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 Received 14th July 2021,
Accepted 1st October 2021

DOI: 10.1039/d1cc03802h

rsc.li/chemcomm

A novel thiol-labile cysteine protecting group for peptide synthesis based on a pyridazinedione (PD) scaffold†

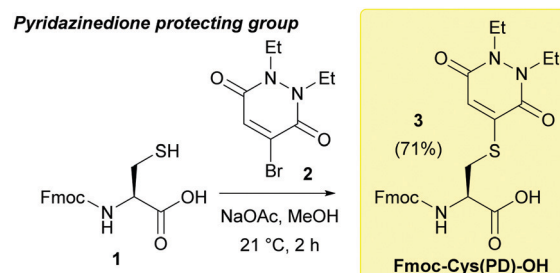
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Herein we report a thiol-labile cysteine protecting group based on an unsaturated pyridazinedione (PD) scaffold. We establish compatibility of the PD in conventional solid phase peptide synthesis (SPPS), showcasing this in the on-resin synthesis of biologically relevant oxytocin. Furthermore, we establish the applicability of the PD protecting group towards both microwave-assisted SPPS and native chemical ligation (NCL) in a model system.

Protecting group strategies for chemoselective protection and deprotection, also referred to as “orthogonal” protecting groups, have proved invaluable in the synthesis of complex biomolecules.¹ In particular, construction of peptides containing multiple disulfide bonds frequently relies on orthogonal cysteine protecting groups to engineer the disulfide linkages in a regioselective manner.² Examples of these include synthesis of hormones such as human insulin³ and bioactive molecules such as conotoxins.⁴ In addition, orthogonal cysteine protecting groups have also been used in protein semisynthesis⁵ and more recently in the synthesis of homogeneous antibody drug conjugates (ADCs) bearing multiple payloads.⁶ The majority of commonly used cysteine protecting groups require acidic conditions and/or toxic heavy metals such as Hg for deprotection, which has thus resulted in a great deal of literature focusing on designing strategies that offer milder deprotection conditions.^{7,8} In particular, thiol labile protecting groups show promise as orthogonal cysteine protecting groups as these can be removed under mild reducing conditions in peptide synthesis. This has primarily been demonstrated with the *tert*-butylsulphenyl/*tert*-butylthio (*StBu*) protecting group.⁹ Deprotection of this group is notably sluggish however, requiring deprotection times of several hours or days depending on peptide sequence.¹⁰ In an effort to address this, the *S*-dimethoxyphenylthio/*S*-trimethoxyphenylthio (*S*-Dmp/*S*-Tmp) groups¹¹ and the *sec*-isoamyl

mercaptan/2-methyloxolane-3-thiol (*SIT*/*MOT*) groups¹² have been reported as replacements for *StBu*, offering significantly faster deprotection times albeit with minor instability towards strongly acidic conditions/piperidine occasionally observed.^{11,12} As evidenced by the development of *S*-Dmp/*S*-Tmp and *SIT*/*MOT*, expansion of the thiol-labile class of cysteine protecting groups would greatly enhance the peptide chemist's current toolkit.

Over the last decade, we have reported pyridazinediones (PDs) as a class of multifunctional molecules for cysteine functionalisation.¹³ We have found that PDs have proved extremely versatile in organic synthesis and chemical biology^{14–16} and can be readily cleaved *via* addition of small molecule thiols in mild, aqueous conditions *i.e.* phosphate buffer (PB) pH 7–8.^{13,17} Given these properties, we anticipated the PD would potentially be well suited to act as a novel, thiol-labile cysteine protecting group in peptide synthesis. We began by validating the use of PDs as cysteine protecting groups in conventional 9-fluorenylmethoxycarbonyl solid phase peptide synthesis (Fmoc SPPS). First, we performed a one-step reaction between Fmoc-Cys-OH **1** and bromopyridazinedione **2** to prepare a suitably protected model Fmoc-Cys(PD)-OH **3** in 71% yield (Fig. 1, see ESI† for full synthesis). Fmoc-Cys(PD)-OH **3** was subsequently incorporated into peptides *via* standard Fmoc SPPS, using 2-(6-chloro-1-*H*-benzotriazole-1-yl)-1,1,3,3-tetramethylammonium hexafluorophosphate (HCTU) and *N,N*-diisopropylethylamine (DIPEA) coupling on a 2-chlorotrityl


 Fig. 1 Synthesis of the protected amino acid Fmoc-Cys(PD)-OH **3**.

