

CrossMark  
click for updatesCite this: *Chem. Sci.*, 2016, 7, 4848

# A widespread bacterial phenazine forms S-conjugates with biogenic thiols and crosslinks proteins†

D. Heine,<sup>a</sup> S. Sundaram,<sup>a</sup> Matthias Beudert,<sup>a</sup> K. Martin<sup>a</sup> and C. Hertweck<sup>\*ab</sup>

Phenazines are redox-active compounds produced by a range of bacteria, including many pathogens. Endowed with various biological activities, these ubiquitous N-heterocycles are well known for their ability to generate reactive oxygen species by redox cycling. Phenazines may lead to an irreversible depletion of glutathione, but a detailed mechanism has remained elusive. Furthermore, it is not understood why phenazines have so many protein targets and cause protein misfolding as well as their aggregation. Here we report the discovery of unprecedented conjugates (panphenazines A, B) of panthetheine and phenazine-1-carboxylic (PCA) acid from a *Kitasatospora* sp., which prompted us to investigate their biogenesis. We found that PCA reacts with diverse biogenic thiols under radical-forming conditions, which provides a plausible model for irreversible glutathione depletion. To evaluate the scope of the reaction in cells we designed biotin and rhodamine conjugates for protein labelling and examined their covalent fusion with model proteins (ketosynthase, carbonic anhydrase III, albumin). Our results reveal important, yet overlooked biological roles of phenazines and show for the first time their function in protein conjugation and crosslinking.

Received 1st February 2016

Accepted 13th April 2016

DOI: 10.1039/c6sc00503a

www.rsc.org/chemicalscience

## Introduction

From an evolutionary perspective, it is most remarkable how small biomolecules have been shaped into privileged structures that fulfil multiple biological functions. This is particularly true for the large family of phenazines, ancient N-heterocycles that are produced by a large variety of bacteria.<sup>1</sup> Initially discovered as colourful bacterial pigments over 150 years ago, to date over one hundred natural phenazines have been characterised, most of which are endowed with highly diverse bioactivities.<sup>2</sup> These findings inspired the chemical synthesis of thousands of phenazine derivatives and propelled the development of antibiotics and anticancer agents.<sup>1–3</sup> However, phenazines are a remarkable example of antibiotic compounds that are actually crucial for microbial interaction and competitive fitness.<sup>4–6</sup> They not only act as signalling molecules in microbial communication,<sup>7–9</sup> mediating biofilm formation and morphogenesis of colonies,<sup>10</sup> but also control gene expression<sup>11</sup> and influence the intracellular redox state<sup>10</sup> as well as exocellular electron transfer.<sup>12</sup> Owing to their redox properties, phenazines play major roles in infection processes of pathogens such as *Pseudomonas*

*aeruginosa*, especially in cystic fibrosis patients.<sup>13</sup> In particular, pyocyanin (PYO, 1) and phenazine-1-carboxylic acid (PCA, 2), the key intermediate in phenazine biosynthesis,<sup>14</sup> represent important virulence factors of *P. aeruginosa* that damage infected tissue (Fig. 1).<sup>15,16</sup> The primary reason for the deleterious effects is phenazine-mediated redox cycling,<sup>1</sup> which leads to high concentrations of reactive oxygen species (ROS). Naturally, ROS is effectively counteracted by reduced glutathione (GSH) as redox buffer, with formation of the oxidised disulfide form (GSSG).<sup>17,18</sup> Despite an impressive body of data on phenazine functions, there are numerous open questions on their mode of action. Specifically, it was observed that phenazines

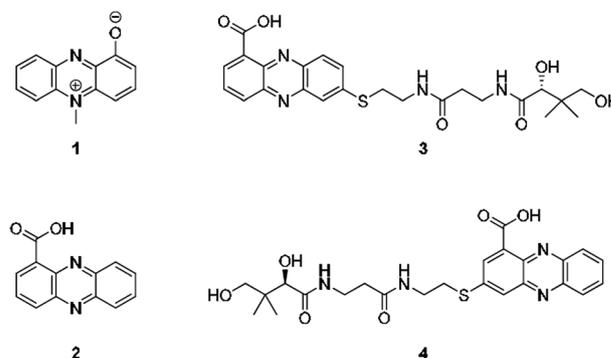


Fig. 1 Structures of pyocyanin (1), PCA (2), and panphenazines A (3) and B (4) discovered in this study.

<sup>a</sup>Leibniz Institute for Natural Product Research and Infection Biology, Hans Knoell Institute, Beutenbergstrasse 11a, 07745 Jena, Germany. E-mail: christian.hertweck@leibniz-hki.de

<sup>b</sup>Friedrich Schiller University, 07737 Jena, Germany

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c6sc00503a





identical with the conjugates **3** and **4** isolated from *Kitasatospora* sp. HKI 714 (Fig. S2†). Furthermore, the photoreaction proved to be a viable route for coupling cysteine and glutathione with PCA, as evidenced by HRMS and MS<sup>n</sup> analyses (Fig. S3 and S4†).

The successful photoreactions and the substitution pattern are highly suggestive for the involvement of resonance-stabilised radicals in thiol conjugation. To interrogate the reaction mechanism, we conducted the coupling reaction of PCA and *N*-acetylcysteamine in the dark, but added azobisisobutyronitrile (AIBN) as a radical initiator. HPLC-HRMS analyses clearly indicated the formation of the corresponding adducts. Likewise, **3** and **4** were formed after incubation of PCA with pantetheine and AIBN (Fig. S5†).

### Mechanistic considerations

The radical mechanism is supported by two other observations: first, addition of an electron source such as triethylamine (TEA) substantially increases product formation (up to tenfold), which is in agreement with the reported mechanism of the formation of the phenazine radical anion.<sup>37</sup> Second, addition of the radical scavenger *t*-butylhydroxytoluene (TBHT) hampered the photoreaction; only traces of conjugates could be detected. Thus, in addition to the observed substitution pattern of the thioconjugates, three lines of experimental evidence support the radical-based reaction mechanism (Fig. 3).

