

Communication

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Diverted Total Synthesis of Promysalin Analogs Demonstrates that an Iron-Binding Motif is Responsible for its Narrow-Spectrum Antibacterial Activity

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Supporting Information Placeholder

Abstract: Promysalin is a species-specific *Pseudomonad* metabolite with unique bioactivity. To better understand the mode of action of this natural product, we synthesized sixteen analogs utilizing diverted total synthesis (DTS). Our analog studies revealed that the bioactivity of promysalin is sensitive to changes within its hydrogen bond network whereby alteration has drastic biological consequences. The DTS library not only yielded three analogs that retained potency but also provided insights that resulted in the identification of a previously unknown ability of promysalin to bind iron. These findings coupled with previous observations hint at a complex multi-faceted role of the natural product within the rhizosphere.

Over the past century, groundbreaking discoveries in the field of antibiotics have resulted in the saving of countless lives. This can be primarily attributed to their broad-spectrum activity that is ideal for the treatment of a wide range of bacterial infections but can also result in significant misuse and unintended side effects.¹ For example, recent work from the Blaser lab strongly suggests that prolonged broad-spectrum antibiotic exposure early in life can lead to an increased likelihood of obesity, allergies, and inflammatory diseases.¹ These findings highlight the effect that certain antimicrobials have on one's commensal population, resulting in an altered community dynamic and leading to undesired outcomes lending credence to the recent call for the development of narrow-spectrum therapies.

To date very few options currently exist for pathogen-specific treatments.² Furthermore, with species-specific compounds in hand one would also have access to tool compounds to aid in deconvoluting the complex multispecies environments present. The recent advances in genetic sequencing, as exemplified by the human microbiome project, would allow for the probing of these environments if the appropriate tools were available.³

Our group has been particularly interested in developing narrow-spectrum agents to combat *Pseudomonas aeruginosa* (PA) infections. PA is a Gram-negative pathogen that is typically found in people with weakened immune systems and/or hospital settings accounting for an estimated 51,000 infections/year in the United States alone.⁴ The community, including our own lab,⁵ has postulated that the rhizosphere, which is defined as the immediate area of soil that is directly influenced by microorganisms and root secretions, is an ideal environment to discover such compounds as the soil is teeming with *Pseudomonads* competing to establish their own niches. This hypothesis has motivated our interest in the unique and highly specific *Pseudomonad* natural product promysalin (**1**, Figure 1).

We recently reported the total synthesis and structural elucidation of promysalin, which is produced by *Pseudomonas putida* (PP), and inhibits the growth of PA at nM concentrations,⁶ while Gram-positive bacteria show no susceptibility.⁷ Furthermore, during our biological studies we found that promysalin inhibits the production of pyoverdine (a

