

Cite this: *Chem. Sci.*, 2022, 13, 7581

All publication charges for this article have been paid for by the Royal Society of Chemistry

Directed evolution of a cyclodipeptide synthase with new activities *via* label-free mass spectrometric screening†

Songya Zhang,^{‡a} Jing Zhu,^{‡a} Shuai Fan,^b Wenhao Xie,^a Zhaoyong Yang^{*b} and Tong Si^{‡a}

Directed evolution is a powerful approach to engineer enzymes *via* iterative creation and screening of variant libraries. However, assay development for high-throughput mutant screening remains challenging, particularly for new catalytic activities. Mass spectrometry (MS) analysis is label-free and well suited for untargeted discovery of new enzyme products but is traditionally limited by slow speed. Here we report an automated workflow for directed evolution of new enzymatic activities *via* high-throughput library creation and label-free MS screening. For a proof of concept, we chose to engineer a cyclodipeptide synthase (CDPS) that synthesizes diketopiperazine (DKP) compounds with therapeutic potential. In recombinant *Escherichia coli*, site-saturation mutagenesis (SSM) and error-prone PCR (epPCR) libraries expressing CDPS mutants were automatically created and cultivated on an integrated work cell. Culture supernatants were then robotically processed for matrix-assisted laser desorption/ionization time-of-flight (MALDI-ToF) MS analysis at a rate of 5 s per sample. The resulting mass spectral data were processed *via* custom computational algorithms, which performed a multivariate analysis of 108 theoretical mass-to-charge (*m/z*) values of 190 possible DKP molecules within a mass window of 115–373 Da. An F186L CDPS mutant was isolated to produce cyclo(L-Phe–L-Val), which is undetectable in the product profile of the wild-type enzyme. This robotic, label-free MS screening approach may be generally applicable to engineering other enzymes with new activities in high throughput.

Received 22nd March 2022

Accepted 1st June 2022

DOI: 10.1039/d2sc01637k

rsc.li/chemical-science

Mimicking random mutagenesis and natural selection in the laboratory,^{1–3} directed enzyme evolution is valuable in improving a variety of properties of biocatalysts (*i.e.*, catalytic activity, stability, substrate specificity, and enantioselectivity).^{4–6} While it is straightforward to generate thousands of enzyme variants using established methods such as error-prone PCR (epPCR), site-saturation mutagenesis (SSM), and DNA shuffling, phenotypic screening remains a major bottleneck due to the substrate and product versatility of biocatalysts.^{7,8} Optical assays are widely applied in high-throughput screening (HTS) during directed evolution, often requiring a surrogate substrate or a coupling assay with a colorimetric or fluorogenic readout.^{9,10} However, it is very challenging to apply such approaches to discover new catalytic activities. On the other hand, mass spectrometry (MS) is label-free and suitable for

untargeted molecular profiling.¹¹ The high sensitivity and chemical resolution of MS further highlight its potential in discovering new products, because new catalytic activities are often weak when first emerging from enzyme promiscuity.¹²

We and others have recently developed a range of high-throughput MS screening methods for directed protein evolution.^{13–16} However, the label-free advantage of MS has not been fully demonstrated in engineering new enzymatic activities, possibly due to the difficulties in untargeted MS screening. Particularly, it requires careful standardization and optimization of sample preparation, MS acquisition, and data processing, which are necessary to minimize experimental noise and spot the weak signals of a new product. On the other hand, biofoundries provide an emerging infrastructure to assist the design–build–test–learn (DBTL) cycles in biological engineering *via* robotic standardization and parallelization.^{17–19} Using an integrated biofoundry, here we report a workflow for unlabeled MS screening of recombinant libraries to rapidly isolate enzyme mutants that catalyze the formation of new products. This new workflow extends our previous matrix-assisted laser desorption/ionization time-of-flight (MALDI-ToF) MS-based screening approach from agar colonies¹⁶ to liquid cultures in industry-standard microplates for better uniformity. Unlike colony

^aCAS Key Lab Laboratory of Quantitative Engineering Biology, Shenzhen Institute of Synthetic Biology, Shenzhen Institute of Advanced Technology, Chinese Academy of Sciences, Shenzhen 518055, China