However, a direct light-induced formation of the thiyl radical can be ruled out. The homolytic cleavage of the SH bond occurs at wavelengths below 300 nm,<sup>38</sup> yet the PCA-thioconjugates are readily produced at 370 nm. Alternatively, it is more plausible that radical formation is initiated by photoexcitation of PCA into the triplet state. Transferring an electron from TEA (or an alternative electron source) leads to the phenazine radical anion (Fig. 3).<sup>37</sup> Subsequent protonation generates a neutral radical, which can react with biogenic thiols (GSH, pantetheine, cysteamine, cysteine) to form a thiyl radical while regenerating PCA. By analogy to the aromatic substitution reaction with thiyl

radicals,<sup>39</sup> one may infer that the thiyl radicals readily attack the electron-rich phenazine core. After H abstraction and re-aromatization, the radical chain reaction continues, which may also lead to multiple substitutions as observed for *N*-acetylcysteamine. On the contrary, conjugate formation with pantetheine seems to appear more selectively, possibly due to an increased steric demand of the thiol. Even though various conjugates could be detected in the HPLC-HRMS trace, only two isomers (**3** and **4**) appear in higher amounts. It should be highlighted that light-independent avenues to the formation of phenazine radicals have been reported in the biological context. It is well known that PYO is chemically reduced through radical intermediates in the cell, and phenazine redox-cycling (intracellular as well as extracellular) may generate PCA radicals,<sup>1</sup> which can react with thiols to form conjugates.

### Phenazine protein adduct formation and crosslinking

Single-electron redox processes yield phenazine radicals in host cells; consequently, these reactive species may not only react with biogenic thiols, but also with SH residues of proteins, an important target of many small molecules.<sup>40,41</sup> Such a process would be reminiscent of non-enzymatic reactions of sugars with proteins, yielding advanced glycation end-products (AGEs), which play important roles in aging and degenerative diseases.<sup>42</sup> Considering the non-selective adduct formation of phenazines and the multitude of thiols in living cells, it is obvious that the corresponding conjugates would barely be detectable *in vivo* as individual entities. Thus we investigated whether PCA radicals form conjugates with individual proteins *in vitro*. For this, we synthesised a chemical probe (**9**) (Fig. S6†) in which the phenazine is fused to biotin *via* a tetraethylene glycol (TEG) linker (Fig. 4A).

First, we incubated **9** and AIBN with a cysteine-rich didomain (ketosynthase-branching, KS-B, 14 cysteines) of a modular polyketide synthase (PKS) module.<sup>43–45</sup> The reaction mixture was loaded onto a streptavidin column, and the biotinylated protein was eluted at 60 °C in buffer with 3 mM biotin after multiple washing steps. Subsequent sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) analysis indicated that the protein was indeed linked to the biotin-labelled probe, unequivocally proving that the phenazine was bound to the PKS component (Fig. 4B). To gain insight into the fate of proteins in the presence of the phenazine and UV light, we also prepared a fluorescent probe (**10**). Although PCA shows fluorescence itself, the respective signal proved to be too weak for the detection of protein conjugates. Therefore we attached PCA to rhodamine B, a fluorophore with an exceptionally high fluorescence quantum yield. The KS-B didomain was subjected to the rhodamine-based probe **10** with irradiation at 370 nm, and the reaction mixture was analysed by SDS-PAGE on a 12% gel. Surprisingly, the fluorescent bands were substantially larger than expected. Instead of the predicted band at 100 kDa, we observed a fluorescent signal that corresponds to a molecular weight of over 200 kDa (Fig. S7†). One possible explanation for this finding is that the probe might have cross-linked two or more equivalents of the

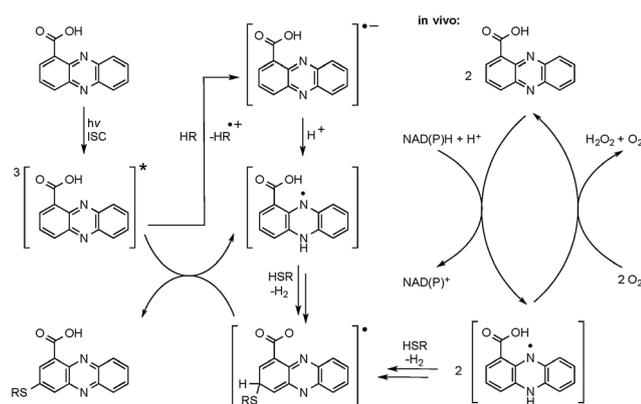
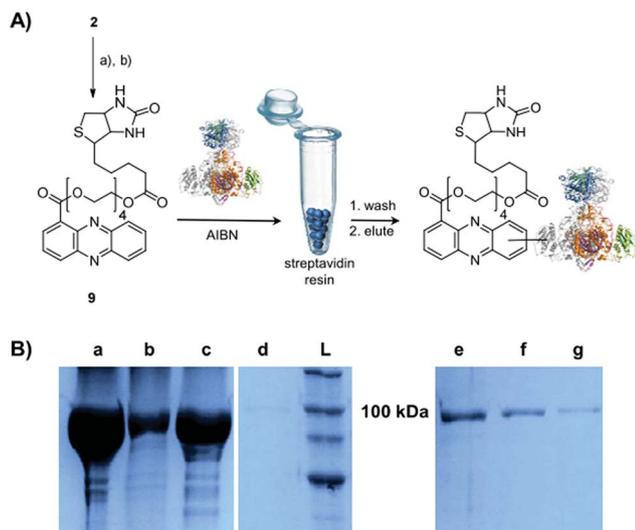


Fig. 3 Proposed reaction mechanism for the radical-induced conjugate addition. Radical formation can proceed *via* a photochemical reaction (left) or by redox cycling inside the cell (right). Only one of the possible intermediates (resonance form) is shown, leading to the derivative substituted at position 7. ISC = intersystem crossing; HR, electron donor (triethylamine, thiol or other); HSR, thiol.