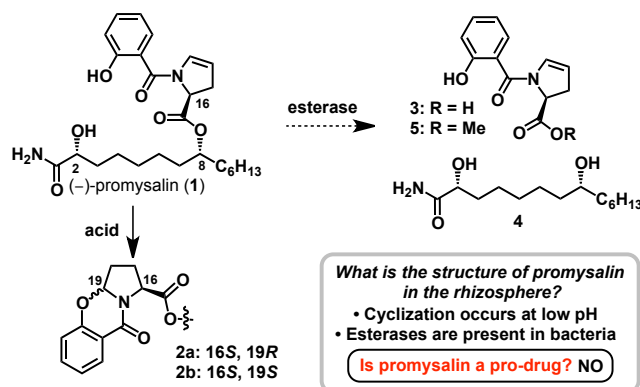
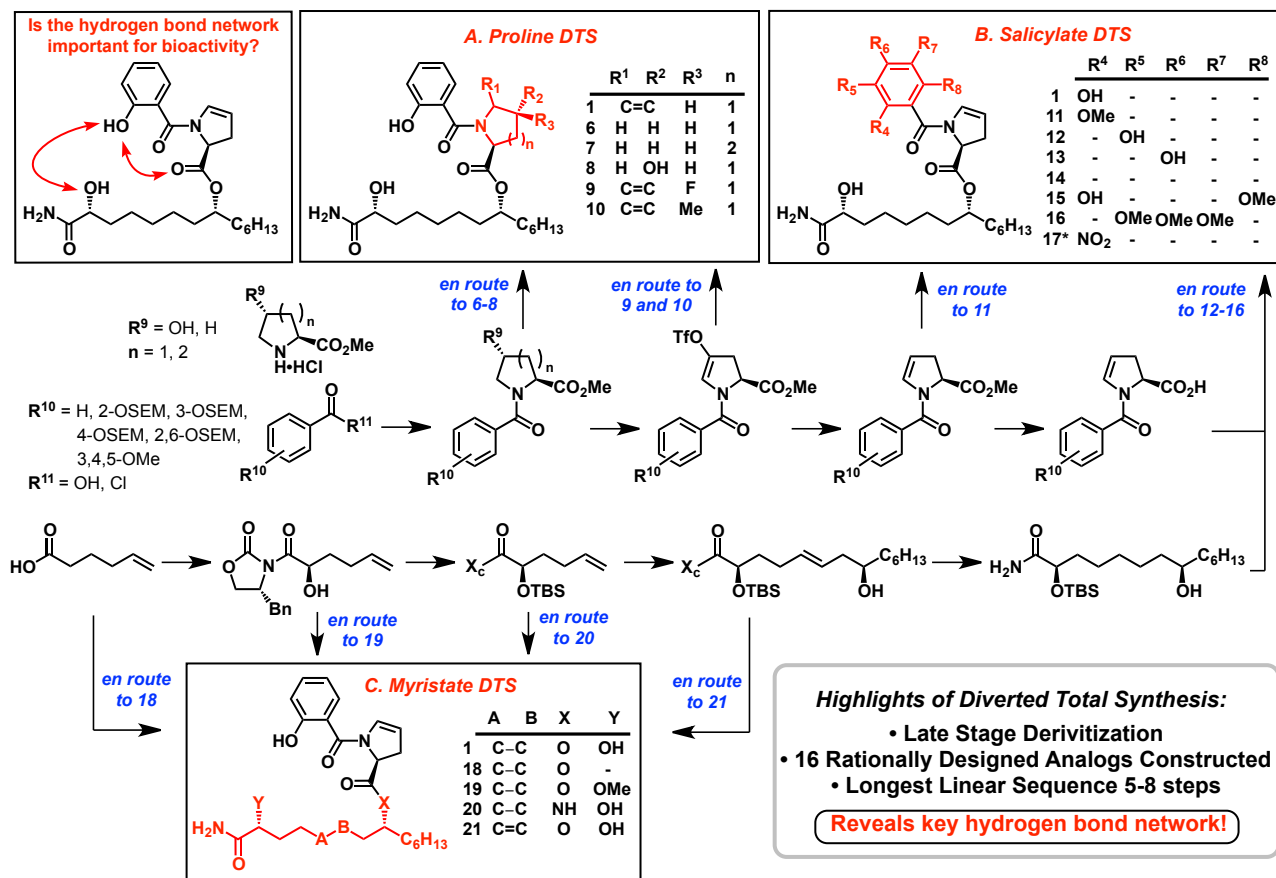


Figure 1. Structure of promysalin (**1**) and compounds **2** - **5** involved in our pro-drug hypothesis.

siderophore) in *P. putida* KT2440, and as expected, only one stereoisomer elicited this phenotype. Siderophores are essential for the growth of *Pseudomonads* as they actively chelate Fe^{3+} for its end use in enzymatic processes. Intriguingly, siderophore production is closely linked to virulence; therefore, inhibiting such a process may yield novel anti-virulence therapies.⁸ These initial discoveries prompted us to initiate a synthetic campaign to elucidate what chemical moieties are responsible for the species-specific nature of promysalin.

