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# Enterobactin- and salmochelin- $\beta$ -lactam conjugates induce cell morphologies consistent with inhibition of penicillin-binding proteins in uropathogenic *Escherichia coli* CFT073†

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The design and synthesis of narrow-spectrum antibiotics that target a specific bacterial strain, species, or group of species is a promising strategy for treating bacterial infections when the causative agent is known. In this work, we report the synthesis and evaluation of four new siderophore- $\beta$ -lactam conjugates where the broad-spectrum  $\beta$ -lactam antibiotics cephalexin (Lex) and meropenem (Mem) are covalently attached to either enterobactin (Ent) or diglycosylated Ent (DGE) *via* a stable polyethylene glycol (PEG<sub>3</sub>) linker. These siderophore- $\beta$ -lactam conjugates showed enhanced minimum inhibitory concentrations against *Escherichia coli* compared to the parent antibiotics. Uptake studies with uropathogenic *E. coli* CFT073 demonstrated that the DGE- $\beta$ -lactams target the pathogen-associated catecholate siderophore receptor IroN. A comparative analysis of siderophore- $\beta$ -lactams harboring ampicillin (Amp), Lex and Mem indicated that the DGE-Mem conjugate is advantageous because it targets IroN and exhibits low minimum inhibitory concentrations, fast time-kill kinetics, and enhanced stability to serine  $\beta$ -lactamases. Phase-contrast and fluorescence imaging of *E. coli* treated with the siderophore- $\beta$ -lactam conjugates revealed cellular morphologies consistent with the inhibition of penicillin-binding proteins PBP3 (Ent/DGE-Amp/Lex) and PBP2 (Ent/DGE-Mem). Overall, this work illuminates the uptake and cell-killing activity of Ent- and DGE- $\beta$ -lactam conjugates against *E. coli* and supports that native siderophore scaffolds provide the opportunity for narrowing the activity spectrum of antibiotics in clinical use and targeting pathogenicity.

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## Introduction

Infections caused by Gram-negative bacteria can be difficult to treat due to the semipermeable outer-membrane (OM) that serves as an efficient barrier against most antibiotics.<sup>1,2</sup> The overuse of broad-spectrum antibiotics in the treatment of such bacterial infections facilitates the selection of resistant strains, which causes antibiotics to lose effectiveness over time.<sup>3,4</sup> Moreover, broad-spectrum antibiotics can damage the commensal microbiota and trigger life-threatening secondary

infections such as those caused by *Clostridioides difficile*.<sup>5,6</sup> Coupled with the scarcity of new antibiotics in the drug pipeline, society is faced with the reality that bacterial infections that were once of minor concern can become lethal.<sup>7</sup> To address this emerging public health crisis, novel therapies that exhibit narrow-spectrum activity must be developed and implemented in combination with rapid diagnostics.<sup>8–11</sup> Such pathogen-selective antibiotics are predicted to reduce the occurrence of antibiotic resistance in microbial populations and have smaller impact on the host microbiome.

To date, various strategies of narrowing the activity spectrum of antibiotics in clinical use have been explored.<sup>9,12–14</sup> In this regard, essential nutrient transporters located in the OM of Gram-negative bacteria provide opportunities for selective recognition and intracellular delivery of antibacterial molecules.<sup>15,16</sup> Iron is an essential nutrient for the vast majority of bacterial species; thus, acquiring adequate levels of iron is important for survival and host colonization.<sup>17–19</sup> To scavenge iron from the host, many bacteria biosynthesize small-molecule Fe<sup>3+</sup> chelators called siderophores. These secondary metabolites are biosynthesized in the cytoplasm, exported to the

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extracellular space to coordinate  $\text{Fe}^{3+}$  and then returned to the bacterial cell *via* specialized OM receptors.<sup>20–23</sup> Several decades of work showed that the siderophores and siderophore uptake machinery of Gram-negative bacteria can be leveraged to deliver toxic cargos into the bacterial periplasm and cytoplasm.<sup>15,24–33</sup> Recent examples demonstrated that the activity of broad-spectrum antibiotics (*e.g.*  $\beta$ -lactams and fluoroquinolones) can be narrowed by targeting select species or a group of strains within a species through covalent attachment to a siderophore.<sup>34–36</sup>

