

# Chem Soc Rev

Chemical Society Reviews

rsc.li/chem-soc-rev



ISSN 0306-0012

**REVIEW ARTICLE**

Ashraf Brik *et al.*

Harnessing the power of transition metals in solid-phase peptide synthesis and key steps in the (semi)synthesis of proteins



Cite this: *Chem. Soc. Rev.*, 2021, **50**, 2367

## Harnessing the power of transition metals in solid-phase peptide synthesis and key steps in the (semi)synthesis of proteins

Shay Laps,  † Gandhesiri Satish  † and Ashraf Brik  \*

Peptides and proteins can be either synthesized using solid-phase peptide synthesis (SPPS) or by applying a combination of SPPS and ligation approaches to address fundamental questions related to human health and disease, among others. The demand for their production either by chemical or biological methods continues to raise significant interests from the synthetic community. In this context, transition metals such as Pd, Ag, Hg, Tl, Au, Zn, Ni, and Cu have also contributed to the field of peptide and protein synthesis such as in peptide conjugation, extending native chemical ligation (NCL), and for regioselective disulfide bonds formation. In this review, we highlight, summarize, and evaluate the use of various transition metals in the chemical synthesis of peptides and proteins with emphasis on recent developments in this exciting research area.

Received 6th September 2020

DOI: 10.1039/d0cs01156h

[rsc.li/chem-soc-rev](http://rsc.li/chem-soc-rev)

### 1. Introduction

#### 1.1 Peptide and protein synthesis

Proteins are often made using molecular biology approaches relying on various strategies that are based on the expression of proteins in yeasts, bacteria, or human cell lines, and cell free

expression systems. These strategies, despite being useful in many systems, are still limited with the ability to produce proteins in high homogeneity and large quantities, which contain a diversity of posttranslational modifications (PTMs) at desired sites and numbers, unnatural functionalities, fluorophores, as well as other unique chemical moieties.<sup>1–3</sup> Such complex proteins, in principle, could be obtained using total or semi chemical synthesis.<sup>4–6</sup> In this approach, peptides derived from the protein sequence are synthesized using SPPS and ligated together by chemoselective reactions. The ligation is

Schulich Faculty of Chemistry, Technion-Israel Institute of Technology, Haifa, 3200008, Israel. E-mail: [abrik@technion.ac.il](mailto:abrik@technion.ac.il)

† These authors contributed equally.



**Shay Laps**

Shay Laps received his teaching certificate and BSc degree in Chemistry from the Hebrew University of Jerusalem in 2011 and MA in Science and Technology Education from Ben-Gurion University of the Negev in 2014. He completed his MSc in Chemistry in the Technion-Israel Institute of Technology in 2018, under the supervision of Prof. Brik. He is currently a PhD student under the Jacobs excellence program also in Prof. Brik's

laboratory at the Schulich Faculty of Chemistry in the Technion, working on the development of Pd chemistries for the synthesis of challenging and inaccessible posttranslationally modified peptides and proteins.



**Gandhesiri Satish**

Gandhesiri Satish received his BSc in Chemistry and MSc in Organic Chemistry from Kakatiya University, India. He obtained his PhD under Prof. Andivelu Ilangovan in 2017 from Bharathidasan University, India. Currently he is a post-doctoral fellow in Prof. Brik's laboratory at the Schulich Faculty of Chemistry, Technion – Israel Institute of Technology. His research work is related to the development of ubiquitin activity-based probes for studying and targeting deubiquitinases and proteolysis targeting chimeras (PROTACs) for protein degradation studies.

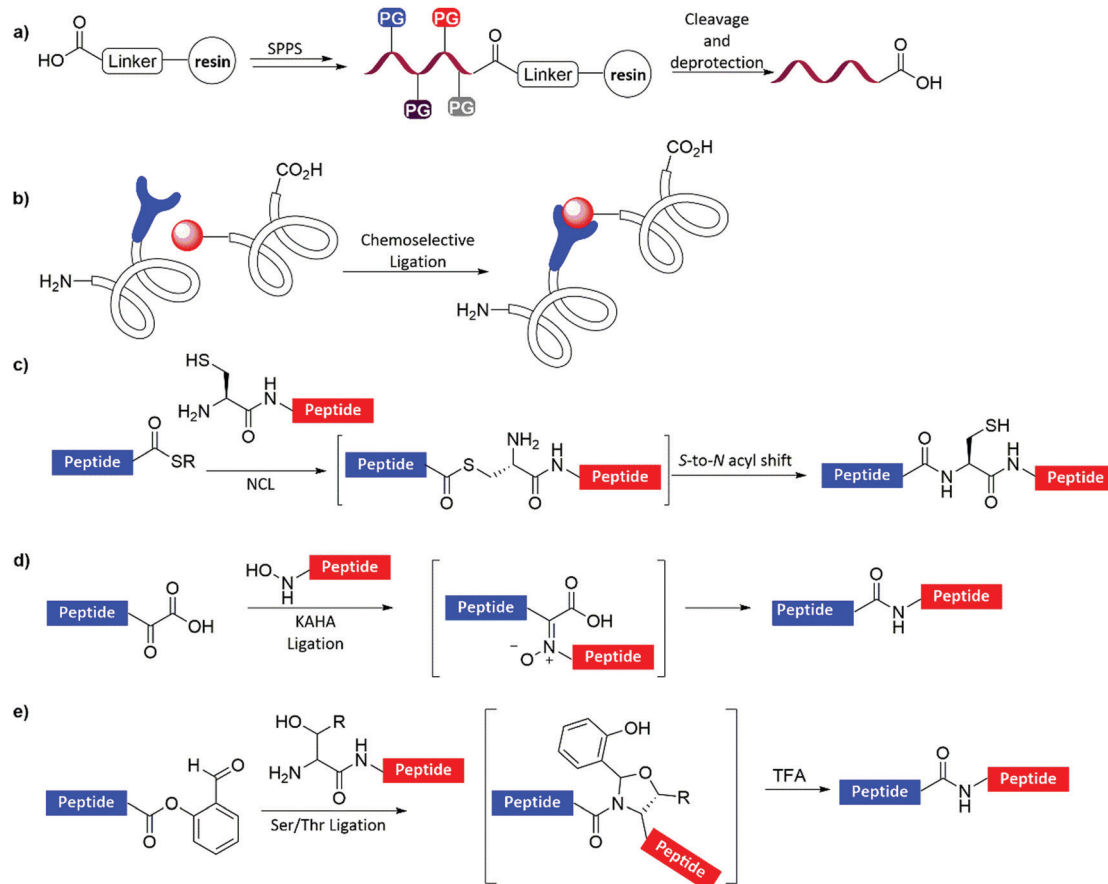


Fig. 1 Chemical synthesis of peptides and proteins: (a) peptide assembly on a solid support (SPPS), (b) general ligation of peptide segments *via* a chemoselective reaction between two functional groups. Examples of chemoselective reactions between two peptides based on: (c) NCL, (d) KAHA ligation, and (e) Ser/Thr ligation, R = H, CH<sub>3</sub>.

based on a chemoselective reaction between two unique functional groups that are introduced into C- and/or the N-termini of peptides.

To simplify peptide synthesis and overcome the limitations of the solution phase synthesis, Bruce Merrifield developed



Ashraf Brik

Ashraf Brik is a professor at the Schulich Faculty of Chemistry in the Technion, Israel Institute of Technology, holding the Jordan and Irene Tark Academic Chair. He received his BSc in Chemistry from the Ben-Gurion University of the Negev and MSc in Chemistry from the Technion. He obtained his PhD in 2001 from the Faculty of Chemistry in the Technion. From 2002 to 2006, he was a Research Associate in the Scripps Research Institute. In 2007, he

joined the Department of Chemistry in the Ben-Gurion University as an assistant professor and was promoted to a full professor in 2012. In 2015, Prof. Brik moved to Technion.

SPPS in 1963.<sup>7</sup> In SPPS, the first amino acid is attached to an insoluble solid support (resin) *via* a special linker (Fig. 1a). Then by repetitive cycles of α-amine deprotection and coupling of the next amino acid, the desired peptide is assembled on the resin. Fluorenylmethyloxycarbonyl (Fmoc)-SPPS,<sup>8</sup> which is the most used strategy relies on α-amine protection with the Fmoc group and can be removed *via* treatment with a base (e.g., piperidine). Protected side chains with acid labile protecting groups (PGs, e.g., trityl (Trt)), are removed during the cleavage from the resin by treatment with trifluoroacetic acid (TFA) and scavengers.<sup>8</sup>

Several chemoselective ligation reactions of peptides have been developed over the years (Fig. 1b) including: (1) NCL, which up to date is the most widely used approach in chemical protein synthesis (Fig. 1c).<sup>4,5,9</sup> In this approach, a peptide bearing an N-terminal Cys reacts in a transthiesterification reaction with another peptide bearing a C-terminal thioester, followed by an N-to-S acyl shift to furnish the amide bond at the ligation junction; (2) the α-keto acid hydroxylamine (KAHA) ligation, which is based on the reaction between an hydroxylamine functionality and α-keto acid to give a native amide bond at the ligation junction (Fig. 1d);<sup>10</sup> (3) Ser or Thr ligation, which is based on the reaction between a peptide with an N-terminal

Ser/Thr and a peptide with a C-terminal *O*-salicylaldehyde ester to form an amide bond (Fig. 1e).<sup>11</sup>

In the process of the peptide synthesis and ligation steps, transition metals have found great utility due to their unique reactivity. This is in addition to the great knowledge we have about their known catalytic nature in a wide scope of organic transformations. Notably, in peptide and protein synthesis, transition metals have been found to be applicable both in organic and aqueous media for various synthetic targets (Fig. 2). This review covers these developments without covering their use in protein modifications (*i.e.*, modification of an expressed form) as some excellent reviews in the literature have already covered this aspect.<sup>12–19</sup>

## 1.2 Protecting groups in peptide and protein synthesis

Protecting the  $\alpha$ -amino group or the side chain of amino acids is crucial for performing SPPS (Fig. 3). In certain cases, protection is necessary also after SPPS and the cleavage from the resin. For example, when NCL is performed between two unprotected peptides, in several cases orthogonal protection of one or more amino acids during the ligation reaction is needed, as is the case in the combination of NCL with desulfurization.<sup>20</sup> Here, native Cys residues have to be protected with orthogonal PGs that are stable to SPPS, cleavage, NCL, and desulfurization conditions, *e.g.*, acetamidomethyl PG (Acm).<sup>21,22</sup> Another example for the requirement of orthogonal protection is when sequential NCL between three peptides or more is desired. In this case, the middle peptide must bear both an N-terminal Cys and C-terminal thioester. In such peptides, the N-terminal Cys has to be protected in order to avoid polymerization or cyclization.<sup>4–6</sup> Apart from this, the applicability of using orthogonal Cys protection was demonstrated in disulfide bond formation and the chemical synthesis of branched proteins, which are discussed in detail in the coming sections.

## 2. Protein synthesis

### 2.1 Silver-mediated peptide ligation

Silver-ion-mediated coupling of peptide bearing a free amine with other peptide bearing a C-terminal thioacid in aqueous DMF was originally reported by Blake in 1981 (Fig. 4a).<sup>23</sup> Later on, Aimoto and co-workers expanded the scope and utility of this method by combining Ag(I) treatment with various activating agents, such as HOBt, to facilitate ligation with the N-terminal amine (Fig. 4a).<sup>24</sup> This interesting strategy, in which the activation of the peptide thioester enables the condensation with an N-terminal amine from the peptide acceptor, was exploited by Hojo and co-workers in their impressive synthesis of a 23 kDa mucin-type glycoprotein, bearing multiple copies of the TN antigen (Fig. 4b).<sup>25</sup> The synthesis involved the initial preparation of a 23-amino acid glycopeptide thioester bearing seven copies of the TN antigen. After being assembled on the solid support, the glycopeptide thioester was cleaved from the resin and then condensed with the trimer peptide (TQT-NH<sub>2</sub>) by the thioester method, which then underwent repetitive ligation cycles *via* silver-mediated activation to form the original repeat sequence.

Later on, Danishefsky and co-workers reported AgCl-assisted phenolic ester-directed amide coupling (PEDAC AgCl) as an extension of this approach using C-terminal *o*-disulfide phenolic ester (Fig. 4c).<sup>26</sup> In this strategy, peptides bearing this moiety on their C-terminus were shown to react rapidly with another peptide bearing an unprotected N-terminal amine under. In this reaction, the authors suspected a dual role of Ag(I), in which silver ions also enhanced disulfide reduction, and therefore they examined the use of a stronger reducing agent, namely tris(2-carboxyethyl)phosphine (TCEP), in place of silver(I) ions to obtain a faster ligation.<sup>26</sup> Nevertheless, the

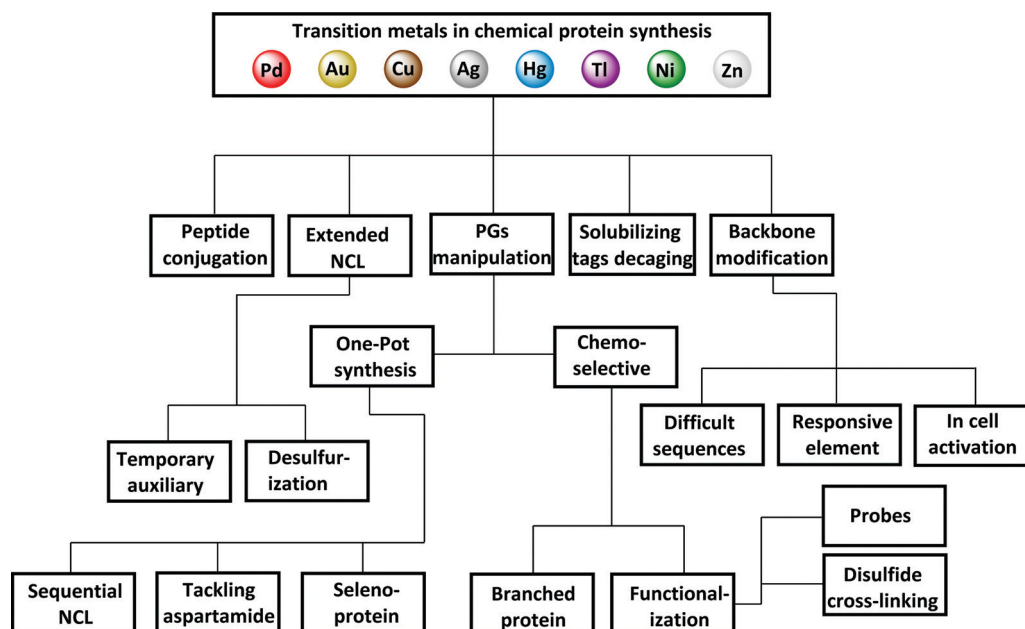


Fig. 2 Transition metals in chemical protein synthesis. PG refers to protecting group and NCL refers to native chemical ligation.

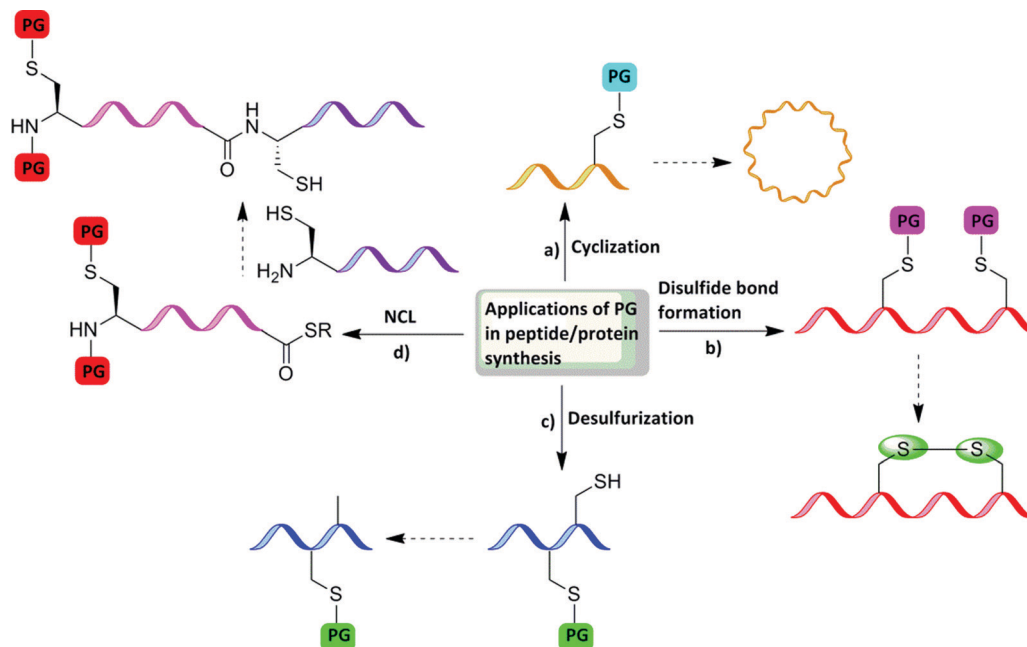


Fig. 3 Applications of PGs in peptide/protein synthesis: (a) cyclization, (b) direct single disulfide bond formation, (c) desulfurization, and (d) ligation.