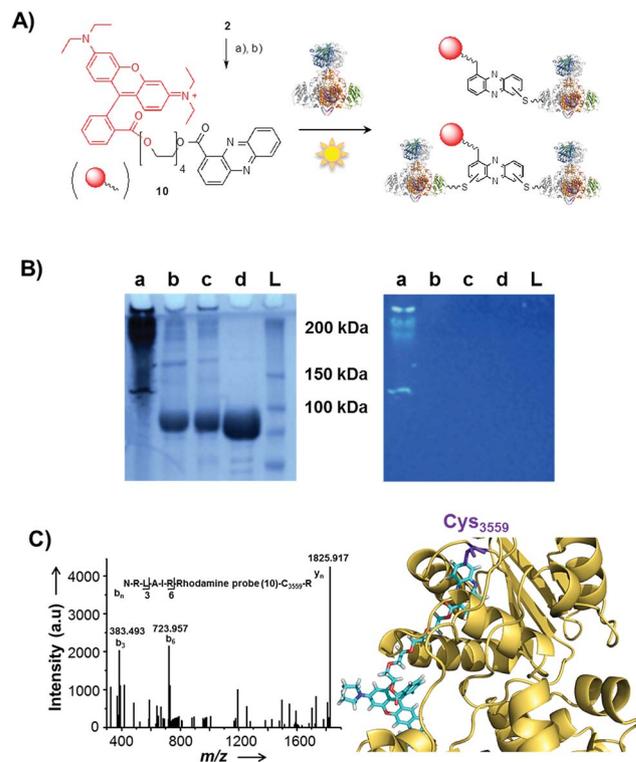




**Fig. 4** Evidence for radical-mediated protein conjugation by PCA. (A) Synthesis of biotin-based probe **9**, thio-adduct formation with keto-synthase (KS-B) and purification with streptavidin resin; (a) tetraethylene glycol, EDC, DMAP, DCM, 86% (b) biotin, EDC, DMAP, DCM, 96%. (B) SDS-PAGE (10%) of KS-B after the incubation with the probe **9** for 24 h in the presence of AIBN. (a) Load, (b) flow-through, (c) first wash, (d) second wash, (e) first elution, (f) second elution, (g) third elution, (L) ladder.

protein. When analysing the products on a lower concentrated gel (10%) for better separation, we detected several protein aggregates of various sizes (Fig. 5B).

By tryptic digestion and MALDI analyses we corroborated that all protein complexes resulting from the incubation of KS-B with **9** and **10** are indeed derived from the KS (Fig. S8.1–S8.6†). In control experiments where the protein was irradiated in the presence of rhodamine B alone no aggregation could be observed (Fig. 5B). MS/MS analysis of the protein (lane a) revealed one of the peptides where **10** was bound to Cys3559 (Fig. 5C). A model was generated to show the proposed binding mode of **10** to this cysteine using Pymol (Fig. 5D). These findings unequivocally showed that the crosslinking activity correlates with the presence of the phenazine moiety and that multiple linkages are formed. To test the scope of the reaction we employed two proteins that are completely unrelated to a PKS, carbonic anhydrase III (5 cysteines) and albumin (35 cysteines). After incubation of carbonic anhydrase III with probe **10** we monitored the time course of its aggregation. Bands corresponding to the native protein slowly disappeared with concomitant formation of extensive protein aggregates bound to the fluorescent probe (Fig. S9†). Using the same conditions, albumin was functionalised with the rhodamine-based probe **10**, yet formation of higher aggregates was negligible (Fig. S10†). Nonetheless, the monoadduct was ideally suited to confirm the binding of the probe to a cysteine side chain by MALDI analysis (Fig. S11 and S12†). Taken together, all selected proteins are derivatised with the phenazine probe, yet they differ in their tendency to form higher aggregates, which may be controlled by steric factors.



**Fig. 5** Photo-induced protein crosslinking by PCA. (A) Synthesis of PCA-rhodamine B conjugate **10** and thio-conjugate formation with KS-B, (a) tetraethylene glycol, EDC, DMAP, DCM, 86%, (b) rhodamine B, PyBOP, Hünig's base, DCM, 63%. (B) SDS-PAGE (10%) of KS-B after the incubation with probe **10** or rhodamine B in water (left picture: under white light; right picture at 370 nm); (contrast adjusted) (a) **10**, 370 nm, 18 h, (b) rhodamine B, 370 nm, 18 h, (c) H<sub>2</sub>O, 370 nm, 18 h, (d) KS-B control; (L) ladder. (C) MALDI-MS/MS spectrum of the rhodamine-based probe (**10**) linked to a peptide fragment of KS-B and model of the rhodamine-based probe (**10**) linked to Cys3559 of KS-B (PDB code 4KC5).

## Conclusions

Considering the large number of studies on phenazines and their biological activities, it is surprising that their ability to capture biogenic thiols and crosslink proteins has so far been overlooked. The discovery of the panphenazines and the mechanistic investigations revealed that phenazine radicals form conjugates with a range of thiols. Thus, we provide a plausible model for irreversible GSH reduction in human endothelial cells.<sup>19</sup> Furthermore, we report for the first time that phenazines form *S*-conjugates with proteins. Protein modification and aggregation mediated by small molecules are the focus of diverse research on biomaterials,<sup>46,47</sup> and these processes are also relevant for protein aging and degenerative diseases.<sup>42</sup> Our findings are significant because PCA is an important biosynthetic building block that is produced in large amounts by many pathogenic pseudomonads during infection processes.<sup>2,48</sup> Since phenazine radicals are readily formed in cells by redox cycling, PCA and congeners may contribute to protein degeneration processes. Specifically, our findings may explain the broad activity profiles of phenazines including protein aggregation in



*C. elegans*.<sup>23</sup> Radical-based thiol conjugation and protein crosslinking by phenazines represent novel modes of action that are an important addition to the body of knowledge on this widespread group of virulence factors.<sup>6</sup>