^bThe Institute of Medicinal Biotechnology, Chinese Academy of Medical Sciences and Peking Union Medical College, Beijing 1000050, China

† Electronic supplementary information (ESI) available. See <https://doi.org/10.1039/d2sc01637k>

‡ These authors contributed equally to this work.



biomass, liquid culture media often contain high concentrations of nonvolatile components, which interfere with MALDI matrix crystallization and cause severe ion suppression during MS analysis. Therefore, sample preparation steps, such as liquid-liquid extraction and solid-phase extraction, need to be incorporated and automated.

Cyclodipeptide synthases (CDPSs) are a family of enzymes that catalyze the formation of a diketopiperazine (DKP) from two aminoacyl-tRNA substrates (aa-tRNAs), which is an important pharmacophore for modern drug development^{20–22} (Fig. 1A and S1†). To produce new DKP derivatives for therapeutic and industrial applications, the catalytic mechanism of CDPSs has been studied^{23,24} to guide engineering. AlbC (239 aa) is the first identified CDPS protein, taking phenylalanyl-tRNA^{Phe} (Phe-tRNA^{Phe}) and leucyl-tRNA^{Leu} (Leu-tRNA^{Leu}) as substrates to synthesize cyclo(L-Phe-L-Leu) (cFL) in its native host *Streptomyces noursei*.^{25,26} Recombinant *E. coli* with AlbC overexpression is able to produce more cyclodipeptide derivatives in addition to cFL.²⁷ To date, eight CDPSs have been structurally characterized,^{26,28–32} and mutagenesis studies reveal that the residues within the P1 and P2 catalytic pockets are the key determinants of substrate specificity. However, the detailed mechanism of substrate recognition and catalysis is still elusive, and CDPSs are recognized as recalcitrant targets for rational engineering.^{33,34} To our knowledge, large-scale screening of CDPS variants has not been reported, possibly due to the lack of applicable HTS assays.

Here we develop and apply untargeted MS screening on a biofoundry for directed evolution of AlbC mutants that form new CDP products. The workflow consists of strain library creation, colony picking, microtiter cultivation, organic solvent extraction, transfer of the sample to a MALDI target and acquisition of mass spectra, followed by data processing and visualization (Fig. 1B). For library creation, AlbC variants were generated by either SSM or epPCR approaches and expressed from a pET28a plasmid under control of a T7 promoter. Upon genetic transformation of *E. coli* Rosetta(DE3), individual clones were transferred by a colony picker into microtiter plates. Subsequent cultivation, inducible production by addition of

isopropyl β-D-1-thiogalactopyranoside (IPTG), ethyl acetate extraction, and MS sample preparation were performed using an integrated robotic workcell in the Shenzhen Synthetic Biology Infrastructure. This workcell integrates common instruments for synthetic biology, including a liquid-handling station, shaking incubators, centrifuge, plate reader, and so on. Organic extracts of the liquid cultures were spotted onto a MALDI target and overlaid with 4-CHCA matrix solution. MS targets were then manually transferred to a stand-alone MALDI mass spectrometer in this study, although a robotic configuration has been previously reported to automate this step.³⁵

We first applied the workflow to analyze the wild-type (WT) AlbC-expressing strain, Rosetta(DE3)/pET28-AlbC(WT). After cell cultivation, inducible production, and sample preparation, MS analysis was performed using an Autoflex MALDI-ToF mass spectrometer in the reflector positive ion mode (Fig. 1). In the resulting MALDI mass spectra, we observed $[M + H]^+$ ion peaks with m/z values corresponding to cFL (m/z 261.17), cFY (m/z 311.15), cFM (m/z 279.12), cYL (m/z 277.16), cFF(cYM) (m/z 295.15), cLL (m/z 227.16) and cMM (m/z 263.12), all of which were absent from the control strain Rosetta(DE3)/pET28 (Fig. S2†). Production of these CDP molecules was further confirmed by examining tandem MS results using LC-MS in the multiple reaction monitoring (MRM) mode (Table S3†), where ion fragmentation patterns were consistent with the literature.^{26,27} Moreover, the observed production profile was also consistent with previous studies.^{26,27} These results validated our workflow to detect CDP products from microplate cultures of AlbC-expressing *E. coli* using MALDI-ToF MS.