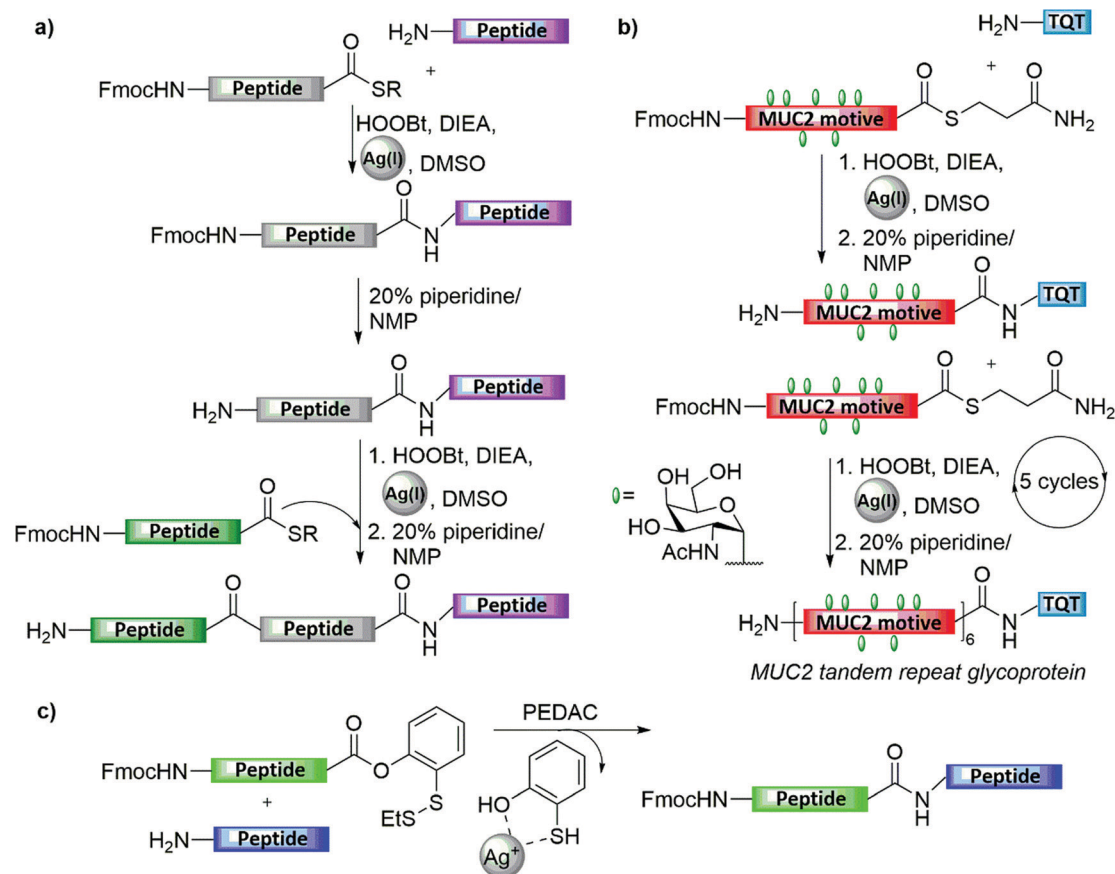


Fig. 4 Silver-mediated peptide ligation: (a) general scheme for silver-mediated sequential ligation, (b) silver-mediated sequential ligation for the synthesis of MUC2, (c) AgCl-assisted phenolic ester-directed amide coupling (PEDAC AgCl).

epimerization of C-terminal residues other than Gly and Pro were obtained in both approaches. Another limitation of this

approach is the necessity to protect the Lys and Cys side chains during the ligation step, to prevent unwanted side reactions.<sup>26</sup>

## 2.2 Desulfurization

A synthetic scheme was also developed for various protein targets that also included a thiol elimination step once the NCL between two or more segments has been accomplished (Fig. 5a).<sup>4–6,27</sup> This combination of NCL and desulfurization extended the utility of protein synthesis for various ligation junctions.<sup>27</sup> The synthesis based on these tools requires the introduction of a thiol functionality at the N-terminal peptide (e.g., N-terminal Cys) to enable the transthioesterification step followed by an *S*-to-*N* acyl shift. Upon polypeptide assembly, the thiol moiety then can be eliminated by employing metal or metal-free desulfurization reactions. Despite an important development emerging from Danishefsky's lab, where a radical-based approach was introduced to achieve rapid desulfurization under homogenous aqueous conditions,<sup>28</sup> transition metals have found great utility in the chemical synthesis of many proteins *via* NCL coupled with desulfurization.

Dawson and co-worker demonstrated the first total synthesis of Microcin J25, determined at that time as a 21 AA cyclic antibiotic peptide, which is naturally produced by a strain of *E. coli*, and which lacks a Cys residue (Fig. 5b).<sup>20</sup> Further structural investigations revealed, however, that Microcin J25 comprises an unusual N-terminus to a Glu8 side-chain amide bond, resulting in an N-terminal ring in a conformation that has been described as a “lasso” type structure.<sup>29–31</sup> In the original synthesis, the linear analogue of microcin J25 was prepared *via* Boc-SPPS. An N-terminal Cys was introduced, replacing natural Ala within the peptide sequence, while the C-terminal of the same peptide was functionalized with a thioester. Cyclization

*via* NCL followed by metal-mediated desulfurization furnished the native microcin J25 (as was known before the structure was further revised).<sup>20</sup> For the desulfurization step, the cyclized product was treated with Pd/Al<sub>2</sub>O<sub>3</sub> (10%) in 20% aqueous acetic acid (AcOH) and exposed to H<sub>2</sub> gas to eliminate the thiol functionality and convert it to the native Ala. In addition, the authors performed a comparison study between various conditions (e.g., Pd/Al<sub>2</sub>O<sub>3</sub>, (10%), Pd/C (10%), Pd/BaSO<sub>4</sub> (10%), PdO and RANEY<sup>®</sup> nickel). The comparison revealed a similar efficiency of Pd/Al<sub>2</sub>O<sub>3</sub> and Pd/C (10%) in 20% aqueous AcOH (>99% yield), while Pd/BaSO<sub>4</sub> and PdO resulted in lowered yields (90% and 30%). RANEY<sup>®</sup> nickel was shown to be as efficient as Pd/Al<sub>2</sub>O<sub>3</sub>; however, the demethylthiolation of Met residues has also been observed.<sup>20</sup>

The Pd-mediated desulfurization enabled the development of sugar-assisted glycopeptide ligation by Brik *et al.*, in which a sugar moiety of a glycopeptide modified with a thiol handle at the C-2 position to form an acetamido thiol handle facilitated the ligation of a Cys-free glycopeptide with a thioester peptide *via* an *S*-to-*N*-acyl shift followed by desulfurization of the ligation product to the corresponding native acetamide moiety on the sugar part.<sup>32,33</sup> The same group developed a strategy for selective Cys desulfurization in the presence of a thiol moiety protected with Acm.<sup>21</sup>

In parallel, Pentelute *et al.* reported a selective desulfurization of Cys in the presence of Cys(Acm) in polypeptides obtained by NCL.<sup>21,22</sup> In this study, the authors aimed to extend the use of NCL followed by desulfurization to enable a more versatile synthesis of Cys-rich peptide and protein targets. The unique and challenging

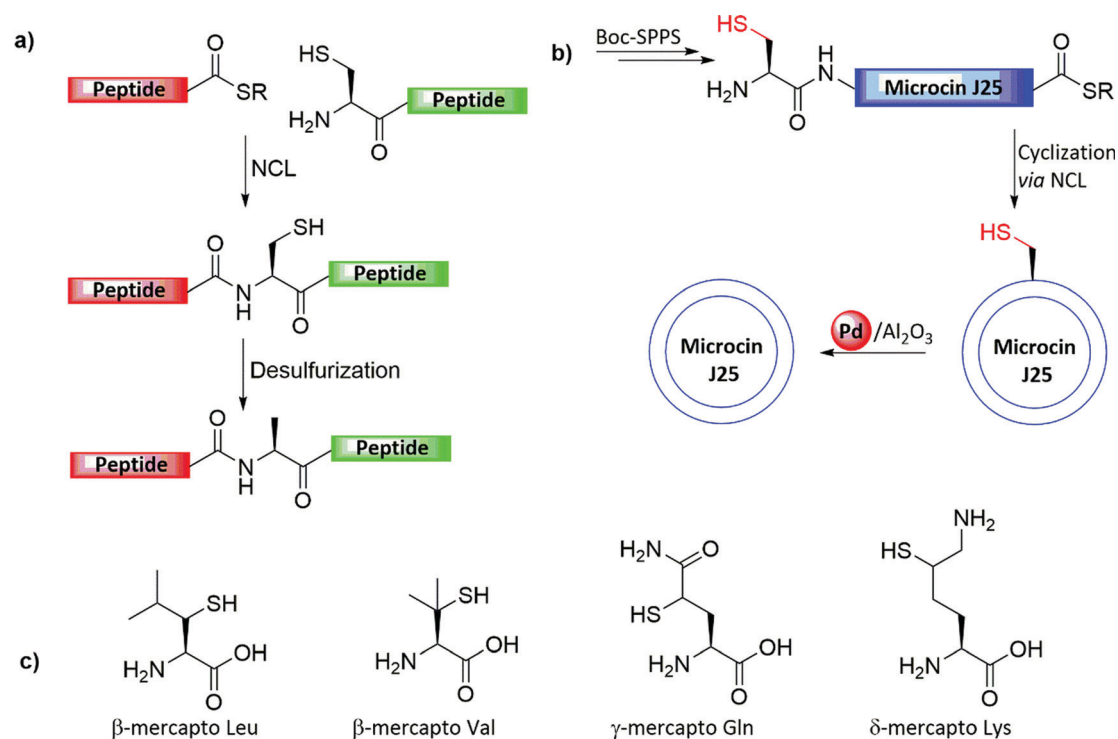


Fig. 5 (a) Strategy for NCL followed by desulfurization to recover native Ala. (b) Synthesis of Microcin J25 cyclic peptide. (c) Selected examples of non-native thiolated amino acids.

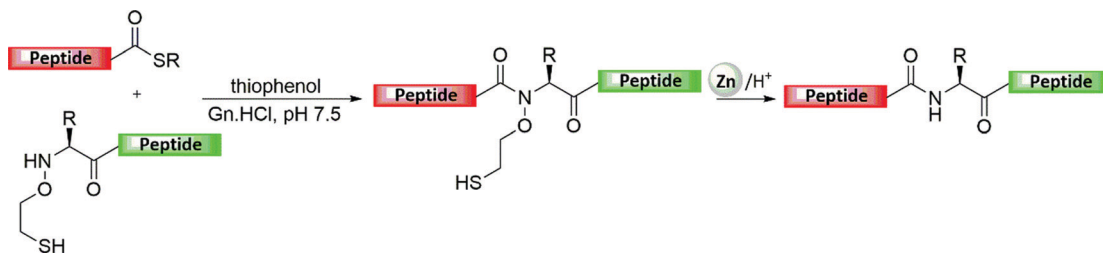


Fig. 6 Representative depiction of a temporary auxiliary-based NCL followed by Zn-mediated removal.

knotted trypsin inhibitor EETI-II protein was chosen as a proof of concept.<sup>22</sup> EETI-II contains 28 AA and six Cys residues that form a highly stable framework *via* three disulfide bonds. The synthetic route involved NCL between a segment bearing a C-terminal thioester and another segment in which Ala17 was mutated to Cys. All the natural six Cys residues were protected with Acm. After NCL of the two segments, selective desulfurization of residue Cys17 to give Ala17 was performed *via* RANEY<sup>®</sup> nickel-mediated reduction, in the presence of the Cys(Acm) residues that remained intact. Following HPLC purification, the Acm PGs were removed and the purified product was subjected to oxidative folding to form the native EETI-II.<sup>22</sup>

These early developments inspired several groups to exploit this concept to develop additional ligation junctions utilizing various artificial AAs bearing temporary thiol handles (such as Lys, Asp, Glu, Val, Trp, and Pro) (Fig. 5c).<sup>4–6,27</sup> These further developments have added considerable flexibility in choosing the ligation junctions when attempting the synthesis of a particular protein.

### 2.3 Zn-Mediated removal of oxyalkyl auxiliary in peptide ligation

Kent and co-workers have sought to extend the applicability of NCL to more junctions beside Cys since the early days of its development. His group reported the development of a cleavable oxyethanethiol-based auxiliary to assist ligation, which was designed to be introduced at the backbone amide of the first AA of the peptide sequence (Fig. 6).<sup>34</sup> In this approach, a C-terminal thioester peptide reacts *via* thiol exchange with another peptide containing the auxiliary, which through a rearrangement involving a six-membered ring gives the amide-linked product.<sup>34</sup> The auxiliary showed stability under HF cleavage conditions yet could be simply removed under treatment with zinc dust to give the native backbone structure of the ligation product. Although this strategy was applied for the syntheses of a few proteins,<sup>35</sup> it has been not widely used mainly due to the slow ligation rates ( $\sim 12$  h for the ligation at Gly junction compared to  $\sim 1$  h *via* Cys-based NCL in the presence of 4-mercaptophenylacetic acid (MPAA) and TCEP).<sup>36</sup>

## 3. Removal of the protecting group with transition metals in organic solvents

### 3.1 Acm PG deprotection

The stability of the Acm group to acidic conditions allows its orthogonal use with other PGs (Fig. 7a and Table 1). It is also

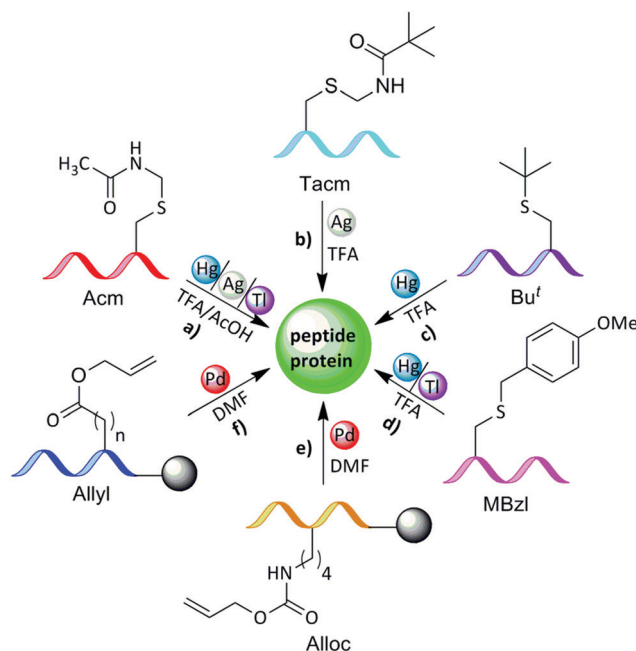


Fig. 7 Transition metal-mediated removal of the PGs in organic solvents for peptide and protein synthesis: (a) removal of Acm PG with Hg, Ag or Tl in TFA or AcOH; (b) removal of *S*-trimethylacetamidelmethyl (Tacm) PG with Ag in TFA; (c) removal of *tert*-butyl PG with Hg in TFA; (d) removal of *p*-methoxybenzyl (MBzl) PG with Hg or Tl in TFA; (e) removal of Alloc PG with Pd in DMF; (f) removal of allyl PG with Pd in DMF.