Enterobactin (Ent) is a tris-catecholate siderophore produced by Gram-negative species including *Escherichia coli*, *Salmonella enterica* and *Klebsiella pneumoniae*.<sup>37,38</sup> The high  $\text{Fe}^{3+}$ -binding affinity of Ent ( $K_a \sim 10^{49} \text{ M}^{-1}$ )<sup>39,40</sup> allows bacteria to scavenge  $\text{Fe}^{3+}$  from the host environment in the form of  $[\text{Fe}(\text{Ent})]^{3-}$ .<sup>41</sup> In *E. coli*,  $[\text{Fe}(\text{Ent})]^{3-}$  is recognized by the OM receptors FepA and IronN, and is transported into the periplasm utilizing the energy provided by the TonB–ExbB–ExbD complex

(Fig. S1†). In addition to producing Ent, a number of Gram-negative pathogens including uropathogenic *E. coli* and *Salmonella enterica* also biosynthesize salmochelins, C-glucosylated analogs of Ent. For instance, salmochelin S4, hereafter named diglucosylated Ent (DGE), is a C5,C5'-diglucosylated analogue of Ent.<sup>42</sup> DGE is deployed by pathogenic bacteria to acquire iron and evade the Ent-scavenging host-defense protein lipocalin-2.<sup>43–46</sup>  $[\text{Fe}(\text{DGE})]^{3-}$  is transported into the periplasm by IronN.<sup>43,47</sup> Whereas all *E. coli* produce Ent and express FepA, the production of salmochelins and expression of IronN are mostly pathogen-associated and require the *iroA* gene cluster.<sup>42,47</sup>

In prior work, we attached the aminopenicillins ampicillin (Amp) and amoxicillin (Amx) to the Ent scaffold *via* a polyethylene glycol (PEG<sub>3</sub>) linker installed at the C5 position of one catechol ring.<sup>34</sup> These Ent- $\beta$ -lactam conjugates afforded 100–1000-fold increases in antimicrobial activity against *E. coli* expressing FepA compared to the unmodified  $\beta$ -lactams. We

