

Small Molecule Inhibitors of the Bacterioferritin (BfrB)–Ferredoxin (Bfd) Complex Kill Biofilm-Embedded *Pseudomonas aeruginosa* Cells

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Cite This: <https://dx.doi.org/10.1021/acsinfecdis.0c00669>



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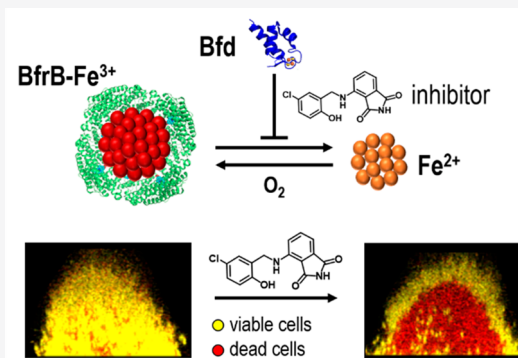
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ABSTRACT: Bacteria depend on a well-regulated iron homeostasis to survive adverse environments. A key component of the iron homeostasis machinery is the compartmentalization of Fe^{3+} in bacterioferritin and its subsequent mobilization as Fe^{2+} to satisfy metabolic requirements. In *Pseudomonas aeruginosa* Fe^{3+} is compartmentalized in bacterioferritin (BfrB), and its mobilization to the cytosol requires binding of a ferredoxin (Bfd) to reduce the stored Fe^{3+} and release the soluble Fe^{2+} . Blocking the BfrB–Bfd complex in *P. aeruginosa* by deletion of the *bfd* gene triggers an irreversible accumulation of Fe^{3+} in BfrB, concomitant cytosolic iron deficiency and significant impairment of biofilm development. Herein we report that small molecules developed to bind BfrB at the Bfd binding site block the BfrB–Bfd complex, inhibit the mobilization of iron from BfrB in *P. aeruginosa* cells, elicit a bacteriostatic effect on planktonic cells, and are bactericidal to cells embedded in mature biofilms.

KEYWORDS: biofilm, *Pseudomonas aeruginosa*, bacterioferritin, iron homeostasis, iron metabolism



Antimicrobial resistant bacteria are a major threat to public health. The World Health Organization (WHO) has released a priority list of pathogens for which antibiotics are urgently needed.¹ Carbapenem-resistant *Acinetobacter baumannii*, *Pseudomonas aeruginosa* and *Enterobacteriaceae* are at the top of the WHO priority list because of their resistance to multiple drug classes and associated mortality. *P. aeruginosa*, one of the leading Gram-negative pathogens associated with nosocomial infections has a propensity to form biofilms, which are populations of bacteria living in organized structures embedded in a self-produced matrix of DNA, proteins, and polysaccharides.^{2–4} Although the preference of bacteria for a biofilm lifestyle was reported almost 80 years ago, the ubiquitous involvement of biofilms in chronic infection has been recognized more recently.^{5,6} Biofilm-related diseases are typically persistent infections that develop relatively slowly, are rarely resolved by the immune system and respond poorly or transiently to antibiotics.⁷ Some examples are chronic wound infection, chronic otitis media, osteomyelitis, recurrent urinary tract infection, endocarditis, and cystic fibrosis-associated infections which can accelerate lung function decay in cystic fibrosis patients.^{8,9} Biofilms can also colonize indwelling catheters, stents, orthopedic implants, endotracheal tubes and urinary catheters,^{10–12} and are ubiquitous in burn wounds and in chronic nonhealing wounds, including diabetic foot ulcers, pressure ulcers and venous leg ulcers.^{13,14} Bacteria in biofilms are thought to be phenotypically and physiologically different

from nonadhered, free living (planktonic) cells.¹⁵ Susceptibility tests have shown that the minimal inhibitory concentration (MIC) and minimal bactericidal concentration (MBC) of antibiotics that are required to challenge bacteria in mature biofilms can be 10- to 100-fold higher than those required for planktonic bacteria.^{16–18} This intrinsic tolerance to antibiotics, which is thought to be an important reason for the persistence of biofilm infections,^{6,10} generates an urgent need to discover novel antibiotics and to validate new targets for combatting the threat posed by multidrug resistant organisms and overcoming the limitations of conventional antibiotics to treat chronic (biofilm) infections.^{19–21}

Iron is an essential element in biology, and a required cofactor for many enzymes that participate in important physiological processes, such as respiration, DNA synthesis, and amino acid synthesis.²² Iron homeostasis in bacteria offers a significant vulnerability because invading pathogens must obtain essential iron from the host, but host-defenses maintain the concentration of free iron at a vanishingly low level ($\sim 10^{-18}$ M).²² Pathogens have evolved mechanisms to “steal”

Received: September 23, 2020

iron from their host, but these depend strongly on well-regulated iron homeostasis.^{23,24} Consequently, we have proposed iron storage proteins as a new and specific target for dysregulating bacterial iron homeostasis.^{23,25,26} Our work has showed that although two ferritin-like molecules coexist in *P. aeruginosa*, bacterioferritin B (BfrB) is the main iron storage protein.^{23,27} Bacterioferritin, which only exists in bacteria, is a spherical protein that can store up to ~3000 Fe³⁺ ions in its interior cavity (~80 Å diameter). Bacterioferritin is unique in that it binds 12 heme groups buried under the external protein surface, which allows the heme propionates to protrude into the interior cavity (Figure 1A).^{28,29} We demonstrated that

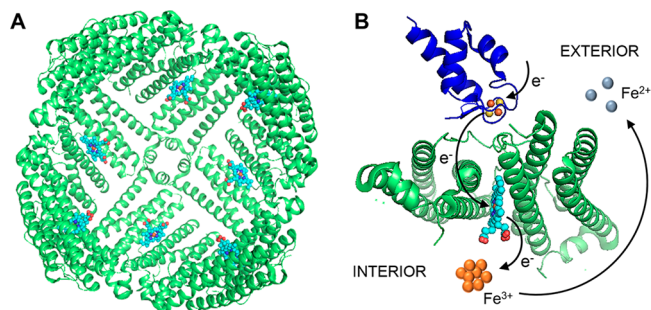


Figure 1. (A) BfrB from *Pseudomonas aeruginosa* (PDB ID 3is7) is a spherical and hollow protein assembly which can store up to 3000 Fe³⁺ in the interior cavity. A heme molecule (cyan) binds between two subunits, where it is buried under the exterior surface with the heme propionates exposed to the interior cavity. (B) Structure of the BfrB-Bfd complex (PDB ID 4E6K) where a Bfd molecule (blue) binds at the interface of two BfrB subunits (green), above a heme molecule (cyan). Electron transfer from the [2Fe-2S] cluster in Bfd to the Fe³⁺ in the interior of BfrB is mediated by the heme. The resultant Fe²⁺ exits the BfrB molecule for incorporation in bacterial metabolism.

mobilization of iron from BfrB requires specific interactions with the bacterioferritin-associated ferredoxin (Bfd).^{30,31} The structure of Bfd revealed a new class of [2Fe-2S] protein³² and the crystal structure of the BfrB-Bfd complex showed that up to 12 Bfd molecules can bind at identical sites on the BfrB surface, at the interface of subunit dimers, above a heme molecule (Figure 1B).³⁰ The heme mediates electron transfer between the [2Fe-2S] cluster of Bfd and the Fe³⁺ mineral stored in the BfrB cavity, enabling mobilization of Fe²⁺ to the cytosol.^{28,30} The Bfd binding sites on BfrB are equivalent and independent, where Bfd binds with a *K*_d of 3 μM.³³

Studies conducted with *P. aeruginosa* PAO1 and a mutant where the *bfd* gene has been deleted (Δbfd) showed that in the absence of Bfd iron is irreversibly accumulated in BfrB, thus demonstrating that Bfd is required to mobilize iron stored in bacterioferritin. These studies also showed that the irreversible accumulation of iron in BfrB is accompanied by depletion of free iron in the cytosol.²³ Given that *P. aeruginosa* requires sufficient environmental and intracellular iron reserves to establish mature biofilms,^{34–37} we reasoned that the iron deficiency that ensues in the cytosol of the Δbfd cells might adversely affect biofilm formation. Studies conducted to pursue this idea showed that the Δbfd cells form poorly developed biofilms and that the biofilm-embedded cells experience cytosolic iron deficiency, even in iron-sufficient culture media.²⁶ These findings, which underscored the inhibition of the BfrB-Bfd complex as a viable target to dysregulate iron homeostasis and possibly develop novel antimicrobial tools,

encouraged us to conduct a fragment-based structure-guided campaign to discover inhibitors of the BfrB-Bfd complex. These efforts allowed us to identify derivatives of 4-aminoisindoline-1,3-dione which bind BfrB selectively at the Bfd binding site,²⁵ block the BfrB-Bfd complex in the *P. aeruginosa* cytosol, and inhibit the mobilization of bacterioferritin-stored iron.²⁵ Challenging planktonic cultures of *P. aeruginosa* with the 4-aminoisindoline-1,3-dione analogues elicited a dose-dependent growth inhibition phenotype, thus providing proof of concept for the usefulness of small molecules designed to inhibit the BfrB-Bfd complex as probes to dysregulate iron homeostasis and weaken bacterial cells.²⁵ Herein we report on improvements made to the 4-aminoisindoline-1,3-dione derivatives which increase the binding affinity for BfrB and the bacteriostatic activity against planktonic *P. aeruginosa* cells. Surprisingly, the 4-aminoisindoline-1,3-dione derivatives are bactericidal to *P. aeruginosa* cells embedded in mature biofilms. The killing effect of the BfrB-Bfd complex inhibitors on biofilm cells, which are normally recalcitrant to several classes of commercial antibiotics, has uncovered a rare weakness that may be exploited to control biofilms.

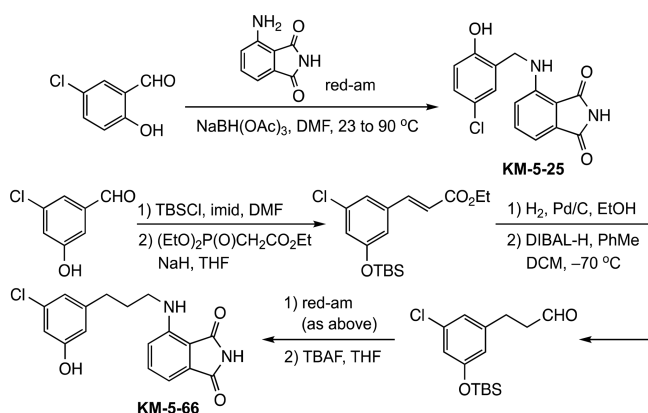
RESULTS AND DISCUSSION

Synthesis of 4-Aminoisindoline-1,3-dione Derivatives. Solving the X-ray structure of the BfrB-Bfd complex³⁰ and identifying the hot-spot residues participating at the protein–protein interface³³ provided a unique molecular platform to study the mechanisms that enable mobilization of bacterioferritin-stored iron.²⁸ The structural information also enabled a fragment based structure-guided strategy to discover inhibitors of the BfrB-Bfd complex. Fragment screening followed by efforts to cocrystallize the fragment hits with BfrB produced the X-ray crystal structure of BfrB in complex with 4-aminoisindoline-1,3-dione (**8**) binding at the Bfd-binding site.²⁵ The cocrystal structure informed a strategy for fragment growth that produced a series of 4-(benzylamino)- and 4-((3-phenylpropyl)amino)isindoline-1,3-dione analogues represented by **11** and **16** (Table 1), which are derivatives of **8** with $-(CH_2)-$ and $-(CH_2)_3-$ linkers, respectively.^{25,38} These analogues were shown to selectively bind BfrB at the Bfd binding site and to elicit a growth defect in *P. aeruginosa* planktonic cell cultures.²⁵ To increase the binding affinity of **11** and **16** for BfrB we looked for information in the available structural data. For example, the crystal structures of **11** and **16** bound to BfrB show clear electron density defining the aminoisindoline-1,3-dione (phthalimide) moieties. The hydroxy-substituted phenyl rings, however, are less well-defined by electron density, suggesting that conformational disorder of the phenyl ring in the cleft formed by L68 and E81 may influence the stability of the complex.²⁵ In an attempt to lower disorder of the phenyl ring, we prepared a set of 4-aminoisindoline-1,3-dione derivatives bearing a relatively bulky halogen atom in the phenyl ring (Table 1). A schematic summary of the representative syntheses for compounds with $-(CH_2)-$ or $-(CH_2)_3-$ linkers is shown in Figure 2. For example, compound **KM-5-25** was prepared from 5-chlorosalicylaldehyde by reductive amination with **8** in *N,N*-dimethylformamide (DMF) solution using sodium triacetoxyborohydride as the reducing agent. Compound **KM-5-66** was prepared from 3-chloro-5-hydroxybenzaldehyde; the phenol was protected with *tert*-butyldimethylsilyl chloride (TBSCl) and the side chain

Table 1. Structure, Binding Affinity, and IC₅₀ of 4-Aminoisoindoline-1,3-dione Derivatives

Analog	Structure	K _d (μM)	IC ₅₀ (μM)	Ref.
8		300 ± 50	not active	25
11		11 ± 1	258 ± 23	25
KM-5-29		6 ± 0.5	nd*	this work
JAG-5-7		7 ± 2	128 ± 26	this work
KM-5-25		4 ± 0.6	69 ± 7	this work
KM-5-30		6 ± 1	156 ± 30	this work
BN-XIV-53		21 ± 3	not active at 120 μM	this work
KM-5-50		11 ± 2	82 ± 16	this work
KM-5-28		1.4 ± 0.2	96 ± 1	this work
JAG-5-6		0.43 ± 0.07	54 ± 9	this work
KM-5-54		0.22 ± 0.04	nd*	this work
16		1.50 ± 0.25	121 ± 4	25
KM-5-57		2.5 ± 0.4	not active at 200 μM	this work
KM-5-66		0.35 ± 0.05	42 ± 6	this work

*Not determined because of low solubility (<30 μM) in PBS buffer.

