



Cite this: *Chem. Commun.*, 2015, 51, 5279

Received 28th October 2014,  
Accepted 12th November 2014

DOI: 10.1039/c4cc08515a

www.rsc.org/chemcomm

# A mild TCEP-based *para*-azidobenzyl cleavage strategy to transform reversible cysteine thiol labelling reagents into irreversible conjugates†

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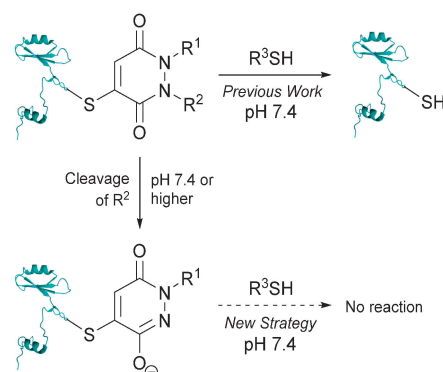
**It has recently emerged that the succinimide linkage of a maleimide thiol addition product is fragile, which is a major issue in fields where thiol functionalisation needs to be robust. Herein we deliver a strategy that generates selective cysteine thiol labelling reagents, which are stable to hydrolysis and thiol exchange.**

Advances in protein modification by chemical means have led to the development of a range of protein bioconjugation methodologies.<sup>1</sup> These methodologies have been successfully applied to a number of fields such as the fluorescent tagging of proteins,<sup>2</sup> and the development of therapeutic protein conjugates<sup>3,4</sup> to treat indications such as HIV,<sup>5</sup> cancer,<sup>6</sup> and malaria.<sup>7</sup> Chemically modified proteins are also utilised as diagnostics.<sup>8</sup>

The use of synthetic methodology to modify proteins has to overcome many major obstacles, the most significant of which is the need for high selectivity, *i.e.* modifying only one amino acid type by discriminating against the other natural amino acids in a protein.<sup>9</sup> As free cysteines are extremely rare in proteins<sup>10</sup> and the thiol side chain has the highest nucleophilicity of all proteinogenic groups at physiological conditions,<sup>11</sup> it is a very popular target for the selective and site-specific modification of proteins.<sup>12</sup> Moreover, with the possibility of facile cysteine introduction by site-directed mutagenesis, cysteine modification is a leading approach. The most popular strategy for labelling the thiol moiety of cysteine residues is by alkylation with maleimides to form thioether-succinimides.<sup>12,13</sup> However, it has recently come to light that such an appendage is sub-optimal owing to issues of hydrolysis, and thiol exchange with reactive thiols in the blood (*e.g.* albumin).<sup>14</sup> This has major implications for biologics that employ a maleimide motif to functionalise a protein thiol for *in vivo* applications. For example, in antibody-drug conjugates (ADCs), where an antibody delivers a toxic payload to cancerous tissue selectively, the use of maleimides to attach cytotoxic

drugs to an antibody is not ideal as thiol exchange onto human serum albumin in the blood results in off-site toxicity.<sup>14</sup> Although recent advances have been made in this area through the use of hydrolysed maleimides and succinimides,<sup>14,15</sup> a strong drive to develop novel reagents for reliable, chemoselective, stable and irreversible thiol labelling remains, and particularly for the construction of ADCs.<sup>16</sup>

Recently, we have described a novel, reversible approach to cysteine bioconjugation through the use of bromomaleimides and bromopyridazinediones.<sup>17</sup> To date, our approach has provided access to complex bioconjugates in high yields, without prior activation of reagents with reliable, reversible conjugation. Owing to the demand for hydrolytically stable and thiol irreversible bioconjugates that react in a chemoselective manner, we naturally sought to explore the use of reagents that would meet these criteria. During the course of developing bromopyridazinediones for reversible cysteine bioconjugation, we became intrigued by the prospect of pyridazinediones (PDs) as irreversible cysteine functionalisation reagents. Previously we have shown that if one of the nitrogen atoms on the PD core is unsubstituted the molecule does not react with thiols at physiological pH or higher.<sup>17a</sup> We postulate that this is a consequence of such a structure existing as its enol tautomer, which is likely to be significantly deprotonated under physiological