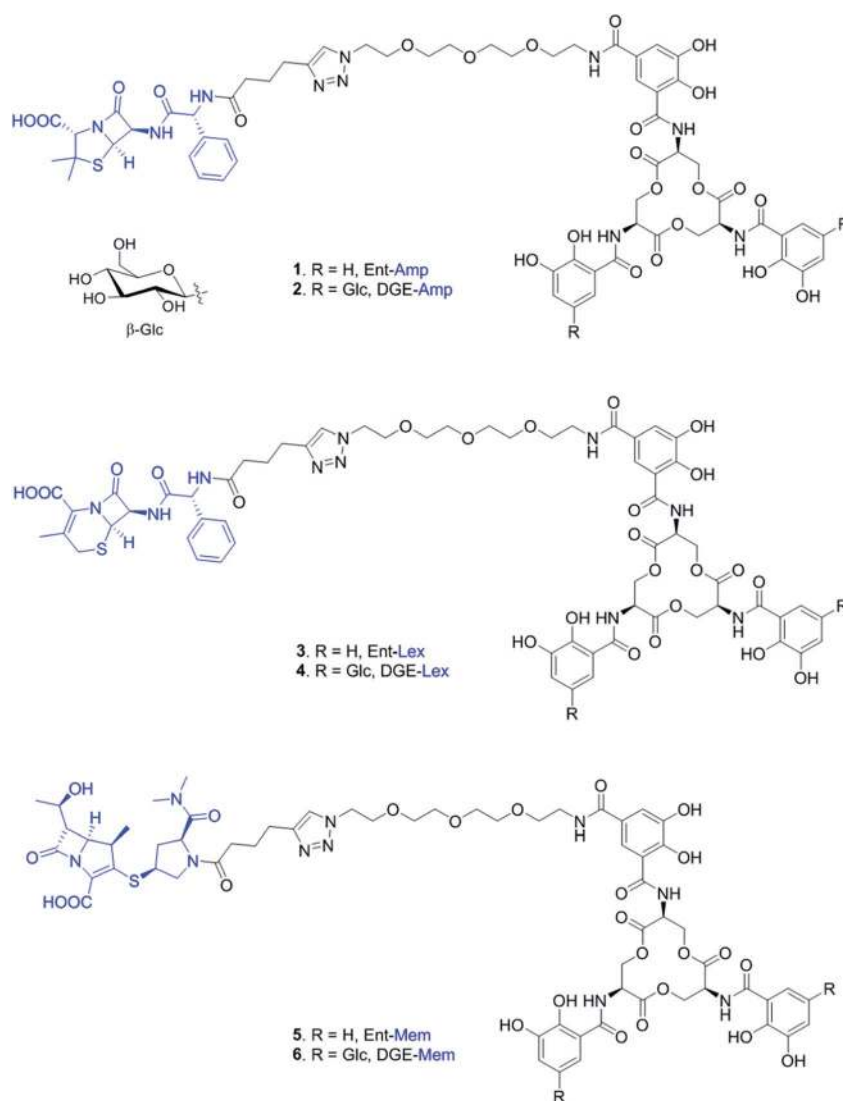


Fig. 1 Chemical structures of the Ent- and DGE- $\beta$ -lactam conjugates with Amp (1, 2), Lex (3, 4), and Mem (5, 6). Glc, glucose. The curved line on the  $\beta$ -glycosidic bond of glucose ( $\beta$ -Glc) represents the point of attachment to the catechol.



then demonstrated that C-glucosylation of the two remaining catechol rings, yielding DGE- $\beta$ -lactam conjugates, afforded selective killing of *E. coli* strains that harbor the *iroA* gene cluster.<sup>35</sup> The antibacterial activity of the Ent/DGE- $\beta$ -lactam conjugates was attributed to the  $\beta$ -lactam cargos because  $\beta$ -lactam hydrolysis of the Ent congeners abrogated this activity.<sup>34</sup>

Although it is accepted that the  $\beta$ -lactam cargo of siderophore- $\beta$ -lactam conjugates is responsible for bacterial cell death, studies addressing the mechanism of antimicrobial activity of siderophore- $\beta$ -lactam conjugates are limited.<sup>48–50</sup> Generally,  $\beta$ -lactam antibiotics enter Gram-negative bacteria *via* OM porins, bind to penicillin-binding proteins (PBPs) in the periplasm and thereby inhibit cell wall (CW) biosynthesis.<sup>51,52</sup> In bacteria, there are numerous PBP enzymes, which are classified according to their enzymatic function. In *E. coli*, class A PBPs such as PBP1a and PBP1b catalyze both transglucosylation (polymerization) of the peptidoglycan and transpeptidation (cross-linking) of the glycan strands. Class B enzymes such as PBP2 and PBP3 are monofunctional transpeptidases with roles in elongation and cell division, respectively. Class C enzymes, which include PBP4, PBP4b, PBP5, PBP6, PBP6b, and AmpH, carry out various peptidase reactions that facilitate maturation, remodeling, and metabolism of PG.<sup>53,54</sup> Consequently, inhibition of essential PBPs by  $\beta$ -lactams induces a diverse array of cellular morphologies, reflecting the roles that the enzymes play in coordinating cell division and elongation. Thus, studying the cellular morphologies that occur as a result of exposing *E. coli* to siderophore- $\beta$ -lactam conjugates is likely to provide insight into the observed antibacterial activity and cellular target(s).

In this work, we report the synthesis of four new siderophore- $\beta$ -lactam conjugates harboring the antibiotics cephalixin (Lex) and meropenem (Mem) (Fig. 1). These conjugates are based on our original Ent/DGE- $\beta$ -lactam design (*e.g.* Ent/DGE-Amp) and extend the scope of antibiotic cargos that can be effectively attached to Ent/DGE. These molecules show siderophore receptor-dependent antibacterial activity against *E. coli*. Importantly, Ent-Mem also retains antibacterial activity against a serine  $\beta$ -lactamase-producing strain. For the first time, we obtain microscopic evidence that Ent/DGE- $\beta$ -lactam conjugates exert antibacterial activity by inhibiting essential PBP enzymes. This study further demonstrates that the Ent and DGE scaffolds