**Figure 2.** Schematic summary of the synthetic process to prepare 4-(benzylamino)isoindoline (i.e., KM-5-25) and 4-((3-phenylpropyl)amino)-4-aminoisoindoline-1,3-dione (i.e., KM-5-66) derivatives of 4-aminoisoindoline-1,3-dione.

elongated by Horner–Wadsworth–Emmons reaction to the α , β -unsaturated ester. Subsequent catalytic reduction of the side chain double bond followed by diisobutylaluminum hydride (DIBAL-H) reduction of the ester under carefully controlled temperature conditions furnished the requisite 3-arylpropionaldehyde. This aldehyde was then linked by reductive amination to **8** as above and deprotected with tetrabutylam-

monium fluoride (TBAF) in THF. Additional details are presented in the [Supporting Information](#).

4-Aminoisoindoline-1,3-dione Derivatives Elicit a Bacteriostatic Effect in Planktonic *P. aeruginosa* Cultures. The relative strength of the association between the new analogues and BfrB was evaluated measuring the dissociation constant K_d (Table 1, Figure S1). The results show that installing a halogen atom in the phenyl ring improves the binding affinity of the derivatives relative to the previously reported analogues **11** and **16**. The K_d values of halogen containing compounds with a $-(CH_2)-$ linker are on average ~ 2 -fold lower when compared to the K_d exhibited by **11**, and the K_d values of halogen-bearing analogues with a $-(CH_2)_3-$ linker are on average ~ 5 -fold lower than the K_d measured for analogue **16**. The relative efficacy of the analogues to inhibit *P. aeruginosa* planktonic growth was evaluated by measuring the half maximal inhibitory concentration (IC₅₀). Inspection of the data in Table 1 shows that all of the halogenated compounds are more active than analogues **11** and **16**; the activity of KM-5-29 and KM-5-54 could not be evaluated because the relatively low aqueous solubility of these compounds prevented measurement of their IC₅₀ values.

Although the compounds synthesized so far do not include all possible substitution isomers, some insights of the governing structure activity relationships have begun to emerge (Table 1): (i) Among the compounds with a $-(CH_2)-$ linker, the data reveal that when the hydroxyl group is at position 2 relative to the linker (KM-5-29, JAG-5-7, KM-5-25, and KM-5-30) a bulkier Cl atom at position 5 (KM-5-25) imparts ~ 2 -fold higher binding affinity for BfrB than a smaller F atom at the same position (JAG-5-7). In line with the nearly 2-fold lower K_d , the IC₅₀ of KM-5-25 is ~ 2 -fold lower than that of JAG-5-7. In comparison, a Cl atom at position 3 (KM-5-30) imparts a K_d similar that of KM-5-25, but an IC₅₀ ~ 2.5 -fold larger, suggesting that KM-5-30 is less efficient at penetrating or accumulating in *P. aeruginosa* cells. Installing a Cl atom at position 6 (KM-5-29) renders the compound poorly soluble in aqueous solution. It is also interesting to compare the two analogues with a hydroxyl group at position 3. The presence of a Cl atom at position 5, KM-5-50, lowers the K_d by a factor of 2 relative to BN-XIV-53 and improves the activity vs planktonic cells significantly. (ii) Examining the compound series with a $-(CH_2)_3-$ linker shows that when the hydroxyl group is at position 3, a Cl atom at position 5 (KM-5-66) decreases the K_d ~ 4 -fold and the IC₅₀ ~ 3 -fold relative to compound **16**. In comparison, the presence of a second hydroxyl group at position 5 (KM-5-57) increases the K_d ~ 1.7 -fold relative to **16** and renders the compound inactive. Given that the K_d measured for KM-5-57 is similar or lower than K_d values measured for other active compounds in Table 1, the immeasurable activity of KM-5-57 suggests that it cannot penetrate or accumulate in *P. aeruginosa* cells. Finally, comparison of compounds where the hydroxyl group is at position 2 (KM-5-28, JAG-5-6, and KM-5-54) also shows that a halogen at position 5 improves binding affinity. A bulkier Cl atom at position 5 increases the binding affinity of KM-5-54 2-fold relative to the compound with a F at the same position (JAG-5-6). Comparing the IC₅₀ values corresponding to KM-5-28 and JAG-5-6 reveals that the ~ 3 -fold lower K_d caused by installing a F atom at position 5 is accompanied by ~ 2 -fold decrease in the IC₅₀. Attempts to determine whether the lower K_d obtained when a Cl atom at position 5 would bring an

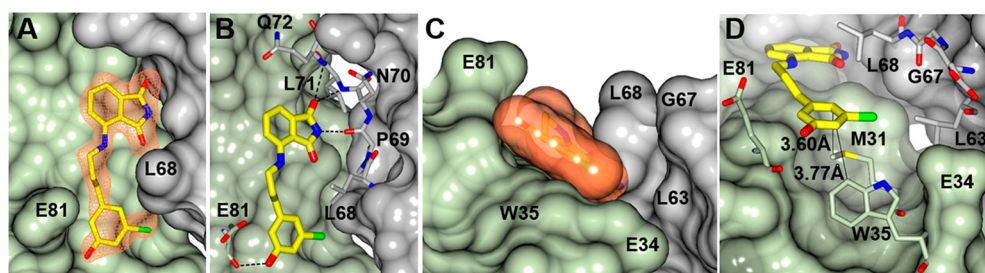


Figure 3. Binding of KM-5-66 to the Bfd binding site on BfrB. Subunits A and B of a BfrB subunit dimer are rendered as surface and colored in gray and green, respectively. (A) $F_o - F_c$ omit map contoured at 3σ (coral mesh) from KM-5-66 bound on BfrB. (B) Hydrogen bonding interactions (dashed lines) between KM-5-66 and BfrB. (C) View along the 3-hydroxyl-5-chloro phenyl ring plane of KM-5-66 (coral) depicted as a surface rendering along with residues surrounding the phenyl ring. (D) View depicting the 3-hydroxyl-5-chloro phenyl ring position relative to residues M31 and W35; pertinent distances are indicated by solid lines. The Cl atom of KM-5-66 is depicted in green.

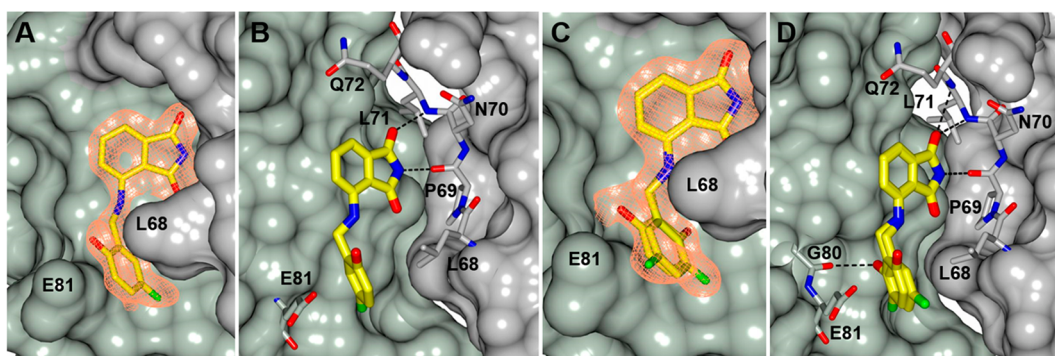


Figure 4. Binding of JAG-5-7 to the Bfd binding site on BfrB. (A,B) $F_o - F_c$ omit map contoured at 3σ (coral mesh) from JAG-5-7 bound at the Bfd binding site in one of the BfrB subunit dimers, and hydrogen bonding interactions (dashed lines). The 2-hydroxyl-5-fluoro phenyl ring is observed in only one conformation at this binding site. (C,D) View of a different binding site where the 2-hydroxyl-5-fluoro phenyl ring is observed in two conformations; in one of the conformations the hydroxyl group is hydrogen bonded to the carbonyl oxygen of G80. The F atom of JAG-5-7 is depicted in green.

additional decrease in the IC_{50} were stymied by the poor water solubility of KM-5-54.

The structure–activity relationships (SAR) information available thus far indicates that a halogen in the aryl ring of the 4-aminoisindoline-1,3-dione derivatives invariably improves binding affinity for BfrB. In compounds with a $-(CH_2)-$ linker, when the hydroxyl group is at position 2, a Cl atom at position 5 imparts favorable properties, a Cl atom at position 3 renders the compound poorly active vs *P. aeruginosa*, despite relatively favorable K_d , and a Cl atom at position 6 imparts poor aqueous solubility. The preparation of compounds with a $-(CH_2)_3-$ linker is more elaborate and accessibility to suitable starting materials at reasonable prices is also more limited. The information available thus far indicates that although a Cl at position 5 is favorable whether the hydroxyl group is at position 2 or 3, poor aqueous solubility impairs the potential biological activity when the hydroxyl is at position 2 (KM-5-54), despite having the most favorable K_d of all compounds in Table 1. Attempts to increase aqueous solubility by installing a hydroxyl group at position 5 succeeded in increasing solubility but rendered a compound (KM-5-57) with lower affinity for BfrB and inactive against *P. aeruginosa* cells. As new compounds become available, similar evaluation of binding affinity and activity against bacterial cells will continue to shed light on the structural requirements that simultaneously enhance target affinity in vitro and activity against *P. aeruginosa* cells. For the purposes of the studies described below, we chose to work with compounds KM-5-25, the most active of the analogues

containing a $-(CH_2)-$ linker, and KM-5-66, the most active of the analogues harboring a $-(CH_2)_3-$ linker.

Structures of Representative 4-Aminoisindoline-1,3-dione Derivatives Bound to BfrB. To gain additional evidence for the selectivity of the 4-aminoisindoline-1,3-dione derivatives for the Bfd binding site on BfrB, we carried out ligand soaking experiments aimed at obtaining cocrystals of compound bound to the bacterioferritin. These experiments resulted in the X-ray crystal structures of KM-5-28, KM-5-66, KM-5-50, and JAG-5-7 bound to BfrB; refinement statistics are summarized in Table S1. In all four structures the 4-aminoisindoline-1,3-dione (phthalimide) moiety binds at the Bfd binding site with a nearly identical pose, forming hydrogen bonding interactions to the O atom of P69 and the backbone N atoms of L71 and N70 (Figures 3A,B, 4A,B, and S2). This binding mode, which is nearly identical with that observed in the previously reported structures of 11, 16, and several other analogues,²⁵ provides strong evidence for the selectivity of the 4-aminoisindoline-1,3-dione derivatives for the Bfd site on BfrB. The linkers and phenyl rings also bind BfrB at the Bfd binding site, where they fit in a cleft formed by the side chains of L68 and E81. As will be discussed below, the details of binding are particular to whether the compound harbors a $-(CH_2)-$ or a $-(CH_2)_3-$ linker and to the substitution pattern on the ring.