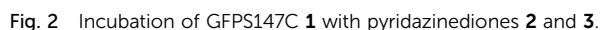


**Fig. 1** A novel strategy for developing thiol-stable pyridazinedione bioconjugates.

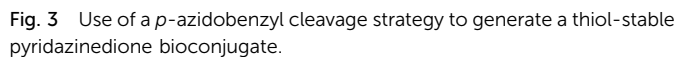
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† Electronic supplementary information (ESI) available: LC-MS, ES-MS and deconvoluted spectra for all reactions with proteins described herein, and <sup>1</sup>H and <sup>13</sup>C NMR spectra for all small molecule constructs. See DOI: 10.1039/c4cc08515a

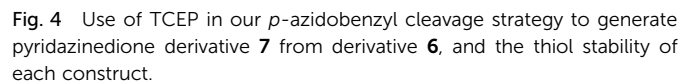


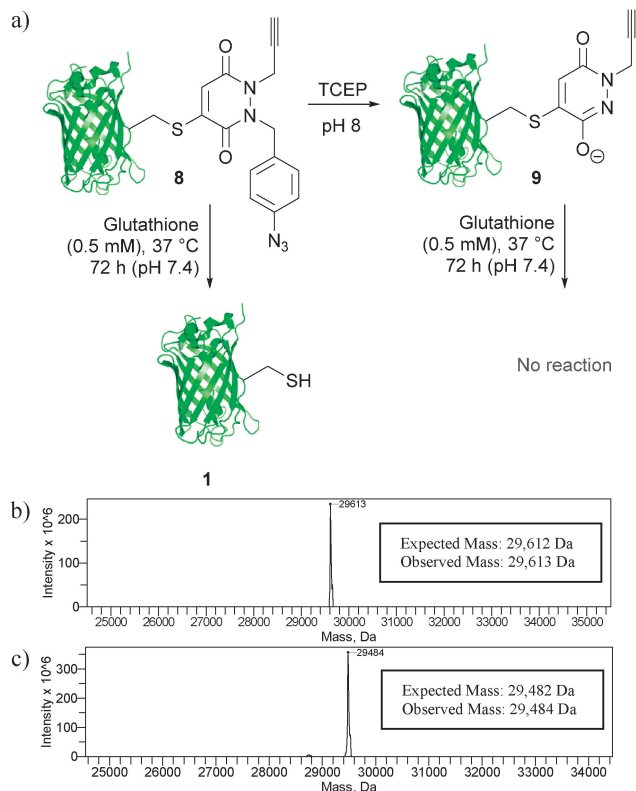


These initial studies paved the way for us to appraise the use of a novel strategy for developing thiol-stable pyridazinedione bioconjugates (see Fig. 1). To do so, we needed to develop a selective method for cleavage of R<sup>2</sup> from the PD core. There are many strategies that could be applied, however, at this juncture we took the opportunity to develop a novel, mild and simple method based on an azide trigger. Our desire to use an azide-based cleavable handle originates from the bioorthogonality of the azide functional group. Taking inspiration from the well-documented work on *p*-aminobenzyloxycarbonyl (PABC) linkers,<sup>19</sup> we set about using a *p*-azidobenzyl cleavage strategy (see Fig. 3).



Following these encouraging results on a small molecule study, we appraised our strategies on a model protein with a single cysteine mutation, GFPS147C **1**. Initially, GFPS147C **1** was incubated with mono-bromo-PD **5** in sodium phosphate buffer (pH 8.0) for 1 h at 37 °C. As expected, this proceeded with complete conversion and afforded GFP-derivative **8**. We next applied our TCEP cleavage strategy, by incubation of this derivative with 10 equivalents of TCEP in phosphate buffer at pH 8.0. Satisfyingly, clean conversion to bioconjugate **9** was observed, which is consistent with our small molecule study. It is also noteworthy that no hydrolysis occurred under these conditions, which is consistent with our previous observations on the PD core being hydrolytically stable.<sup>17a</sup>