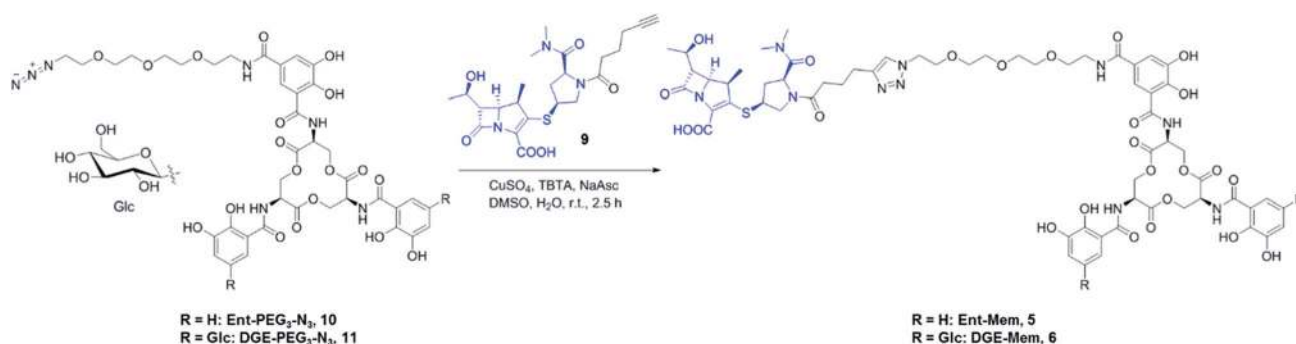
can render broad-spectrum antibiotics selective towards pathogenic bacteria on the basis of siderophore-uptake machineries.

## Results and discussion

### Design and synthesis of Ent/DGE- $\beta$ -lactam conjugates harboring Lex and Mem

We prepared and evaluated the Ent/DGE conjugates 3–6 carrying two additional broad-spectrum  $\beta$ -lactam antibiotics, Lex and Mem (Fig. 2A). Lex is a first-generation cephalosporin analog of Amp, whereas meropenem is a carbapenem with enhanced resistance to extended-spectrum  $\beta$ -lactamases and cephalosporinases.<sup>55–60</sup> From a structural standpoint, Amp, Lex and Mem differ by the size and nature of the ring fused with the  $\beta$ -lactam core: thiazolidine (Amp), dihydrothiazine (Lex), and thiazoline (Mem) (Fig. 1). Compared to Amp, the 4,6-fused ring system of Lex provides enhanced stability to acid hydrolysis,<sup>55,61</sup> whereas the 4,5-ring system of Mem, in which the endocyclic S atom of the 5-membered ring is replaced by C, confers superior resistance to serine  $\beta$ -lactamases.<sup>58–60</sup> Additionally, these structural differences (i) determine the rate with which Amp, Lex, and Mem diffuse through bacterial porins<sup>62</sup> and (ii) induce variations in  $\beta$ -lactam reactivity.<sup>61</sup>

The syntheses of conjugates 3–6 were carried out in one step from either Ent-PEG<sub>3</sub>-N<sub>3</sub> **10** or DGE-PEG<sub>3</sub>-N<sub>3</sub> **11** and one of the acylated  $\beta$ -lactams 7–9, using the methods developed for the assembly of Ent/DGE-Amp/Amx (Schemes 1 and S1†).<sup>34,35</sup> Each  $\beta$ -lactam was acylated with hex-5-ynoyl chloride to provide the alkyne functionality for the coupling reactions with azides **10** and **11** (Scheme S2†). The preparation of Lex-alkyne **8** was similar to that developed for Amp-alkyne **7** due to the comparable hydrophilicities of Amp and Lex ( $\log P_{\text{Amp}} = 1.35$ ,  $\log P_{\text{Lex}} = 0.65$ ),<sup>63–65</sup> which allows for an efficient extraction of **7** and **8** from acidified solution. Because Mem is more hydrophilic and difficult to extract from an aqueous phase ( $\log P = -0.6$ ),<sup>66</sup> the acylation reaction of Mem was optimized to avoid acid-catalyzed hydrolysis of Mem-alkyne **9** during extraction from an acidic solution. Thus, the water–acetone system employed for preparation of Amp/Lex-alkyne **7**, **8** was replaced by MeOH. Removal of MeOH under reduced pressure eliminated the need for



**Scheme 1** Synthesis of Ent/DGE-Mem **5**, **6**. The curved line on the  $\beta$ -glycosidic bond of glucose ( $\beta$ -Glc) represents the point of attachment to the catechol. Syntheses for the Amp and Lex conjugates are provided in Scheme S1.†