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/d1cc03802h



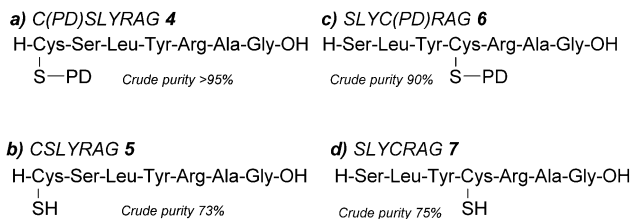


Fig. 2 Peptides synthesised (a) C(PD)SLYRAG 4, (b) CSLYRAG 5, (c) SLYC(PD)RAG 6 and (d) SLYCRAG 7.

resin with Fmoc deprotections carried out using 20% piperidine in DMF (5×2 min treatments). We envisaged that, upon completion of on-resin synthesis, TFA-mediated cleavage of the peptide from the resin (TFA:TIS:H₂O, 95:2.5:2.5) would give the corresponding PD protected peptide, whereas the free thiol-containing peptide would be obtained instead if the resin was treated with a suitable thiol-containing cocktail prior to resin cleavage. Initially, we synthesised on-resin the protected 7-mer peptide Cys(PD)-Ser(*t*Bu)-Leu-Tyr(*t*Bu)-Arg(Pbf)-Ala-Gly, which was cleaved with TFA:TIS:H₂O (95:2.5:2.5). The corresponding PD protected peptide H-Cys(PD)-Ser-Leu-Tyr-Arg-Ala-Gly-OH (C(PD)SLYRAG 4 (Fig. 2a) was obtained as judged by liquid-chromatography mass spectrometry (LCMS), suggesting the PD protecting group was stable to both the acidic and basic conditions used during Fmoc SPPS. Furthermore, crude peptide 4 was obtained in >95% purity as judged by high-performance liquid chromatography (HPLC) analysis. Next, we investigated conditions for on-resin deprotection of the PD protecting group to synthesise the corresponding CSLYRAG peptide CSLYRAG 5 (Fig. 2b). We have previously established that the unsaturated PD motif in protein-PD bioconjugates can be cleaved by addition of excess thiols.¹⁸ With this in mind, we attempted on-resin PD deprotection with 1,4-dithiothreitol (DTT) due to its capacity to remove thiol-labile cysteine protecting groups and its comparative lack of toxicity/stench compared to other thiols such as β -mercaptoethanol (BME) and 1,2-ethanedithiol (EDT). Taking inspiration from the previously described *S*-Dmp/*S*-Tmp protecting groups,¹¹ we initially treated resin-bound peptide (following the final Fmoc deprotection cycle) with a 5% DTT in DMF, 0.1 M *N*-methylmorpholine (NMM) deprotection solution (3×5 min treatments). Following peptide cleavage from the resin, we obtained the CSLYRAG peptide CSLYRAG 5 with minimal deprotection of the PD observed by LCMS (see ESI[†]). In an effort to increase this conversion, we modified the deprotection solution used to comprise an aqueous component, given that our previous work involving deconjugation of protein-PD bioconjugates has primarily been performed in aqueous buffer systems. We therefore treated portions of resin-bound PD protected peptide (3×5 min treatments) with deprotection solutions containing either 5% or 10% DTT w/v with differing ratios of DMF and 5 mM PB pH 8.0, followed by peptide cleavage with TFA:TIS:H₂O (95:2.5:2.5, with 5% DTT w/v to ensure the liberated thiols were kept in the reduced form). Improved conversions (as judged by LCMS) to the corresponding peptide 5 were observed, particularly when using a 5% or