The structure of KM-5-66 (Figure 3) shows the $-(CH_2)_3-$ linker wedged at the “floor” of the cleft formed by the side chains of L68 and E81 on BfrB (Figure 3A,B), where it engages in hydrophobic packing with the several residues forming the

cleft floor and with the hydrophobic portion of the L68 and E81 side chains. The 3-hydroxyl-5-chloro phenyl ring, which is nearly parallel with the cleft floor, enables the 3-hydroxyl group to interact with the E81 carbonyl oxygen via H-bonding, and the Cl atom to interact with the side chains of L63 and L68 (Figure 3A,B). The phenyl ring, at its closest point, is also 3.60 Å from the M31 C_E atom, a favorable interaction that is also probable to hinder rotation of the ring toward a perpendicular position relative to the cleft floor (Figure 3C,D). It is also interesting to note that the 3-hydroxyl-5-chloro phenyl ring is positioned to form a pseudo edge-to-face interaction with the indole ring of W35; the phenyl ring, at its closest position, is 3.77 Å from the indole ring C_{Z2} atom, the distance between the centroids of the phenyl and the indole 6-member ring is 5.92 Å, and the angle between the planes of these rings is 67.4°, suggesting a weak edge-to-face interaction.

Despite significant effort, we could not obtain good diffraction data from BfrB crystals soaked in solutions containing **KM-5-25**. However, we were able to obtain the structure of the fluorinated analogue **JAG-5-7** (Figure 4). Strong electron density consistent with **JAG-5-7** was observed in 8 of the 12 subunits. The shorter $-(\text{CH}_2)-$ linker places the 2-hydroxyl-5-fluoro phenyl ring well within the cleft formed by the side chains of L68 and E81, with the ring nearly perpendicular to the cleft floor. The electron density from the 2-hydroxyl-5-fluoro phenyl ring is consistent with the ring experiencing one orientation in certain subunits (Figure 4A,B) and with the ring in two conformations in other subunits (Figure 4C,D). In one of these orientations the hydroxyl group forms a H-bond with the carbonyl oxygen of G80, whereas the fluorine atom packs against the side chain of L68. The structure of **KM-5-50** (Figure S2), also places the 3-hydroxyl-5-chloro phenyl ring within the cleft formed by L68 and E81, in a similar nearly perpendicular conformation relative to the cleft floor, except that no H-bonding was observed for the 3-hydroxyl group.

Taken together, the structures of the 4-aminoisoinndoline-1,3-dione derivatives reported here and those reported previously²⁵ provide important insights. The linkers and the phenyl rings of all the compounds containing a $-(\text{CH}_2)_3-$ linker adopt nearly identical binding modes, with the phenyl rings oriented parallel to the cleft floor (Figure S3) and positioned to engage the 6 member ring of the indole in W35. Superposing the structures of analogue **16** and **KM-5-66** (Figure S4A) shows that the phenyl ring in **16** is notably pitched relative to the phenyl ring in **KM-5-66** (the angle between the mean planes of both rings is 22.6°), and the 3-hydroxyl group is not engaged in H-bonding interactions. In comparison, the Cl atom at position 5 in **KM-5-66** appears to induce a nearly parallel orientation of the ring relative to the cleft floor and a conformation that places the 3-hydroxyl within H-bonding distance of the carbonyl oxygen of E81. Together, the H-bonding engagement of the 3-hydroxyl group, the packing of the Cl atom with the side chains of L63 and L68, and the more extensive packing of the phenyl ring against the cleft floor residues are probably responsible for the higher affinity of **KM-5-66** for BfrB relative to **16**. It is also interesting to note that in the structures of all the compounds with a shorter $-(\text{CH}_2)-$ linker the phenyl ring is nearly perpendicular to the cleft floor. This is illustrated by superposing the structures of **KM-5-66** and **JAG-5-7** (Figure S4B), which shows that the phenyl rings of both compounds adopt a nearly perpendicular angle (74°) relative to one another. The

structural information currently available suggests that the shorter linker and the relatively less efficient packing of the phenyl ring against the hydrophobic portions of the L68 and E81 side chains may contribute to the higher K_d values of these compounds relative to those with a $-(\text{CH}_2)_3-$ linker.

Planktonic *P. aeruginosa* Cells Treated with 4-Aminoisoinndoline-1,3-dione Derivatives Overproduce Pyoverdine. Previous studies directed at evaluating the repercussions of blocking the BfrB-Bfd complex in *P. aeruginosa* cells relied on deleting the *bfd* gene (Δbfd). These investigations showed that blockade of the BfrB-Bfd complex in planktonic Δbfd cells causes an irreversible accumulation of iron in BfrB and iron deficiency in the cytosol. The resultant phenotype is hyperproduction of pyoverdine relative to the wild type cells.²³ Pyoverdine is a siderophore produced by *P. aeruginosa* when the cells experience iron limitation.³⁹ A similar pyoverdine overproduction phenotype was observed when wild type *P. aeruginosa* cells were treated with small molecule inhibitors of the BfrB-Bfd complex (**11** and **16**).²⁵ Therefore, to determine that compounds **KM-5-25** and **KM-5-66** inhibit iron mobilization from BfrB in the *P. aeruginosa* cytosol, we investigated whether cells treated with these compounds express the characteristic pyoverdine hyperproduction phenotype. To this end, planktonic cells were cultured in the presence of **KM-5-25** (70 μM) or **KM-5-66** (50 μM) for 27 h in M63 media and the content of the secreted pyoverdine in the cell-free spent media was analyzed by measuring the fluorescence intensity at 460 nm. Normalizing the intensity of pyoverdine fluorescence to CFU/mL shows that as expected, cells treated with **KM-5-25** or **KM-5-66** secrete ~5-fold more pyoverdine than the untreated control (Figure S5), an overproduction level similar to that observed with the Δbfd mutant.²³ These observations indicate that both analogues bind BfrB in the *P. aeruginosa* cytosol, block the BfrB-Bfd interaction and inhibit iron mobilization from BfrB, resulting in cytosolic iron limitation that is manifested in a pyoverdine hyperproduction phenotype. The cytosolic iron limitation caused by treating planktonic cultures with **KM-5-25** or **KM-5-66** exerts a bacteriostatic effect on the cells, as indicated by the IC₅₀ values in Table 1. In stark contrast, when the same compounds are used to treat *P. aeruginosa* biofilms, a bactericidal effect is observed. The results from these experiments are discussed below.

4-Aminoisoinndoline-1,3-dione Derivatives Kill *P. aeruginosa* Cells in Mature Biofilms. A characteristic of biofilms is their high tolerance to antimicrobial agents. Tolerance is a physiological condition which does not involve mutation and enables bacteria to survive in the presence of antibiotics.^{40–43} The persistent biofilm phenotype is thought to arise from several factors, including restricted penetration of antibiotic molecules due to interactions with components of the biofilm matrix, slow cell metabolism in the biofilm, differential expression of specific genes, and the presence of persister cells. In addition, biofilms are composed of distinct subpopulations that exhibit different physiological activity; cells in the biofilm interior exhibit low metabolic activity, distinct from the high metabolism of cells near the surface.^{40,44,45} The dissimilar metabolic activity is thought to result from a concentration gradient of O₂ and nutrients, which are high at the biofilm surface and low in the deeper layers of the biofilm.^{45,46} Commercial antibiotics that interfere with cell replication (e.g., ciprofloxacin), or protein translation (e.g., tobramycin), preferentially kill the metabolically active bacteria

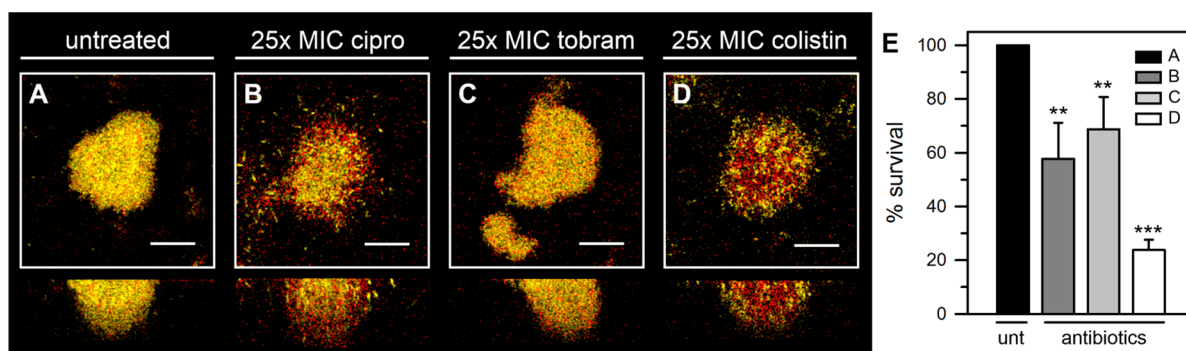


Figure 5. Biofilms cultured in flow cells are tolerant to ciprofloxacin and tobramycin and susceptible to colistin. EYFP-expressing *P. aeruginosa* PAO1 biofilms were cultured for 3 days by flowing AB media supplemented with 15 μ M Fe and then treated for 24 h by flowing the same media containing antibiotic. Biofilms were counterstained with Sytox Red and imaged with the aid of CLSM. Top-down views (x - y plane) are depicted with side views (x - z plane) at the bottom. Viable cell mass is in yellow and dead cells and extracellular DNA in red. (A) Untreated (DMSO) control, (B) 25 \times MIC ciprofloxacin (19 μ M), (C) 25 \times MIC tobramycin (27 μ M), (D) 25 \times MIC colistin (20 μ M). (E) The % survival obtained from viable biomass calculated with the aid of COMSTAT software. The scale of the bars represents 20 μ m. $p < 0.01$ denoted by ** and $p < 0.001$ by *** relative to untreated.

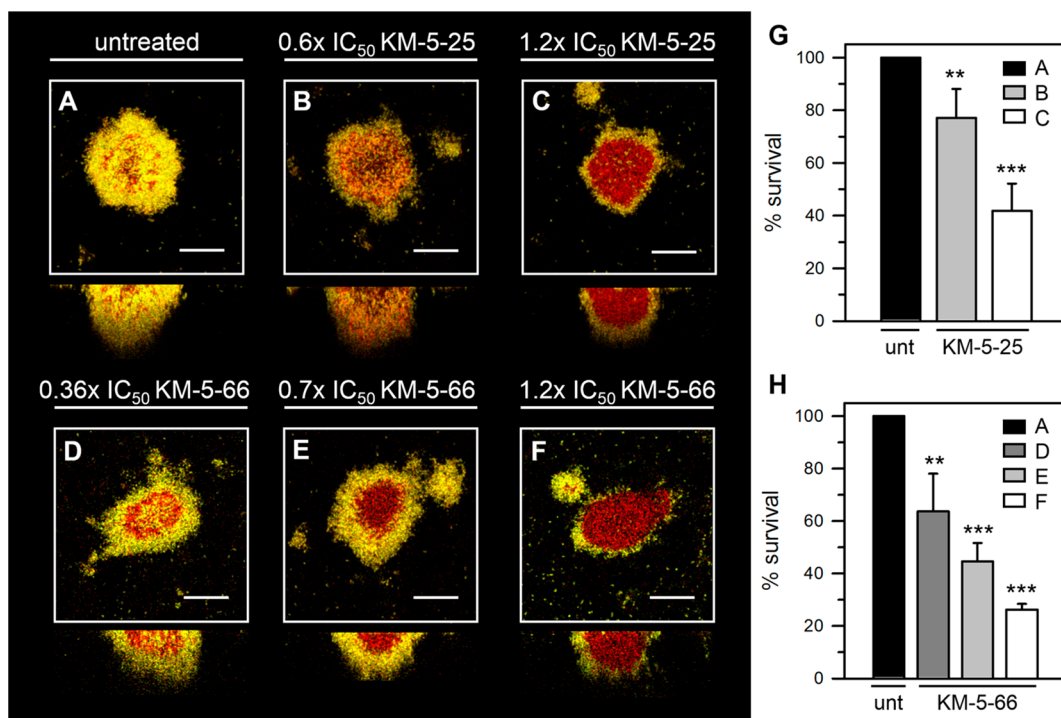


Figure 6. *P. aeruginosa* cells embedded in mature biofilms grown in flow cells are susceptible to 4-aminoisoindoline-1,3-dione analogues. EYFP-expressing *P. aeruginosa* PAO1 biofilms were cultured for 3 days by flowing AB media supplemented with 15 μ M Fe and then treated for 24 h with 4-aminoisoindoline-1,3-dione analogue. Biofilms were counterstained with Sytox Red and imaged with the aid of CLSM. Top-down views (x - y plane) are depicted with side views (x - z plane) at the bottom. Viable cell mass is in yellow and dead cells and extracellular DNA in red. (A) Untreated (DMSO) control, (B) 0.6 \times IC₅₀ KM-5-25 (40 μ M), (C) 1.2 \times IC₅₀ KM-5-25 (80 μ M), (D) 0.36 \times IC₅₀ KM-5-66 (15 μ M), (E) 0.7 \times IC₅₀ KM-5-66 (30 μ M), and (F) 1.2 \times IC₅₀ KM-5-66 (50 μ M). (G,H) The % survival obtained from viable biomass calculated with the aid of COMSTAT software for cells treated with KM-5-25 and KM-5-66, respectively. The scale of the bars represents 20 μ m. $p < 0.01$ denoted by ** and $p < 0.001$ by *** relative to untreated.