## Experimental

### General analytical procedures

All NMR experiments were performed on Bruker AVANCE II 300, AVANCE III 500 or a 600 MHz spectrometer, equipped with a Bruker Cryo Platform. The chemical shifts are reported in parts per million (ppm) relative to the solvent residual peak of chloroform- $d_1$  ( $^1\text{H}$ : 7.24 ppm, singlet;  $^{13}\text{C}$ : 77.00 ppm, triplet), methanol- $d_4$  ( $^1\text{H}$ : 3.30 ppm, quintet;  $^{13}\text{C}$ : 49.00 ppm, septet) or DMSO- $D_6$  ( $^1\text{H}$ : 2.50 ppm, quintet;  $^{13}\text{C}$ : 39.50 ppm, septet). HR ESI-MS was carried out on an Accela UPLC-system (Thermo Scientific) combined with an Exactive mass spectrometer (Thermo Scientific) equipped with an electrospray ion source. Solid phase extraction was carried out using Chromabond  $\text{C}_{18}$  ec Cartridges filled with 2000 mg of octadecyl-modified silica gel (Macherey-Nagel). HPLC-MS measurements were performed on a HPLC 1100 System connected to a 1100 Series LC/MSD Trap using a Zorbax Eclipse XDB-C8,  $4.6 \times 150$  mm column, particle size  $5 \mu\text{m}$  (Agilent Technologies). For thin layer chromatography TLC aluminium sheets silica gel 60  $\text{F}_{254}$  (MERCK) were used. Open column chromatography was performed using silica gel 60; particle size 0.015–0.04 mm (Macherey-Nagel) and on Sephadex® LH-20 (Sigma-Aldrich). Semi preparative HPLC was performed on an 1260 Infinity System (Agilent Technologies) using the following columns: Zorbax Eclipse XDB-C8,  $9.5 \times 250$  mm, particle size  $5 \mu\text{m}$  (Agilent Technologies), Zorbax Eclipse XDB-C18,  $9.5 \times 250$  mm, particle size  $5 \mu\text{m}$  (Agilent Technologies) and EC 250/4 Nucleodur Sphinx RP,  $10 \times 250$  mm, particle size  $5 \mu\text{m}$  (Macherey-Nagel). All solvents for analytical and preparative HPLC were obtained commercially in gradient grade and were filtered prior to use. To avoid microbial growth 0.1% formic acid was added to the water, used for analytical and preparative HPLC. UV-spectra were obtained using a UV-spectrometer UV-1800 (Shimadzu). IR-spectra were recorded on an FT/IR-4100 ATR spectrometer (Jasco). For irradiation experiments an UV-lamp TL MINI 6 Watt 33-640 (Philips) and an UV-lamp F 6W T5 G5 129 3000K 280 lm (Sylvania) were used.

### Strain cultivation and isolation of panphenazines

The cultivation of *Kitasatospora* sp. HKI 714 and the fractionation of the crude extract were performed in exactly the same way as reported before.<sup>27</sup> Compounds **1** and **2** were isolated by semipreparative HPLC (Zorbax Eclipse XDB-C18,  $9.5 \times 250$  mm,  $5 \mu\text{m}$ , flow rate  $4 \text{ mL min}^{-1}$ , gradient starting from  $\text{H}_2\text{O}/\text{MeOH}$  90/10, in 2.5 min to 50/50, in 15.5 min to 46/54, in 7 min to 42/58, in 5 min to 0/100).

**Compound 3.** Yellow solid, for a full list of NMR signals see Table S1† in the SI; IR (film): 3308, 2923, 2854, 1700, 1652, 1421, 1265, 1053,  $759 \text{ cm}^{-1}$ ; UV/vis (MeOH):  $\lambda_{\text{max}}$  nm (log  $\epsilon$ ): 236 (6.22), 274 (6.36), 381 (5.87), 427 (5.78) nm; HR ESI-MS:  $m/z = 501.1807$  [ $\text{M} + \text{H}$ ]<sup>+</sup> (calculated 501.1809 for  $\text{C}_{24}\text{H}_{29}\text{O}_6\text{N}_4\text{S}$ ).

**Compound 4.** Yellow solid, for a full list of NMR signals see Table S1† in the SI; IR (film): 3297, 2929, 1651, 1541, 1410, 1045,  $757 \text{ cm}^{-1}$ ; UV/vis (MeOH):  $\lambda_{\text{max}}$  nm (log  $\epsilon$ ): 261 (6.05), 372 (5.52), 408 (5.41) nm; HR ESI-MS:  $m/z = 501.1808$  [ $\text{M} + \text{H}$ ]<sup>+</sup> (calculated 501.1809 for  $\text{C}_{24}\text{H}_{29}\text{O}_6\text{N}_4\text{S}$ ).

### General synthetic procedures

All reagents were obtained from commercial suppliers (Sigma Aldrich and Acros Organics) and were used without further purification. Carbonic anhydrase III (29.5 kDa) was obtained from GE Healthcare UK Limited and albumin (66 kDa) was obtained from Carl Roth GmbH. Chemical solvents were obtained at least in HPLC grade. Methanol and chloroform were distilled prior to use. Reactions were performed under inert atmosphere (Argon) using the Schlenk technique.

### Pantetheine

A solution of D-pantetheine (200 mg, 0.36 mmol) and dithiothreitol (100 mg, 0.65 mmol) in a methanolic solution (4 mL; methanol : water; 1 : 1; v/v) was stirred for 2 days in an inert atmosphere (argon). The solvent was removed under reduced pressure and the product purified by column chromatography using silica gel (chloroform/methanol: 98/2: v/v) to give 195 mg of a colourless liquid. Yield 97%.  $^1\text{H}$  NMR (300 MHz, MeOD):  $\delta$  0.9 (6H, s), 2.4 (2H, t), 2.6 (2H, t), 3.3–3.6 (6H, m), 3.9 (1H, s) ppm.  $^{13}\text{C}$  NMR (75.5 MHz, MeOD):  $\delta$  20.9, 21.3, 24.5, 36.4, 40.4, 43.9, 70.3, 77.2, 79.5, 173.9, 176.1 ppm.