10% DTT w/v in DMF:5 mM phosphate buffer (8:2) deprotection solution (3×5 min treatments), of which the best results were observed when using 10% DTT w/v. Incorporation of Fmoc-Cys(PD)-OH 3 at an internal position within a peptide sequence was then investigated. In a similar manner to the synthesis of 4 and 5, we synthesised peptides SLYC(PD)RAG 6 (Fig. 2c) and SLYCRAG 7 (Fig. 2d) which were obtained following resin cleavage and confirmed by LCMS, with crude purities of 90% and 75% respectively as judged by HPLC. Additionally, we found that no racemisation from L-Cys to D-Cys had occurred in the synthesis of SLYCRAG 7 when using the Cys(PD) protecting group strategy (see ESI[†]). As a direct comparison for the lability of PD protecting group vs. the *t*Bu protecting group, we also synthesised SLYC(*t*Bu)RAG 8 in addition to SLYC(PD)RAG 6 and attempted on-resin synthesis of SLYCRAG 7 (Fig. 3a), with Cys deprotection performed using a 10% DTT w/v in DMF:5 mM phosphate buffer (8:2) deprotection solution (3×5 min treatments) and resin cleavage achieved using TFA:TIS:H₂O (95:2.5:2.5, forgoing 5% DTT w/v addition in the cocktail for both peptides; this was done to avoid unintended *t*Bu deprotection unrelated to prior DTT treatment). Synthesis of SLYC(*t*Bu)RAG 8 proceeded similarly to that of SLYC(PD)RAG 6, whereas attempts to synthesise the deprotected peptide 7 using the Cys(*t*Bu) protecting group strategy gave a mixture of deprotected 7/protected peptide 8 (60% deprotected) as judged by HPLC, as opposed to near quantitative deprotection achieved using the Cys(PD) protecting group strategy (>90% deprotected, Fig. 3b). Overall, the PD protecting group shows faster deprotection times than those reported for the *t*Bu protecting group¹⁰ and similar to *S*-Dmp/*S*-Tmp¹¹ and SIT/MOT.¹²

a) Pyridazinedione vs *tert*-butylthio

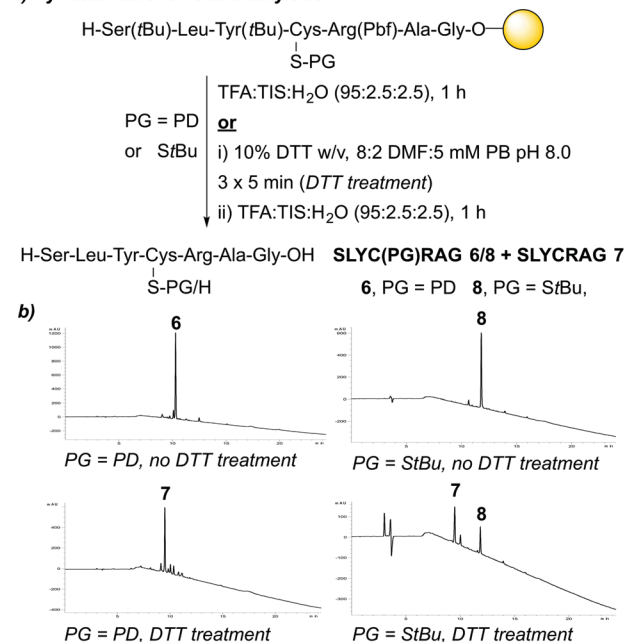


Fig. 3 (a) Outline of synthesis of peptides 6, 7, and 8. (b) HPLC data obtained for attempted synthesis of peptides 6 and 8, along with attempted synthesis of 7.



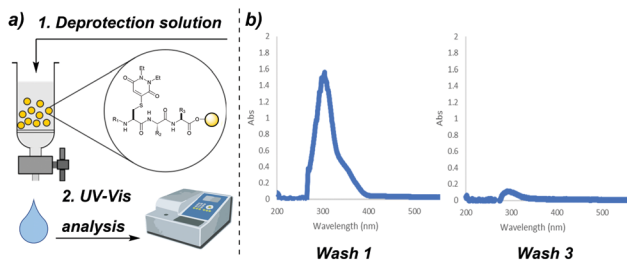


Fig. 4 (a) Outline monitoring on-resin peptide deprotection by UV-vis analysis of wash solutions. (b) Example UV-vis traces of wash solutions in the synthesis of SLYCRAG 7.