To engineer AlbC mutants that synthesize new products, we first focused on the key residues in the substrate-binding pockets. AlbC employs a “ping-pong” catalytic mechanism: the initial step transfers the aminoacyl moiety of the first aa-tRNA onto a conserved serine, leading to the formation of an aminoacyl enzyme intermediate; then, the aminoacyl enzyme reacts with the aminoacyl moiety of the second aa-tRNA to form a dipeptidyl enzyme.³² Previous biochemical experiments demonstrated that two binding pockets P1 and P2 of AlbC accommodate the aminoacyl moieties of the two aa-tRNA during the biosynthesis²⁴ (Fig. 2). Only a limited number of mutations were examined on P1 and P2 residues,^{25,26} possibly due to the lack of HTS assays for large-scale analysis. Instead, we applied our workflow to create and screen the SSM libraries of select residues, including 10 residues in the binding pockets (L33, V65, L119, L185, L200, M152, M159, I204, T206 and P207) and 4 residues (R99, R101, R102 and D205) outside the pockets which have interactions of the tRNA moiety reported in the literature (Fig. 2A).

For SSM libraries, we adopted the “22c-trick” strategy to design degenerative primers, and 94 library clones were randomly picked for each residue library to reach a >98.6% probability of full library coverage of the 20 canonical amino acids.³⁶ In addition to library variants, the WT and control strains were also included in the same 96-well plate. Overall, the abovementioned workflow (Fig. 1B) takes approximately 5 s for sample preparation and MS analysis for each mutant culture. Tentative ion peaks of cyclodipeptides were assigned based on

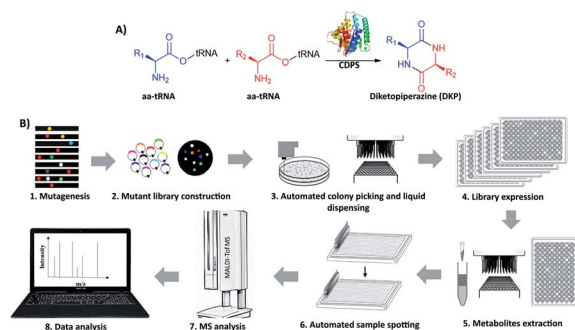


Fig. 1 (A) Scheme of diketopiperazine (DKP) biosynthesis catalyzed by cyclodipeptide synthase (CDPS). (B) Workflow of the matrix-assisted laser desorption/ionization time-of-flight mass spectrometry (MALDI-ToF MS)-based high-throughput screening method for directed evolution of AlbC protein in *E. coli*.



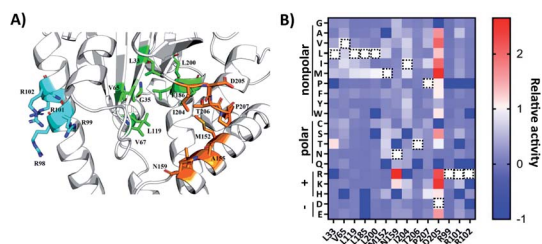


Fig. 2 (A) Structural analysis of the AlbC (PDB ID, 3OQV). Enlarged view of the catalytically active pocket. The possible catalytic residues are shown in green (pocket-1) and orange (pocket-2) and basic residues on helix α_4 are labelled in cyan. (B) Heatmap of the relative activity change in the cFL production levels of mutation of 14 amino acid residues in AlbC. WT residues are labeled with a black dashed rectangle. Dark blue boxes indicate that a specific mutant was not covered in randomly picked clones (activity assigned as -1).

theoretical m/z values (ESI Section 1†) using the MetaboAnalyst webserver.³⁷ For each ion peak, one-way analysis of variance (ANOVA) was used to evaluate statistical differences between the mean peak areas of a library variant and those of the control group. AlbC gene mutations were revealed by Sanger sequencing for all 14 SSM library members.