in the outer biofilm layers, whereas cells in the biofilm interior survive,^{44,47–49} despite the ability of both antibiotics to diffuse into the inner regions of the biofilm.^{44,50} In contrast, some antimicrobials that affect membrane structure, such as colistin, a “last-line” therapy to treat multidrug resistant infections,^{51–53} can kill cells in the deeper biofilm layers.⁴⁹

We tested the susceptibility of mature biofilms to treatment with analogues of 4-aminoisoindoline-1,3-dione using two platforms, biofilms cultured at the solid–liquid interface (flow

cell biofilms) and biofilms cultured at the air–liquid interface (pellicles). Biofilms of *P. aeruginosa* cells expressing an enhanced yellow fluorescent protein (EYFP) were cultured in flow cells using AB minimal media supplemented with 15 μ M Fe. 3-Day old biofilms were treated for 24 h with commercial antibiotics or with 4-aminoisoindoline-1,3-dione analogues by flowing AB media containing analogue or commercial antibiotic, 0.025% hypromellose (HPMC), 1.5% DMSO and 15 μ M Fe. In most experiments the concentration

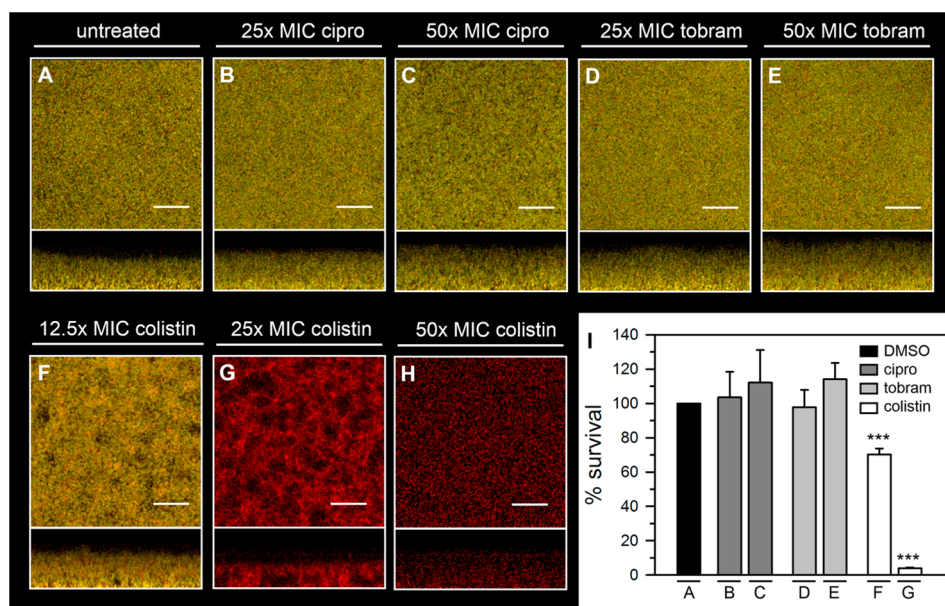


Figure 7. Pellicle biofilms are tolerant to ciprofloxacin and tobramycin, and susceptible to colistin. Pellicles of *P. aeruginosa* PAO1 expressing EYFP were cultured in PI media supplemented with 20 μM Fe for 48 h, and then treated with antibiotics for 24 h. Pellicles were counterstained with Sytox Red and imaged with the aid of CLSM. Images depict top-down views (squares) and side views (rectangles) where viable cells are shown in yellow and dead cells and extracellular DNA in red. (A) Untreated (DMSO) control, (B) 25 \times MIC ciprofloxacin (19 μM), (C) 50 \times MIC ciprofloxacin (38 μM), (D) 25 \times MIC tobramycin (27 μM), (E) 50 \times MIC tobramycin (54 μM), (F) 12.5 \times MIC colistin (10 μM), (G) 25 \times MIC colistin (20 μM), (H) 50 \times MIC colistin (40 μM). (I) The % survival obtained from viable biomass calculated with the aid of COMSTAT software. The scale of the bars represents 20 μm . $p < 0.001$ denoted by *** relative to untreated.

of commercial antibiotics used was 25 \times or 50 \times MIC, and the concentration of 4-aminoisoindoline-1,3-dione analogues was between 0.36 \times and 1.2 \times the IC_{50} . The treated biofilms were counterstained with the cell impermeable fluorescent nucleic acid dye Sytox Red and then imaged with the aid of confocal laser scanning microscopy (CLSM). Figure 5A depicts a representative image of the untreated control, illustrating yellow-fluorescent viable cells and red-stained dead cells and extracellular DNA. Figures 5B and 5C show representative images depicting 4-day old biofilms tolerant to 24 h treatment with ciprofloxacin or tobramycin, respectively, and Figure 5D shows 4-day old biofilms susceptible to 24 h treatment with colistin. In agreement with previously reported observations,^{49,54} treatment with ciprofloxacin preferentially kills cells at the biofilm surface, leaving the interior of the biofilm almost unaffected (Figure 5B). In contrast, treatment with colistin preferentially kills bacteria in the biofilm interior, leaving the biofilm surface less affected (Figure 5D). COMSTAT software was used to attempt a quantitative comparison of the biofilm biomass by estimating the biovolume, which is calculated as the overall volume/substratum area ($\mu\text{m}^3/\mu\text{m}^2$).⁵⁵ Comparing the biofilm biomass as the ratio of (untreated biomass)/(treated biomass) expressed as % survival (Figure 5E) shows that ~20% of biomass remains viable (yellow fluorescent) after treatment with colistin at 25 \times MIC. In comparison, ~60% and 70% of biomass remains viable after treatment with 25 \times MIC ciprofloxacin or tobramycin, respectively.

Having established that the biofilms are susceptible to colistin and significantly tolerant to ciprofloxacin and tobramycin, we treated similarly cultured 3-day old biofilms with KM-5-25 and KM-5-66 for 24 h. Compounds KM-5-25 and KM-5-66 are soluble in aqueous media to ~110 and ~80 μM , respectively. To ensure that the compounds remain

soluble during the 24 h treatment period, KM-5-25 was used at concentrations 40 and 80 μM , equivalent to 0.6 \times and 1.2 \times the IC_{50} , and KM-5-66 was used at concentrations 15, 30, and 50 μM , equivalent to 0.36 \times , 0.7 \times , and 1.2 \times the IC_{50} (Table 1). Representative CLSM images obtained after treating 3-day old biofilms with each of the analogues for 24 h show that both compounds kill biofilm cells in a concentration dependent manner (Figure 6). Treatment with 50 μM KM-5-66 elicits a similar level of killing as treatment with 80 μM KM-5-25. The higher efficacy exhibited by KM-5-66 agrees with its higher binding affinity for BfrB and lower IC_{50} . Inspection of the images obtained upon treatment with the higher concentrations of KM-5-25 or KM-5-66 clearly shows that the inhibitors of the BfrB-Bfd complex kill the cells in the interior of the biofilm, leaving most of the viable cells located at the biofilm surface (Figure 6C,F). This pattern of killing is reminiscent of previously reported observations showing that treatment of biofilms with Ga^{3+} preferentially kills cells in the inner portion of the biofilm.⁵⁶ The same authors concluded that cells in the biofilm interior are more sensitive to Ga^{3+} because this population experiences a more pronounced iron starvation. Therefore, we speculate that in biofilms treated with KM-5-25 or KM-5-66 the internal biofilm population is more susceptible to iron limitation caused by the nearly irreversible accumulation of iron in BfrB. It is important to underscore that although the mechanisms whereby iron starvation contribute to cell death in the biofilm interior are not yet understood, the fact remains that perturbation of iron homeostasis, either by systemic replacement of Fe^{3+} with Ga^{3+} , or by selective inhibition of the BfrB-Bfd complex, leads to bacterial cell death.

To expand the observations made with flow cell biofilms into a second biofilm model we also studied the susceptibility of biofilms grown at the air–liquid interface (pellicles).^{57,58}

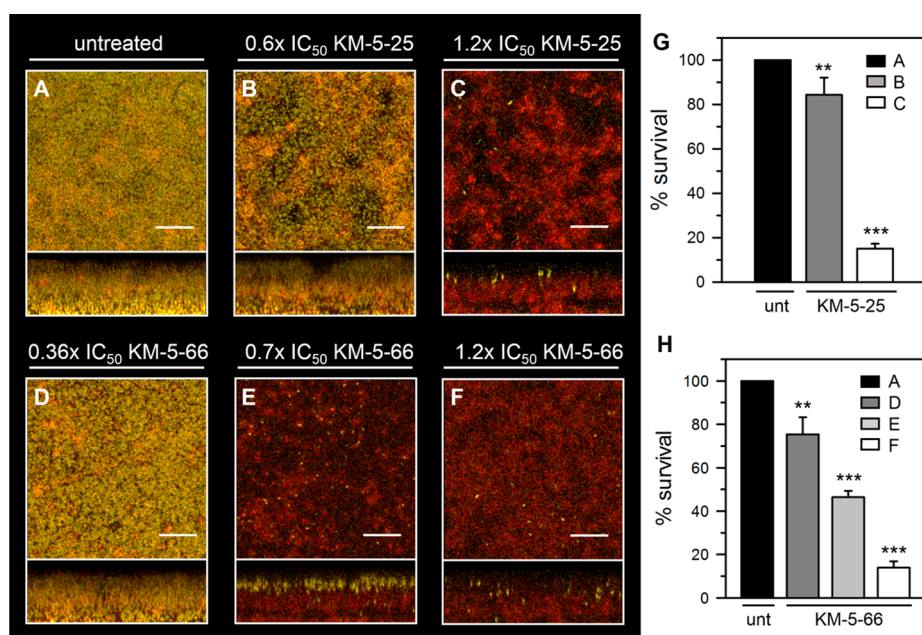


Figure 8. 4-Aminoisoindoline-1,3-dione analogues kill *P. aeruginosa* cells embedded in pellicle biofilms. Pellicles of *P. aeruginosa* PAO1 cells expressing EYFP were cultured for 48 h in PI media supplemented with 20 μ M Fe, and then treated with KM-5-25 or KM-5-66 for 24 h. Pellicles were counterstained with Sytox Red and imaged with the aid of CLSM. Images depict top-down views (squares) and side views (rectangles) where viable cells are shown in yellow and dead cells and extracellular DNA in red. (A) Untreated (DMSO) control, (B) 0.6 \times IC₅₀ KM-5-25 (40 μ M), (C) 1.2 \times IC₅₀ KM-5-25 (80 μ M), (D) 0.36 \times IC₅₀ KM-5-66 (15 μ M), (E) 0.7 \times IC₅₀ KM-5-66 (30 μ M), and (F) 1.2 \times IC₅₀ KM-5-66 (50 μ M). (G,H) The % survival obtained from viable biomass calculated with the aid of COMSTAT software for pellicles treated with KM-5-25 and KM-5-66, respectively. The scale of the bars represents 20 μ m. $p < 0.01$ denoted by ** and $p < 0.001$ by *** relative to untreated.

