4). Importantly, the Cys-selectivity of the arylation reaction was confirmed by LC-MS/MS analysis of **12** following trypsin digestion (see ESI† for details). As with the MUC1 VNTR, we found that the arylation chemistry and CuAAC could be performed in one-pot without purification of intermediate **12**. Specifically, arylation followed by CuAAC with azido-PEG<sub>3</sub>-biotin in the same pot furnished the affibody conjugate **13** (94% conversion over the two steps as judged by UPLC-MS), which was purified by HPLC and characterised by ESI-MS, analytical UPLC, and MALDI-ToF MS (Scheme 4, see ESI† for synthetic details). Circular dichroism (CD) spectroscopy of functionalised zEGFR affibodies **11–13** showed that the characteristic  $\alpha$ -helical secondary structure of the proteins was maintained following conjugation (see ESI† for details).



**Scheme 5** (a) Arylation of histone protein **15** to afford **16** for subsequent oxime-ligation reactions to furnish **17**, **18**, and **19**; (b) characterisation of the oxime ligated H2A conjugates **17**, **18**, and **19** by MALDI-ToF MS. Structure of H2A from PDB (ID: 4QYL).



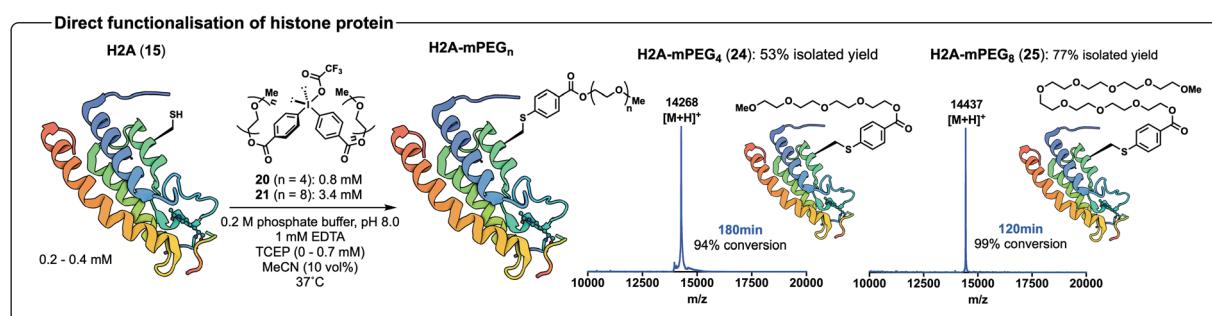


**Scheme 6** Arylation of affibody **11** with PEGylated diaryliodonium salts **20** and **21** to yield PEGylated proteins **22** and **23**, respectively; MALDI-ToF MS spectra of purified **22** and **23**. Structure of zEGFR from PDB (ID: 2KZI).

Having demonstrated that alkyne-derived diaryliodonium salts can be used for the efficient functionalisation of peptides and proteins *via* CuAAC chemistry, we next sought to generate a ketone-containing diaryliodonium salt to access single-site-functionalised proteins primed for diversification with an alternative reaction manifold, namely the oxime-ligation.<sup>44</sup> To this end, we synthesised diaryliodonium salt **14** (ref. 45) (see ESI†) and moved to assess the efficiency of oxime ligation chemistry on a more complex protein system, namely the histone H2A, one of the four proteins found in the octameric complex that make nucleosomes (Scheme 5a).<sup>46</sup> The 129 residue and 14 kDa protein was expressed as the T120C mutant **15** by site-directed mutagenesis in *Escherichia coli* and purified by HPLC (see ESI† for details).<sup>36</sup> It should be noted that the aqueous buffer used in these reactions was changed from HEPES to phosphate due to side-product formation when arylation of **15** was performed in HEPES (see ESI† for details). Gratifyingly, treatment of H2A **15** with diaryliodonium salt **14** in phosphate buffer led to the generation of the desired arylated conjugate **16** in just 60 min with 98% conversion. Purification by HPLC afforded **16** in 54% isolated yield (Scheme 5a, see ESI† for ESI-MS and analytical HPLC data). We also confirmed the Cys-selectivity of the arylation reaction by LC-MS/MS analysis of trypsin-digested **16** (see ESI† for details). The ketone-containing protein conjugate **16** was successfully functionalised with a variety of hydroxylamines using 5-methoxyanthranilic acid (5MA) as a nucleophilic catalyst<sup>47,48</sup> (Scheme 5a). The resulting histone protein conjugates, functionalised with biotin (**17**),

TAMRA (**18**), and the cell-penetrating peptide ‘penetratin’ (**19**),<sup>49</sup> were each generated cleanly following oxime ligation in 1–20 h (83–93% conversions). Each of the constructs was purified by HPLC to afford the oxime conjugates **17–19** in 82–83% isolated yields (Scheme 5b, see ESI† for characterisation data).