At the outset, we hypothesized two potential mechanisms of action. First, promysalin may be a "prodrug", whereby it would be activated by either an enzyme (i.e. an esterase) unique to PA or by the physical environment surrounding PA itself. A second alternative is that the chemical architecture of the molecule is exquisitely selective for a target harbored by PA but not present in other bacteria. Our earlier synthetic studies hinted that the exact three-dimensional shape of the molecule was key to its biological activity as we observed a 100-200-fold decrease in activity by altering either stereocenter on the fatty acid fragment. To address these questions, we constructed a library of sixteen hypothesis-driven analogs via diverted total synthesis (DTS).⁹

Figure 1 depicts the structure of promysalin (**1**) and the structures involved in our hypotheses. Enamines are quite susceptible to acid hydrolysis, and we were curious about the



Scheme 2. The diverted total synthesis of promysalin analogs to probe the hydrogen-bonding network. Blue wording depicts the branching points from our previously reported synthesis (for reaction conditions see ref. 6 and SI). Dashes in tables denote where hydrogens are present. *Compound 17 was prepared by a different route (See SI).

stability of promysalin in low pH environments. Somewhat surprisingly, we (and others)¹⁰ noticed the instability of promysalin under mildly acidic conditions, which we were able to attribute to the formation of cyclized products **2a/b**. We wondered if this cyclization event was biologically relevant, whereby the compound would be dispensed by PP and would undergo cyclization in the known acidic environment present around PA.¹¹ However, the cyclized compounds showed no biological activity. Similarly, it is well known that bacteria possess enzymes capable of hydrolyzing esters. Therefore, we questioned if the hydrolyzed fragments (**3** and **4**) would be active. We synthesized methyl ester **5** in lieu of the acid, for cell permeability purposes. Methyl ester **5** and diol **4** were both inactive up to concentrations of 250 μ M. Additionally, we have also recently shown that appending the acid fragment of promysalin to another myristate-derived natural product, lyngbic acid, yielded compounds devoid of activity against PA.¹² Taken in sum, it appears that the cyclization and hydrolysis reactions are synthetic artifacts and not biologically relevant.

Based on these results we next sought to leverage DTS to access a library of analogs to build a structure-activity relationship (SAR) profile and further test the importance of the dehydropyrrolidine heterocycle. DTS has proven useful in previous natural product mechanistic studies,¹³ and in some cases has provided therapeutically useful analogs, exemplified by the development of fludelone.¹⁴ It should be noted that the analogs accessed from DTS presented here are inaccessible by

enzymatic or chemical manipulation of the natural product directly, thus showcasing the power of organic synthesis.

We initially hypothesized that the enamide moiety of promysalin was responsible for the bioactivity wherein it would covalently interact with its biological target; therefore, we postulated that any modification would render the molecule inactive. We utilized DTS to access five specific dehydropyrrolidine derivatives (Scheme 1, Box A). The synthesis of the proline, piperidine, and hydroxyproline analogs **6** - **8** was straightforward using standard coupling reactions and protecting group manipulations. We next exploited the inherent reactivity of our triflate intermediate via Pd-mediated coupling reactions to provide access to analogs **9** and **10**.

Initial computational models of promysalin highlighted an intricate intramolecular hydrogen-bonding network resulting in a rigidified molecular framework (*vide infra*). We hypothesized that this network, composed of the phenol, hydroxyl, and ester moieties, were therefore critical for activity. To test this hypothesis we designed analogs varying these key functionalities (Scheme 1, Box B). The salicylate fragment was systematically altered in three specific ways: 1) the position (**12** and **13**) and presence (**14**) of the phenol, 2) the substitution of the phenol with either the corresponding methyl ether (**11**) or nitro group (**17**), or 3) by increasing the electron density within the ring (**15** and **16**). In a similar fashion we sought to probe the role of ester linkage and secondary alcohol (Scheme 1, Box C). For example, we envisioned converting the ester to the corresponding amide (**20**) to create a hydrolytically more stable analog, albeit one

possessing a drastically altered hydrogen-bond network. Alkene analog **21** would add rigidity to the scaffold and potentially stabilize the active conformation. Finally, the importance of the hydroxyl group was probed by either methylation (**19**) or omission (**18**). The detailed synthetic route for the analogs library is presented in the Supporting Information (Scheme S1 and S2).

With a library of sixteen analogs in hand, we evaluated all of the compounds for inhibitory activity against *P. aeruginosa* strains PAO1 and PA14. The data for the active analogs ($IC_{50} < 250 \mu M$) is shown in Table 1 along with the inhibitory data for the less potent diastereomers of promysalin (for numbering see Figure 1). The inhibition data supported our initial hypothesis that the conformation of promysalin is exquisitely linked to its inhibition of PA. Of our modifications to the proline structure, fluorination (**9**) – being the smallest steric perturbation, was the only compound with equipotency to that of promysalin. Methylation (**10**) was slightly tolerated, while the piperidine (**7**) and hydroxyproline (**8**) derivatives were inactive. To our surprise, proline derivative **6** retained modest activity, which may hint at an inhibitory mechanism that involves both structural recognition and covalent binding.