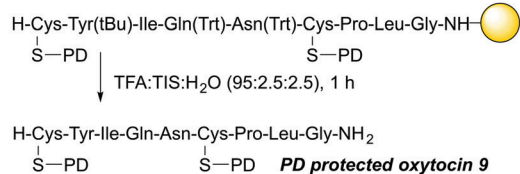
Previously published work featuring PDs reports that the unsaturated motif typically displays an absorption maximum at *ca.* $\lambda = 300\text{--}350\text{ nm}$, *e.g.* $\epsilon_{345} = 9100\text{ cm}^{-1}$ for a dithiopyridazinedione.¹⁹ We hypothesised that on-resin PD deprotection would yield thiol-PD adducts that could be monitored by performing UV-vis analysis of wash solutions post incubation with a given resin-bound, PD protected peptide (Fig. 4a). To test this hypothesis, we collected the wash solutions from resin-bound peptides that had been treated with our deprotection solutions and performed UV-vis analysis. We observed a large absorbance readout that decreased with each washing of the resin; for example, in the synthesis of SLYCRAG 7 (Fig. 4b). This decrease in absorbance was also found to coincide with efficient PD deprotection as judged by LCMS (following resin cleavage). We found that UV-vis analysis of wash solutions collected after treating the resin with deprotection solution in this manner proved to be a rapid and reliable way of tracking PD deprotection during peptide synthesis without the need to cleave portions of the resin-bound peptide to analyse deprotection by HPLC or LCMS. We therefore continued to use this method of analysis throughout the remainder of our studies of the PD protecting group.

We next investigated the applicability of the PD protecting group towards the synthesis of peptides containing more than one cysteine. To this end, we focused on synthesising oxytocin, a disulfide containing 9-mer peptide which can be used for various therapeutic applications such as induction of labour.²⁰

We performed on-resin peptide synthesis on a Rink Amide resin *via* Fmoc SPPS, with Fmoc-Cys(PD)-OH 3 incorporated at both positions 1 and 6 of the peptide. Following the final coupling/Fmoc deprotection step and subsequent peptide cleavage from the resin with TFA:TIS:H₂O (95:2.5:2.5), successful synthesis of the crude PD protected linear oxytocin 9 was confirmed by LCMS analysis and obtained in 87% purity as judged by HPLC. Next, we performed stepwise PD deprotection and subsequent thiol oxidation for the on-resin synthesis of oxytocin. PD deprotection of the resin-bound protected linear oxytocin was performed using a 10% w/v DTT in DMF:5 mM phosphate buffer (8:2) deprotection solution (7 × 5 min treatments, deprotection tracked by UV-vis analysis). Following quantitative PD deprotection (as judged by HPLC/UV-vis analysis), on-resin oxidation of the resin-bound, protected, free thiol-containing linear oxytocin peptide was carried out using *N*-chlorosuccinimide (NCS) as reported previously by Postma and Albericio.²¹ Following thiol oxidation and peptide cleavage from the resin with TFA:TIS:H₂O (95:2.5:2.5), the crude material was subsequently purified by semi-preparative HPLC, with purified oxytocin 10 confirmed by LCMS (Fig. 5a-d).

Finally, we sought to investigate the applicability of the PD protecting group within two different aspects of peptide synthesis as a proof of concept. We first focused on the compatibility of the PD protecting group with microwave-assisted peptide synthesis. We synthesised SLYC(PD)RAG 6 on-resin using microwave-assisted conditions (see ESI† for details) for Fmoc-amino acid couplings (Fig. 6a). As a test of stability, upon completion of on-resin synthesis we also subjected the resin-bound peptide to 20% piperidine in DMF under microwave-assisted conditions. The desired PD protected peptide was obtained in a crude purity of 82% (Fig. 6b), suggesting the PD protecting group was compatible with microwave-assisted conditions. Next, we turned our attention to native chemical ligation (NCL),²² a well-documented strategy for the synthesis of large peptides/proteins through reaction of an N-terminal cysteine-containing peptide fragment and a

a) PD protected oxytocin synthesis



b) Oxytocin synthesis

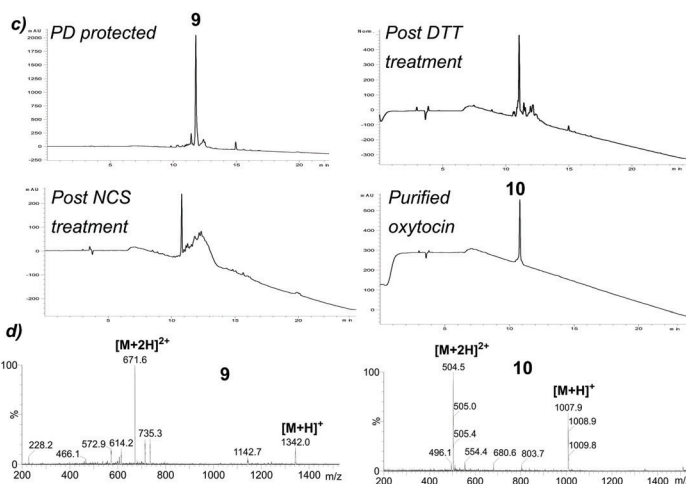
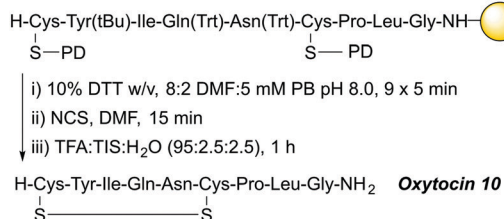