Having thus far used our arylation chemistry to install a reactive handle for subsequent functionalisation, we next sought to employ the method to directly functionalise proteins (Scheme 6). PEGylation of proteins is known to improve their efficacy and half-life, and we therefore selected this system to test direct functionalisation chemistry.<sup>11</sup> Towards this end, we first synthesised diaryliodonium salts equipped with mPEG<sub>4</sub> (**20**) and mPEG<sub>8</sub> (**21**) chains (Scheme 6, see ESI† for synthetic details). With the PEGylated diaryliodonium salts in hand we next investigated their efficiency in the late-stage modification of the affibody **11** (Scheme 6). Arylation of the affibody **11** with **20** proceeded smoothly to 90% conversion in 180 min in phosphate buffer. The PEGylated protein was subsequently purified by HPLC to afford **22** in 32% yield (Scheme 6, see ESI† for details). Likewise, we were delighted to find that treatment of **11** with **21** in phosphate buffer delivered the protein conjugate **23** with 90% conversion in 120 min and could be isolated by HPLC in 21% yield (Scheme 6, see ESI† for details). The functionalised affibodies were characterised by ESI-MS, MALDI-ToF MS, analytical UPLC, and CD spectroscopy (see ESI† for data). The Cys-selectivity of the arylation reactions was verified by LC-MS/MS analysis of the trypsin digested proteins (see ESI† for details).



**Scheme 7** Arylation of histone protein **15** with **20** and **21** to yield protein conjugates **24** and **25**, respectively; accompanied by MALDI-ToF MS spectra of crude **24** and **25** before HPLC purification. Structure of H2A from PDB (ID: 4QYL).



Having validated our direct arylation chemistry on the affibody **11** we next moved to the histone protein **15**. Gratifyingly, arylation of **15** with **20** progressed to 94% conversion in 180 min and purification by HPLC provided **24** in 53% isolated yield (Scheme 7, see ESI† for ESI-MS and analytical UPLC data). Likewise, we were pleased to find that treatment of the protein **15** with the diaryliodonium salt **21** led to 99% conversion to **25** in 120 min as judged by MALDI-ToF MS analysis, and subsequent HPLC purification provided the PEGylated protein **25** in 77% isolated yield (Scheme 7, see ESI† for ESI-MS and analytical UPLC data). The Cys-selectivity of each of the arylation reactions were unequivocally demonstrated by tryptic digestion and LC-MS/MS analysis of the proteolysed fragments (see ESI†).

## Conclusions

In summary, we have shown that diaryliodonium salts are both mild and robust Cys selective reagents for the late-stage synthesis of a variety of stable peptide and protein conjugates. The power of this chemistry was highlighted through the installation of a variety of different functionalities in aqueous buffer that is compatible with proteins. The chemoselectivity of the methodology was also exemplified through the functionalisation of the affibody zEGFR and protein H2A that possessed each of the 20 proteinogenic amino acids. Given the overall simplicity and efficiency of the chemistry, we anticipate that the Cys functionalisation methodology reported here will find widespread application in the synthesis of high value peptide and protein conjugates in the future.

## Data availability

All experimental procedures, analytical, and spectroscopic data are provided in the ESI.† The mass spectrometry proteomics data have been deposited to the ProteomeXchange Consortium *via* the PRIDE partner repository with the dataset identifier PXD028382, username: reviewer\_pxd028382@ebi.ac.uk, password: 3UCckTu3.

## Author contributions

SB and RJP designed the project, SB and LC synthesised the diaryliodonium salts, SB synthesised the arylated model peptide, SB synthesised the MUC1 VNTR conjugates, SB synthesised the zEGFR conjugates, SB and DF synthesised the H2A conjugates, MJB synthesised azido-PEG<sub>3</sub>-biotin and azido-PEG<sub>3</sub>-TAMRA, MJB synthesised zEGFR, LC conducted the NMR analysis on the arylated model peptide, LC and DJF synthesised the alkoxyamines, YZ and JPM produced the H2A protein, CF conducted the CD spectroscopy, ML conducted LC-MS/MS experiments and data analysis, SB and RJP prepared the manuscript with assistance from all authors. RJP obtained funding for the project.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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