Table 1 Transition metals-mediated removal of PGs in organic solvents

Amino acid	Protecting group	Compatibility	Transition metal	Solvent
Cys	Acm	Boc, Fmoc	Hg(II)	AcOH
			Ag(I)	TFA, TFE
			Ag(II)	AcOH
			Tl(III)	TFA + CH <sub>3</sub> CN
				TFA
Cys	TAcM	Boc, Fmoc	Hg(II)	TFA
			Ag(I)	TFA
			Hg(II)	TFA, AcOH
			Tl(III)	TFA, AcOH
Cys	MBzl	Boc, <sup>a</sup> Fmoc	Hg(II)	TFA
			Tl(III)	TFA
Lys	Alloc	Boc, Fmoc	Pd(0)	DCM
			Pd(0)	DCM

<sup>a</sup> Except for repetitive treatments.

stable to the applied conditions for the generation of hydrazides from esters and azides from hydrazides.<sup>37</sup> The main advantage

of the Acm group comes from its stability under the conditions commonly used in peptide synthesis (Fmoc or Boc) in solution or in the solid-phase method. Acm can be used as a PG during segment condensation,<sup>38</sup> sequential ligation,<sup>21,39–41</sup> desulfurization,<sup>21,22,42</sup> and disulfide bond formation.<sup>43</sup>

The protection of thiol group of Cys by the Acm group was introduced by Hirshmann.<sup>37</sup> In this report, mercury(II) acetate [Hg(OAc)<sub>2</sub>] in AcOH was used for the deprotection of Acm in octa-5-Acm-octahydro-5-protein to give S-protein, with further air oxidation to enable the synthesis of ribonuclease-S. The mercuric ion was removed by treatment with mercaptoethanol for a long time (19 h) by producing acetamido methanol as a byproduct.

Yajima and co-workers utilized Acm-protected Cys residues to form a single disulfide bond in oxytocin peptide using thalium(III)trifluoroacetate [Tl(OTf)<sub>3</sub>] in combination with TFA for the removal step.<sup>44</sup> However, under these conditions, Trp and Met were found to be unstable and formed some unidentified side products. Hence, these functionalities must be protected during the Tl(OTf)<sub>3</sub> treatment. To overcome these issues, another catalytic system involving silver trifluoromethanesulfonate (AgOTf) in TFA and anisole was developed.<sup>45</sup> Under these conditions, Trp, Met, and Tyr were less prone to undergo side reactions, and other Cys PGs, like *S*-*t*-butyl (Bu<sup>t</sup>) and 4-methylbenzyl, were not affected.

Surprisingly, when these catalytic systems [Tl(OTf)<sub>3</sub> and Hg(OAc)<sub>2</sub>] were employed for the synthesis of an amino terminal loop of the snake  $\alpha$ -neurotoxin, the Acm group was shifted from the Cys residue to the Ser or Thr residue.<sup>46</sup> However, the addition of glycerol (trivalent alcohol) as a scavenger in the presence of both metal complexes prevented this shift.

As discussed in Section 2.1,<sup>24</sup> Aimoto and co-workers selectively achieved peptide ligation and Acm removal in the presence of AgNO<sub>3</sub> by changing the nucleophile. Here, the nucleophilic character of the additive HOOBt was enough to attack the thioester activated with a silver ion, but was too weak to attack the thioether of the Acm, eventually leading to a coupling reaction. In another study, however, the same conditions (AgNO<sub>3</sub>, HOOBt) led to the removal of the Acm group when the synthesis of a reaper polypeptide was attempted.<sup>47</sup> The reason for this could be explained by the influence of the amino acid residue in the immediate vicinity of the Cys(Acm), which greatly influences the stability of the Acm group toward different silver ions. Hence, changing the silver salt, such as by using AgCl instead of AgNO<sub>3</sub>, for the condensation step provided the reaper as well as the molt-inhibiting hormone (Prc-MIH).

Kent and co-workers reported the synthesis of crambin by three peptide segments, which gave a 10-fold increase in yield compared to from two segments (Fig. 8).<sup>48</sup> Here, the N-terminal Cys of the middle segment was temporally protected with the Acm group to prevent undesired cyclization or polymerization. Following the ligation step, the Acm group was subjected to removal with Hg(OAc)<sub>2</sub>; however, the reaction was less efficient. Changing the metal to Ag(OAc)<sub>2</sub>, however, enabled the efficient removal of Acm to facilitate the subsequent ligation. Similarly, the total synthesis of a range of proteins, such as erythropoietin (EPO),<sup>38</sup> sortase A,<sup>49</sup> interferon- $\beta$ -1a (IFN- $\beta$ -1a),<sup>50</sup> glycosylated

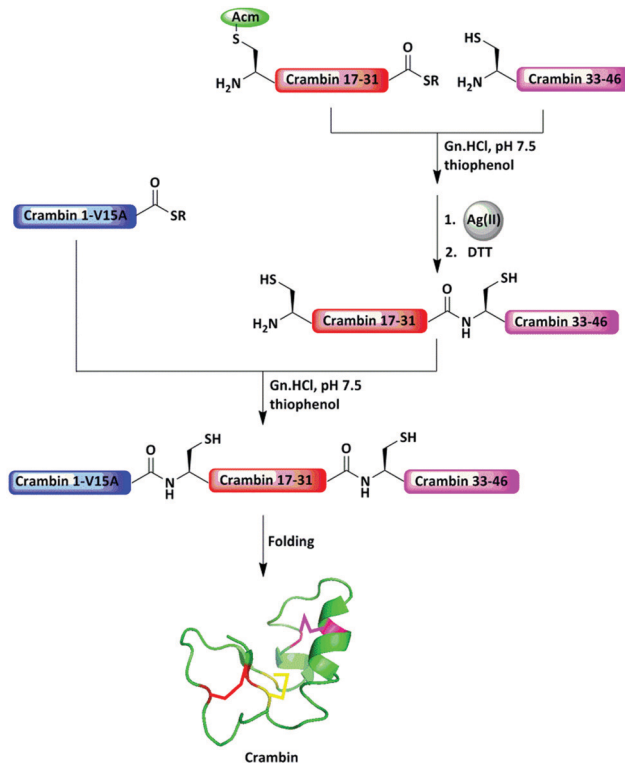


Fig. 8 Protection of the middle fragment with an Acm group to facilitate sequential ligation for the synthesis of crambin.

granulocyte-macrophage colony-stimulating factor (GM-CSF),<sup>43</sup> designed ankyrin repeat protein (DARPin) PE59,<sup>41</sup> bacillus amyloliquefaciens RNase (Barnase),<sup>41</sup> and HIV-1 Rev protein,<sup>51</sup> were achieved employing Acm as a PG.

Our group utilized Acm as an N-terminal Cys PG of the middle fragment during isopeptide chemical ligation (ICL) for site-specific ubiquitination at the lysine 34 position of H2B as well as the  $\delta$ -mercaptolysine of the  $\alpha$ -synuclein-thioester fragment to enable expressed protein ligation (EPL) with the recombinant Cys fragment (Fig. 9),<sup>52</sup> and later, the same target was prepared by a sequential strategy.<sup>40</sup> Similarly, other ubiquitinated proteins, like ubiquitinated  $\alpha$ -synuclein,<sup>53</sup> H3K4me3,<sup>54</sup> and H2K120Ub,<sup>42</sup> were also obtained.

Kiso and co-workers developed the *S*-trimethylacetamidomethyl (TAcM) group as a thiol PG of Cys (Fig. 7b).<sup>55</sup> This PG has similar chemical properties as Acm, but is less susceptible to air oxidation. The utility of this group was demonstrated by the synthesis of porcine brain natriuretic peptide and somatostatin.<sup>55</sup>

### 3.2 Tertiary-butyl (Bu<sup>t</sup>) PG deprotection

The Cys *t*-butyl (Bu<sup>t</sup>) group is completely stable under the regular conditions of Fmoc-SPPS and its deprotection has been less explored (Fig. 7c and Table 1). Fujino and co-workers reported the removal of the *S*-Bu<sup>t</sup> group under Hg(OAc)<sub>2</sub> in TFA or Hg(OTf)<sub>2</sub> in AcOH using anisole as a scavenger.<sup>56</sup> In the subsequent step, the free thiol was regenerated quantitatively by treatment with hydrogen sulfide. Other heavy metal ions, including Ag<sup>+</sup>, Cu<sup>2+</sup>, Zn<sup>2+</sup>, Ni<sup>2+</sup>, and Pb<sup>2+</sup>, were completely ineffective for the removal of the *S*-Bu<sup>t</sup> group.<sup>56</sup> The same



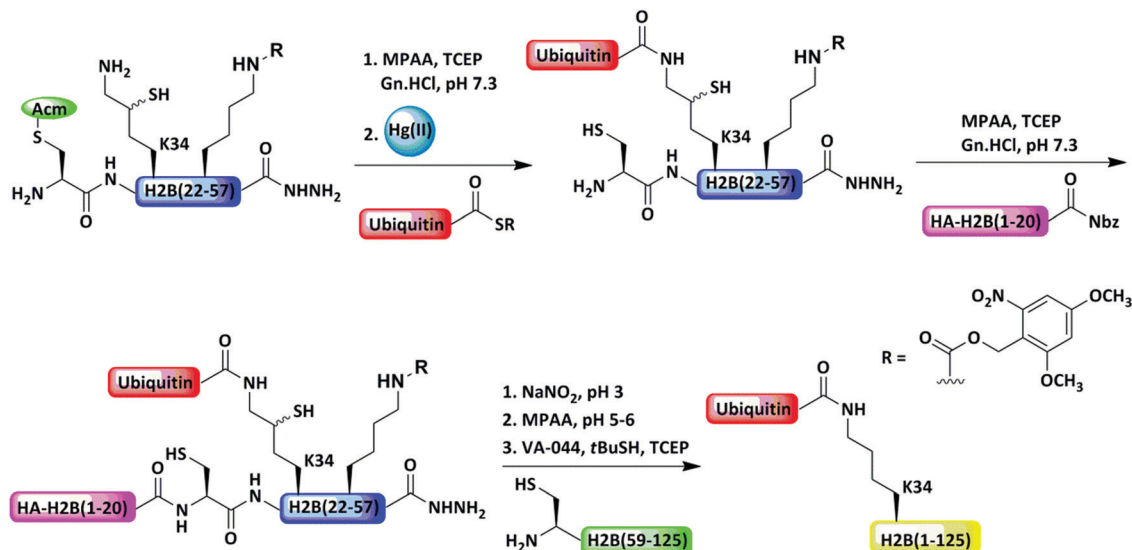


Fig. 9 Protection of the N-terminal Cys with Acm for ICL in the synthesis of ubiquitinated proteins.

cleavage conditions for S-Bu<sup>t</sup> were successfully applied for the removal of another thiol-PG, namely *p*-methoxybenzyl (MBzl),<sup>56</sup> with an even faster rate and higher yields.

### 3.3 *p*-Methoxybenzyl (MBzl) PG deprotection

Yazima and co-workers found that the treatment of Cys (MBzl) with Tl(OTf)<sub>3</sub> as a soft Lewis acid under acidic conditions resulted in effective MBzl cleavage (Fig. 7d and Table 1).<sup>44</sup> However, under these conditions, Trp gave several unidentified products and Met was partially oxidized. Hence, unprotected Trp and Met are not compatible with Tl(OTf)<sub>3</sub> treatment. The utility of this method was demonstrated in the syntheses of a urotensin II peptide bearing a single disulfide bond and a Trp, which was temporally protected and upon formation of the disulfide bond, was then removed with TMSOTf/TFA in the presence of diphenyl sulfide.

### 3.4 Allyloxy carbonyl (Alloc) PG deprotection

The Alloc group was first utilized by Stevens and Watanabe for the protection of the N-terminal group of D,L-amino acids (Fig. 7e, f and Table 1).<sup>57</sup> This PG is widely used mainly due to its compatibility with either Boc- or Fmoc-SPPS, and as it can be cleaved efficiently under mild and neutral conditions. The cleavage of the Alloc group was demonstrated by catalytic hydrogenolysis either with Pt or Pd catalysts. Later, Kunz and Unverzagt developed effective conditions for the removal of the Alloc group by Pd(PPh<sub>3</sub>)<sub>4</sub> in tetrahydrofuran *via* an allyl group transfer.<sup>58</sup> In these reactions, dimedone and others, such as phenylsilane, were used as scavengers to capture the released allyl group.<sup>59</sup> Several research groups employed Alloc/allyl as PGs to the side chains of Lys/Glu residues for the synthesis of cyclic peptides on resin through a coupling of the head to tail/side chain to side chains.<sup>60,61</sup> Apart from this, the Alloc group served as an orthogonal PG for the thiazolidine (Thz) on resin in the total synthesis of ubiquitinated  $\alpha$ -globin.<sup>62</sup>

## 4. Removal of the protecting group with transition metals in aqueous solvents

### 4.1 N-terminal Cys protection

Thz is a widely used N-terminal-protected form of Cys when the sequential ligation of three or more fragments is applied (Table 2).<sup>63</sup> Nevertheless, the unmasking step with methoxylamine takes a long time, which slows down the entire synthetic process of the target protein. In addition, thioester peptide with C-terminal Gly has been reported to be modified irreversibly with the methoxylamine.<sup>12,64–66</sup> In addition, Thz was found to be unstable during the oxidative switching of hydrazide peptide.<sup>67</sup> Our laboratory addressed these challenges and found that Pd(II) complexes can effectively open the Thz ring to give the free unprotected N-terminal Cys, within minutes, under NCL conditions (Fig. 10a).<sup>68</sup> This approach has been applied for the synthesis of Lys34-ubiquitinated H2B among other proteins.<sup>69</sup> We also reported that the propargyloxycarbonyl group (Proc) at the N-terminus Cys undergoes deprotection in an effective manner *via* exposure to PdCl<sub>2</sub> under NCL conditions, as demonstrated in the first synthesis of neddlyated peptides (Fig. 10b).<sup>68</sup> In this study, the addition of Mg<sup>2+</sup>

Table 2 Transition metals-mediated removal of PGs in aqueous solvents

Amino acid	Protecting group	Transition metal	Solvent	Introduction to the sequence	
Cys	Acm	Pd(II)	6 M Gn-HCl	Fmoc-SPPS	
		Cu(II)	6 M Gn-HCl	Fmoc-SPPS	
	Thz	Pd(II)	6 M Gn-HCl	Fmoc-SPPS	
		Cu(II)	6 M Gn-HCl	Fmoc-SPPS	
		Pd(II)	6 M Gn-HCl	Fmoc-SPPS	
	Proc	Pd(II)	6 M Gn-HCl	Fmoc-SPPS	
	Alloc	Pd(0)	6 M Gn-HCl	Fmoc-SPPS	
Bu <sup>t</sup>	Pd(II)	Urea/Tris buffer	Fmoc-SPPS		
Lys	Succinimide	Pd(II)	6 M Gn-HCl	Solution phase	
		Au(I)	6 M Gn-HCl	Fmoc-SPPS	
	Asp/Glu	Allyl	Pd(II)	6 M Gn-HCl	Fmoc-SPPS
		Allyl	Pd(0)	6 M Gn-HCl	Fmoc-SPPS
Asp	Allyl	Pd(0)	6 M Gn-HCl	Fmoc-SPPS	
Sec	Sez	Cu(II)	6 M Gn-HCl	Fmoc-SPPS	

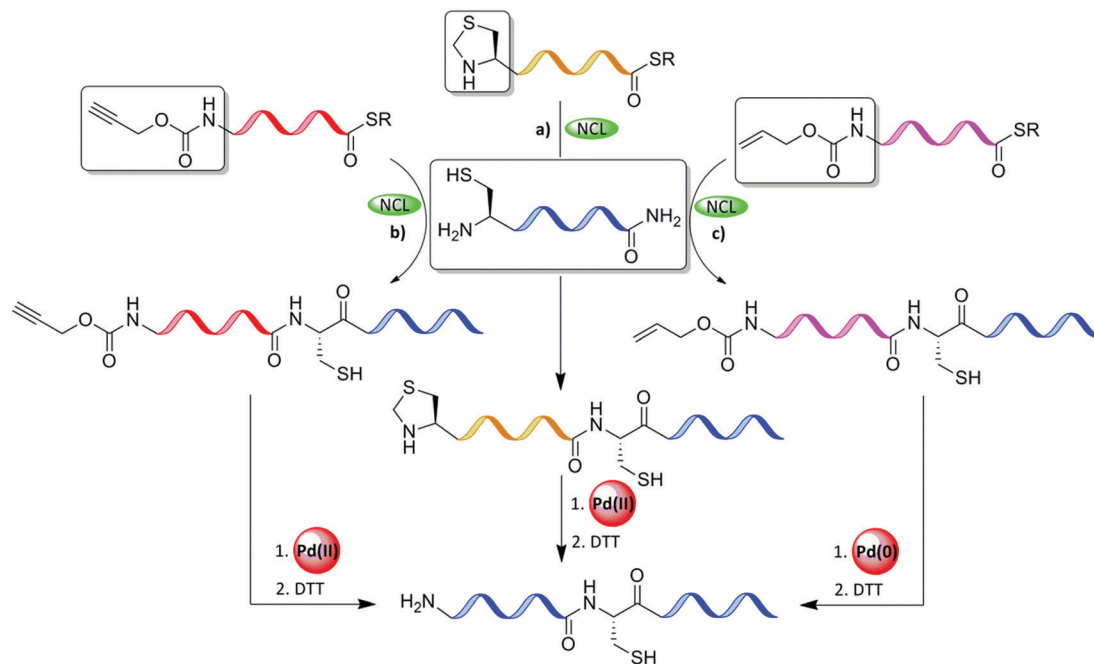


Fig. 10 N-terminal Cys protection of the middle fragments (a) Thz, (b) Proc, and (c) Alloc during sequential ligation and removal with Pd complexes.

was found to facilitate the efficient one-pot Pd(II)-mediated Thz/Proc deprotection under NCL conditions.<sup>68</sup>

Inspired by our findings, Okamoto's laboratory applied Pd(0) complexes to remove the Alloc PG from N-terminal Cys (Fig. 10c and Table 2). In this study, MPAA was used as an enhancement additive for ligation and as a scavenger for the  $\pi$ -allyl complexes, and for quenching Pd.<sup>70</sup> This approach was demonstrated in the chemical synthesis of histone H2AX.