To visualize the MALDI-ToF MS screening results, a heatmap was generated based on the cFL production of library members relative to that of WT (Fig. 2B), and we observed a general consistency between our results and literature data (Table 1). For example, the L200N mutation almost abolished cFL production (Table 1) but increased cYL (m/z 277.16, $[M + H]^+$), cYY (m/z 327.16, $[M + H]^+$), and cYM (m/z 295.18 $[M + H]^+$) production by about 1.2-fold, 5-fold, and 3-fold (Fig. 3A), respectively, which was similar to a previous study.²⁶ Also, mutagenesis of basic residues (R99, R101 and R102) outside the pockets led to the decline of cFL production in the mutants, although R101 was more tolerant to mutations than R99 and R102. Furthermore, the substitution of D205 in the β_6 - α_8 loop to alanine enhanced AlbC catalytic activity as observed previously.²⁵ On the other hand, some previously unnoticed

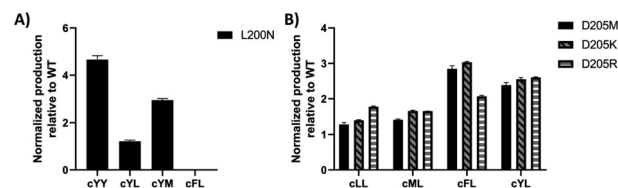


Fig. 3 Cyclodipeptide product analysis of AlbC mutants. (A) LC-MS characterization of the L200N mutant; (B) LC-MS characterization of select mutants in the D205 saturation mutagenesis library. Normalized production levels relative to WT (set as 1) are reported. Error bars represent the standard deviations of three biological replicates.

phenomena were observed. For example, in the D205 SSM library, relative ion intensities of cFL (m/z 261.17) in many mutants are higher than that of WT (Fig. S3†). The top 3 mutants, D205M, D205K and D205R, were subsequently analyzed by LC-MS/MS. The results not only confirmed augmented synthesis of cFL, but also revealed increased production of other leucine-containing products including cYL, cLL and cML (Fig. 3B). Also interestingly, the discovery that most mutations of the T206 residue in the P2 pocket abolished the biosynthesis of the main product cFL was unprecedented (Fig. S3 and ESI Section 2†). When examining MALDI mass spectra in detail, we noticed that although the T206F mutation greatly reduced cFL production, cFF and cFY products were largely not affected. Replacing the small threonine side chain with a bulky aromatic phenylalanine side chain could affect the activity of WT, because the phenyl ring structure increased the steric hindrance of T206F, thus impairing binding with its substrate and causing loss of its enzyme activity for cFL (Fig. 5A). These results suggested that T206 is also a key residue for the selectivity of the second Leu-tRNA substrate binding in the pocket. Unfortunately, we did not observe any new CDP molecules produced from the 14 SSM library mutants.

The failure of isolating AlbC mutants with new substrate specificities from the SSM libraries highlights the limitation of semi-rational approaches that target manually selected

Table 1 Impact of AlbC mutations on the main product cFL

Target residue	Mutants and production levels relative to WT (literature)	Mutants and production levels relative to WT (this study)
	WT: cFL (set as 100%)	WT: cFL (set as 100%)
Within binding pockets		
L33/L185D	L33Y/L185D: cFL (0) ²⁶	L33Y: cFL (n.d.) ^a
L200	L200N: cFL (<10%) ²⁶	L200N: cFL (n.d., ^a 5.6%) ^b
N159	N159A: cFL (45%) ²⁵	N159A: cFL (60%) ^a
Outside the pockets		
R98	R98A: cFL (20%) ²⁶	
R99	R99A: cFL (<10%) ²⁶	R99A: cFL (n.d.) ^a
	R98A/R99A: cFL (0)	
R101/R102	R101A/R102A: cFL (20%) ²⁶	R101A: cFL (>50%) ^a
		R102A: cFL (23%) ^a
D205	D205A: cFL (>100%) ²⁵	D205A: cFL (>100%) ^a

^a Calculated from MALDI-TOF MS data. ^b Calculated from LC-MS/MS data.