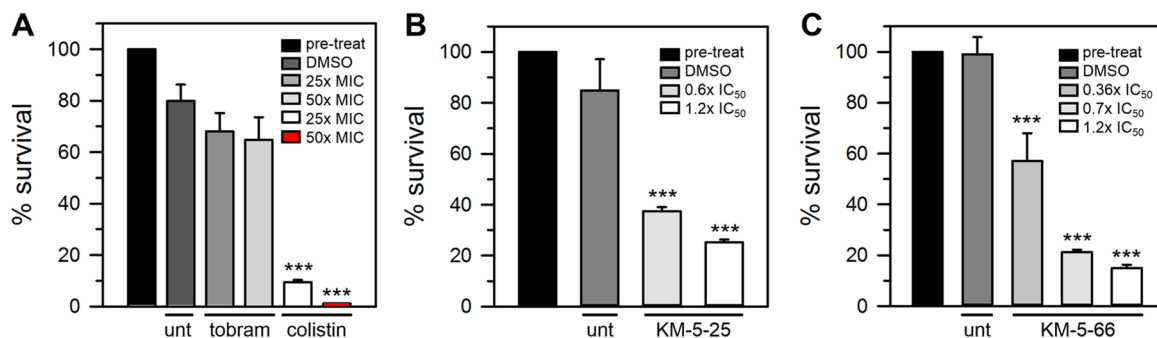


Figure 9. Assessment of cell survival in pellicle biofilms by dispersing and counting viable cells. EYFP-expressing *P. aeruginosa* PAO1 cells embedded in two-day old pellicles treated for 24 h with antibiotic or 4-aminoisoindoline-1,3-dione derivatives were dispersed for enumeration of viable culturable cells (CFU/mL). The % survival is expressed as the ratio CFU/mL_(after treatment)/CFU/mL_(pretreatment). Pellicles were treated for 24 h with (A) tobramycin (27 or 54 μ M) or colistin (20 or 40 μ M), concentrations equivalent to 25 \times and 50 \times the corresponding MIC, (B) compound KM-5-25 (40 and 80 μ M) concentrations equivalent to 0.6 \times and 1.2 \times IC₅₀, and (C) compound KM-5-66 (15, 30, and 50 μ M), concentrations equivalent to 0.36 \times , 0.7 \times , and 1.2 \times IC₅₀. $p < 0.001$ denoted by *** relative to untreated.

Pellicle biofilms (henceforth pellicles) are an attractive alternative platform to study biofilms because pellicles are amenable to imaging by CLSM and to harvesting, which can be desirable for additional biofilm analysis.²⁶ To determine the susceptibility of pellicles to antibiotics or inhibitors of the BfrB-Bfd complex, we cultured 2-day old pellicles of EYFP-expressing *P. aeruginosa* cells in PI media supplemented with 20 μ M Fe, as indicated in the Methods. The pellicles were transferred onto glass coverslips by allowing the surface of the coverslip to contact a pellicle. The coverslip-adhered pellicles were subsequently exposed to treatment solution (AB media supplemented with 15 μ M Fe, 0.025% HPMC, 1.5% DMSO, and antibiotic or analogue) for 24 h prior to staining with Sytox Red and imaging with the aid of CLSM. The pellicle biofilms are tolerant to ciprofloxacin and tobramycin at

concentrations 25 \times and 50 \times the MIC (Figure 7B–E), as is evident by the yellow fluorescence and near complete absence of red-stained dead cells. In contrast, the pellicle biofilms are susceptible to colistin at concentrations above 10 \times MIC (Figure 7F–H). Analysis of the images with COMSTAT, which allowed a more quantitative comparison of cell survival upon treatment with each of the antibiotics (Figure 7I), confirms tolerance to ciprofloxacin and tobramycin, but sensitivity to colistin. Note that when the concentration of colistin is 50 \times MIC the fluorescence signal from viable cells expressing EYFP is undetectable. When pellicle biofilms are challenged with compound KM-5-25 or KM-5-66 bacterial cell death occurs in a concentration dependent manner (Figure 8). These results agree with the idea that the 4-aminoisoindoline-1,3-derivatives penetrate the bacterial cell and bind to their

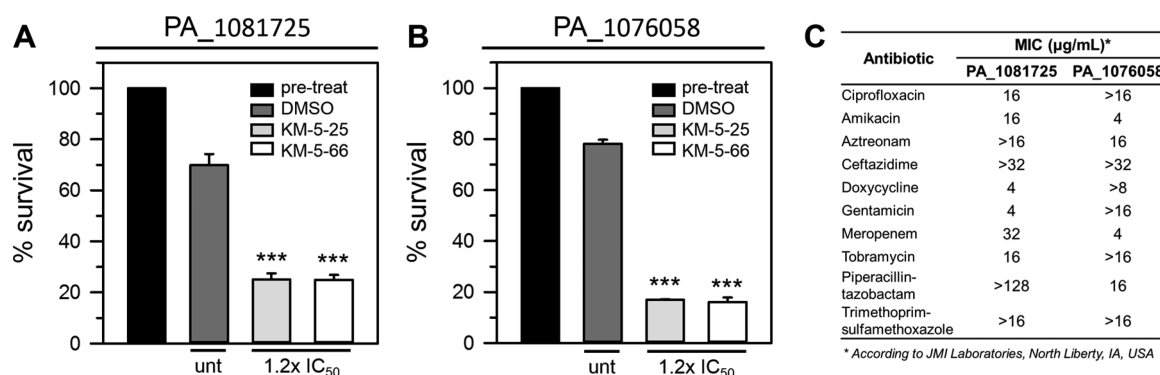


Figure 10. 4-Aminoisoindoline-1,3-dione analogues are likely broadly active against *P. aeruginosa* strains. Two-day old pellicle biofilms formed by *P. aeruginosa* clinical isolates (PA_1081725 and PA_1076058) were challenged for 24 h with 4-aminoisoindoline-1,3-dione derivatives prior to dispersing the cells for enumeration of viable culturable cells (CFU/mL). The % survival is expressed as the ratio of CFU/mL_(after treatment)/CFU/mL_(pretreatment). Pellicles of (A) PA_1081725 or (B) PA_1076058 were treated with concentrations equivalent to 1.2× IC₅₀ of KM-5-25 (80 μM), or KM-5-66 (50 μM). (C) Antibiotic susceptibility of clinical isolates PA_1081725 and PA_1076058. *p* < 0.001 denoted by *** relative to untreated.

target in the *P. aeruginosa* cytosol. Inspection of the images and analysis with COMSTAT (Figure 8G and H) shows that compound KM-5-66 is more efficacious than KM-5-25, observations that are consistent with the lower *K_d* and IC₅₀ values measured for KM-5-66. It is also important to note that when compounds KM-5-25 and KM-5-66 are used at a concentration of 80 μM and 50 μM, respectively (Figure 8G,H), which correspond approximately to 1.2× IC₅₀, nearly 85% of the cells in the pellicle are killed. This efficacy is similar that observed with colistin when used at 20 μM, equivalent to 25× MIC (Figure 7I).

To assess the efficacy of antibiotics and compounds with an approach complementary to imaging with CLSM, we resorted to dispersing biofilm cells for subsequent enumeration of viable cells (CFU/mL). To this end, we cultured pellicle biofilms for 48 h in PI media containing 20 μM Fe and exposed them to AB media containing 15 μM Fe and antibiotic or compound for 24 h, as described above. The biofilms were then harvested, and the cells dispersed into sterile PBS by vortexing in the presence of zirconia beads, prior to plating the cell suspensions for subsequent enumeration of CFU/mL. The results from these experiments are summarized in the plots of Figure 9 which show the % cell survival of pellicle-embedded cells after challenges with antibiotic or compound, calculated from the fraction CFU/mL_(after treatment)/CFU/mL_(pretreatment). When colistin is used to treat the pellicles at concentrations equivalent to 25× and 50× MIC, the treated biofilms exhibit ~10% and ~1% cell survival, respectively, relative to the pretreated biofilm (Figure 9A), corroborating the sensitivity of the pellicles to colistin. In contrast, challenging the pellicles with tobramycin (25× and 50× MIC) results in ~70% survival relative to the pretreated biofilm, and nearly identical cell survival relative to the untreated (DMSO control) pellicles (Figure 9A). These observations, which are in good agreement with those made with the aid of CLSM (Figure 7D–H) corroborate that the pellicles are tolerant to tobramycin and sensitive to colistin. Interestingly, attempts to enumerate cells after challenging the pellicles with ciprofloxacin (25× or 50× MIC) resulted in extremely low CFU/mL, findings which at first glance appear to be in conflict with the tolerance of the biofilms to ciprofloxacin observed in the CLSM images (Figure 7B,C). To reconcile these seemingly discrepant observations it is important to consider that several stressors, including ciprofloxacin, can induce a dormancy state in bacterial cells

known as the viable but not culturable (VBNC) state. A characteristic of cells in the VBNC state is their inability to develop into colonies on routine culture media, even though the cells remain viable for long periods of time.^{59,60} Evidence that bacterial cells can enter the VBNC has been obtained by several distinct methods,⁶⁰ one of which is the utilization of bacteria engineered to constitutively express bioluminescent proteins, and using the bioluminescence as a reporter of metabolic activity. Studies conducted with *P. aeruginosa* showed that following treatment with ciprofloxacin the bioluminescence emitted by *P. aeruginosa* cells decreased significantly less than the viable cell counts (CFU/mL). The perceived reduction in viable cell counts, which did not correlate with the relatively high metabolic activity reported by the small decrease in bioluminescence, indicated that challenges with ciprofloxacin induce *P. aeruginosa* cells to enter a VBNC state.^{61,62} Our observations suggest a similar situation. Imaging the pellicles with CLSM following the 24 h challenge with ciprofloxacin (Figure 7B,C) shows that most of the cells are metabolically active (yellow fluorescent), but dispersion of the cells from the pellicles for enumeration of CFU/mL shows a large reduction in culturable cells relative to the untreated control. These observations strongly suggest that treating the pellicles with ciprofloxacin induces the cells to enter the VBNC state, thus rendering them tolerant to the antibiotic. These findings, which highlight the complexities associated with biofilm embedded cells, also underscore the importance of resorting to more than one platform to study the efficacy of antibiofilm agents.

Enumeration of CFU/mL was also carried out after challenging pellicles with 4-aminoisoindoline-1,3-dione derivatives. Treating the pellicles with KM-5-25 at concentrations equivalent to 0.6× and 1.2× IC₅₀ results in ~38% and ~25% survival relative to cells in the pretreated biofilm (Figure 9B), while treating with KM-5-66 at concentrations equivalent to 0.3×, 0.6×, or 1.2× IC₅₀ results in approximately 57%, 21%, and 15% survival relative to cells in the pellicles prior to treatment (Figure 9C). These observations, which are in good agreement with the efficacy of the compounds evaluated by COMSTAT analysis of the CLSM images, corroborate the bactericidal activity of the compounds against *P. aeruginosa* biofilms, and provide additional evidence indicating that KM-5-66 used at 50 μM (1.2× IC₅₀) exhibits nearly the same efficacy as colistin used at 20 μM (25× MIC). We also used

the strategy of dispersing and counting viable cells to compare the relative efficacy of analogues **11**, **16**, **KM-5-25**, and **KM-5-66**. To this end, pellicles formed by *P. aeruginosa* PAO1 cells were treated (24 h) with each of the compounds at a concentration of 50 μM . The results (Figure S6) show that the compound activity against biofilm (**KM-5-66** > **KM-5-25** > **16** > **11**) track with the K_d and IC_{50} values (Table 1). Taken together, these findings suggest that additional modifications made to the structures of **KM-5-25** or **KM-5-66** which improve the binding affinity for BfrB have the potential to produce molecules with similar or better efficacy toward biofilms than colistin.