#### 4.2 Internal Cys protection

The harsh removal conditions of the AcM PG, which were discussed in Section 3.1, often requires long reaction times, prevents one-pot ligation and/or desulfurization processes, and necessitates additional purification steps before and after its removal. Our laboratory discovered that Pd(II) complexes can effectively unmask Cys(AcM) in peptides and proteins, within 5 min, under aqueous NCL/desulfurization conditions (Fig. 11a and Table 2).<sup>71</sup> Notably, PdCl<sub>2</sub> dissolved in 6 M Gn·HCl more efficiently than in water to form the corresponding complexes, probably due to the coordination of the guanidinium group to Pd(II).<sup>72</sup> We applied these mild AcM removal conditions to construct proteins containing native Cys residues by NCL and desulfurization.<sup>12,71,73</sup> As a proof of concept, the ubiquitin-like protein (UBL-5) was synthesized by applying this chemistry.<sup>71</sup> UBL-5 is a 73 AAs protein containing two Cys residues, which are not located in strategic positions suitable for NCL (6, 18). Consequently, the sequence was dissected into two segments: UBL-5N(1–32) bearing two Cys protected with AcM and a C-terminal thioester, and a second segment of UBL-5C(33–73), while a mutation of Ala33 into Cys was devised (Fig. 11b). A one-pot operation of NCL followed by desulfurization and AcM removal was successfully conducted, demonstrating the power of the developed tools for efficient protein synthesis.<sup>69,71,74</sup>

Inspired by our work, Wang and co-workers reported a method for the site-specific incorporation of selenocysteine (Sec) into proteins in *E. coli* and mammalian cells.<sup>75</sup>

The Sec incorporation required idiosyncratic translational machinery, which makes it challenging to recombinantly prepare selenoproteins with high Sec specificity. Here, the authors reported an orthogonal tRNA<sup>Pyl</sup>-ASecRS to site-specifically incorporate allyl-protected Sec. Subsequently, the allyl group was removed using Pd chemistry under mild conditions. This method provides the opportunity to introduce site-specific Sec to proteins and enables their biochemical study.<sup>75</sup>

Very recently, Otaka and co-workers reported the removal of AcM from Cys (Table 2).<sup>76</sup> The deprotection was established *via* treatment with Cu(II) in the presence of free N-terminal Cys or an external source of 1,2-aminothiols, which was required in order to avoid severe side reactions. The reaction took 3 h compared to 5 min when using Pd, highlighting the differences in the used transition metal in these transformations. In their proposed mechanism, Cu was involved in the removal of AcM and the formation of Cys to the S–OH intermediate, which could then be reacted with EDTA or DTT to give the oxidized or reduced peptide, respectively.<sup>76</sup>

#### 4.3 C-terminal Asp protection

As discussed in Section 1.1, NCL based on XX-Cys (XX represents any of the 20 canonical amino acids) sites has been shown to proceed with different kinetics.<sup>77</sup> However, the carboxylate group side-chain of Asp and Glu are known to form the cyclic C-terminal anhydride through intramolecular cyclization, which can further be attacked by the Cys-peptide to form the  $\beta$ - or  $\gamma$ -linked product as well as the native product. (Fig. 12).<sup>78,79</sup> When Boc chemistry<sup>79</sup> or *N*-acylbenzimidazolone (Nbz)-based

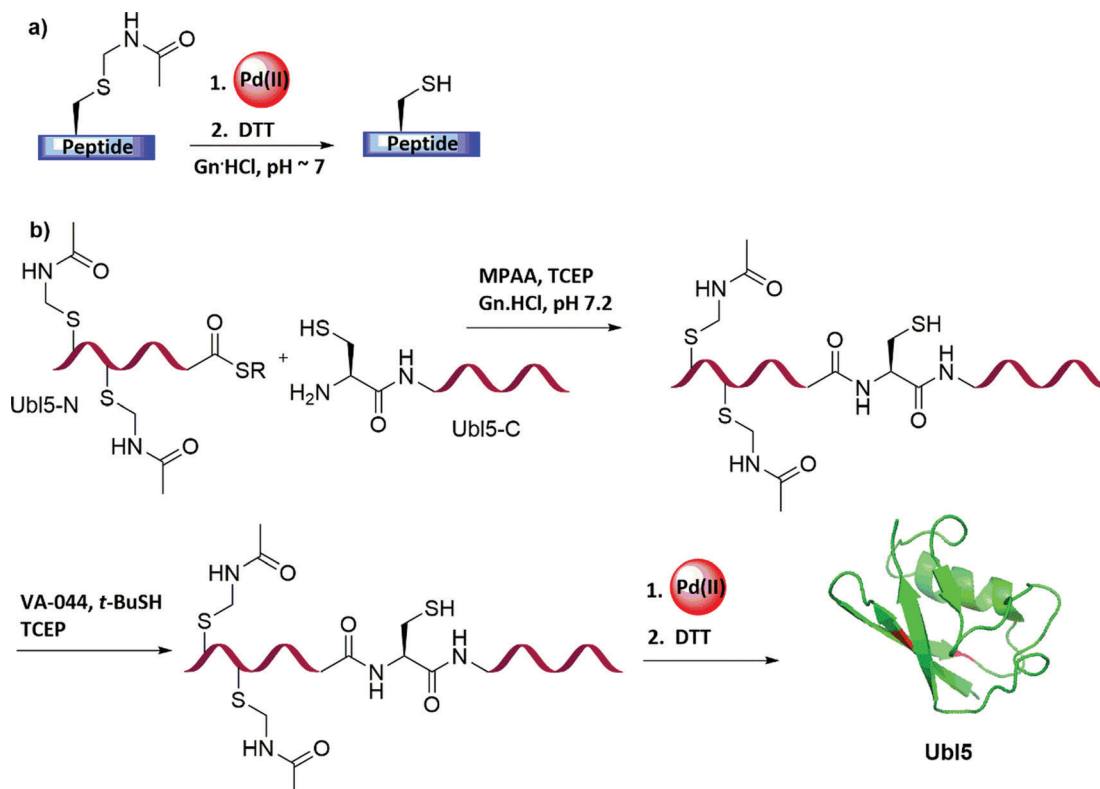


Fig. 11 Transition metal-mediated removal of AcM in aqueous medium for peptide and protein syntheses. (a) General scheme for the removal of AcM with Pd(II). (b) One-pot synthesis of Ubl5 employing AcM.

methods<sup>80</sup> were employed for the generation of C-terminal Asp thioesters, the formation of carboxylic anhydride was observed in 30% of the overall reaction products. However, in the case of the C-terminal hydrazide, the side-chain carboxyl group of Asp is prone to C-terminal cyclization upon oxidative switching, rendering it completely inactive in the ligation reaction.<sup>81</sup> Therefore, the carboxylate side-chain of Asp and Glu must be protected during ligation to avoid undesired product formation.

Botti and co-workers developed the PG 1-methyl-2-oxo-2-phenylethyl ester (OMop) for the side chain of Asp and Glu, which was introduced during SPPS.<sup>78</sup> This PG was found to be completely stable during HF cleavage and the ligation reaction. After ligation, the PG was removed by Zn in AcOH. In another report, Hojo's group protected the side chain of C-terminal Asp with a 2-(4-pyridyl)-2-propyl (PyP) group to prevent side reactions during NCL. Upon ligation, this PG could be selectively removed after 30 min by Zn in the presence of 3-mercaptopropionic acid (MPA).<sup>82</sup> This method was applied to the synthesis of human histone H4.

Our group recently reported that the protection of the side chain of the Asp and Glu with allyl functionality could serve as a temporal PG to prevent undesired cyclization of the C-terminal hydrazide (Fig. 12).<sup>83</sup> Upon ligation at these sites, the allyl group was rapidly (10 min) removed by [Pd(allyl)Cl]<sub>2</sub> and GSH under aqueous conditions. In this case, GSH was found to enhance the Pd reactivity and acted as scavenger for the released allyl group (Fig. 12 and Table 2).<sup>83</sup>

Parallel to our study, Okamoto and co-workers demonstrated NCL at the Asp–Cys junction, while protecting the Asp residues with an allylic group. (Fig. 13 and Table 2).<sup>84</sup> The removal step of the allylic moiety required less than 10 min upon the addition of Pd/TPPTS complex under the ligation conditions. Here, an excess amount of MPAAs was used as a scavenger for the  $\pi$ -allyl complex and Mg<sup>2+</sup> was added to speed up the deprotection reaction. This approach was demonstrated by the preparation of a fluorescein-labeled polypeptide, PEP-19.

## 5. Transition metal-mediated removal of reversible tags in protein synthesis and manipulations

### 5.1 Side-chain solubilizing tags

Chemists often encounter difficult sequences derived from proteins, which are problematic to make (*e.g.*,  $\beta$ -amyloid peptides) and to handle at the purification step or in ligation reactions (*e.g.*, membrane proteins).<sup>85</sup> Dissecting a polypeptide into segments to enable protein synthesis could lead to the formation of insoluble peptides in aqueous media.<sup>4,8</sup> To overcome these challenges, the concept of solubilizing tags has emerged, where a highly soluble tag is installed into the peptide fragments through a cleavable linker.<sup>86</sup> The linker should be stable to SPPS, compatible to ligation approaches, and capable of being removed in full aqueous media.

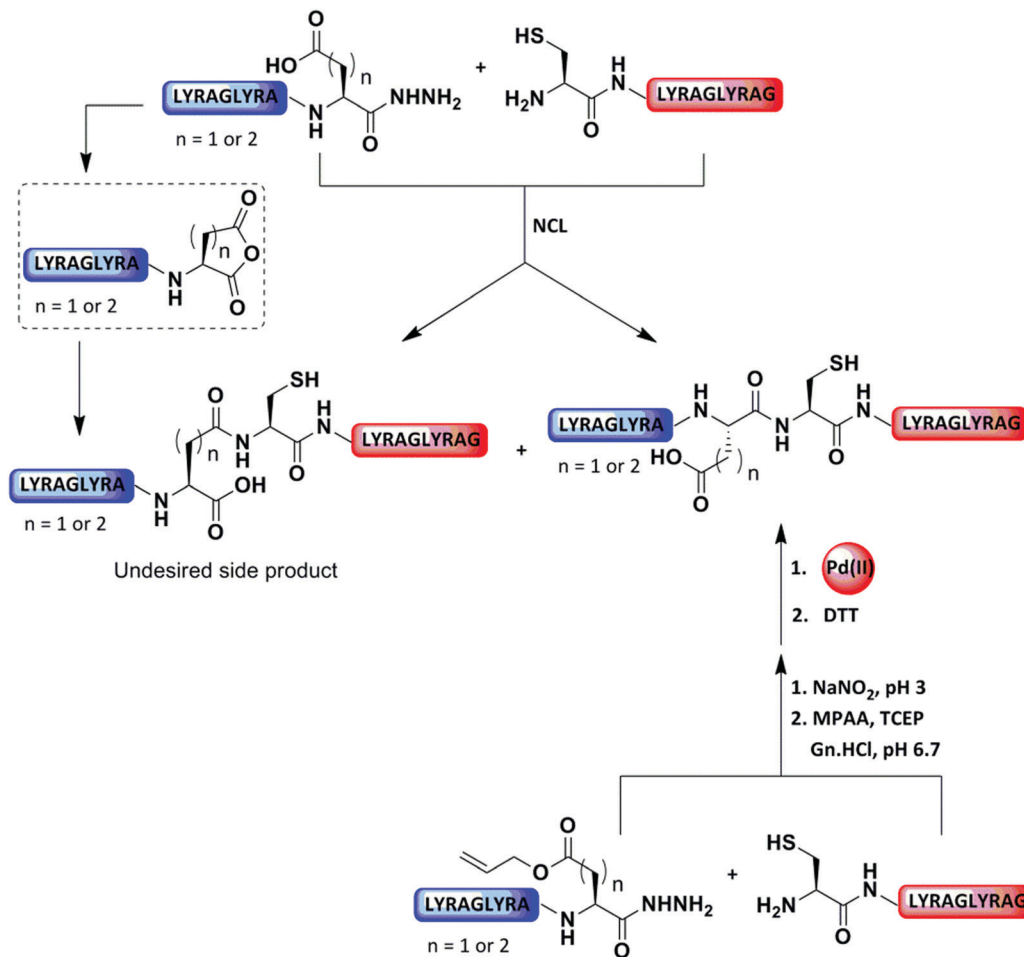


Fig. 12 Pd(II)-Mediated removal of allyl PG after the ligation at Asp/Glu for peptide synthesis.