Then, we investigated possible mechanisms underlying new substrate specificity of the F186L variant from structural aspects using computational modeling. In the substrate binding pocket, the cFL substrate formed three hydrogen bonds with two amino acid residues (N40 and E182) in the WT AlbC. The phenyl side chain of cFL forms a π - π stacking interaction with F186 (Fig. 5B and S5 $\dagger$). However, when the residue at position 186 was mutated to leucine, this π - π stacking interaction was abolished. This leads to a change in the conformation of the Phe1 ring and thereby the hydrogen bond between Phe1 and N40 is abolished (Fig. 5C and S5 $\dagger$), which ultimately results in outward movement of loop G33-S44 (Fig. S6 $\dagger$). Therefore, the volume of the substrate binding cavity, which was measured to be 197 Å³, increased up to 288 Å³ when the residue at position 186 of WT was mutated to leucine (Table S4 $\dagger$). Overall, it is possible that an increase in the production of cFV occurs precisely due to the larger volume of the substrate-binding cavity in F186L. Moreover, cFL and the new derivative cFV were separately docked into the binding pockets of the WT and F186L variant for 100 ns MD simulations (Fig. S5 $\dagger$). The results showed that the binding energy of F186L was significantly increased when docking with the substrate cFV (Table 2). The root-mean-square deviations (RMSD) were calculated in 100 ns to investigate the stability of the WT/F186L-L-lysine complexes (Fig. S5 $\dagger$). Both WT and F186L reached the equilibrium state from an early stage.

In conclusion, we developed a robotic assay for directed evolution of AlbC, a model CDPS, using MALDI-ToF MS for label-free screening. Compared with conventional LC-MS (typically more than 5 min per sample), our HTS workflow represents a two-magnitude reduction of analytical time (5 s per sample). Contrary to previous reports that only study limited mutations of select residues, 14 SSM libraries were created and profiled for sequence-activity profiling, which not only confirmed the impact of known mutations (Table 1), but also revealed new specificity-modulating mutations (*i.e.*, T206F reduced preference towards Leu-tRNA as the second substrate). Notably, by creating and screening ~4500 epPCR library members within a week using unlabeled MS on an integrated, robotic workcell, an AlbC mutant producing a new cyclopeptide product was identified, revealing a previously unknown residue (F186) that exerts a substantial impact on substrate specificity. Other MS modalities, such as droplet microfluidics coupled with ESI-MS,^{38,39} may also serve as label-free HTS assays, but further development is needed to address ion suppression issues caused by direct infusion of complex culture media into a mass spectrometer. On the other hand, the scarcity of positive hits in the single-residue SSM and epPCR libraries in this study confirmed AlbC as a difficult target for evolving new activities. Therefore, it is desirable to create and screen new AlbC mutant libraries that are comprehensive (*i.e.*, deep mutational scanning⁴⁰), combinatorial (*i.e.*, combinatorial active site saturation test/iterative saturation mutagenesis, CAST/ISM⁴¹), or data-driven (*i.e.*, machine learning-assisted directed evolution, MLDE^{42,43}) in the future. Together, we envision that the label-

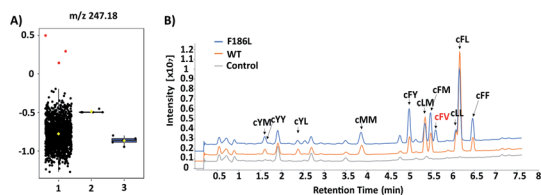


Fig. 4 (A) Box-whisker plots of 1152 samples at m/z 247.18 based on ANOVA analysis. Group 1, epPCR variants; 2, Rosetta(DE3)/pET28a-AlbC (WT); 3, Rosetta(DE3)/pET28a (control). Three clones containing F186L were labeled in red. ANOVA, analysis of variance. (B) LC-MS traces of WT and F186L mutant.