The results from experiments aimed at determining the efficacy of the 4-aminoisindoline-1,3-dione derivatives presented so far have been conducted with the reference strain *P. aeruginosa* PAO1. To investigate whether the compounds are also active against other strains of *P. aeruginosa*, we cultured pellicles of several clinical isolates from JMI Laboratories. Isolates PA_1081725 and PA_1076058 were chosen for additional testing because these strains exhibit relatively high MIC values for several antibiotics (Figure 10C) and form robust pellicles under the same culture conditions used to grow pellicle biofilms of *P. aeruginosa* PAO1. Challenging the pellicles formed by PA_1081725 and PA_1076058 with analogues **KM-5-25** or **KM-5-66** at concentrations equivalent to $1.2 \times \text{IC}_{50}$ elicits approximately 80% reduction of viable cells (Figure 10A,B), indicating that the activity of the 4-aminoisindoline-1,3-dione analogues is not unique to biofilms formed by the *P. aeruginosa* PAO1 strain. To gain a broader understanding of the potential activity spectrum of the BfrB-Bfd inhibitors against *P. aeruginosa* strains, BLASTp⁶³ was used to find homologues of BfrB and Bfd sequences in the >4400 *P. aeruginosa* genomes in the Pseudomonas Genome Database.⁶⁴ The results reveal two important facts: (i) The *bfrB* and *bfd* genes are adjacent to one another in all the *P. aeruginosa* strains. (ii) There is an extremely high level of conservation among the *bfd* and *bfrB* sequences. These findings allow us to predict that the structures of the BfrB and Bfd proteins, as well as the structures of the BfrB-Bfd complexes, are highly conserved across *P. aeruginosa* strains, thus suggesting that the 4-aminoisindoline-1,3-dione derivatives can be expected to be broadly active against *P. aeruginosa*.

4-Aminoisindoline-1,3-dione Derivatives Inhibit Iron Mobilization from BfrB in *P. aeruginosa* Cells. The results presented above demonstrate that the 4-aminoisindoline-1,3-dione derivatives can kill cells in mature biofilms. To demonstrate that the bactericidal activity is likely a result of the compounds engaging BfrB in the *P. aeruginosa* cytosol, inhibiting the BfrB-Bfd complex and blocking iron mobilization from the bacterioferritin in the *P. aeruginosa* cytosol, we carried out experiments aimed at visualizing the iron stored in BfrB. These experiments capitalize on a strategy we reported previously to demonstrate that the Δbfd mutant of *P. aeruginosa* irreversibly accumulates iron in BfrB²³ and to show that analogue **16** inhibits iron mobilization from BfrB in planktonic cells.²⁵ To visualize BfrB-stored iron in biofilm-embedded cells, 2-day old pellicles of *P. aeruginosa* PAO1 cells were treated for 24 h with analogue **KM-5-25** (40 μM) or **KM-5-66** (20 μM) at a concentration predicted to kill approximately 50% of the cells in the biofilm. The treated pellicles were dispersed in sterile PBS and the cell suspension was harvested by centrifugation after a small aliquot had been sampled to enumerate viable cells. To visualize iron stored in BfrB the

harvested cells were lysed, the lysate solution supernatant was clarified by centrifugation and then loaded onto native PAGE gels for separation and subsequent staining with Ferene S, which reacts with iron to develop a blue color. Since the viable cell count dispersed from the pellicle biofilms treated with **KM-5-25** or **KM-5-66** was 42% and 37% of the cells in the untreated pellicle (Figure 11A), the clarified lysate super-

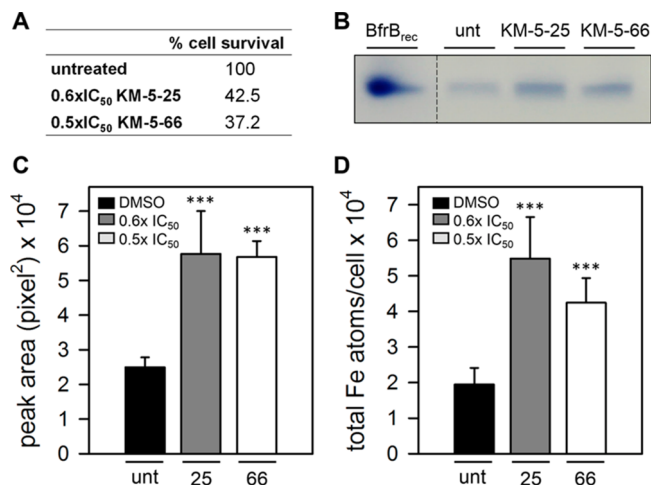


Figure 11. 4-Aminoisindoline-1,3-dione analogues penetrate the *P. aeruginosa* cell, bind BfrB, and inhibit mobilization of BfrB-stored iron. (A) Treating pellicles with 0.6× IC_{50} **KM-5-25** (40 μM) or 0.5× IC_{50} **KM-5-66** (20 μM) for 24 h reduces the number of viable cells to <50%. (B) The iron stored in BfrB in the viable cells was visualized with the aid of native PAGE gels stained with Ferene S, which stains the iron in the interior cavity of BfrB. Recombinant BfrB (BfrB_{rec}) was used as a standard for the electrophoretic mobility of BfrB. The lane corresponding to untreated control was loaded with 0.5× the volume of the lanes loaded with lysates from treated pellicles to account for the ~2-fold larger number of viable cells in the untreated pellicles. Lanes loaded with treated pellicle lysates show greater accumulation of iron in BfrB relative to untreated cells. (C) Peak areas obtained from densitometry analysis (ImageJ) of the bands in the native PAGE gel of panel B indicate that there is ~3-fold more iron stored in BfrB in the treated cells relative to the untreated control. (D) Analysis of total intracellular iron levels normalized to CFU/mL indicates ~2.5-fold higher iron levels in the pellicle-embedded cells treated with the 4-aminoisindoline-1,3-dione analogues relative to untreated control. Panel B shows results from a representative experiment from 3 biological replicates. Panels A, C, and D show the average of 3 biological replicates. $p < 0.001$ is denoted by *** relative to untreated.

natants from the untreated control were diluted approximately 2-fold prior to loading the native gels. Results obtained with a representative gel are shown in Figure 11B, where it can be observed that lanes loaded with lysate solutions from pellicles treated with analogues **KM-5-25** or **KM-5-66** exhibit significantly higher Ferene S stain intensity than the lane loaded with lysate solution from the untreated pellicle. To enable quantitative comparison we measured the relative intensities of the Ferene S-stained bands with the aid of ImageJ. Comparing the resultant peak areas (Figure 11C) shows that BfrB from the cells treated with **KM-5-25** or **KM-5-66** has ~3-fold more iron relative to BfrB from cells in the untreated pellicles. These findings provide strong evidence indicating that compounds **KM-5-25** and **KM-5-66** bind BfrB in the *P. aeruginosa* cytosol, inhibit the formation of the BfrB-Bfd complex required to mobilize iron from BfrB, and lead to nearly irreversible iron accumulation in BfrB. Consistent with

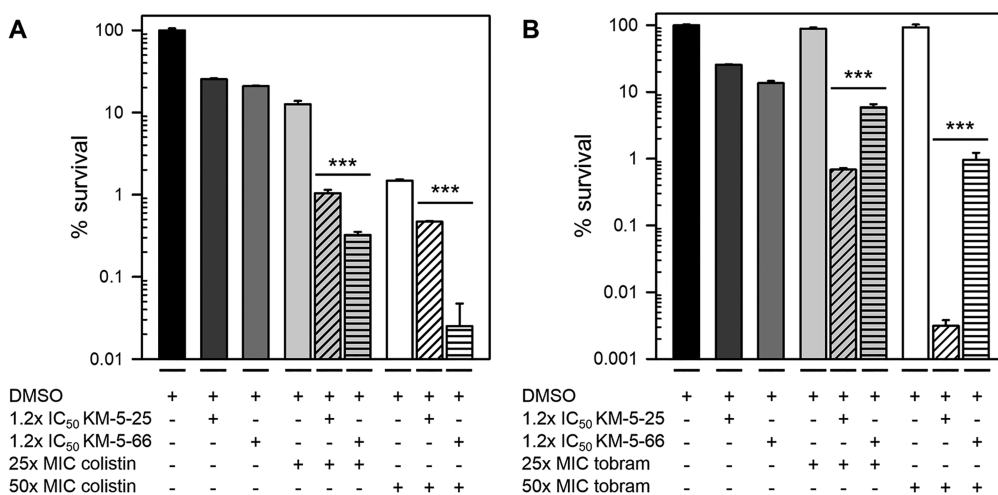


Figure 12. 4-Aminoisoindoline-1,3-dione derivatives enhance the efficacy of colistin and tobramycin against *P. aeruginosa* biofilms. Two-day old pellicles of EYFP-expressing *P. aeruginosa* PAO1 were treated for 24 h with (A) colistin alone 25× MIC (20 μM), or 50× MIC (40 μM), KM-5-25 (80 μM) or KM-5-66 (50 μM) alone, equivalent to 1.2× IC₅₀, or a combination of colistin and KM-5-25 or KM-5-66, and (B) tobramycin alone 25× MIC (27 μM) or 50× MIC (40 μM), KM-5-25 or KM-5-66 alone at a concentration equivalent to 1.2× IC₅₀, or a combination of tobramycin and KM-5-25 or KM-5-66. The % survival is expressed as the ratio CFU/mL_(after treatment)/CFU/mL_(pretreatment). $p < 0.001$ is denoted by *** in the combination treatment relative to treatment with antibiotic alone.

the nearly irreversible accumulation of iron in BfrB in cells treated with KM-5-25 or KM-5-66, quantification of the total intracellular iron and normalizing the values to viable cell counts demonstrates that *P. aeruginosa* cells dispersed from pellicles treated with KM-5-25 or KM-5-66 harbor ~2.5-fold more intracellular iron than cells obtained from untreated biofilms (Figure 11D). Taken together, these observations support the idea that 4-aminoisoindoline-1,3-derivatives dysregulate iron homeostasis by inhibiting the BfrB-Bfd complex, causing the accumulation of unusable iron in the bacterial cell.

4-Aminoisoindoline-1,3-dione Derivatives Enhance the Efficacy of Colistin and Tobramycin against Biofilm-Embedded Cells. As demonstrated above and in previous reports,^{65,66} mature biofilms formed by *P. aeruginosa* cells are susceptible to colistin and tolerant to tobramycin. Since these biofilms are also susceptible to the 4-aminoisoindoline-1,3-dione inhibitors of the BfrB-Bfd complex, we asked whether these compounds would enhance the efficacy of colistin and tobramycin. To answer this question, we cultured 2-day old pellicle biofilms of EYFP-expressing *P. aeruginosa* PAO1 and treated them for 24 h with colistin alone, compound alone (KM-5-25 or KM-5-66), and with a combination of colistin and compound. In the combination treatment experiments, the concentration of compound was kept constant (1.2× IC₅₀), while colistin was used at two different concentrations, equivalent to 25× and 50× MIC (Figure 12A). Treatment with each of the compounds or with colistin alone caused a reduction of viable cells similar to that reported in Figure 9A. Challenging the pellicles with a combination of colistin and compound, however, causes a significant additional reduction in the number of viable cells (Figure 12A): the combination treatment with KM-5-66 results in ~0.3% survival when colistin is present at 25× MIC, and ~0.02% survival when colistin is used at 50× MIC, which correspond to approximately 1.7 log and 1.9 log reduction of viable cells relative to treatment with colistin alone. In comparison, the combination treatment with KM-5-25 results in ~1% survival when colistin is used 25× MIC and ~0.5%

survival when colistin is present at 50× MIC, which correspond to nearly 1 log and 0.7 log reduction of viable cells, respectively, compared to colistin alone. Encouraged by these results we tested whether the 4-aminoisoindoline-1,3-dione derivatives can also enhance the bactericidal activity of tobramycin. The results are shown in Figure 12B: the combination treatment with KM-5-25 results in ~0.5% survival when tobramycin is used at 25× MIC and ~0.003% survival when tobramycin is present at 50× MIC, which correspond to approximately 2.5 log and 4.7 log reduction of viable cells when compared to treatment with tobramycin alone. The combination treatment with KM-5-66 results in 5% survival when tobramycin is present at 25× MIC and 1% survival when tobramycin is used at 50× MIC, which correspond to approximately 1.5 log and 2 log reduction in viable cells relative to treatment with tobramycin alone. It is interesting to note that compound KM-5-66 is more effective at enhancing the efficacy of colistin (Figure 12A), whereas compound KM-5-25 is more effective at enhancing the efficacy of tobramycin (Figure 12B). Additional studies are clearly required to understand the underlying reasons.