Based on the Acm functionality, our group designed the synthesis of an Alloc-Phacm linker and introduced this to SPPS to aid the solubility of histone H4 C-terminal fragment (Fig. 14a).<sup>87</sup> This linker allowed the addition of several Arg (RRR-Phacm) as a solubilizing tag. Following the ligation step with the histone H4 N-terminal fragment, the reversible RRR-Phacm tag was completely removed using PdCl<sub>2</sub>, within 20 min, from the crude peptide under aqueous conditions. A final desulfurization step enabled the synthesis of the full-length histone protein H4 in a one-pot manner.<sup>74,87</sup>

In another work, our group attempted to synthesize ISGylated-Ub, which turned out to be extremely challenging due to its hydrophobic nature and high tendency for its aggregation as well as its fragments during SPPS (Fig. 14b).<sup>88</sup> We envisioned that the introduction of a cleavable solubilizing tag could aid the solubility. Consequently, we designed a new reversible Acm solubilizing tag in which the  $\alpha$ -carbon of the Acm was modified with a dimethylated amine moiety (Acm-NMe<sub>2</sub>).<sup>88</sup> The key benefits of this tag were the positively charged amino groups lowered the aggregation tendency and there was less steric hindrance caused by the Acm-NMe<sub>2</sub> group compared to the Ph-Acm-based tag. This solubilizing tag efficiently assisted the SPPS, assembly of the peptide segments by ligation,

and the HPLC purification throughout the synthetic process. Upon the ISGylation of Ub through Lys 29, this temporal tag could be liberated by applying PdCl<sub>2</sub> under aqueous conditions. Finally, a desulfurization step followed by folding afforded the target protein.<sup>88</sup>

## 5.2 Backbone modifications

Interest in the introduction of a responsive element to an external stimulus in peptides and proteins for various purposes has led to the use of different strategies, such as the expression of unnatural amino acids<sup>89,90</sup> and the (semi)chemical synthesis of proteins.<sup>4,91,92</sup>

We recently reported the backbone modification of a synthetic protein, namely ubiquitin (Ub)-aldehyde attached with a cell-penetrating peptide (CPP) through the Thz cleavable linker as a backbone (Fig. 15).<sup>93,94</sup> We then delivered the engineered protein into live cells (DU-145 cells) with the help of CPP, and subsequently the Thz was cleaved by Pd as an external stimulus. Using this approach, we were able to generate the Ub C-terminal aldehyde as an electrophile, which resulted in an enhanced inhibition of the cellular deubiquitinating enzyme (DUB), USP2a. The inhibition of USP2a in the cellular environment

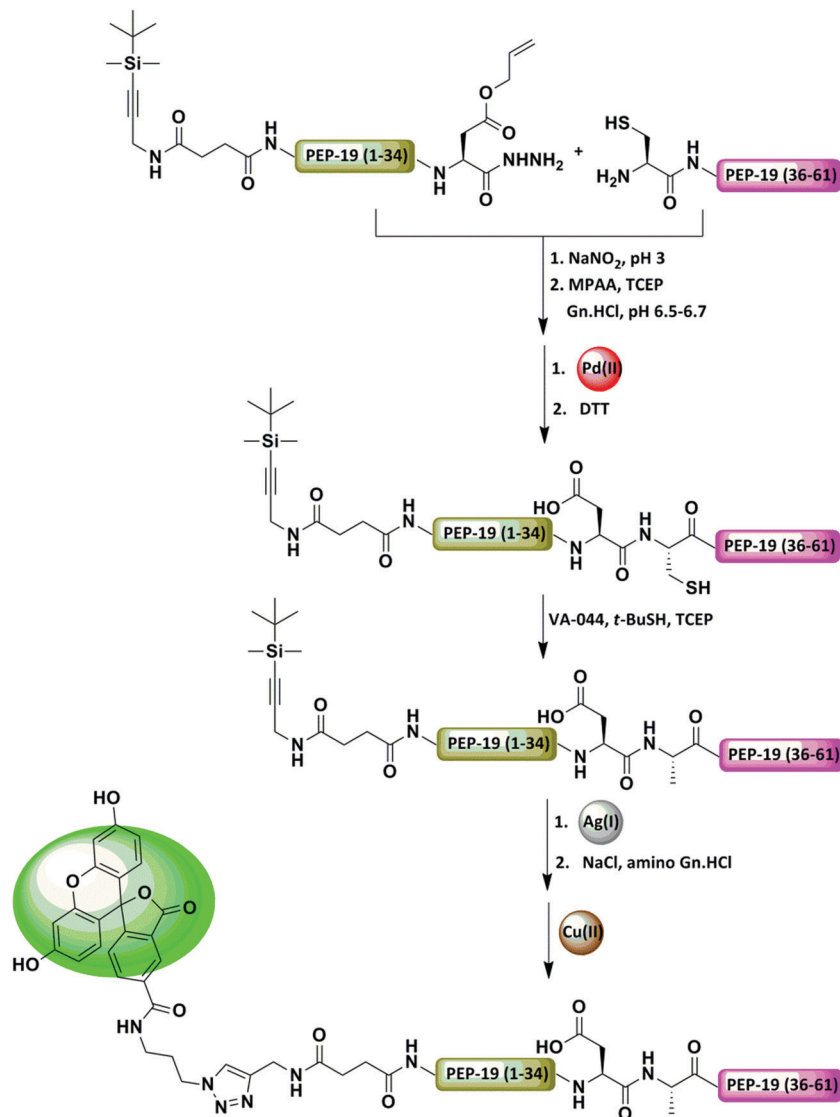


Fig. 13 Pd(II)-Mediated removal of allyl PG to facilitate PEP-19 synthesis.

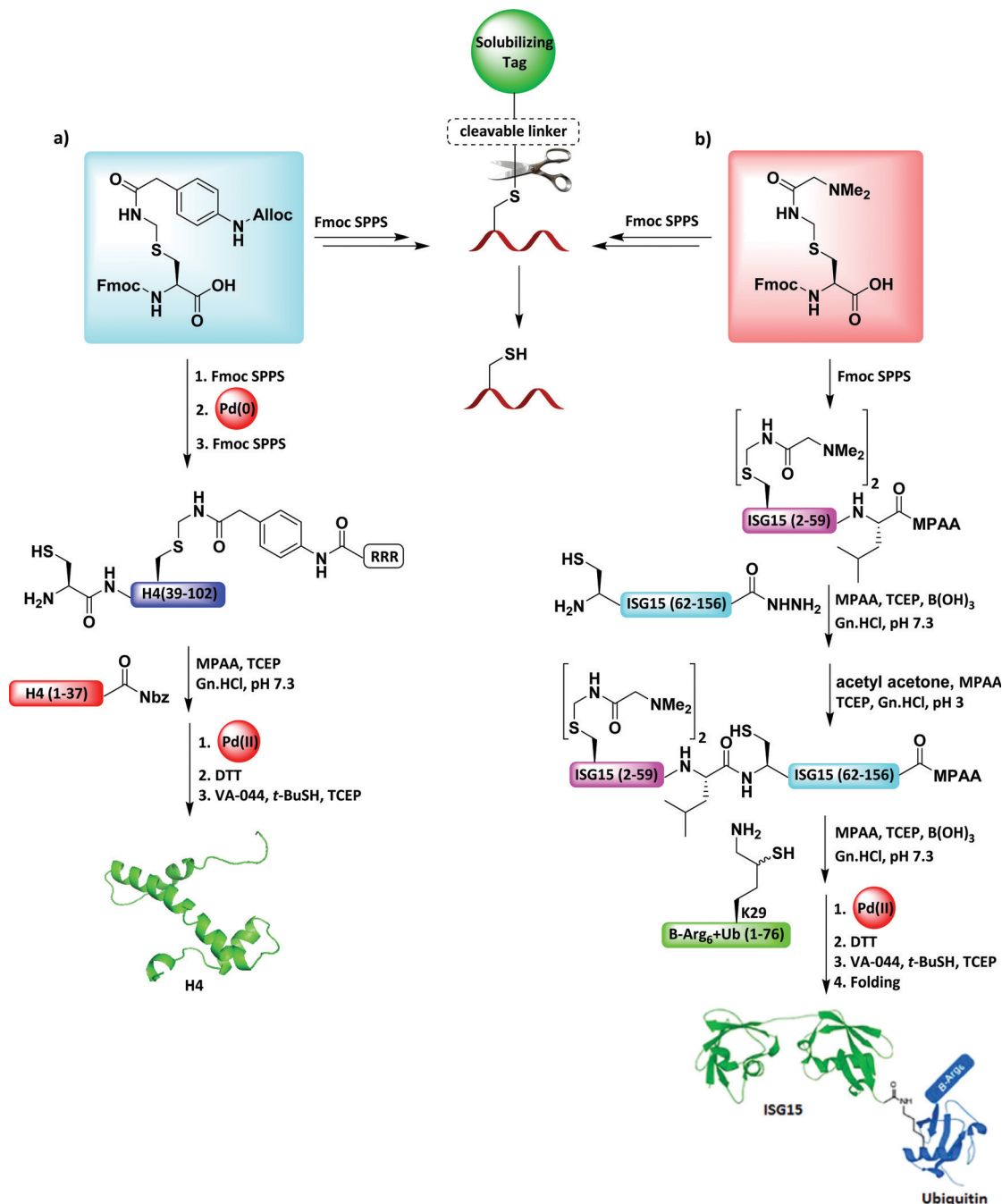
was supported by western blot, using the antibody against MDM2, which is a known substrate for USP2a. The fluorescence quenching pair sulfo-cyanine 3/5 (Cy3/Cy5) were used to monitor the entire process.

The incorporation of reversible backbone modification is a useful strategy to interrupt the secondary structures forming during SPPS or after cleavage, therefore assisting in avoiding peptide aggregation on the resin or in solution.<sup>95,96</sup> The pseudoproline dipeptide introduced by Mutter and co-workers is restricted to particular amino acids (*e.g.*, Ser) and often cleaved using TFA.<sup>97</sup> Our group recently reported the depropargylation of amides in aqueous media utilizing gold [Au(I)] (Fig. 16 and Table 2).<sup>98</sup> The incorporation of temporary propargylated amides in the sequence of NEDD8 protein enabled its efficient synthesis due to the effect of backbone modification on the peptide solubility. We also discovered a rapid site-selective amide bond cleavage, which depended on the position of the propargylation site (Gly-XX).<sup>98</sup>

## 6. Orthogonal removal of different PGs with transition metals

### 6.1 Orthogonal removal of Acm vs. Thz

The incorporation of orthogonal PGs is extremely important for the synthetic scheme of branched proteins and/or to guide the synthetic direction (*e.g.*, in the case of sequential NCL and late-stage functionalization). Recently, we reported the Pd-mediated on-demand deprotection of Thz and Acm groups in a fully aqueous medium (Fig. 17a).<sup>99</sup> We also showed the efficient removal of *t*-Bu PG using Pd and demonstrated its stability under the removal conditions of Thz and Acm (Fig. 17b).<sup>99</sup> This chemistry has been applied for multiple selective functionalizations of peptides containing Cys residues and in the challenging total chemical synthesis of the copper storage protein 1 (Fig. 17c).<sup>99</sup> Utilizing these synthetic tools, we also accomplished the chemical synthesis of an activity-based probe (ABP). The ABP was composed of histone H2A, which was



**Fig. 14** Pd(II)-Mediated removal of Acm-derived solubilizing tags to facilitate protein synthesis. (a) Histone H4 total synthesis. (b) ISGylated ubiquitin total synthesis (B-Arg<sub>6</sub> refers to biotinylated six Arg).

specifically ubiquitinated at Lys119 and contained dehydroalanine (DHA), at the junction between H2A and Ub, as a Cys proteases trap. This ABP reacted selectively with the Calypso/ASX heterodimer DUB, *via* a Michael addition reaction, to furnish a nucleosome-enzyme complex stabilized by a covalent C–S bond.<sup>99</sup>

## 6.2 Orthogonal removal of Acm vs. Alloc

Okamoto's group developed another selective strategy relying on the different properties of Pd complexes for the selective removal of the Alloc PG for N-terminal Cys, in the presence of internal Cys

protected with Acm, (Fig. 18).<sup>100</sup> In this approach, the Alloc PG was cleaved *via* a Pd(0)-mediated reaction, while the Acm was effectively unmasked with Pd(II).<sup>100</sup> The applicability of this chemistry has been demonstrated in the total chemical synthesis of histone H3 bearing a trimethylated Lys9 (K9me<sub>3</sub>) through the orthogonal removal of Alloc and Acm groups.<sup>100</sup>

## 6.3 Orthogonal removal of Thz vs. maleimide

As discussed in Sections 6.1 and 6.2, most of the PGs (*e.g.*, Acm) employed in peptide and protein synthesis need to be installed

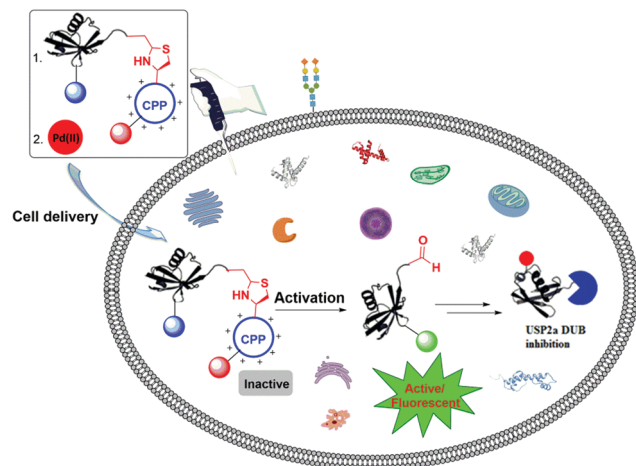


Fig. 15 Pd(II)-Mediated cleavage of a Ub activity-based probe with a Thz-modified backbone in live cells.

during the synthesis of peptide fragments on SPPS. Our group recently developed maleimide as a PG, which could be attached or detached on demand to the Cys residues of unprotected peptides or proteins in aqueous conditions (Fig. 19).<sup>101</sup> The maleimide group was efficiently removed by employing PdCl<sub>2</sub>/MgCl<sub>2</sub> and dithiothreitol (DTT). To demonstrate the synthetic applicability, the maleimide group was employed as a stable linker during solid-phase chemical ligation for the synthesis of Ub-ABPs. To elongate the polypeptide on the solid support, the N-terminal-protected Thz was removed selectively against the maleimide group by using [Pd(allyl)Cl]<sub>2</sub>/GSH.<sup>101</sup> After multiple ligations and desulfurization, the elongated polypeptide was

then liberated efficiently from the solid support by cleaving the maleimide handle using PdCl<sub>2</sub>/MgCl<sub>2</sub> (Fig. 19 and Table 2). In the final step, the liberated polypeptide was subjected with bisamide to form the corresponding Ub-ABPs.<sup>101</sup>

## 7. Pd in the synthesis of peptides and proteins bearing disulfide bonds

For over six decades, tremendous efforts have been invested to synthesize peptides and proteins bearing multiple disulfide bonds, yet no effective and/or general synthetic scheme has been devised to date, despite the great potential of these products for various applications.<sup>102</sup> Unfortunately, performing studies with isolated or biologically expressed peptides and proteins with disulfide bonds is very challenging for various targets, which has prompted the search for efficient synthetic approaches for their production. Two main strategies were adopted along this long journey: (1) the metal-free approach, which is based on oxidative folding and depends on the folding pathway of the particular target. However, this strategy suffers mainly from long reaction times (up to several days), low yields, and a lack of success in many cases; (2) a strategy based on forming disulfide bonds, which builds on the stepwise thiol deprotection and oxidation. In this approach, orthogonal PGs are applied for the side-chain Cys, such as of S-Trt, Acm, and *t*-Bu groups.<sup>102,103</sup> Nevertheless, the removal conditions of these PGs and the requirement for purification before and after these steps reduce the practicability of this approach due to the long processing time, significant side reactions, and extremely

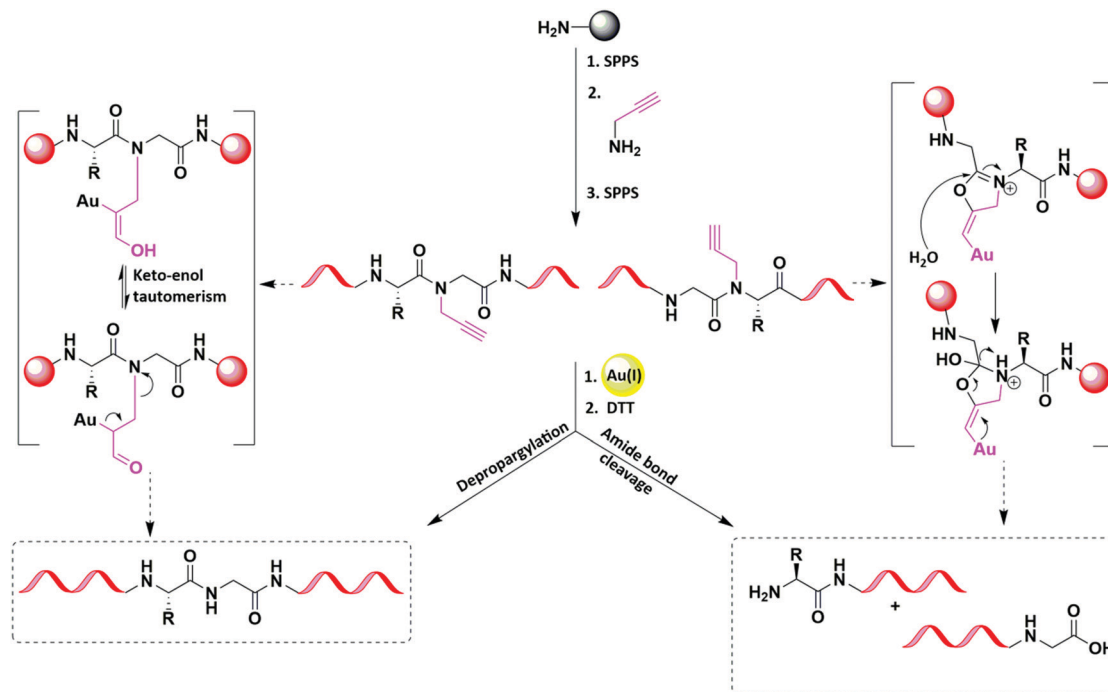


Fig. 16 Au(I)-Mediated removal of the backbone facilitated the chemical synthesis of the protein and the cleavage reaction when reversing the residue order.