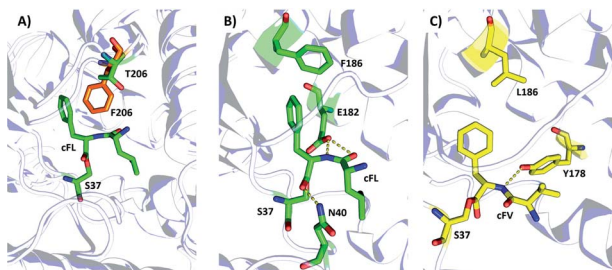


Fig. 5 Structural illustration of the substrate binding pocket of the wildtype AlbC and the mutants F186L and T206. Note: Panel (A) Superimposition of the WT with T206F. Panels (B and C) Schematic representation of the hydrogen-bond and hydrophobic interaction between cFL-WT (B) and cFV-F186L (C). Dashed lines indicate hydrogen bond (yellow) interactions. The key amino acid residues of WT, F186L, and T206F are drawn as sticks in green, yellow and orange, respectively.

Table 2 Binding energy of ligand for AlbC WT and F186L

	Binding energy ^a (kcal mol ⁻¹) of WT	Binding energy ^b (kcal mol ⁻¹) of F186L	Binding energy ^c (kcal mol ⁻¹) of F186L
ΔG_{VDW}^d	-35.01	-23.67	-34.27
ΔG_{Et}^e	-31.95	-8.70	-29.37
ΔG_{polar}^f	48.71	31.08	47.60
ΔG_{apolar}^g	-5.60	-4.08	-5.50
$\Delta G_{binding}^h$	-23.85	-5.37	-21.54

^a WT-cFL. ^b F186L-cFL. ^c F186L-cFL. ^d van der Waals energy. ^e Electrostatic energy. ^f Polar-solvation energy. ^g Nonpolar solvation energy. ^h $\Delta G_{binding} = \Delta G_{VDW} + \Delta G_{Et} + \Delta G_{polar} + \Delta G_{apolar}$.

free MS screening method should be generally applicable to engineering other enzymes with new activities.

Data availability

The datasets supporting this article have been uploaded as part of the ESI.†

Author contributions

T. S., J. Z. and S. Y. Z. conceived the project and experimental design; J. Z. and S. Y. Z. conducted all the wet experimental work; S. Y. Z., S. F. and Z. Y. Y. performed the molecular docking and the dynamic simulation calculation; W. H. X. assisted with sample preparation for MALDI-ToF MS analysis. S. Y. Z., J. Z., S. F. and T. S. prepared the manuscript, and all authors refined the draft.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by the National Key R&D Program of China (2021YFA0910800), National Natural Science Foundation of China (32071428 to T. S., 82104044 to S. Y. Z. and 81872782 to Z. Y. Y.), and the Chinese Postdoctoral Grant (2019M663193 to S. Y. Z.). We thank Zhiyu Zhang and Yunquan Lan at the Shenzhen Synthetic Biology Infrastructure for help with robotic workflow development.