### Synthesis of panphenazines (3 and 4)

Phenazine-1-carboxylic acid (**2**, 3 mg, 0.013 mmol) was dissolved in DMSO (300  $\mu\text{L}$ ) and pantetheine (4 mg, 0.013 mmol) was added. The stirred mixture was incubated at 370 nm for 24 h. The solvent was removed under reduced pressure and the crude product dissolved in MeOH. HPLC-HRMS proved the identity of one product with compound **4**. Compound **3** was only produced in minor amounts under these conditions. The yield of **4** was quantified by integration of its peak area at a detector wavelength of 430 nm. Under these conditions yield was determined to be 2%. Repeating the reaction with the addition of TEA (1.9  $\mu\text{L}$ , 0.014 mmol) increased the yield to 24%. In contrast, addition of TBHT instead of TEA hampered the reaction and no corresponding signal for **4** could be detected in the UV trace.

### Synthesis of phenazine *N*-acetylcysteamine conjugates (5–8)

Phenazine-1-carboxylic acid (**2**, 10 mg, 0.045 mmol) and *N*-acetylcysteamine (5 mg, 0.042 mmol) were dissolved in DMSO and incubated at 370 nm for 24 h. The solvent was removed under reduced pressure and the residue was subjected to semipreparative HPLC (Zorbax Eclipse XDB-C8,  $9.5 \times 250$  mm,  $5 \mu\text{m}$ , flow rate  $4 \text{ mL min}^{-1}$ , gradient starting from  $\text{H}_2\text{O}/\text{ACN}$  90/10, in 3 min to 80/20, in 25 min to 60/40, in 2 min to 0/100). Yields of monosubstituted phenazines: **5** (10%), **6** (8%), **7** (5%), **8** (6%) (yellow solids).

**Compound 5.**  $^1\text{H}$  NMR (500 MHz, DMSO- $D_6$ ):  $\delta$  1.58 (3H, s), 3.15 (2H, t,  $^3J = 6.7$  Hz), 3.22–3.27 (2H, m), 7.92–7.98 (2H, m),



8.03 (1H, d,  $^3J = 9.2$  Hz), 8.12 (1H, d,  $^3J = 9.2$  Hz), 8.16–8.21 (1H, m), 8.21–8.25 (1H, m) ppm.  $^{13}\text{C}$  NMR (125 MHz, DMSO- $D_6$ ):  $\delta$  22.3, 32.7, 38.8, 128.2, 129.3, 131.0, 131.3, 132.4, 139.8, 141.3, 142.3, 142.5, 163.1<sup>#</sup>, 168.2, 169.3 ppm.

**Compound 6.**  $^1\text{H}$  NMR (600 MHz, DMSO- $D_6$ ):  $\delta$  1.82 (3H, s), 3.28 (2H, t,  $^3J = 6.8$  Hz), 3.36–3.44 (2H, m)\*, 7.58 (1H, m), 7.87–7.94 (3H, m), 8.16 (1H, dd,  $^3J = 8.4$  Hz,  $^4J = 1.4$  Hz), 8.19 (1H, d,  $^3J = 8.4$  Hz), 8.23 (1H, t,  $^3J_1 = 5.5$  Hz, NH) ppm.  $^{13}\text{C}$  NMR (150.9 MHz, DMSO- $D_6$ ):  $\delta$  22.6, 30.4, 37.7, 120.7, 127.0, 128.9, 129.4, 130.2, 131.2, 139.2, 141.2, 141.3, 142.6, 143.2, 164.3, 168.5, 169.6 ppm.

**Compound 7.**  $^1\text{H}$  NMR (600 MHz, DMSO- $D_6$ ):  $\delta$  1.83 (3H, s), 3.22 (2H, t,  $^3J = 6.9$  Hz), 3.40–3.44 (2H, m), 7.75 (1H, d,  $^3J = 7.2$  Hz), 7.86 (1H, d,  $^3J = 7.2$  Hz), 7.95–8.01 (2H, m), 8.20–8.29 (3H, m) ppm.  $^{13}\text{C}$  NMR (150.9 MHz, DMSO- $D_6$ ):  $\delta$  22.6, 29.6, 37.5, 124.0, 128.6, 129.2, 129.6, 131.2, 131.4, 133.6, 140.1, 140.6, 140.7, 142.1, 143.8, 165.4, 169.5 ppm.

**Compound 8.**  $^1\text{H}$  NMR (600 MHz, DMSO- $D_6$ ):  $\delta$  1.82 (3H, s), 3.30 (2H, t,  $^3J = 6.9$  Hz), 3.39–3.43 (2H, m), 7.73 (1H, d,  $^3J = 6.6$  Hz), 7.77 (1H, dd,  $^3J = 9.2$  Hz,  $^4J = 1.5$  Hz), 7.80 (1H, dd,  $^3J_1 = 8.6$  Hz,  $^3J_2 = 6.6$  Hz), 7.99 (1H, s), 8.04 (1H, d,  $^3J = 8.6$  Hz), 8.09 (1H, d,  $^3J = 9.2$  Hz), 8.28–8.33 (1H, m, NH) ppm.  $^{13}\text{C}$  NMR (150.9 MHz, DMSO- $D_6$ ):  $\delta$  22.5, 30.5, 37.6, 121.9, 125.4, 127.0, 127.5, 129.3, 130.3, 130.9, 140.4, 141.1, 142.0, 142.5, 165.5, 169.6, 169.8 ppm. \*Signal partly obscured, #signal only detectable from HMBC data.