The observations above, which indicate that the iron limitation induced by inhibitors of the BfrB-Bfd complex can increase the efficacy of colistin and tobramycin against biofilms, are in good agreement with previous studies showing that the Fe chelator HBDE is an effective colistin adjunct against *P. aeruginosa*,⁶⁷ and the iron chelators deferoxamine and deferisirox increase the efficacy of tobramycin against *P. aeruginosa* biofilms.⁶⁸ When taken together, the observations made in the presence of HBDE, deferoxamine, deferisirox or 4-aminoisoindoline-1,3-dione derivatives, strengthen the idea that inducing intracellular iron limitation is probably a viable strategy to enhance the efficacy of colistin or tobramycin against biofilms. Since colistin is often used as one of the very few therapeutic options available to combat multidrug resistant Gram-negative organisms such as *Acinetobacter baumannii*, *Pseudomonas aeruginosa*, and *Klebsiella pneumoniae*,^{69,70} it is encouraging that small molecule inhibitors of the BfrB-Bfd complex are capable of increasing the effectiveness of colistin

against *P. aeruginosa*. In this context, it is noteworthy that the *bfr* and *bfd* genes in *A. baumannii* and *K. pneumoniae* strains are contiguous, as is the case in the >4000 *P. aeruginosa* genomes currently available in the Pseudomonas Genome Data Base, because it suggests that the function of the Bfr-Bfd complex in *P. aeruginosa* is conserved in *A. baumannii* and *K. pneumoniae*. Moreover, amino acid sequence alignment with the aid of Clustal Omega⁷¹ reveals very high conservation in the amino acid sequences of Bfr and Bfd proteins from *P. aeruginosa*, *A. baumannii* and *K. pneumoniae* (Figure S7–S8). Importantly, the amino acids identified as hot spot residues in the Bfr-Bfd complex of *P. aeruginosa*³³ are highly conserved in the Bfr and Bfd sequences of *A. baumannii* and *K. pneumoniae*, suggesting that the 4-aminoisindoline-1,3-dione derivatives designed to inhibit the Bfr-Bfd interaction in *P. aeruginosa* are likely to act similarly in *A. baumannii* and *K. pneumoniae*.

CONCLUDING REMARKS

Bacterial iron metabolism is a vulnerability that may be exploited in the development of new antimicrobial therapies. The field has centered on four major avenues: (i) The development of siderophore-antibiotic conjugates aims at capitalizing on siderophore uptake receptors to enhance antibiotic cell penetration.^{72,73} (ii) The sequestration of iron with the aid of chelators is also directed at depleting intracellular iron, and whereas iron chelation has been shown to lead to biofilm dispersion or to prevent biofilm maturation in vitro,^{35,37,48,68,74} a potential concern is the probability of secondary infections fueled by the chelated iron.^{75,76} (iii) The perturbation of heme uptake or heme degradation aims at denying pathogens from using heme, a rich iron source during infection.^{77,78} (iv) The perturbation of iron homeostasis with Ga³⁺ aims at inducing iron limitation by replacing Fe³⁺ in the active site of enzymes with Ga³⁺ which cannot support the rich redox chemistry typical of iron.^{79,80} Systemic Ga³⁺ treatment has been shown to improve lung function in patients with chronic *P. aeruginosa* infection.⁵⁶ Our work, which aims at adding an entirely new strategy for exploiting the bacterial iron metabolism vulnerability, is directed at inhibiting the mobilization of iron from bacterioferritin. Blocking iron mobilization from BfrB by deletion of the *bfd* gene has been shown to elicit a breakdown in iron homeostasis due to the irreversible accumulation of unusable iron in BfrB, which causes cytosolic iron deficiency.²³ Because of the relatively high iron requirement necessary to support the biofilm lifestyle, the intracellular iron deficiency induced in the Δbfd cells has been shown to result in poorly developed biofilms.²⁶ Encouraged by these findings our laboratories have pursued drug discovery strategies to discover small molecule inhibitors of the BfrB-Bfd complex, which led us to a series of 4-aminoisindoline-1,3-dione derivatives capable of penetrating the *P. aeruginosa* cell envelope, inhibiting the BfrB-Bfd complex and eliciting a cell growth defect in planktonic cells.²⁵ Additional structural manipulation of these 4-aminoisindoline-1,3-dione derivatives led us to analogues with increased affinity for BfrB and enhanced bacteriostatic activity against planktonic *P. aeruginosa* cells (Table 1). Testing these new derivatives against mature biofilms revealed that these small molecules inhibit iron mobilization from BfrB in biofilm-embedded cells (Figure 11) and kill *P. aeruginosa* cells in mature biofilms cultured in flow cells (Figure 6) or in pellicles cultured at the air–liquid interface (Figure 8). These findings demonstrate the potential of small molecules developed to inhibit the BfrB-Bfd complex

in *P. aeruginosa* as important tools to expose a rare weakness of biofilms. It is also encouraging that the small molecule inhibitors of the BfrB-Bfd complex increase the effectiveness of tobramycin and colistin because the latter is often used as one of the very few therapeutic options available to combat multidrug resistant Gram-negative organisms.^{69,70} The high level of conservation in the Bfr and Bfd sequences from *P. aeruginosa*, *A. baumannii*, and *K. pneumoniae* (Figure S7–S8) suggests that the compounds developed to block the BfrB-Bfd complex in *P. aeruginosa* may exert similar activity in *A. baumannii* and *K. pneumoniae* cells.

METHODS

Chemicals, Bacterial Strains, and Growth Media.

Chemicals were purchased from Fisher Scientific unless otherwise stated. *P. aeruginosa* PAO1 was obtained from the University of Washington Genome center.⁸¹ The PAO1 strain expressing enhanced yellow fluorescent protein (EYFP) was prepared previously.²⁶ Clinical isolates of *P. aeruginosa* were purchased from JMI Laboratories (North Liberty, IA, USA). IC₅₀ determinations were carried out in defined media (50 mM KH₂PO₄ (Sigma-Aldrich) 7.5 mM (NH₄)₂SO₄ (Sigma, Life Sciences), 0.1% (w/v) glucose (Acros Organics, 99+%), 0.5 mM MgSO₄·7H₂O (Sigma-Aldrich, 99+%), 5% v/v nonessential amino acids (Gibco), 2% v/v essential amino acids (Gibco), 4 μ M (NH₄)₂Fe(SO₄)₂, and 0.025% (w/v) hypromellose (HPMC, Sigma-Aldrich), pH 7.0. The media was filter-sterilized by passing through a 0.2 μ m cellulose acetate membrane syringe filter (VWR). Starter cultures of *P. aeruginosa* PAO1 in 5 mL LB media were grown for 13 h in 50 mL conical tubes at 37 °C and 220 rpm. For biofilm experiments, a EYFP-expressing *P. aeruginosa* strain was routinely grown in Pseudomonas Isolation (PI) media (20 g L⁻¹ peptone, 0.3 g L⁻¹ MgCl₂·6H₂O, 10 g L⁻¹ K₂SO₄, 25 mg L⁻¹ irgasan, and 20 mL L⁻¹ glycerol, pH 7.0). Starter cultures were grown from a single colony at 37 °C and shaking at 220 rpm for 14 h in 5 mL PI media supplemented with 10 μ M Fe. Pellicle biofilms were cultured for 48 h at 30 °C in PI media supplemented with 20 μ M Fe. Surface-attached biofilms were cultured in AB minimal media⁸² supplemented with trace metals [0.15 μ M (NH₄)₂MoO₄, 3 μ M CuSO₄, 2 μ M Co(NO₃)₂, 9.4 μ M Na₂B₄O₇, and 7.6 μ M ZnSO₄], 3 mM glucose and 15 μ M Fe. Iron supplementation was carried out by addition of a small volume of filter-sterilized 10 mM (NH₄)₂Fe(SO₄)₂ (pH ~ 2.0) solution. The antibiotics ciprofloxacin, colistin and tobramycin were used at concentrations equivalent to 25 \times and 50 \times the reported MIC:⁸³ ciprofloxacin MIC = 0.25 μ g/mL = 0.75 μ M; tobramycin MIC = 0.5 μ g/mL = 1.07 μ M; colistin MIC = 1 μ g/mL = 0.79 μ M. Compound stock solutions (100 mM or 10 mM) in DMSO (Sigma-Aldrich) were prepared weekly and stored at 4 °C. Solutions used to treat biofilms or planktonic cells include 0.025% (w/v) HPMC, and 1.5% or 2% DMSO (Sigma-Aldrich) to prevent aggregation of the analogues in aqueous solution.

Synthesis and Preparation of 4-Aminoisindoline-1,3-dione Derivatives. Experimental details of the synthetic protocols developed to prepare the 4-aminoisindoline-1,3-dione derivatives to be tested as inhibitors of the BfrB-Bfd complex, as well as ¹H, ¹³C NMR and MS data, are presented in the Supporting Information.

Measurement of Dissociation Constant (K_d). Dissociation constants for the interaction between BfrB and 3-

aminoisindoline-1,3-dione derivatives (Table 1, Figure S1) were measured in vitro with a fluorescence polarization method based on the intrinsic fluorescence of the isindoline-1,3-dione moiety, as described previously.²⁵

Measurement of Half Maximal Inhibitory Concentration (IC₅₀). IC₅₀ values (Table 1, Figure S1) were determined as reported previously²⁵ with small modifications. In brief: precultures of *P. aeruginosa* PAO1 (5 mL) were grown in LB media for 13 h at 37 °C and 220 rpm in 50 mL conical tubes (VWR International, PA). The cells were centrifuged for 5 min at 4000 rpm and 4 °C, washed two times and then diluted in buffer (100 mM KH₂PO₄ and 15 mM (NH₄)₂SO₄) to an optical density at 600 nm (OD₆₀₀) of 0.1. A small volume of compound stock solution (10 mM) was transferred to a microcentrifuge tube, initially diluted with DMSO to 20 μL, and then diluted to 1 mL with preculture cell suspension in defined media with OD₆₀₀ = 0.0001, so the final DMSO concentration is 2%. The resultant cell suspension (200 μL) was transferred to a clear-bottom polystyrene 96-well plate (VWR) covered with a lid and incubated at 35 °C and 205 cpm for 24 h in a Synergy H1 microplate reader (Biotek Instruments Inc., Vermont). The cell cultures were serially diluted and then plated on PI Agar (PIA; BD biosciences) plates for enumeration of viable cells (CFU/mL). The % growth was calculated from the ratio CFU/mL_(treated)/CFU/mL_(untreated control). To calculate the IC₅₀ values, the % growth was plotted as a function of compound concentration, expressed as log[compound] (μM), and fitted to the 4-parameter logistic model describing the sigmoid-shaped response pattern (eq 1),⁸⁴ where *b* is the slope factor, max is the upper asymptote (plateau), and min is the lower asymptote. Values are the average and standard deviation from three independent experiments.

$$\% \text{ growth} = \min + \frac{\max - \min}{1 + 10^{(\log \text{IC}_{50} - x)b}} \quad (1)$$

Analysis of Secreted Pyoverdine in Planktonic Cultures. These experiments were conducted in 96 well plates as described above for the determination of IC₅₀, except that the cells were cultured in M63 media (2 g/L (NH₄)₂SO₄, 13.6 g/L KH₂PO₄ (Sigma-Aldrich), 2 g/L glucose, 4 g/L citric acid, 5 g/L technical grade casamino acids (BD Scientific), 0.24 g/L MgSO₄ (Alfa Aesar), and 0.05% (w/v) HPMC, pH 7.0 adjusted with KOH). Cultures of *P. aeruginosa* PAO1 treated with KM-5-25 (70 μM) or KM-5-66 (50 μM) were grown for 27 h prior to diluting the contents of each well in PBS (pH 7.4) and plating the cells on PIA plates for enumeration of CFU/mL. The 500-fold diluted solution was clarified by centrifugation and the pyoverdine in the cell-free supernatant was analyzed by acquiring fluorescence emission spectra (430–550 nm) with excitation at 400 nm (10 nm slit width) and emission at λ_{max} = 460 nm (10 nm slit width) using a PerkinElmer LS50B spectrophotometer.

Flow Cell Biofilm Assays. Surface-attached biofilms of *P. aeruginosa* PAO1 cells expressing EYFP were grown on flow cells with an 800 μm channel depth (μ-slide 1^{0.8} Luer, Ibidi) using an automated perfusion system (Ibidi, Munich, Germany), as described previously.²⁶ Briefly, the flow cell was inoculated with 200 μL of an overnight culture diluted to OD₆₀₀ = 0.5, followed by 1 h incubation at 30 °C to allow bacterial cell attachment. The μ-slide was connected to the Ibidi Pump System and the biofilms were cultured for 3 days at 30 °C while flowing AB minimal media containing 15 μM Fe.