Notes and references

- 1 F. H. Arnold, *Angew. Chem., Int. Ed. Engl.*, 2018, **57**, 4143–4148.
- 2 Y. Wang, P. Xue, M. Cao, T. Yu, S. T. Lane and H. Zhao, *Chem. Rev.*, 2021, **121**, 12384–12444.
- 3 G. Qu, A. Li, C. G. Acevedo-Rocha, Z. Sun and M. T. Reetz, *Angew. Chem., Int. Ed. Engl.*, 2020, **59**, 13204–13231.
- 4 H. Y. Choi, H. S. Lim, K. H. Park, J. Kim and W. G. Kim, *Appl. Microbiol. Biotechnol.*, 2021, **105**, 4599–4607.
- 5 B. Morawski, S. Quan and F. H. Arnold, *Biotechnol. Bioeng.*, 2001, **76**, 99–107.
- 6 R. Carr, M. Alexeeva, A. Enright, T. S. Eve, M. J. Dawson and N. J. Turner, *Angew. Chem., Int. Ed. Engl.*, 2003, **42**, 4807–4810.
- 7 H. Yu, J. Li, D. Zhang, Y. Yang, W. Jiang and S. Yang, *Appl. Microbiol. Biotechnol.*, 2009, **82**, 279–285.
- 8 M. L. Geddie and I. Matsumura, *J. Biol. Chem.*, 2004, **279**, 26462–26468.
- 9 N. N. Rosic, *Methods Mol Biol.*, 2013, **987**, 205–224.
- 10 V. Tarallo, K. Sudarshan, V. Nosek and J. Misek, *Chem. Commun.*, 2020, **56**, 5386–5388.
- 11 T. X. Si, P. K. Choe, H. Zhao and J. V. Sweedler, in *Protein Engineering*, 2021, pp. 57–79.
- 12 T. de Rond, J. Gao, A. Zargar, M. de Raad, J. Cunha, T. R. Northen and J. D. Keasling, *Angew. Chem., Int. Ed. Engl.*, 2019, **58**, 10114–10119.
- 13 P. Xue, T. Si, S. Mishra, L. Zhang, K. Choe, J. V. Sweedler and H. Zhao, *Biotechnol. Bioeng.*, 2020, **117**, 2131–2138.
- 14 C. Yan, F. Parmeggiani, E. A. Jones, E. Claude, S. A. Hussain, N. J. Turner, S. L. Flitsch and P. E. Barran, *J. Am. Chem. Soc.*, 2017, **139**, 1408–1411.
- 15 A. J. Pluchinsky, D. J. Wackelin, X. Huang, F. H. Arnold and M. Mrksich, *J. Am. Chem. Soc.*, 2020, **142**, 19804–19808.
- 16 T. Si, B. Li, T. J. Comi, Y. Wu, P. Hu, Y. Wu, Y. Min, D. A. Mitchell, H. Zhao and J. V. Sweedler, *J. Am. Chem. Soc.*, 2017, **139**, 12466–12473.
- 17 N. Hillson, M. Caddick, Y. Cai, J. A. Carrasco, M. W. Chang, N. C. Curach, D. J. Bell, R. Le Feuvre, D. C. Friedman, X. Fu, N. D. Gold, M. J. Herrgard, M. B. Holowko, J. R. Johnson, R. A. Johnson, J. D. Keasling, R. I. Kitney, A. Kondo, C. Liu, V. J. J. Martin, F. Menolascina, C. Ogino, N. J. Patron, M. Pavan, C. L. Poh, I. S. Pretorius, S. J. Rosser, N. S. Scrutton, M. Storch, H. Tekotte, E. Travník, C. E. Vickers, W. S. Yew, Y. Yuan, H. Zhao and P. S. Freemont, *Nat. Commun.*, 2019, **10**, 2040.
- 18 J. Zhang, Y. Chen, L. Fu, E. Guo, B. Wang, L. Dai and T. Si, *Curr. Opin. Biotechnol.*, 2021, **67**, 88–98.
- 19 R. Chao, S. Mishra, T. Si and H. M. Zhao, *Metab. Eng.*, 2017, **42**, 98–108.
- 20 A. D. Borthwick, *Chem. Rev.*, 2012, **112**, 3641–3716.
- 21 N. Canu, M. Moutiez, P. Belin and M. Gondry, *Nat. Prod. Rep.*, 2020, **37**, 312–321.
- 22 A. Ortiz and E. Sansinenea, *Curr. Med. Chem.*, 2017, **24**, 2773–2780.
- 23 I. B. Jacques, M. Moutiez, J. Witwinowski, E. Darbon, C. Martel, J. Seguin, E. Favry, R. Thai, A. Lecoq, S. Dubois,