## Synthesis of phenazine-biotine and -rhodamine probes

### Tetraethyleneglycol phenazine-1-carboxylate (11)

Phenazine-1-carboxylic acid **2** (10 mg, 0.044 mmol) was dissolved in dichloromethane (5 mL) and cooled to 0 °C. Tetraethylene glycol (26 mg, 0.13 mmol, 3 eq.), DMAP (5 mg, 0.04 mmol) and EDC (8.4 mg, 0.05 mmol) were added. After 24 h, water was added and the aqueous phase extracted with dichloromethane (2 × 10 mL). The combined organic fractions were dried over sodium sulfate and concentrated under reduced pressure. Column chromatography over silica gel (chloroform/methanol: 98/2, v/v) gave the product (15.3 mg) as yellow oil. Yield: 86%.  $^1\text{H}$  NMR (600 MHz, MeOD):  $\delta$  3.47 (2H, t,  $^3J = 4.9$  Hz), 3.54 (2H, m), 3.58 (2H, t,  $^3J = 4.9$  Hz), 3.61 (2H, m), 3.66 (2H, m), 3.73 (2H, m), 3.93 (2H, t,  $^3J = 4.7$  Hz), 4.65 (2H, t,  $^3J = 4.7$  Hz), 7.96–7.99 (3H, m), 8.24–8.27 (1H, m), 8.31 (1H, dd,  $^3J = 6.8$  Hz,  $^4J = 1.3$  Hz), 8.33 (1H, m), 8.40 (1H, dd,  $^3J = 8.8$  Hz,  $^4J = 1.3$  Hz) ppm.  $^{13}\text{C}$  NMR (150.9 MHz, MeOD):  $\delta$  62.2, 66.0, 70.2, 71.4, 71.6, 71.7, 73.6, 79.5, 130.3, 130.9, 131.0, 132.7, 132.8, 132.9, 133.4, 134.0, 142.0, 143.8, 144.5, 144.9, 168.1 ppm; IR (film): 1119, 1214, 1237, 1274, 1350, 1421, 1523, 1727, 2874, 3018  $\text{cm}^{-1}$ , HRMS: (ESI<sup>+</sup>):  $m/z$  calculated for  $\text{C}_{21}\text{H}_{25}\text{O}_6\text{N}_2$  [M + H]<sup>+</sup> 401.1707, found: 401.1710.

### Biotin-labelled phenazine-1-carboxylic acid (9)

Tetraethylene-glycol phenazine-1-carboxylate (**7** mg, 0.017 mmol) **11** and biotin (8.5 mg, 0.035 mmol, 2 eq.) were dissolved in dry dichloromethane (2 mL) and cooled to 0 °C. DMAP (2 mg, 0.017 mmol) and EDC (8 mg, 0.044 mmol, 2.5 eq.) were added. Water (5 mL) was added after 20 h and the aqueous phase

extracted with dichloromethane (2 × 5 mL). The combined organic fractions were dried over sodium sulfate and concentrated under reduced pressure. Column chromatography over silica gel (chloroform/methanol: 95/5, v/v) gave 10.5 mg of a yellow solid. Yield: 96%.  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ ):  $\delta$  1.56–1.70 (6H, m), 2.28–2.32 (2H, m), 2.67–2.73 (1H, m), 2.81–2.88 (1H, m), 3.06–3.13 (1H, m), 3.57–3.65 (10H, m), 3.72 (2H, dd,  $^3J_1 = 5.4$  Hz,  $^3J_2 = 4.1$  Hz), 3.91 (2H, t,  $^3J_1 = 4.8$  Hz), 4.14–4.18 (2H, m), 4.23–4.28 (1H, m), 4.43–4.48 (1H, m), 4.65 (2H, t,  $^3J_1 = 4.8$  Hz), 7.82–7.86 (3H, m), 8.20–8.23 (1H, m), 8.23 (1H, dd,  $^3J_1 = 7.1$  Hz,  $^4J_2 = 1.3$  Hz), 8.27–8.30 (1H, m), 8.37 (1H, dd,  $^3J_1 = 8.8$  Hz,  $^4J_2 = 1.3$  Hz) ppm.  $^{13}\text{C}$  NMR (125 MHz, MeOD):  $\delta$  ppm.  $^{13}\text{C}$  NMR (150.9 MHz,  $\text{CDCl}_3$ ):  $\delta$  24.6, 28.1, 28.2, 33.7, 40.4, 55.4, 60.1, 61.9, 63.3, 64.6, 69.0, 69.1, 70.4, 70.5, 70.6, 70.6, 129.1, 129.3, 130.3, 131.0, 131.2, 131.3, 132.1, 133.2, 141.0, 142.5, 143.1, 143.6, 166.4, 173.5, 174.1 ppm; IR (film): 747, 1038, 1119, 1179, 1236, 1264, 1455, 1522, 1700, 2867, 2925, 3236  $\text{cm}^{-1}$ , HRMS: (ESI<sup>+</sup>):  $m/z$  calculated for  $\text{C}_{31}\text{H}_{39}\text{O}_8\text{N}_4\text{S}$  [M + H]<sup>+</sup> 627.2483, found: 627.2477.

### Rhodamine-labelled phenazine-1-carboxylic acid (10)

To a solution of rhodamine B (1.9 mg, 0.0043 mmol), tetraethyleneglycol phenazine-1-carboxylate **11** (1.7 mg, 0.0043 mmol) and PyBop (2.7 mg, 0.0052 mmol, 1.2 eq.) in anhydrous dichloromethane (1 mL) was added Hünigs base (2.9  $\mu\text{L}$ , 4 eq.) at 0 °C. The reaction was allowed to warm to room temperature and stirred for 24 h. The solution was concentrated under reduced pressure and the residue was subjected to column chromatography over silica gel (gradient chloroform/MeOH: 100/0–95/5, v/v). The crude product was subjected to semi-preparative HPLC using a Zorbax Eclipse XDB-C8 (acetonitrile- $\text{H}_2\text{O}$ , 0.1% formic acid; starting conditions: 20% acetonitrile, 40% in 4 min, 60% in 12.5 min, 100% in 0.5 min, 100% for 5 min, 20% in 1 min) to give 2.2 mg of a white solid. Yield: 63%.  $^1\text{H}$  NMR ( $\text{CDCl}_3$ , 600 MHz)  $\delta$  = 1.26–1.37 (12H, m), 3.45–3.49 (3H, m), 3.50–3.59 (9H, m), 3.61–3.64 (2H, m), 3.64–3.68 (1H, m), 3.69–3.72 (2H, m), 3.73–3.79 (1H, m), 3.90 (2H, t,  $^3J = 4.8$  Hz), 4.12 (2H, t,  $^3J = 4.7$  Hz), 4.63 (2H, t,  $^3J = 4.8$  Hz), 6.78–6.81 (3H, m), 7.02 (1H, s), 7.04 (1H, s), 7.28 (1H, dd,  $^3J_1 = 7.6$  Hz,  $^4J_2 = 0.7$  Hz), 7.61–7.82 (3H, m), 7.85–7.91 (3H, m), 8.25 (1H, dd,  $^3J_1 = 6.9$  Hz,  $^4J_2 = 1.1$  Hz), 8.26 (1H, dd,  $^3J_1 = 8.0$  Hz,  $^4J_2 = 1.0$  Hz), 8.28–8.32 (2H, m), 8.28 (1H, d,  $^3J_1 = 8.8$  Hz) ppm.  $^{13}\text{C}$  NMR ( $\text{CDCl}_3$ , 600 MHz)  $\delta$  = 12.6, 12.6, 12.6, 13.7, 46.0, 46.0, 47.4, 64.7, 64.8, 68.6, 69.2, 70.4, 70.5, 70.6, 70.7, 96.4, 113.6, 114.0, 119.0, 128.7, 129.0, 129.6, 129.8, 130.2, 130.3, 131.2, 131.3, 131.3, 131.4, 131.4, 131.7, 132.3, 132.8, 132.9, 133.0, 133.6, 141.1, 141.9, 142.5, 143.8, 153.8, 155.5, 157.8, 158.8, 165.0, 166.3 ppm. IR (film): 1077, 1126, 1183, 1342, 1410, 1462, 1590, 1651, 1722, 2353, 2860, 2923, 3318  $\text{cm}^{-1}$ , HRMS: (ESI<sup>+</sup>):  $m/z$  calculated for  $\text{C}_{49}\text{H}_{53}\text{O}_8\text{N}_4$  [M]<sup>+</sup> 825.3858, found: 825.3868.