Fig. 5 (a) SPPS of protected linear oxytocin 9. (b) SPPS of oxytocin 10. (c) HPLC data for the synthesis of oxytocin 10. (d) LCMS data for 9 and purified 10.



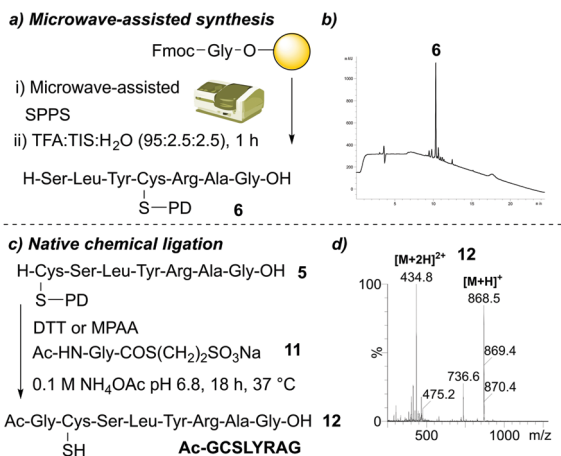


Fig. 6 (a) Outline of microwave-assisted synthesis of SLYC(PD)RAG **6**. (b) HPLC data obtained. (c) Outline of Ac-GCSLYRAG **12** synthesis. (d) LCMS data of crude reaction containing peaks corresponding to **12**.

C-terminal thioester-containing fragment.²³ We envisaged that addition of thiol would act to deprotect the PD protecting group to yield the N-terminal cysteine required for NCL and that this could be done in a one pot manner along with NCL in the presence of a suitable thioester fragment. We attempted a one-pot PD deprotection/NCL procedure by incubating C(PD)SLYRAG **4** with a model *N*-acetylglycine thioester **11** in the presence of either DTT or 4-mercaptophenylacetic acid (MPAA) in ammonium acetate (NH₄OAc) pH 6.8 for 18 h at 37 °C (Fig. 6c). Total consumption of the PD protected peptide **4** was observed, along with conversion to the anticipated NCL product Ac-GCSLYRAG-COOH **12** as judged by LCMS analysis (Fig. 6d).

In conclusion, we have explored the use of pyridazinediones in peptide synthesis, establishing the motif as a thiol-labile protecting group compatible with Fmoc SPPS. We found on-resin deprotection of the PD could be achieved through addition of thiols and monitored by UV-vis analysis. We have demonstrated the applicability of the PD protecting group within on-resin synthesis of disulfide-containing oxytocin, microwave-assisted synthesis and peptide coupling *via* a one pot deprotection/NCL procedure under aqueous conditions. We anticipate the ability to track thiol deprotection by UV-vis analysis, as well as the potential to diversify the ethyl chains of the PD scaffold with moieties such as affinity/solubility tags and clickable handles will assist in enabling more complex peptide syntheses. Furthermore, given its compatibility with aqueous conditions, we anticipate the PD protecting group could be applicable within research focused on designing more environmentally-friendly, “greener” peptide synthesis.²⁴

We gratefully acknowledge the Leverhulme Trust (RPG-2017-288, 176274 and RPG-2020-010) for funding R. J. S. and C. J. B. (respectively), the EU (projects 859458 and 838703 respectively)

for funding L. N. C. R. and F. T. (respectively), and the EPSRC (EP/T517793/1 and EP/R034621/1) for funding I. A. T. and N. F. (respectively). We also acknowledge the UCL Chemistry Mass Spectrometry (MS) Facility (Dr Kersti Karu).