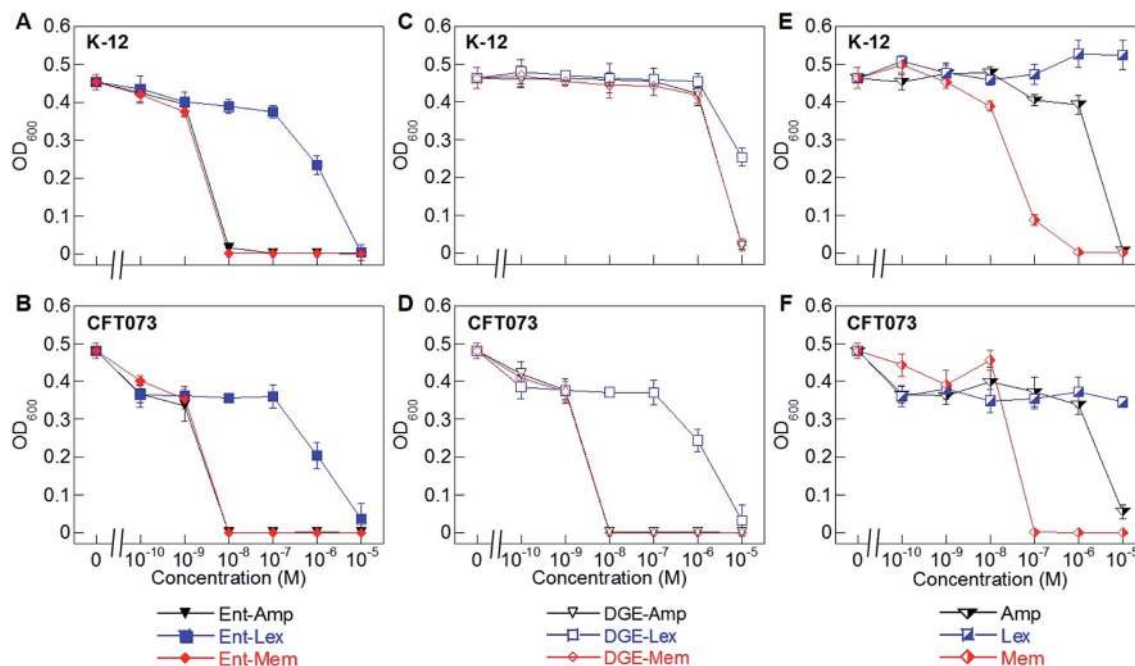


Fig. 2 Antibacterial activity of Ent/DGE- $\beta$ -lactam conjugates 1–6 against *E. coli* K-12 and CFT073. (A) and (B) Activity of Ent-Amp/Lex/Mem 1, 3, and 5; (C) and (D) activity of DGE-Amp/Lex/Mem 2, 4, and 6; (E) and (F) activity of the parent antibiotics Amp, Lex and Mem. All assays were performed in modified M9 medium (20 h, 30 °C; mean  $\pm$  standard deviation,  $n = 3$ ). Data for *E. coli* UTI89 are presented in Fig. S2† A summary of all MIC values for conjugates 1–6 is provided in Table S4.†

extraction from acidified solvent, which avoided hydrolysis of Mem-alkyne **9** during workup.

The resulting  $\beta$ -lactam alkynes **7–9** were coupled to Ent- or DGE-PEG<sub>3</sub>-N<sub>3</sub> **10** and **11** via copper-catalyzed azide-alkyne cycloaddition (Schemes 1 and S1†). TBTA (tris[(1-benzyl-1*H*-1,2,3-triazol-4-yl)methyl]amine) was used in all cycloaddition reactions to chelate copper and thereby prevent copper-mediated decomposition of the  $\beta$ -lactam ring.<sup>34,35</sup> The azide-alkyne cycloaddition reactions yielded 40–70% Ent/DGE- $\beta$ -lactam conjugates on a milligram scale.