### Phenazine-protein binding assays

The biotinylated probe **9** was directly mixed with the protein (KS-B – 104.7 kDa) in 20 mM Tris-HCl (pH 8.0). Final concentrations of the probe and protein were 0.5 mM and 20  $\mu\text{M}$  respectively. The reaction mixture was incubated for 24 h under



the following conditions: (a) at 254 nm (b) at 370 nm or (c) at 22 °C in the presence of 0.5 mM azobisisobutyronitrile (AIBN). For purification of the biotinylated KS-B, the reaction mixture was diluted twice with 50 mM Tris, 0.2 M NaCl, 0.1% SDS (pH 8.0) and incubated with 1 mL of streptavidin beads (Thermo Scientific) at 22 °C for 1 h with shaking at 400 rpm. The beads were washed twice with 50 mM Tris, 0.2 M NaCl, 2% SDS (pH 8.0) after incubation at 60 °C for 10 min. Elution of the bound protein was achieved by heating the sample thrice at 60 °C for 30 min in 50 mM Tris, 0.2 M NaCl, 2% SDS, 3 mM D-biotin (pH 8.0).

The rhodamine-based probe **10** was incubated with the protein (KS-B, carbonic anhydrase III, albumin) in water or in DMSO. Final concentrations of the probe and the protein were 10 μM and 20 μM respectively. The reaction mixture was incubated for varying times between 15 min and 20 h under UV-light at 254 nm, at 370 nm, at 22 °C in the presence of 0.5 mM AIBN in the dark or at sunlight (specific times and conditions for each experiment are indicated in the respective figure legend). After the assay the protein samples were analysed by PAGE, on a 10% or 12% SDS-gel for KS-B. The structures of KS-B and albumin were visualised using Pymol. The probe **10** was generated as a fragment and added specifically to the Cys residue of the proteins by replacing the hydrogen atom in Pymol.

## Acknowledgements

We thank H. Heinecke and A. Perner for NMR and HRMS measurements, M. Poetsch for MALDI experiments, K. Martin and K. Perlet for strain handling, and K.-D. Menzel and M. Steinacker for support in large-scale fermentation and downstream processing. We gratefully acknowledge the Studienstiftung des Deutschen Volkes (to D. H.) and the International Leibniz Research School for Microbial and Biomolecular Interactions (ILRS, to S. S.) for financial support.