- J. L. Pernodet, M. Gondry and P. Belin, *Nat. Chem. Biol.*, 2015, **11**, 721–727.
- 24 M. Gondry, I. B. Jacques, R. Thai, M. Babin, N. Canu, J. Seguin, P. Belin, J. L. Pernodet and M. Moutiez, *Front. Microbiol.*, 2018, **9**, 46.
- 25 M. Moutiez, J. Seguin, M. Fonvielle, P. Belin, I. B. Jacques, E. Favry, M. Arthur and M. Gondry, *Nucleic Acids Res.*, 2014, **42**, 7247–7258.
- 26 L. Sauguet, M. Moutiez, Y. Li, P. Belin, J. Seguin, M. H. Le Du, R. Thai, C. Masson, M. Fonvielle, J. L. Pernodet, J. B. Charbonnier and M. Gondry, *Nucleic Acids Res.*, 2011, **39**, 4475–4489.
- 27 M. Gondry, L. Sauguet, P. Belin, R. Thai, R. Amouroux, C. Tellier, K. Tuphile, M. Jacquet, S. Braud, M. Courcon, C. Masson, S. Dubois, S. Lautru, A. Lecoq, S. Hashimoto, R. Genet and J. L. Pernodet, *Nat. Chem. Biol.*, 2009, **5**, 414–420.
- 28 M. W. Vetting, S. S. Hegde and J. S. Blanchard, *Nat. Chem. Biol.*, 2010, **6**, 797–799.
- 29 L. Bonnefond, T. Arai, Y. Sakaguchi, T. Suzuki, R. Ishitani and O. Nureki, *Proc. Natl. Acad. Sci. U. S. A.*, 2011, **108**, 3912–3917.
- 30 G. Bourgeois, J. Seguin, M. Babin, P. Belin, M. Moutiez, Y. Mechulam, M. Gondry and E. Schmitt, *J. Struct. Biol.*, 2018, **203**, 17–26.
- 31 C. J. Harding, E. Sutherland, J. G. Hanna, D. R. Houston and C. M. Czekster, *RSC Chem. Biol.*, 2021, **2**, 230–240.
- 32 M. Moutiez, E. Schmitt, J. Seguin, R. Thai, E. Favry, P. Belin, Y. Mechulam and M. Gondry, *Nat. Commun.*, 2014, **5**, 5141.
- 33 K. Brockmeyer and S. M. Li, *J. Nat. Prod.*, 2017, **80**, 2917–2922.
- 34 T. Yao, J. Liu, Z. Liu, T. Li, H. Li, Q. Che, T. Zhu, D. Li, Q. Gu and W. Li, *Nat. Commun.*, 2018, **9**, 4091.
- 35 S. Lin, S. Dikler, W. D. Blincoe, R. D. Ferguson, R. P. Sheridan, Z. Peng, D. V. Conway, K. Zawatzky, H. Wang, T. Cernak, I. W. Davies, D. A. DiRocco, H. Sheng, C. J. Welch and S. D. Dreher, *Science*, 2018, 361.
- 36 S. Kille, C. G. Acevedo-Rocha, L. P. Parra, Z. G. Zhang, D. J. Opperman, M. T. Reetz and J. P. Acevedo, *ACS Synth. Biol.*, 2013, **2**, 83–92.
- 37 Z. Pang, J. Chong, G. Zhou, D. A. de Lima Morais, L. Chang, M. Barrette, C. Gauthier, P. E. Jacques, S. Li and J. Xia, *Nucleic Acids Res.*, 2021, **49**, W388–W396.
- 38 X. W. Diefenbach, I. Farasat, E. D. Guetschow, C. J. Welch, R. T. Kennedy, S. Sun and J. C. Moore, *ACS Omega*, 2018, **3**, 1498–1508.
- 39 D. J. Steyer and R. T. Kennedy, *Anal. Chem.*, 2019, **91**, 6645–6651.
- 40 D. M. Fowler and S. Fields, *Nat. Methods*, 2014, **11**, 801–807.
- 41 G. Li, M. Furst, H. R. Mansouri, A. K. Ressmann, A. Ilie, F. Rudroff, M. D. Mihovilovic, M. W. Fraaije and M. T. Reetz, *Org. Biomol. Chem.*, 2017, **15**, 9824–9829.
- 42 B. J. Wittmann, Y. Yue and F. H. Arnold, *Cell Syst.*, 2021, **12**, 1026–1045.
- 43 Z. Wu, S. B. J. Kan, R. D. Lewis, B. J. Wittmann and F. H. Arnold, *Proc. Natl. Acad. Sci. U. S. A.*, 2019, **116**, 8852–8858.