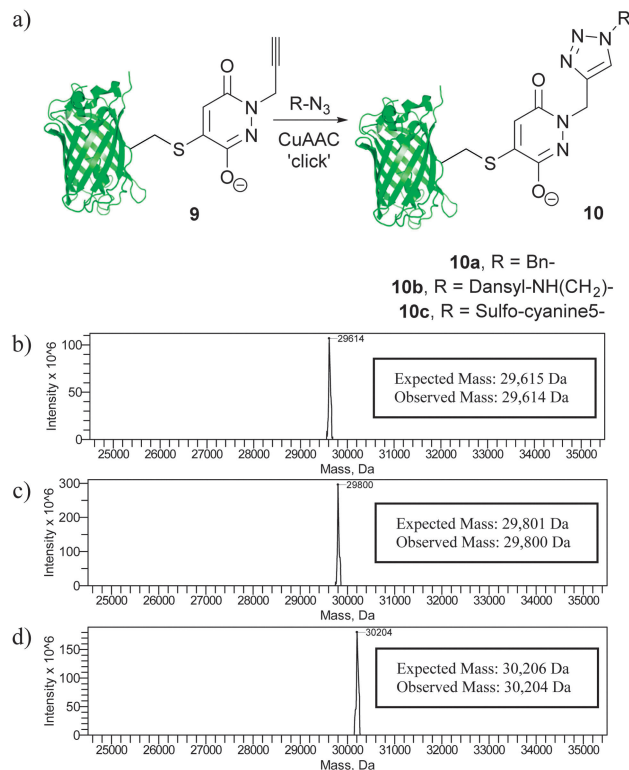


**Fig. 5** (a) Translation of our thiol cleavage and thiol stability strategies onto protein bioconjugates **8** and **9**, and deconvoluted MS data for (b) bioconjugate **8** and (c) bioconjugate **9**.

Having established, using mass spectrometry, that our cleavage strategy is applicable on a protein, we next compared the thiol stability of **8** and **9** by incubation with glutathione (0.5 mM) for 72 h at pH 7.4 and 37 °C. Gratifyingly, GFP-derivative **9** was completely stable under the reaction conditions, whereas derivative **8** showed complete thiol exchange with glutathione. This therefore established proof of concept for both our strategies on a model protein scaffold. Moreover, this work also highlights the versatility of the PD platform with a facile shift from reversible to irreversible constructs achieved under mild conditions (Fig. 5).

Following our work on developing a novel *p*-azidobenzyl cleavage strategy and obtaining a thiol stable construct, we set about functionalising protein scaffold **9** by the use of 'click' chemistry. If successful, this would result in a facile method for functionalising the thiol-stable bioconjugate. A number of 'click' conditions were trialled using benzyl azide as our model azide. The most promising conditions were the use of Cu(I)Br as copper source and THPTA as ligand. These conditions gave complete conversion of starting material alkyne **9** to triazole bioconjugate **10a**. Moreover, these conditions also allowed for clean reaction of the alkyne derivative with a dansyl azide and a sulfo-cyanine5 azide to afford **10b** and **10c**, respectively (Fig. 6).

In conclusion, we have developed, *via* a novel *p*-azidobenzyl cleavage strategy, a route to thiol stable cysteine-bioconjugates that has a clear advantage over conventional maleimide chemistry. The strategy has been demonstrated on both a small molecule



**Fig. 6** (a) The use of 'click' chemistry to functionalise bioconjugate **9**, and deconvoluted MS data for (b) bioconjugate **10a**, (c) bioconjugate **10b** and (d) bioconjugate **10c**.

system and on a model protein, GFPS147C. Owing to the plethora of fields where thiol functionalisation needs to be robust and irreversible, *e.g.* in antibody-drug conjugates (ADCs), imaging and theranostics, we believe this work will find use in a variety of domains. We hope to deliver on the application of our platform in a range of contexts, including ADCs, in the near future.

The authors gratefully acknowledge the EPSRC, Ramsay Memorial Trust and UCL for support of our programme.

## Notes and references

‡ MarvinSketch and its calculator plugins were used for  $pK_a$  prediction, MarvinSketch 14.11.3.0, ChemAxon (<http://www.chemaxon.com>).

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