The structure of the salicylate moiety was largely unforgiving in terms of the position (**12**, **13**), substitution (**17**), or exclusion of the phenol (**14**). In addition, adding methoxy substituents in the presence (**15**) or absence (**16**) of the *o*-phenol was not tolerated. The only active salicylate analog was the methyl ether (**11**), albeit with of an order of magnitude less potency. In contrast, the side chain analogs all retained some, if not all activity, with the one exception being the amide (**20**). Methylation of the hydroxyl group (**19**) resulted in a decrease in potency on par with the methylated-phenol (**11**). Rigidifying the side chain by including the alkene (**21**) led to an equipotent analog. However, when the secondary alcohol is removed, providing compound **18**, biological activity is fully retained. At first this result was particularly surprising, as we postulated that the alcohol was integral to the hydrogen-bonding network, as evidenced by the difference in activity between **1** and **1b/c** (10-100 fold decrease in potency). However, when considering our proposed macrocyclic structure, the implications of epimerization at a hydroxyl group can lead to drastic bond angle changes, in contrast, **18** can adopt a similar conformation simply by substituting the amide carbonyl as a Lewis base in place of the alcohol. Furthermore, the synthesis of **18** requires two less overall steps, and is more atom-economical, than that of promysalin as the use of a protecting group and chiral auxiliary is avoided.

As mentioned earlier, *Pseudomonads* are well known to produce a variety of siderophores in the rhizosphere. A cursory look at the literature reveals two specific siderophores with known salicyamide iron-binding motifs, reminiscent of promysalin: pyochelin¹⁵ and pseudomonine¹⁶ (Figure 2A). PQS, the *Pseudomonas aeruginosa* 4-Quinolone Signaling Molecule, which regulates virulence in PA, has also been shown to bind iron.¹⁷ Intriguingly, ferrocin,¹⁸ another *Pseudomonad* siderophore that bears a fatty acid moiety, has been shown to possess antimicrobial activity specifically against Gram-negative bacteria, and in particular PA. Based on these findings, and in light of our bioactivity and modeling results, we speculated that promysalin might be capable of binding iron. This hypothesis was reinforced by earlier findings demonstrating that promysalin is capable of

promoting swarming in PP and inhibiting pyoverdine production. With these questions in mind, we tested the ability of promysalin to bind iron. Indeed, promysalin tested positive for Fe^{3+} chelation on CAS agar at concentrations ranging from 6-100 mM, albeit with reduced affinity when compared to the known iron chelator EDTA (Figure 2B). Furthermore, Fe^{3+} chelation was confirmed down to 1 mM using an Iron-CAS shuttle assay (see Supporting Information).¹⁹ These results hint at the ability of promysalin to act as a siderophore in various *Pseudomonads*; however, additional studies are needed to confirm this role and are currently ongoing in our laboratory.

	PAO1	PA14
1 (2R,8R)	4.1	0.067
1b (2R,8S)	46	6.6
1c (2S,8S)	90	22
1d (2S,8R)	33	4.3
6	111	28
9	7.7	0.019
10	32	12
11	57	6.7
18	5.8	0.035
19	38	11
21	8.3	0.067

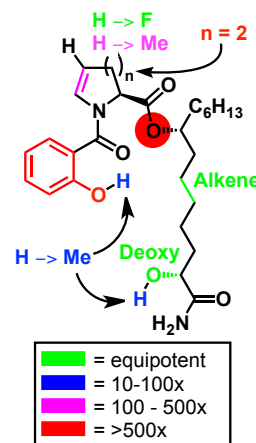


Table 1. IC_{50} values of promysalin diastereomers and active compounds in μM (compounds with $IC_{50} > 250 \mu M$ are not shown). Values are averages of three independent experiments (see SI for graphs and experimental details). A color-coded SAR is provided depicting the fold-decrease in activity (right).