### Conjugation to Ent/DGE enhances the antimicrobial activity of parent $\beta$ -lactams and affords selectivity towards pathogenic *E. coli* independent of $\beta$ -lactam structure

To evaluate the antimicrobial activity (AMA) of the Ent/DGE- $\beta$ -lactam conjugates against *E. coli*, we employed three *E. coli* strains: K-12, CFT073, and UTI89. *E. coli* K-12 (ref. 67) is a non-pathogenic laboratory strain that lacks the *iroA* gene cluster and relies on FepA for Ent-mediated iron uptake. Uropathogenic *E. coli* CFT073 (ref. 68–70) and UTI89 (ref. 71) harbor the *iroA* gene cluster and thus biosynthesize DGE and express IronN.<sup>72–75</sup> CFT073 also harbors the *iha* gene, which encodes an outer membrane receptor, Iha, that has been shown to transport [Fe(Ent)]<sup>3+</sup>.<sup>76,77</sup>

We performed AMA assays using a 10-fold dilution series to compare the AMA of Ent/DGE-Amp/Lex/Mem 1–6 and the parent antibiotics Amp/Lex/Mem (Fig. 2A–F and S2†). In these assays, the bacteria were treated with the apo siderophore- $\beta$ -lactam conjugates, which can scavenge iron from the culture

medium. These assays were conducted in modified M9 medium that provides iron-limiting conditions ( $\sim 0.3 \mu\text{M}$  Fe, Table S2†), causing *E. coli* to express the Ent/DGE uptake machineries.

Under iron limitation, the three *E. coli* strains were more susceptible to Ent-Amp/Lex/Mem (Fig. 2A, B and S2†) than to the corresponding parent antibiotics (Fig. 2E and F). Ent-Amp **1** showed minimum inhibitory concentration (MIC) values that were lower than Amp ( $10^{-5}$  M) by 1000-fold for both K-12 and CFT073, and by 100-fold for UTI89 (Fig. S2†). These findings are in overall agreement with our prior results obtained in MHB medium supplemented with the iron chelator 2,2'-bipyridine (Bpy).<sup>34</sup> Ent-Mem **5** afforded an MIC value of  $10^{-8}$  M in K-12, CFT073 and UTI89 (Fig. 2A, B and S2†), which represents a 10-fold reduction when compared to Mem ( $10^{-7}$  M) (Fig. 2E and F). In contrast to the Ent-Amp and -Mem conjugates, Ent-Lex **3** exhibited a relatively modest enhancement in the AMA compared to parent Lex (Fig. 2A, B and S2†). This fact may be attributed to the lower potency of Lex against the strains employed in this study.<sup>78</sup> It is also possible that the cell filamentation readily induced by Lex (*vide infra*) may impact the relationship between OD<sub>600</sub> and cell viability.

The AMA of the DGE- $\beta$ -lactams **2**, **4** and **6** paralleled that of the corresponding Ent- $\beta$ -lactams against *E. coli* CFT073 (Fig. 2D) and UTI89 (Fig. S2†). For instance, in CFT073 and UTI89, DGE-Mem **6** afforded an MIC value of  $10^{-8}$  M, and the MIC values of DGE-Amp **2** were  $10^{-8}$  M in CFT073 and  $10^{-7}$  M in UTI89, whereas DGE-Lex **4** showed a modest MIC value ( $10^{-5}$  M) consistent with that of Ent-Lex **3** (Fig. 2B, D and S2†). In contrast, the DGE- $\beta$ -lactam conjugates, especially DGE-Amp **2** and DGE-Mem **6**, exhibited attenuated AMA against K-12



(Fig. 2C). This result agrees with our prior observations for DGE-Amp/Amx<sup>35</sup> and, taken with the results for CFT073, indicates that expression of *IroN* is required for DGE- $\beta$ -lactam conjugates to exert AMA. We attribute the growth inhibition observed at  $10^{-5}$  M DGE-Amp/Lex/Mem to iron limitation that results from DGE- $\beta$ -lactams sequestering  $\text{Fe}^{3+}$  in the growth medium.