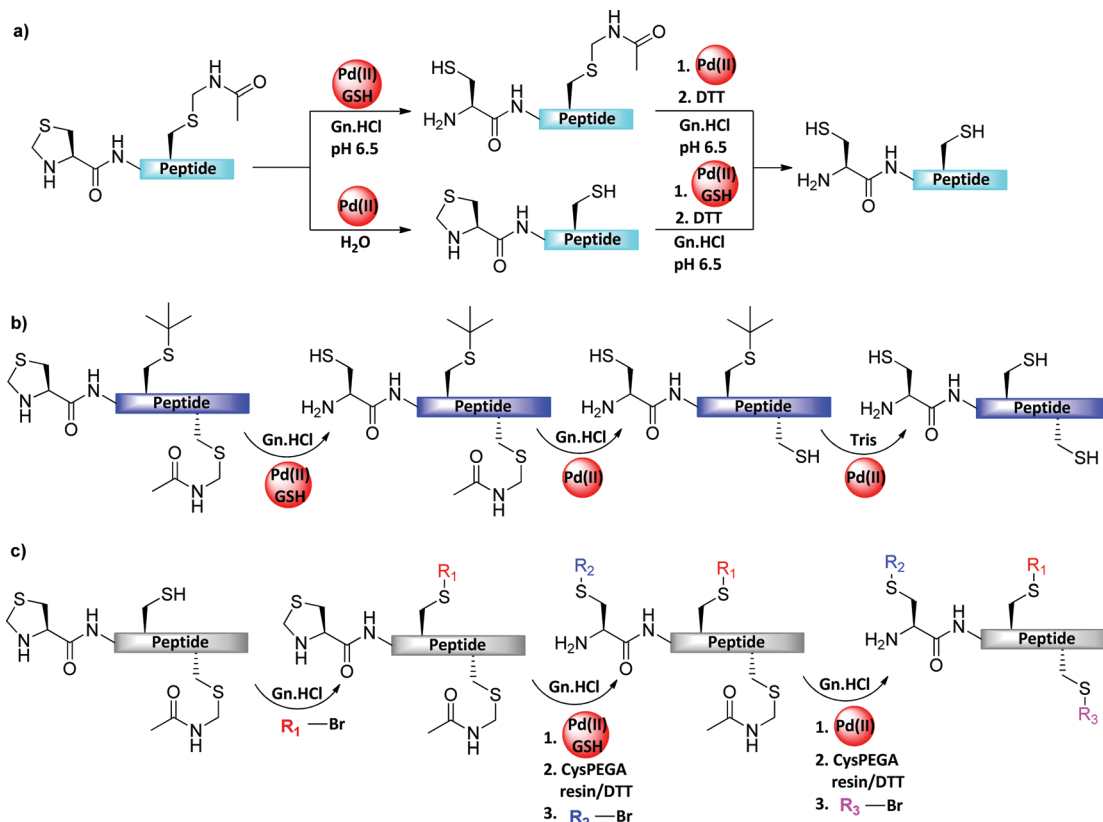


Fig. 17 Cys deprotection on demand: (a) Pd(II)-mediated selective removal of AcM and Thz groups, (b) Pd(II)-mediated selective removal of AcM, Thz, and *t*-butyl groups, (c) Pd(II)-mediated site-specific and differential modification of a peptide.

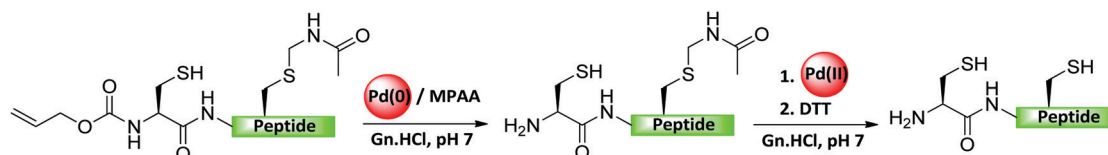


Fig. 18 Pd-Mediated chemoselective removal of Alloc and AcM groups.

low yields, meaning millions of potentially therapeutic active targets still remain chemically inaccessible.<sup>104,105</sup>

### 7.1 Pd-Mediated single disulfide bond formation

Recently, our group developed a method for accomplishing single disulfide bond formation between two Cys(AcM) in a one-pot process using NCL, Pd chemistry, and diethyldithiocarbamate (DTC) (Fig. 20a).<sup>106</sup> The crucial role of using DTC over DTT in this reaction was attributed to its Pd scavenging property and not its reducing property, like for DTT, which cleaves the disulfide bond. The applicability of this method was demonstrated by the synthesis of oxytocin and the first total chemical synthesis of thioredoxin-1 (Trx-1) enzyme *via* NCL, desulfurization, and a one-pot disulfide bond formation.<sup>106</sup> Nevertheless, under these conditions, all attempts to create more than one disulfide bond in a regioselective manner failed and resulted in significant reshuffling, probably due to the formation of a Pd-S intermediate that destabilizes the already existing disulfide bonds *via* a yet unknown pathway.

Similarly, the maleimide group was utilized as a PG of Cys residues for intein-based expressed protein fragment [Trx-1(2–93)-Mesna] to enable (EPL)/desulfurization and subsequent direct single disulfide bond formation to give the full sequence of Trx-1 (Fig. 20b).<sup>101</sup> Here, PdCl<sub>2</sub>/MgCl<sub>2</sub> assisted the removal of the maleimide groups and DTC was used to efficiently recover the final product.

### 7.2 Ultrafast regioselective formation of multiple disulfide bonds in one pot

Very recently, we reported a novel synthetic strategy for the ultrafast, high-yielding formation of multiple disulfide bonds in peptides and proteins.<sup>107</sup> Analysis of the current approaches and their main limitations prompted us to design a synthetic pathway in which: (1) the total process will be completed as fast as possible to avoid reshuffling and side reactions on sensitive AAs (*e.g.*, Met and Trp); (2) the solvent will desaturate the peptide/protein (preferably aqueous) to overcome kinetically



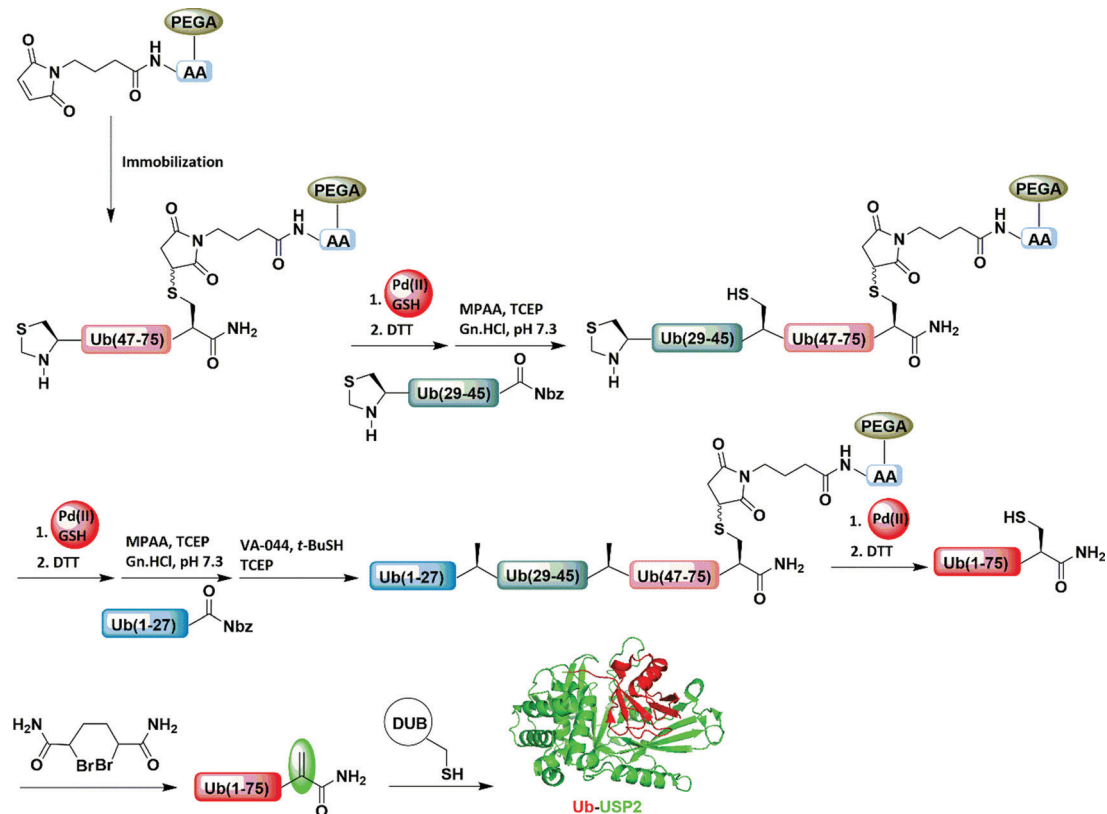


Fig. 19 Pd(II)/GSH-mediated chemoselective removal of Thz against maleimide.

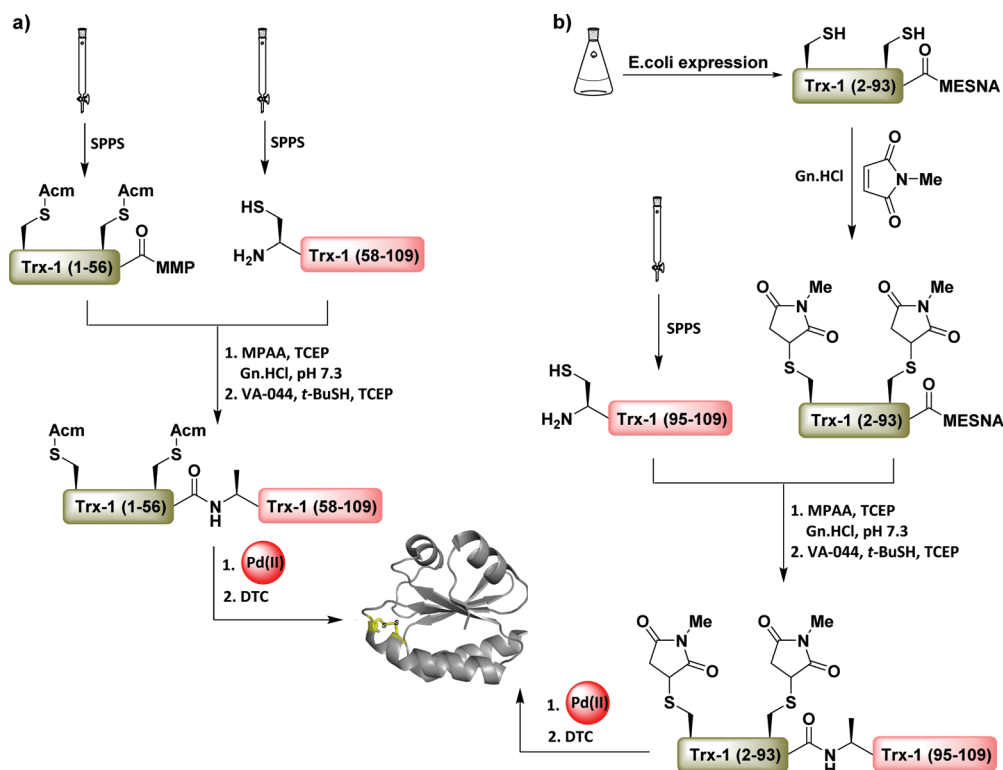


Fig. 20 Pd(II)-Mediated one-pot disulfide bond formation for the total (semi) synthesis of Trx-1 employing Cys (Acm) (a) and Cys (maleimide) (b).

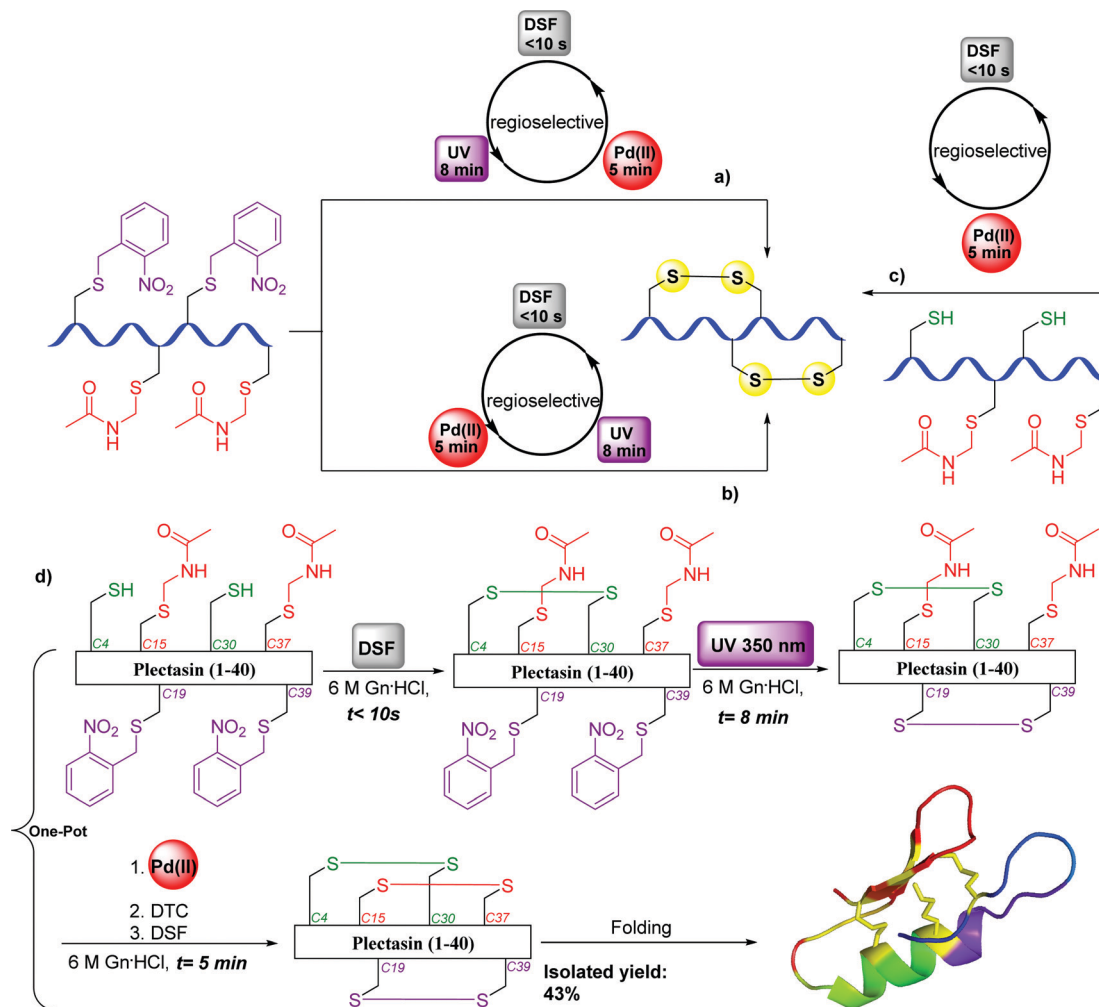


Fig. 21 One-pot formation of disulfide bonds: (a) general scheme for the regioselective formation of two disulfide bonds via chemoselective UV-light irradiation in the presence of DSF followed by Pd(II) treatment in one pot; (b) regioselective formation of two disulfide bonds via chemoselective Pd(II) treatment in the presence of DSF followed by UV-light irradiation in one pot; (c) regioselective formation of two disulfide bonds via exposure to DSF followed by Pd(II) treatment in one pot; (d) regioselective synthesis of the antimicrobial plectasin peptide bearing three disulfide bonds.