The experimental shear stress was 0.14 dyn/cm² (shear rate = 14 s⁻¹, pressure = 7.1 mbar, flow rate = 0.4 mL/min) and the switch time was set to 540 s. The biofilms were treated for 24 h by flowing AB minimal media supplemented with 15 μM Fe, 0.025% HPMC, 1.5% DMSO and commercial antibiotics or 4-aminoisindoline-1,3-dione derivatives in the concentrations indicated in the corresponding figure captions. During biofilm growth and challenge with antibacterial, the culture medium in the reservoirs was removed every 12 h and replaced with fresh prewarmed medium. Prior to imaging with the aid of CLSM the biofilms were stained with 4 mL of 2.5 nM Sytox Red (Invitrogen Life Technologies), a cell impermeable fluorescent nucleic acid dye that stains dead cells and extracellular DNA,⁸⁵ for 20 min (switch time = 200 s) and then washed with AB media for 20 min to remove excess fluorescent dye. The biofilms were imaged with the aid of a Leica TCS SP8 confocal microscope (Leica Microsystems, Germany) using a HC PL apo CS2 63 × /1.4 oil objective. For detecting the EYFP fluorescence the laser line was set at 506 nm and the emission range to 520–610 nm. Sytox Red fluorescence was detected with excitation at 631 nm and emission range 637–779 nm. Image stacks were acquired with a z-step size of 0.3 μm at randomly chosen positions. The Leica Application Suite X (LAS-X) software was used for image stack processing.⁵⁵ Quantitative analysis of biofilm biomass was performed using the COMSTAT computer program⁵⁵ and the Otsu method of automatic thresholding.⁸⁶

Pellicle Biofilm Assays. Pellicle biofilms of EYFP-expressing *P. aeruginosa* PAO1 or clinical isolates were grown in PI media supplemented with 20 μM Fe. Starter cultures were diluted to OD₆₀₀ = 0.001 in 4 mL media, placed in 35 × 10 mm Petri dishes and incubated statically at 30 °C for 48 h. The pellicles were transferred onto circular (1.5 cm diameter) glass coverslips by gently allowing the surface of a coverslip to contact a pellicle. The pellicle-adhered coverslip was washed in PBS and then deposited on top of 1.5 mL AB challenge media contained in a well of a 12-well microplate, with the pellicle exposed to the media. Challenge media consists of AB minimal media supplemented with 15 μM Fe, 0.025% HPMC, 1.5% DMSO, and commercial antibiotic or 4-aminoisindoline-1,3-dione derivative, used in the concentrations specified in the figure captions. The 4-aminoisindoline-1,3-dione derivatives were prepared as 10 mM stock solutions in DMSO and then diluted in culture media to the appropriate concentrations. The coverslip-adhered pellicles were exposed to challenge media at 30 °C for 24 h, changing the challenge media every 12 h by transferring the pellicle-adhered coverslip to a new plate containing prewarmed challenge media.

Prior to imaging with the aid of CLSM, pellicles formed by EYFP-expressing PAO1 cells were washed with PBS and then stained by placing the coverslip-adhered pellicles in 1 mL of PBS containing 2.5 nM Sytox Red (20 min). Excess fluorescent dye was washed with PBS, the coverslip was mounted on a glass slide using 5 μL SlowFade (Invitrogen Life Technologies) and the edges sealed with fingernail polish. CLSM image stacks (z-step size of 0.3 μm) were acquired with the aid of a Leica TCS SP8 microscope, as described above. Quantitative analysis was performed by determination of pellicle biomass using COMSTAT⁵⁵ and the Otsu method of automatic thresholding.⁸⁶

Determination of Viable Cells in Pellicle Biofilms. Pellicle biofilms were grown as described above. Planktonic and loosely attached cells were washed (3 times) by immersing

the coverslip-adhered pellicles (biofilm facing up) into a well of a 12-well plate containing 3 mL PBS, and incubating (5 min) with gentle rocking. To remove the pellicle from the coverslip, break the extracellular matrix and release cells from the biofilm, the coverslip-adhered pellicle was placed in a 50 mL conical tube containing a 2 mL suspension of zirconia beads (0.1 mm diameter, BioSpec Products), 10 mL PBS, 0.2 $\mu\text{g/mL}$ alginate lyase and 0.2 $\mu\text{g/mL}$ DNase. The resultant mixture was incubated at room temperature for 3 min, followed by vigorous vortexing for 4 min. After sedimentation of the zirconia beads, a 100 μL aliquot was used for serial dilution and plating on PIA plates for subsequent enumeration of viable cells (CFU/mL).

Imaging of Iron Stored in BfrB and Analysis of Total Intracellular Iron in Biofilm-Embedded Cells. Pellicle biofilms were grown for 48 h as described above. The pellicles were transferred onto square (2×2 cm) glass coverslips by gently contacting the pellicle with the coverslip. The pellicle-adhered coverslip was washed in PBS and then placed in a 50 mL conical tube containing 2 mL suspension of zirconia beads, 15 mL PBS, 0.2 $\mu\text{g/mL}$ alginate lyase and 0.2 $\mu\text{g/mL}$ DNase; the resultant mixture was incubated (3 min) at room temperature, and then vortexed vigorously for 4 min. After the zirconia beads had sedimented, a 100 μL aliquot was sampled from the cell suspension for plating and enumeration of viable cells and a 14 mL sample was used to harvest the cells by centrifugation (20 min, 400 rpm). The cell pellet was resuspended in 1 mL PBS, transferred to a 1.5 mL microcentrifuge tube, centrifuged for 10 min at 12 500 rpm at 4 °C and the cell pellet frozen at -80 °C. The frozen cells were subjected to three freeze–thaw cycles and then lysed by addition of 200 μL of lysis buffer (50 mM Tris-HCl buffer (pH 8.0) containing 10% (v/v) glycerol, 20 mg/mL lysozyme, 0.2 mg/mL DNase, 0.1 M NaCl, 1 mM MgSO_4 and 1% (v/v) Triton-X100) and incubated at ambient temperature (30 min) and at 37 °C (30 min). Imaging of iron in BfrB was carried out as previously reported:²³ lysate suspensions were clarified by centrifugation (10 min at 12 500 rpm), mixed with 10 μL loading dye (5.9 mL deionized water, 0.5 mL glycerol, 0.4 mL β -mercaptoethanol, 0.4 mL 1% (w/v) bromophenol blue, and 0.5 mL 1 M Tris-HCl (pH 6.8), and loaded onto 1.5 mm-thick native PAGE gels (4% stacking gel, 8% resolving gel). Electrophoresis was carried out at 60 V and 4 °C for 9 h, and the gels were stained in the dark by immersion (10 min) in a solution containing 0.049 g Ferene S, 250 μL thioglycolic acid, 2.4 mL acetic acid and 100 mL deionized water. Levels of total intracellular iron were determined as reported previously:^{23,87} The cell pellets were treated with 500 μL of freshly prepared digestion reagent (0.6 N HCl, 2.25% (w/v) KMnO_4 in water), thoroughly mixed by vortexing, and then incubated at 65 °C for 3.5 h. The resultant solutions were cooled to ambient temperature, treated with 100 μL of iron detection reagent (6.5 mM Ferene S, 15.4 mM neocuproine, 2 M ascorbic acid, and 5 M ammonium acetate), incubated for 30 min at ambient temperature and centrifuged for 5 min at 12 500 rpm. The iron concentration was measured from the absorbance of the Fe^{2+} –Ferene S complex ($\epsilon_{593} = 34.5 \text{ mM}^{-1} \text{ cm}^{-1}$),⁸⁸ normalized by the viable cell counts and reported as Fe atoms per cell.

Statistical Analysis. Statistical significance between the means and standard deviation of values obtained in experiments comparing results from untreated vs treated with antibiotic or analogue conditions was determined using one-way ANOVA followed by Tukey's multiple post hoc test, with the aid of SigmaPlot (Systat Software, Inc. CA).

X-ray Crystallography. Crystallization screening of wild type BfrB was conducted in Compact 300 (Rigaku Reagents) sitting drop vapor diffusion plates at 20 °C using equal volumes of protein and crystallization solution equilibrated against 75 μL of the latter. Crystals displaying a prismatic morphology were obtained in 1–2 days from the Cryo 1–2 HT screen (Rigaku Reagents) condition H6 (30% (v/v) PEG 200, 100 mM Na acetate pH 4.5, 100 mM NaCl). To prepare the ligand complexes, stock solutions (100 mM) of ligand in DMSO were mixed with the crystallization solution to obtain a 10 mM ligand soaking solution. Crystals were transferred to the soaking solution and incubated for 3 h before harvesting directly from the drop and storing in liquid nitrogen. Due to the low solubility of KM-5-66, a multicomponent mixture⁸⁹ composed of 25% (v/v) dioxane, 25% (v/v) glycerol, 25% (v/v) diethylene glycol and 25% (v/v) ethylene glycol (SM3) was prepared to improve the ligand solubility. The ligand stock solution (100 mM in DMSO) was mixed in a 1:1 (v/v) ratio with SM3; 4 μL of the resultant solution was mixed with 5 μL of 2 \times crystallant solution and 1 μL of 10 \times buffer (1 M HEPES pH 7.5) to give a 20 mM ligand solution. Crystals were transferred to the soaking solution and incubated for 3 h before harvesting directly from the drop and storing in liquid nitrogen. Data were collected at the Advanced Photon Source beamline IMCA-CAT beamline 17-ID. Intensities were integrated using XDS⁹⁰ via Autoproc⁹¹ and the Laue class analysis and data scaling were performed with Aimless.⁹² Structure solution was conducted by molecular replacement using the previously determined structure of wt holo-BfrB (PDB: 5D8O)³³ as the search model with Phaser.⁹³ Structure refinement and manual model building were conducted with Phenix⁹⁴ and Coot,⁹⁵ respectively. Structure validation was conducted with Molprobit.⁹⁶

■ ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acsinfectdis.0c00669>.

Quantification of K_d and IC_{50} for KM-5-25 and KM-5-66, superposition of structures of 4-aminoisindoline-1,3-dione derivatives with a $-(\text{CH}_2)_3-$ linker bound to BfrB, structure of KM-5-50 bound to BfrB, pyoverdine release from planktonic cells treated with KM-5-25 and KM-5-66, comparison of the efficacy of analogues 11, 16, KM-5-25, and KM-5-66 against pellicle biofilms, multiple sequence alignments of representative Bfr and Bfd proteins, X-ray diffraction data refining statistics table, synthetic procedures for the preparation of 4-aminoisindoline-1,3-dione derivatives, ^1H , ^{13}C NMR spectra and mass spectra of synthesized compounds (PDF)

Accession Codes

Coordinate and structures factors for the following inhibitor complexes with BfrB were deposited to the Worldwide Protein Data Bank with the accession codes 7K5E (JAG-5-7), 7K5G (KM-5-28), 7K5F (KM-5-50), and 7K5H (KM-5-66).

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Author Contributions

AS: Performed the pellicle and surface biofilm studies, tested the susceptibility of biofilms to antibiotics and 4-aminoisindoline-1,3-dione derivatives, and measured the total and stored iron in bacterioferritin. HY: Measured the K_d values, the susceptibility of planktonic cells to 4-aminoisindoline-1,3-dione derivatives, and the pyoverdine release. ANDPH: Screened 4-aminoisindoline-1,3-dione derivatives and contributed to evaluate the susceptibility of planktonic cells. KM and JKA-G: Synthesized and purified 4-aminoisindoline-1,3-dione derivatives. RAB: Directed the synthesis of 4-aminoisindoline-1,3-dione-derivatives. KPB: Collected X-ray diffraction data. SL: Solved X-ray crystal structures. MR: Directed the biology and analytical biochemistry studies and wrote the manuscript. AS, HY, RAB, and SL edited the manuscript.

Notes

The authors declare no competing financial interest.

ACKNOWLEDGMENTS

This study was supported by a grant from the National Science Foundation (MCB1837877) and a grant from the National Institutes of Health (AI125529) to MR. Use of the IMCA-CAT beamline 17-ID at the Advanced Photon Source was supported by the companies of the Industrial Macromolecular Crystallography Association through a contract with Hauptman-Woodward Medical Research Institute. Use of the Advanced Photon Source was supported by the U.S. Department of Energy, Office of Science, Office of Basic Energy Sciences under contract no. DE-AC02-06CH11357.

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