Conflicts of interest

V. C. and J. R. B. are Directors of the spin-out ThioLogics, but there are no competing financial interests to declare.

References

- 1 A. Isidro-Llobet, M. Álvarez and F. Albericio, *Chem. Rev.*, 2009, **109**, 2455–2504.
- 2 R. He, J. Pan, J. P. Mayer and F. Liu, *ChemBioChem*, 2020, **21**, 1101–1111.
- 3 K. Akaji, K. Fujino, T. Tatsumi and Y. Kiso, *J. Am. Chem. Soc.*, 1993, **115**, 11384–11392.
- 4 A. Cuthbertson and B. Indrevoll, *Org. Lett.*, 2003, **5**, 2955–2957.
- 5 M. Jbara, S. K. Maity, M. Seenaiyah and A. Brik, *J. Am. Chem. Soc.*, 2016, **138**, 5069–5075.
- 6 M. R. Levensgood, X. Zhang, J. H. Hunter, K. K. Emmerton, J. B. Miyamoto, T. S. Lewis and P. D. Senter, *Angew. Chem., Int. Ed.*, 2017, **56**, 733–737.
- 7 R. J. Spears, C. McMahon and V. Chudasama, *Chem. Soc. Rev.*, 2021, **50**, 11098–11155.
- 8 S. Laps, G. Satish and A. Brik, *Chem. Soc. Rev.*, 2021, **50**, 2367–2387.
- 9 R. Eritja, J. P. Ziehler-Martin, P. A. Walker, T. D. Lee, K. Legesse, F. Albericio and B. E. Kaplan, *Tetrahedron*, 1987, **43**, 2675–2680.
- 10 M. Góngora-Benitez, J. Tulla-Puche, M. Paradís-Bas, O. Werbitzky, M. Giraud and F. Albericio, *Biopolymers*, 2011, **96**, 69–80.
- 11 T. M. Postma, M. Giraud and F. Albericio, *Org. Lett.*, 2012, **14**, 5468–5471.
- 12 A. Chakraborty, A. Sharma, F. Albericio and B. G. de la Torre, *Org. Lett.*, 2020, **22**, 9644–9647.
- 13 V. Chudasama, M. E. B. Smith, F. F. Schumacher, D. Papaioannou, G. Waksman, J. R. Baker and S. Caddick, *Chem. Commun.*, 2011, **47**, 8781–8783.
- 14 J. C. F. Nogueira, K. Paliashvili, A. Bradford, F. Di Maggio, D. A. Richards, R. M. Day and V. Chudasama, *Org. Biomol. Chem.*, 2020, **18**, 2215–2218.
- 15 C. Bahou, R. J. Spears, A. E. Aliev, A. Maruani, M. Fernandez, F. Javaid, P. A. Szijj, J. R. Baker and V. Chudasama, *Chem. Commun.*, 2019, **55**, 14829–14832.
- 16 A. Maruani, P. A. Szijj, C. Bahou, J. C. F. Nogueira, S. Caddick, J. R. Baker and V. Chudasama, *Bioconjugate Chem.*, 2020, **31**, 520–529.
- 17 M. Fernández, A. Shamsabadi and V. Chudasama, *Chem. Commun.*, 2020, **56**, 1125–1128.
- 18 C. Bahou, P. A. Szijj, R. J. Spears, A. Wall, F. Javaid, A. Sattikar, E. A. Love, J. R. Baker and V. Chudasama, *Bioconjugate Chem.*, 2021, **32**, 672–679.
- 19 A. Maruani, M. E. B. Smith, E. Miranda, K. A. Chester, V. Chudasama and S. Caddick, *Nat. Commun.*, 2015, **6**, 6645.
- 20 J. M. Shyken and R. H. Petrie, *Clin. Obstet. Gynecol.*, 1995, **38**, 232–245.
- 21 T. M. Postma and F. Albericio, *Org. Lett.*, 2013, **15**, 616–619.
- 22 P. E. Dawson, T. W. Muir, I. Clark-Lewis and S. B. Kent, *Science*, 1994, **266**, 776–779.
- 23 V. Agouridas, O. El Mahdi, V. Diemer, M. Cargoët, J.-C. M. Monbaliu and O. Melynk, *Chem. Rev.*, 2019, **119**, 7328–7443.
- 24 O. Al Musaimi, B. G. de la Torre and F. Albericio, *Green Chem.*, 2020, **22**, 996–1018.