trapped folded intermediates and/or aggregations; and (3) purifications/isolations during the steps will be avoided (one-pot manner) to increase the synthesis yield. To achieve this, we designed an approach based on a combination of the small molecule and extremely active disulfiram (DSF), UV-light at 350 nm, and Pd(II) under pH tuning for the chemo- and regioselective activation of multiple Cys residues.<sup>107</sup> Applying these conditions enabled the efficient one-pot syntheses of representative examples (in only 5 min for two- and 13 min for three disulfide bonds) from various peptide/protein families (Fig. 21), tackling a decades old standing problem in peptide and protein chemistry.<sup>107</sup>

## 8. Synthesis of selenoprotein

The biological function of many Sec-containing proteins remains unclear or poorly studied.<sup>108</sup> The incorporation of Sec into selenoproteins was achieved co-translationally by the suppression of the UGA stop codon.<sup>109,110</sup> However, this requires unusual

tRNA, a Sec insertion sequence element, a Sec-specific elongation factor, and other factors.<sup>111–113</sup> Therefore it is highly difficult to obtain the desired amounts of wild-type selenoproteins by the recombinant expression method, hence chemical (semi)synthesis has emerged as an elegant approach to solve this issues.<sup>114</sup>

The protection of Sec is highly necessary when aiming to perform the sequential ligations to enable selenoprotein synthesis. Recently, Metanis' group developed selenazolidine (Sez) as an easily accessible N-terminal-protected form of Sec for the chemical synthesis of proteins.<sup>115</sup> However, the generally used methoxylamine for the conversion of Sez-to-Sec requires a long reaction time similar to Thz-to-Cys conversion, and results in unwanted side products after reacting with the C-terminal thioester.<sup>115</sup> To address these issues, Metanis' group developed a Cu-mediated Sez deprotection for the synthesis of challenging proteins (Fig. 22a and Table 2).<sup>116</sup> In this study, the group established the chemical synthesis of peptides and proteins using Cu(II)-mediated Sez deprotection, followed by NCL and deselenization, in a one-pot manner.<sup>116</sup> The group also reported that among other metals

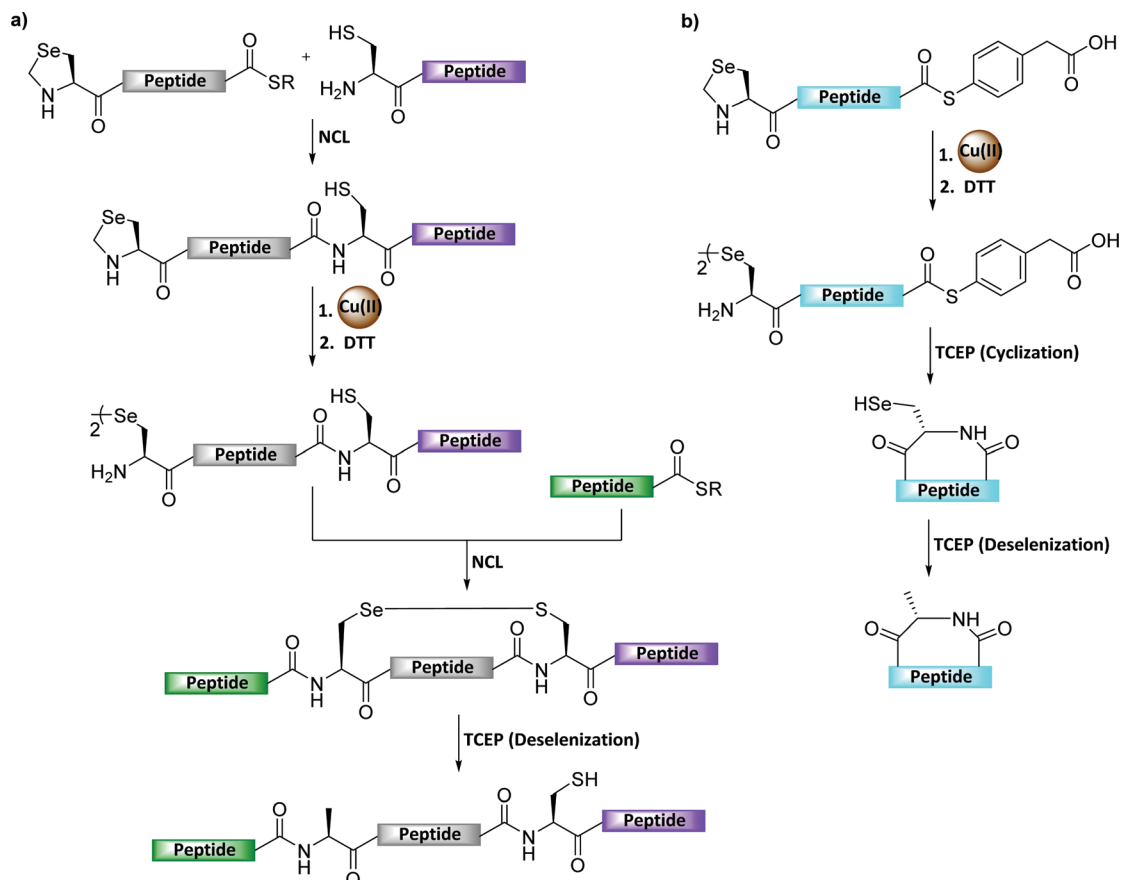


Fig. 22 (a) Cu-Mediated Sez deprotection in the chemical synthesis of seleno-peptides. (b) Cu-mediated Sez deprotection in the chemical synthesis of cyclic peptides.

the most efficient additive for Sez unmasking was Cu(II), which could complete the process within 30 min. The utility of this method was demonstrated in the syntheses of human thiosulfate:glutathione sulfurtransferase (TSTD1) and an analogue bearing Sec. In addition, this chemistry was extended for the preparation of cyclic peptides by sequential Sez deprotection using Cu and intramolecular cyclization in a one-pot manner (Fig. 22b).<sup>117</sup> The usefulness of this method was also demonstrated by the synthesis of cyclotide Se-Kalata S, in which the Sez-linear peptide-MPAA was synthesized using Fmoc-SPPS followed by the deprotection of Sez by CuCl<sub>2</sub>.<sup>117</sup>

In a closely related study by Otaka's group, they investigated the application of the Cu-mediated ring opening of the Thz to practical peptide synthesis. Their study revealed that the opening of Thz with CuSO<sub>4</sub> in the presence of ascorbate proceeded within a few hours. In addition, they observed the opening of Sez under similar conditions.<sup>118</sup> These tools have been applied for the chemical synthesis of 77-mer CXCL14 protein. In this study, the opening of Sez in a model peptide was also achieved with the use of a lower amount of Cu salt than for the Thz opening and therefore this process was applicable for one-pot/sequential reactions.<sup>118</sup>

## 9. Summary and outlook

In this review, we discussed the chemical synthesis of peptides and proteins assisted by transition metals as effective reagents

for resolving various issues that arise with conventional methods during their syntheses. Using transition metals in peptide and protein synthesis enabled the reactions to be performed under mild conditions, with high selectivity and in a short time and as a one-pot process, and gave the products in high yields.

With the increasing demand on the synthesis of complex peptides (*e.g.*, with disulfide bonds) and proteins (ubiquitinated proteins),<sup>119</sup> it became clear that the synthetic scheme requires PGs.<sup>4</sup> On the other hand, these PGs must be removed under mild conditions, in a selective and orthogonal manner while being unreactive to other functional groups present in the sequence. In this regard, transition metals, especially Pd, chemistry offer the chance for the one-pot synthesis of complex protein targets and orthogonal cleavage of different PGs.<sup>69,99</sup> This chemistry was also extended to the efficient formation of single disulfide bonds directly from the protected form of the Cys, *e.g.*, Cys(Acm).<sup>106</sup> Recently, Cu chemistry was also found to assist the removal of selenazolidine during the synthesis of selenoproteins.<sup>116,118</sup>

Pd chemistry played a crucial role in achieving the cleavage of temporally incorporated solubilizing tags, which were applied to aid the solubility of hydrophobic peptide fragments. Using this approach, several proteins with difficult sequences have been synthesized.<sup>74,87,88,99</sup> In another direction, Au assisted also in the removal of backbone modifications of peptides and proteins that were applied to improve their synthesis.<sup>98</sup> It is worth

mentioning that transition metal chemistry has been extended to the cellular environment, where it helps to degrade the temporal PGs or cleave the backbone elements of engineered peptides and proteins to aid studying the biochemical properties in the native environment.<sup>93,94</sup> Despite the relative toxicity of some of the transition metals (e.g., Tl, Hg) being well established in various examples,<sup>120</sup> the use of appropriate metal scavengers could afford highly purified and active proteins. For example, inductively coupled plasma (ICP) measurements for Pd traces after scavenging with DTT showed that the amount of Pd detected was below 0.07% of the protein.<sup>87,98,101,106</sup> Moreover, metals such as Pd, Cu, and others have been shown to be fully compatible with the live cell environment.<sup>94,121–125</sup>

Despite possessing several advantages, many aspects remain to be addressed regarding the use of transition metals in peptide and protein chemistry. Can highly abundant and low-cost metals such as Fe, Ni be used instead of expensive or toxic metals? The newly evolved chemistry must be selective, must work under mild conditions, and ideally should work in catalytic amounts and with the possibility to work in the cellular environment. Such new inventions could further bring forward the field of the chemical synthesis of peptides and proteins and their modifications.

Future directions in this exciting research area include the employment of transition metal nanoparticles<sup>126</sup> and metal-loaded water-compatible resin for peptide and protein synthesis,<sup>127</sup> which have been proved to be excellent catalytic systems in modern organic synthesis. In addition, further studies should be performed in order to understand the mechanism of these transformations under aqueous media and in the presence of the various additives that are normally used in ligation chemistry, such as MPAA, TCEP, and guanidine. Such an understanding could have great implications in the design of new complexes or catalytic reactions for peptides and proteins as well as for other biomolecules (e.g., DNA). These developments might also inspire the scientific community to perform small molecule reactions in aqueous conditions or to use the newly discovered complexes in new chemical transformations.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

A. Brik holds The Jordan and Irene Tark Academic Chair. We would like to thank the European Research Council (ERC) under the European Union's Horizon 2020 research and innovation program for funding (grant agreement no. 831783). S. L. thanks the Technion for the Jacobs fellowship for excellence.

## References

- 1 T. S. Young and P. G. Schultz, *J. Biol. Chem.*, 2010, **285**, 11039–11044.
- 2 Y. Zhang, J. L. Ptacin, E. C. Fischer, H. R. Aerni, C. E. Caffaro, K. San Jose, A. W. Feldman, C. R. Turner and F. E. Romesberg, *Nature*, 2017, **551**, 644–647.
- 3 J. W. Chin, *Nature*, 2017, **550**, 53–60.
- 4 S. Bondalapati, M. Jbara and A. Brik, *Nat. Chem.*, 2016, **8**, 407–418.
- 5 S. S. Kulkarni, J. Sayers, B. Premdjee and R. J. Payne, *Nat. Rev. Chem.*, 2018, **2**, 0122.
- 6 V. Agouridas, O. El Mahdi, V. Diemer, M. Cargoët, J. C. M. Monbaliu and O. Melnyk, *Chem. Rev.*, 2019, **119**, 7328–7443.
- 7 R. B. Merrifield, *J. Am. Chem. Soc.*, 1963, **85**, 2149–2154.
- 8 I. Coin, M. Beyermann and M. Bienert, *Nat. Protoc.*, 2007, **2**, 3247–3256.
- 9 S. B. H. Kent, P. E. Dawson, T. W. Muir and I. Clark-Lewis, *Science*, 1994, **266**, 776–779.
- 10 J. W. Bode, R. M. Fox and K. D. Baucom, *Angew. Chem., Int. Ed.*, 2006, **45**, 1248–1252.
- 11 Y. Zhang, C. Xu, H. Y. Lam, C. L. Lee and X. Li, *Proc. Natl. Acad. Sci. U. S. A.*, 2013, **110**, 6657–6662.
- 12 M. Jbara, S. K. Maity and A. Brik, *Angew. Chem., Int. Ed.*, 2017, **56**, 10644–10655.
- 13 M. Jbara, *Pure Appl. Chem.*, 2021, DOI: 10.1515/pac-2020-0504.
- 14 P. G. Isenegger and B. G. Davis, *J. Am. Chem. Soc.*, 2019, **141**, 8005–8013.
- 15 C. Zhang, E. V. Vinogradova, A. M. Spokoiny, S. L. Buchwald and B. L. Pentelute, *Angew. Chem., Int. Ed.*, 2019, **58**, 4810–4839.
- 16 A. Brik, *Adv. Synth. Catal.*, 2008, **350**, 1661–1675.
- 17 E. Latocheski, G. M. Dal Forno, T. M. Ferreira, B. L. Oliveira, G. J. L. Bernardes and J. B. Domingos, *Chem. Soc. Rev.*, 2020, 7710–7729.
- 18 J. L. Mascarenas, P. Destito, C. Vidal and F. López, *Chem. Eur. J.*, 2021, DOI: 10.1002/chem.202003927.
- 19 W. Wang, M. M. Lorion, J. Shah, A. R. Kapdi and L. Ackermann, *Angew. Chem., Int. Ed.*, 2018, **57**, 14700–14717.
- 20 L. Z. Yan and P. E. Dawson, *J. Am. Chem. Soc.*, 2001, **123**, 526–533.
- 21 Y. Y. Yang, S. Ficht, A. Brik and C. H. Wong, *J. Am. Chem. Soc.*, 2007, **129**, 7690–7701.
- 22 B. L. Pentelute and S. B. H. Kent, *Org. Lett.*, 2007, **9**, 687–690.
- 23 J. Blake, *Int. J. Pept. Protein Res.*, 1981, **17**, 273–274.
- 24 S. Aimoto, N. Mizoguchi, H. Hojo and S. Yoshimura, *Bull. Chem. Soc. Jpn.*, 1989, **62**, 524–531.
- 25 H. Hojo, E. Haginoya, Y. Matsumoto, Y. Nakahara, K. Nabeshima, B. P. Toole and Y. Watanabe, *Tetrahedron Lett.*, 2003, **44**, 2961–2964.
- 26 G. Chen, Q. Wan, Z. Tan, C. Kan, Z. Hua, K. Ranganathan and S. J. Danishefsky, *Angew. Chem., Int. Ed.*, 2007, **46**, 7383–7387.
- 27 P. E. Dawson, *Isr. J. Chem.*, 2011, **51**, 862–867.
- 28 Q. Wan and S. J. Danishefsky, *Angew. Chem., Int. Ed.*, 2007, **46**, 9248–9252.
- 29 K. J. Rosengren, R. J. Clark, N. L. Daly, U. Göransson, A. Jones and D. J. Craik, *J. Am. Chem. Soc.*, 2003, **125**, 12464–12474.
- 30 K. A. Wilson, M. Kalkum, J. Ottesen, J. Yuzenkova, B. T. Chait, R. Landick, T. Muir, K. Severinov and S. A. Darst, *J. Am. Chem. Soc.*, 2003, **125**, 12475–12483.