These findings may allude to how the natural product can elicit its species-specific activity against PA and, furthermore, the increased selectivity of PA14 over that of PAO1. One possible mechanism of action of promysalin is through the inhibition of siderophore transport pathways thereby severely limiting or inhibiting the organism's ability to acquire iron. This particular mechanism is especially intriguing since PA14 significantly differs from PAO1 in its ability to acquire iron. Rahme and coworkers have recently shown that ybtQ, a yersiniabactin ABC-transporter (yersiniabactin is a phenolate-thiazoline siderophore of similar chemical structure to that of pyochelin), is found in PA14 and is primarily responsible for its virulence.²⁰

Chemists, and in particular the Miller group, have beautifully leveraged siderophore transport pathways using "sideromimetic" compounds whereby antibiotics are coupled to known siderophores for uptake.²¹ Conversely, in Nature, there are only a few reports of biosynthesized iron-binding molecules that possess antibacterial activity and none have been conclusively shown to inhibit siderophore transport systems. However, it seems increasingly likely that within the rhizosphere this type of process is indeed taking place, resulting in narrow-spectrum Gram-negative agents. As mentioned earlier, ferrocin is a compound produced by the soil-dwelling bacteria *Pseudomonas fluorescens* YK-310 and has been shown to inhibit Gram-negative bacteria. In addition to binding iron, both compounds possess amphiphilic features that may play a role in their potent activities. Nevertheless, the promise of highly potent and selective agents against

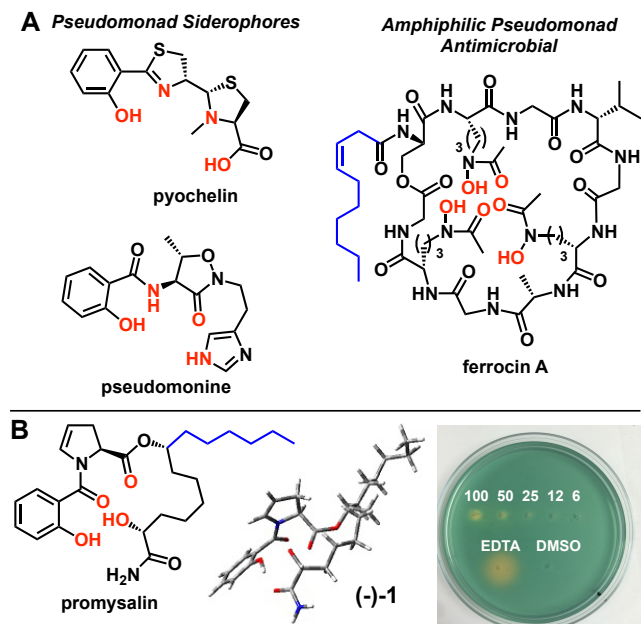


Figure 2. A) Structures of PA siderophores pyochelin and pseudomonine, and antimicrobial siderophore ferrocin. Putative iron-binding atoms are colored red and fatty acid tails in blue. B) Structure of promysalin with proposed iron contacts shown in red (left), minimized calculated structure (middle), CAS agar plate (right). Numbers refer to the concentration (mM) of 10 μ L of promysalin used. EDTA (10 μ L of 6 mM) and DMSO (10%) are used as controls.

Pseudomonads warrants further investigation.

In conclusion, we have leveraged the power of DTS to access a sixteen-membered library of rationally designed synthetic promysalin analogs. The structural diversity has shed light on the key structural features responsible for the bioactivity and highlights the importance of the key functionality within the hydrogen-bonding network (which are presumably responsible for binding iron). Further, these findings have led to the discovery of the iron-binding ability of promysalin, hinting at a secondary role for the natural product as a rhizosphere siderophore. In light of these results, a potential mechanism of action via the inhibition of siderophore transport seems feasible. Current work in our laboratory is now focused on determining if in fact promysalin is a siderophore and unequivocally identifying its biological target and will be reported in due course.

ASSOCIATED CONTENT

Supporting Information

Experimental procedures, characterization data, NMR spectra and supporting figures. The Supporting Information is available free of charge on the ACS Publications website at DOI:xxxx

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Notes

The authors declare no competing financial interest

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