## References

- S. Chincholkar and L. Thomashow, *Microbial Phenazines*, Springer, Berlin, 2013.
- J. B. Laursen and J. Nielsen, *Chem. Rev.*, 2004, **104**, 1663–1686.
- A. Cimmino, A. Evidente, V. Mathieu, A. Andolfi, F. Lefranc, A. Kornienko and R. Kiss, *Nat. Prod. Rep.*, 2012, **29**, 487–501.
- J. Davies and K. S. Ryan, *ACS Chem. Biol.*, 2012, **7**, 252–259.
- A. Price-Whelan, L. E. P. Dietrich and D. K. Newman, *Nat. Chem. Biol.*, 2006, **2**, 71–78.
- L. S. I. Pierson and E. A. Pierson, *Appl. Microbiol. Biotechnol.*, 2010, **86**, 1659–1670.
- D. Romero, M. F. Traxler, D. Lopez and R. Kolter, *Chem. Rev.*, 2011, **111**, 5492–5505.
- D. V. Mavrodi, J. A. Parejko, O. V. Mavrodi, Y. S. Kwak, D. M. Weller, W. Blankenfeldt and L. S. Thomashow, *Environ. Microbiol.*, 2013, **15**, 1462–2920.
- L. E. Dietrich, A. Price-Whelan, A. Petersen, M. Whiteley and D. K. Newman, *Mol. Microbiol.*, 2006, **61**, 1308–1321.
- L. E. Dietrich, C. Okegbe, A. Price-Whelan, H. Sakhtah, R. C. Hunter and D. K. Newman, *J. Bacteriol.*, 2013, **195**, 1371–1380.
- L. E. Dietrich, T. K. Teal, A. Price-Whelan and D. K. Newman, *Science*, 2008, **321**, 1203–1206.
- A. J. Stams, F. A. de Bok, C. M. Plugge, M. H. van Eekert, J. Dolging and G. Schraa, *Environ. Microbiol.*, 2006, **8**, 371–382.
- C. C. Caldwell, Y. Chen, H. S. Goetzmann, Y. Hao, M. T. Borchers, D. J. Hassett, L. R. Young, D. Mavrodi, L. Thomashow and G. W. Lau, *Am. J. Pathol.*, 2009, **175**, 2473–2488.
- M. Mentel, E. G. Ahuja, D. V. Mavrodi, R. Breinbauer, L. S. Thomashow and W. Blankenfeldt, *ChemBioChem*, 2009, **10**, 2295–2304.
- R. C. Hunter, V. Klepac-Ceraj, M. M. Lorenzi, H. Grotzinger, T. R. Martin and D. K. Newman, *Am. J. Respir. Cell Mol. Biol.*, 2012, **47**, 738–745.
- H. Ran, D. J. Hassett and G. W. Lau, *Proc. Natl. Acad. Sci. U. S. A.*, 2003, **100**, 14315–14320.
- K. Apel and H. Hirt, *Annu. Rev. Plant Biol.*, 2004, **55**, 373–399.
- Y. Q. O'Malley, K. J. Reszka, D. R. Spitz, G. M. Denning and B. E. Britigan, *Am. J. Physiol.: Lung Cell. Mol. Physiol.*, 2004, **287**, L94–L103.
- M. Muller, *Free Radical Biol. Med.*, 2002, **33**, 1527–1533.
- M. Muller, *Free Radical Biol. Med.*, 2011, **50**, 971–977.
- R. Cheluvappa, R. Shimmon, M. Dawson, S. N. Hilmer and D. G. Le Couteur, *Acta Biochim. Pol.*, 2008, **55**, 571–580.
- M. Muller and N. D. Merrett, *Chem. Biol. Interact.*, 2015, **232**, 30–37.
- A. Ray, C. Rentas, G. A. Caldwell and K. A. Caldwell, *Neurosci. Lett.*, 2015, **584**, 23–27.
- A. J. McFarland, S. Anoopkumar-Dukie, A. V. Perkins, A. K. Davey and G. D. Grant, *Arch. Toxicol.*, 2012, **86**, 275–284.
- B. Rada and T. L. Leto, *Trends Microbiol.*, 2013, **21**, 73–81.
- J. M. Winter, S. Behnken and C. Hertweck, *Curr. Opin. Chem. Biol.*, 2011, **15**, 22–31.
- D. Heine, K. Martin and C. Hertweck, *J. Nat. Prod.*, 2014, **77**, 1083–1087.
- M. Wagner, W. M. Abdel-Mageed, R. Ebel, A. T. Bull, M. Goodfellow, H.-P. Fiedler and M. Jaspars, *J. Nat. Prod.*, 2014, **77**, 416–420.
- M. L. Gilpin, M. Fulston, D. Payne, R. Cramp and I. Hood, *J. Antibiot.*, 1995, **48**, 1081–1085.
- A. Braunshausen and F. P. Seebeck, *J. Am. Chem. Soc.*, 2011, **133**, 1757–1759.
- D. H. Scharf, P. Chankhamjon, K. Scherlach, T. Heinekamp, K. Willing, A. A. Brakhage and C. Hertweck, *Angew. Chem., Int. Ed.*, 2013, **52**, 11092–11095.
- D. H. Scharf, N. Remme, A. Habel, P. Chankhamjon, K. Scherlach, T. Heinekamp, P. Hortschansky, A. A. Brakhage and C. Hertweck, *J. Am. Chem. Soc.*, 2011, **133**, 12322–12325.
- S. Coyne, C. Chizzali, M. N. Khalil, A. Litomska, K. Richter, L. Beerhues and C. Hertweck, *Angew. Chem., Int. Ed.*, 2013, **52**, 10564–10568.



- 34 E. Sasaki, X. Zhang, H. G. Sun, M. Y. Lu, T. L. Liu, A. Ou, J. Y. Li, Y. H. Chen, S. E. Ealick and H. W. Liu, *Nature*, 2014, **510**, 427–431.
- 35 K. V. Goncharenko, A. Vit, W. Blankenfeldt and F. P. Seebeck, *Angew. Chem., Int. Ed.*, 2015, **54**, 2821–2824.
- 36 B. Li, J. P. Yu, J. S. Brunzelle, G. N. Moll, W. A. van der Donk and S. K. Nair, *Science*, 2006, **311**, 1464–1467.
- 37 T. Ogata, Y. Yamamoto, Y. Wada, K. Murakoshi, M. Kusaba, N. Nakashima, A. Ishida, S. Takamuku and S. Yanagida, *J. Phys. Chem.*, 1995, **99**, 11916–11922.
- 38 G. A. Amaral, F. Ausfelder, J. G. Izquierdo, L. Rubio-Lago and L. Banares, *J. Chem. Phys.*, 2007, **126**, 024301.
- 39 S. Naumov and C. Schoneich, *J. Phys. Chem. A*, 2009, **113**, 3560–3565.
- 40 M. Gersch, J. Kreuzer and S. A. Sieber, *Nat. Prod. Rep.*, 2012, **29**, 659–682.
- 41 T. Böttcher, M. Pitscheider and S. A. Sieber, *Angew. Chem., Int. Ed.*, 2010, **49**, 2680–2698.
- 42 P. Ulrich and A. Cerami, *Recent Prog. Horm. Res.*, 2001, **56**, 1–22.
- 43 T. Bretschneider, J. B. Heim, D. Heine, R. Winkler, B. Busch, B. Kusebauch, T. Stehle, G. Zocher and C. Hertweck, *Nature*, 2013, **502**, 124–128.
- 44 D. Heine, T. Bretschneider, S. Sundaram and C. Hertweck, *Angew. Chem., Int. Ed.*, 2014, **53**, 11645–11649.
- 45 D. Heine, S. Sundaram, T. Bretschneider and C. Hertweck, *Chem. Commun.*, 2015, **51**, 9872–9875.
- 46 S. Adolph, V. Jung, J. Rattke and G. Pohnert, *Angew. Chem., Int. Ed.*, 2005, **44**, 2806–2808.
- 47 N. Reddy, R. Reddy and Q. Jiang, *Trends Biotechnol.*, 2015, **33**, 362–369.
- 48 G. V. Bloembergen and B. J. J. Lugtenberg, *New Phytol.*, 2003, **157**, 503–523.