- 31 M. J. Bayro, J. Mukhopadhyay, G. V. T. Swapna, J. Y. Huang, L. C. Ma, E. Sineva, P. E. Dawson, G. T. Montelione and R. H. Ebright, *J. Am. Chem. Soc.*, 2003, **125**, 12382–12383.
- 32 A. Brik, Y. Y. Yang, S. Ficht and C. H. Wong, *J. Am. Chem. Soc.*, 2006, **128**, 5626–5627.
- 33 A. Brik, S. Ficht, Y. Y. Yang, C. S. Bennett and C. H. Wong, *J. Am. Chem. Soc.*, 2006, **128**, 15026–15033.
- 34 L. E. Canne, S. J. Bark and S. B. H. Kent, *J. Am. Chem. Soc.*, 1996, **118**, 5891–5896.
- 35 C. E. Weller, A. Dhall, F. Ding, E. Linares, S. D. Whedon, N. A. Senger, E. L. Tyson, J. D. Bagert, X. Li, O. Augusto and C. Chatterjee, *Nat. Commun.*, 2016, **7**, 1–10.
- 36 E. C. B. Johnson and S. B. H. Kent, *J. Am. Chem. Soc.*, 2006, **128**, 6640–6646.
- 37 D. F. Veber, J. D. Milkowski, S. L. Varga, R. G. Denkwalter and R. Hirschmann, *J. Am. Chem. Soc.*, 1972, **94**, 5456–5461.
- 38 S. Liu, B. L. Pentelute and S. B. H. Kent, *Angew. Chem., Int. Ed.*, 2012, **51**, 993–999.
- 39 M. T. Weinstock, M. T. Jacobsen and M. S. Kay, *Proc. Natl. Acad. Sci. U. S. A.*, 2014, **111**, 11679–11684.
- 40 M. Seenaiiah, M. Jbara, S. M. Mali and A. Brik, *Angew. Chem., Int. Ed.*, 2015, **54**, 12374–12378.
- 41 S. K. Mong, A. A. Vinogradov, M. D. Simon and B. L. Pentelute, *ChemBioChem*, 2014, **15**, 721–733.
- 42 Y. David, M. Vila-Perelló, S. Verma and T. W. Muir, *Nat. Chem.*, 2015, **7**, 394–402.
- 43 Q. Zhang, E. V. Johnston, J. H. Shieh, M. A. S. Moore and S. J. Danishefsky, *Proc. Natl. Acad. Sci. U. S. A.*, 2014, **111**, 2885–2890.
- 44 N. Fujii, A. Otaka, S. Funakoshi, K. Bessho, T. Watanabe, K. Akaji and H. Yajima, *Chem. Pharm. Bull.*, 1987, **35**, 2339–2347.
- 45 N. Fujii, A. Otaka, T. Watanabe, A. Okamachi, H. Tamamura, H. Yajima, Y. Inagaki, M. Nomizu and K. Asano, *J. Chem. Soc., Chem. Commun.*, 1989, 283–284.
- 46 H. Lamthanh, C. Roumestand, C. Deprun and A. Ménez, *Int. J. Pept. Protein Res.*, 1993, **41**, 85–95.
- 47 T. Kawakami, S. Yoshimura and S. Aimoto, *Tetrahedron Lett.*, 1998, **39**, 7901–7904.
- 48 S. B. H. Kent, *J. Am. Chem. Soc.*, 2004, **126**, 1377–1383.
- 49 F. K. Deng, L. Zhang, Y. T. Wang, O. Schneewind and S. B. H. Kent, *Angew. Chem., Int. Ed.*, 2014, **53**, 4662–4666.
- 50 I. Sakamoto, K. Tezuka, K. Fukae, K. Ishii, K. Taduru, M. Maeda, M. Ouchi, K. Yoshida, Y. Nambu, J. Igarashi, N. Hayashi, T. Tsuji and Y. Kajihara, *J. Am. Chem. Soc.*, 2012, **134**, 5428–5431.
- 51 P. Siman, O. Blatt, T. Moyal, T. Danieli, M. Lebendiker, H. A. Lashuel, A. Friedler and A. Brik, *ChemBioChem*, 2011, **12**, 1097–1104.
- 52 P. Siman, S. V. Karthikeyan, M. Nikolov, W. Fischle and A. Brik, *Angew. Chem., Int. Ed.*, 2013, **52**, 8059–8063.
- 53 M. Hejjaoui, M. Haj-Yahya, K. S. A. Kumar, A. Brik and H. A. Lashuel, *Angew. Chem., Int. Ed.*, 2011, **50**, 405–409.
- 54 J. Li, Y. Li, L. Liu, Y. Li, Q. He and H. Li, *Org. Biomol. Chem.*, 2014, **12**, 5435–5441.
- 55 Y. K. M. Yoshida, T. Tatsumi, Y. Fujiwara, S. Iinuma, T. Kimura and K. Akaji, *Chem. Pharm. Bull.*, 1990, **38**, 1551–1557.
- 56 O. Nishimura, C. Kitada and M. Fujino, *Chem. Pharm. Bull.*, 1978, **26**, 1576–1585.
- 57 C. M. Stevens and R. Watanabe, *J. Am. Chem. Soc.*, 1950, **72**, 725–727.
- 58 H. Kunz and C. Unverzagt, *Angew. Chem., Int. Ed. Engl.*, 1984, **23**, 436–437.
- 59 G. B. F. Giube and O. Dangles, *Tetrahedron Lett.*, 1986, **27**, 2365–2368.
- 60 J. Alsina, F. Rabanal, E. Giralt and F. Albericio, *Tetrahedron Lett.*, 1994, **35**, 9633–9636.
- 61 P. Grieco, P. M. Gitu and V. J. Hruby, *J. Pept. Res.*, 2001, **57**, 250–256.
- 62 H. P. Hemantha, S. N. Bavikar, Y. Herman-Bachinsky, N. Haj-Yahya, S. Bondalapati, A. Ciechanover and A. Brik, *J. Am. Chem. Soc.*, 2014, **136**, 2665–2673.
- 63 D. Bang and S. B. H. Kent, *Angew. Chem., Int. Ed.*, 2004, **43**, 2534–2538.
- 64 M. Seenaiiah, M. Jbara, S. M. Mali and A. Brik, *Angew. Chem., Int. Ed.*, 2015, **54**, 12374–12378.
- 65 E. C. B. Johnson, E. Malito, Y. Shen, D. Rich, W. J. Tang and S. B. H. Kent, *J. Am. Chem. Soc.*, 2007, **129**, 11480–11490.
- 66 C. Piontek, D. Varón Silva, C. Heinlein, C. Pöhner, S. Mezzato, P. Ring, A. Martin, F. X. Schmid and C. Unverzagt, *Angew. Chem., Int. Ed.*, 2009, **48**, 1941–1945.
- 67 G.-M. Fang, J.-X. Wang and L. Liu, *Angew. Chem., Int. Ed.*, 2012, **51**, 10347–10350.
- 68 M. Jbara, S. K. Maity, M. Seenaiiah and A. Brik, *J. Am. Chem. Soc.*, 2016, **138**, 5069–5075.
- 69 S. K. Maity, M. Jbara, G. Mann, G. Kamnesky and A. Brik, *Nat. Protoc.*, 2017, **12**, 2293–2322.
- 70 N. Kamo, G. Hayashi and A. Okamoto, *Angew. Chem., Int. Ed.*, 2018, **57**, 16533–16537.
- 71 S. K. Maity, M. Jbara, S. Laps and A. Brik, *Angew. Chem., Int. Ed.*, 2016, **55**, 8108–8112.
- 72 S. Li, Y. Lin, J. Cao and S. Zhang, *J. Org. Chem.*, 2007, **72**, 4067–4072.
- 73 M. Jbara, N. Guttmann-Raviv, S. K. Maity, N. Ayoub and A. Brik, *Bioorg. Med. Chem.*, 2017, **25**, 4966–4970.
- 74 S. Bondalapati, E. Eid, S. M. Mali, C. Wolberger and A. Brik, *Chem. Sci.*, 2017, **8**, 4027–4034.
- 75 J. Liu, F. Zheng, R. Cheng, S. Li, S. Rozovsky, Q. Wang and L. Wang, *J. Am. Chem. Soc.*, 2018, **140**, 8807–8816.
- 76 D. Kobayashi, N. Naruse, M. Denda, A. Shigenaga and A. Otaka, *Org. Biomol. Chem.*, 2020, **18**, 8638–8645.
- 77 T. M. Hackeng, J. H. Griffin and P. E. Dawson, *Proc. Natl. Acad. Sci. U. S. A.*, 1999, **96**, 10068–10073.
- 78 M. Villain, H. Gaertner and P. Botti, *Eur. J. Org. Chem.*, 2003, 3267–3272.
- 79 B. Dang, T. Kubota, K. Mandal, F. Bezanilla and S. B. H. Kent, *J. Am. Chem. Soc.*, 2013, **135**, 11911–11919.
- 80 J. B. Blanco-Canosa, B. Nardone, F. Albericio and P. E. Dawson, *J. Am. Chem. Soc.*, 2015, **137**, 7197–7209.
- 81 G. M. Fang, Y. M. Li, F. Shen, Y. C. Huang, J. Bin Li, Y. Lin, H. K. Cui and L. Liu, *Angew. Chem., Int. Ed.*, 2011, **50**, 7645–7649.
- 82 Y. Asahina, T. Kawakami and H. Hojo, *Chem. Commun.*, 2017, **53**, 2114–2117.

- 83 M. Jbara, E. Eid and A. Brik, *Org. Biomol. Chem.*, 2018, **16**, 4061–4064.
- 84 N. Kamo, G. Hayashi and A. Okamoto, *Chem. Commun.*, 2018, **54**, 4337–4340.
- 85 T. Kobori, S. Iwamoto, K. Takeyasu and T. Ohtani, *Biopolymers*, 2007, **85**, 392–406.
- 86 J. Bin, Li, S. Tang, J. S. Zheng, C. L. Tian and L. Liu, *Acc. Chem. Res.*, 2017, **50**, 1143–1153.
- 87 S. K. Maity, G. Mann, M. Jbara, S. Laps, G. Kamnesky and A. Brik, *Org. Lett.*, 2016, **18**, 3026–3029.
- 88 E. Eid, G. N. Boross, H. Sun, M. Msallam, S. K. Singh and A. Brik, *Bioconjugate Chem.*, 2020, **31**, 889–894.
- 89 C. H. Kim, J. Y. Axup and P. G. Schultz, *Curr. Opin. Chem. Biol.*, 2013, **17**, 412–419.
- 90 J. W. Chin, *Annu. Rev. Biochem.*, 2014, **83**, 379–408.
- 91 J. Li and P. R. Chen, *Nat. Chem. Biol.*, 2016, **12**, 129–137.
- 92 T. W. Muir, *Annu. Rev. Biochem.*, 2003, **72**, 249–289.
- 93 M. Jbara, S. Laps, S. K. Maity and A. Brik, *Chem. – Eur. J.*, 2016, **22**, 14851–14855.
- 94 G. Mann, G. Satish, R. Meledin, G. B. Vamisetti and A. Brik, *Angew. Chem., Int. Ed.*, 2019, **58**, 13540–13549.
- 95 A. Isidro-Llobet, X. Just-Baringo, M. Álvarez and F. Albericio, *Biopolymers*, 2008, **90**, 444–449.
- 96 J. S. Zheng, Y. He, C. Zuo, X. Y. Cai, S. Tang, Z. A. Wang, L. H. Zhang, C. L. Tian and L. Liu, *J. Am. Chem. Soc.*, 2016, **138**, 3553–3561.
- 97 T. Wöhr, F. Wahl, A. Nefzi, B. Rohwedder, T. Sato, X. Sun and M. Mutter, *J. Am. Chem. Soc.*, 1996, **118**, 9218–9227.
- 98 M. Jbara, E. Eid and A. Brik, *J. Am. Chem. Soc.*, 2020, **142**, 8203–8210.
- 99 M. Jbara, S. Laps, M. Morgan, G. Kamnesky, G. Mann, C. Wolberger and A. Brik, *Nat. Commun.*, 2018, **9**, 3154.
- 100 N. Kamo, G. Hayashi and A. Okamoto, *Org. Lett.*, 2019, **21**, 8378–8382.
- 101 G. B. Vamisetti, G. Satish, S. Prasad, G. Mann, M. Glickman and A. Brik, *J. Am. Chem. Soc.*, 2020, **142**, 19558–19569.
- 102 M. Góngora-Benítez, J. Tulla-Puche and F. Albericio, *Chem. Rev.*, 2014, **114**, 901–926.
- 103 A. Isidro-Llobet, M. Álvarez and F. Albericio, *Chem. Rev.*, 2009, **109**, 2455–2504.
- 104 S. J. De Veer, M. W. Kan and D. J. Craik, *Chem. Rev.*, 2019, **119**, 12375–12421.
- 105 A. H. Jin, M. Muttenthaler, S. Dutertre, S. W. A. Himaya, Q. Kaas, D. J. Craik, R. J. Lewis and P. F. Alewood, *Conotoxins: Chemistry and Biology*, American Chemical Society, 2019, vol. 119.
- 106 S. Laps, H. Sun, G. Kamnesky and A. Brik, *Angew. Chem., Int. Ed.*, 2019, **58**, 5729–5733.
- 107 S. Laps, F. Atamleh, G. Kamnesky, H. Sun and A. Brik, *ChemRxiv*, 2020, DOI: 10.26434/chemrxiv.13065278.v1.
- 108 R. Mousa, R. Notis Dardashti and N. Metanis, *Angew. Chem., Int. Ed.*, 2017, **56**, 15818–15827.
- 109 L. Flohé, *Biochim. Biophys. Acta*, 2009, **1790**, 1389–1403.
- 110 S. Yoshizawa and A. Böck, *Biochim. Biophys. Acta*, 2009, **1790**, 1404–1414.
- 111 S. Chiba, Y. Itoh, S. ichi Sekine and S. Yokoyama, *Mol. Cell*, 2010, **39**, 410–420.
- 112 S. Y. Y. Itoh, M. J. Brocker, S. Sekine, G. Hammond, S. Suetsugu and D. Söil, *Science*, 2013, **340**, 75–79.
- 113 K. Caban and P. R. Copeland, *Selenium*, 2012, **5**, 61–72.
- 114 L. Dery, P. S. Reddy, S. Dery, R. Mousa, O. Ktorza, A. Talhami and N. Metanis, *Chem. Sci.*, 2017, **8**, 1922–1926.
- 115 P. S. Reddy, S. Dery and N. Metanis, *Angew. Chem., Int. Ed.*, 2016, **55**, 992–995.
- 116 Z. Zhao and N. Metanis, *Angew. Chem., Int. Ed.*, 2019, **58**, 14610–14614.
- 117 Z. Zhao and N. Metanis, *J. Org. Chem.*, 2020, **85**, 1731–1739.
- 118 N. Naruse, D. Kobayashi, K. Ohkawachi, A. Shigenaga and A. Otaka, *J. Org. Chem.*, 2020, **85**, 1425–1433.
- 119 H. Sun and A. Brik, *Acc. Chem. Res.*, 2019, **52**, 3361–3371.
- 120 R. R. Crichton, *RSC Metallobiol.*, 2017, 1–23.
- 121 J. Li, J. Yu, J. Zhao, J. Wang, S. Zheng, S. Lin, L. Chen, M. Yang, S. Jia, X. Zhang and P. R. Chen, *Nat. Chem.*, 2014, **6**, 352–361.
- 122 S. Li, L. Wang, F. Yu, Z. Zhu, D. Shobaki, H. Chen, M. Wang, J. Wang, G. Qin, U. J. Erasquin, L. Ren, Y. Wang and C. Cai, *Chem. Sci.*, 2017, **8**, 2107–2114.
- 123 M. Yang, J. Li and P. R. Chen, *Chem. Soc. Rev.*, 2014, **43**, 6511–6526.
- 124 R. M. Yusop, A. Unciti-Broceta, E. M. V. Johansson, R. M. Sánchez-Martín and M. Bradley, *Nat. Chem.*, 2011, **3**, 239–243.
- 125 J. Clavadetscher, S. Hoffmann, A. Lilienkampf, L. Mackay, R. M. Yusop, S. A. Rider, J. J. Mullins and M. Bradley, *Angew. Chem., Int. Ed.*, 2016, **55**, 15662–15666.
- 126 B. Van Vaerenbergh, J. Lauwaert, J. W. Thybaut, P. Vermeir and J. De Clercq, *Chem. Eng. J.*, 2019, **374**, 576–588.
- 127 P. Mastroilli, M. M. Dell'anna, A. Rizzuti, M. Mali, M. Zapparoli and C. Leonelli, *Molecules*, 2015, **20**, 18661–18684.