### Ent- $\beta$ -lactam conjugates retain $\beta$ -lactamase stability of parent $\beta$ -lactams

Hydrolysis of  $\beta$ -lactams by  $\beta$ -lactamases is a major mechanism of resistance to these antibiotics.<sup>51,79–81</sup> Of the three  $\beta$ -lactams considered in this work, Mem is the most resistant to  $\beta$ -lactamases as it is only a substrate for carbapenemases (metallo- $\beta$ -lactamases).<sup>59,60,82–84</sup> In contrast, Amp and Lex are readily hydrolyzed by serine  $\beta$ -lactamases.<sup>85,86</sup> To determine whether conjugation to the Ent scaffold affects the susceptibility of the  $\beta$ -lactam cargos to hydrolysis by a serine  $\beta$ -lactamase, we evaluated the AMA of Ent-Amp/Lex/Mem 1, 3, 5 against *E. coli* ATCC 35218, a strain that expresses a class A serine  $\beta$ -lactamase (Fig. 3A and B). As expected, neither Amp nor Lex displayed AMA against *E. coli* ATCC 35218, whereas Mem retained its activity (MIC =  $10^{-7}$  M) (Fig. 3B). In agreement with prior work,<sup>34</sup> Ent-Amp 1 exhibited negligible AMA against ATCC 35218 (Fig. 3A). Similarly to Ent-Amp 1, Ent-Lex 3 showed no AMA against *E. coli* ATCC 35218 (MIC >  $10^{-5}$  M) (Fig. 3A and B). In contrast, Ent-Mem 5 retained potent AMA against *E. coli* ATCC 35218, showing a 10-fold enhancement of the MIC value relative to Mem ( $10^{-8}$  M and  $10^{-7}$  M, respectively) (Fig. 3A and B). Thus, conjugation of Mem to Ent has negligible effect on the stability of the warhead to a class A serine  $\beta$ -lactamase. A genome search indicated that *E. coli* ATCC 35218 does not harbor *iroN*. In agreement with this analysis, none of the DGE- $\beta$ -lactams inhibited the growth of this strain over the concentration range tested, presumably due to lack of uptake (Table S4†). Taken together with the potent AMA of the Ent/DGE-Mem conjugates 5, 6, we conclude that Mem would be the most promising broad-spectrum  $\beta$ -lactam cargo for further *in vitro* and *in vivo* studies.

### Ent/DGE- $\beta$ -lactam conjugates are transported across the OM by Ent and salmochelin receptors

To ascertain the contributions of the Ent and salmochelin OM transporters for the uptake of the Ent/DGE- $\beta$ -lactam conjugates, we examined the AMA of Ent/DGE- $\beta$ -lactam conjugates 1–6 against *E. coli* CFT073. We selected *E. coli* CFT073 as a case study because this strain exhibits high susceptibility to all six conjugates and expresses *FepA*, *IroN*, and *Iha*.<sup>76,77</sup> We took a genetic approach using six OM and IM receptor mutants. Namely, mutants in *fepA*, *iroN*, *fepA iroN*, and *fepA iroN iha* were employed to probe conjugate recognition and transport across the OM (Fig. 4 and S3†). A *fepC* mutant (ATPase) and a *fepDG* double mutant (IM translocase) were used to investigate whether transport across the IM affects the AMA of the Ent/DGE- $\beta$ -lactams (Fig. S4†).

AMA assays under iron limitation revealed that single deletion of either *fepA* or *iroN* has negligible effect on the AMA of Ent-Mem 5 (Fig. 4A). This result is in agreement with our expectations and is explained by uptake of the Ent-based conjugate through both *FepA* and *IroN*.<sup>87</sup> In the case of DGE-Mem 6, deletion of *fepA* had no effect on the MIC value, whereas deletion of *iroN* attenuated the AMA by 1000-fold (Fig. 4B). This result indicates that *IroN* is essential for transport of the DGE- $\beta$ -lactam conjugates. Overall, these results are in agreement with prior siderophore competition studies that indicated that both *FepA* and *IroN* provide transport of Ent-Amp/Amx into the periplasm, whereas DGE-Amp/Amx can only be transported through *IroN*.<sup>85</sup>

The AMA of both Ent- and DGE-Mem conjugates 5 and 6 (Fig. 4A and B) was attenuated against a *fepA iroN* double mutant as well as a *fepA iroN iha* triple mutant. Moreover, the similar susceptibility of both mutants to the conjugates suggests that *Iha* is not involved in conjugate transport. The origin of the residual AMA for Ent/DGE-Mem is unclear. One possible explanation is that these mutant strains are more susceptible to conjugate-mediated  $\text{Fe}^{3+}$  sequestration from the growth medium compared to the parent strain because they lack multiple siderophore receptors. Alternatively, *E. coli* CFT073 might express an additional OM receptor that can

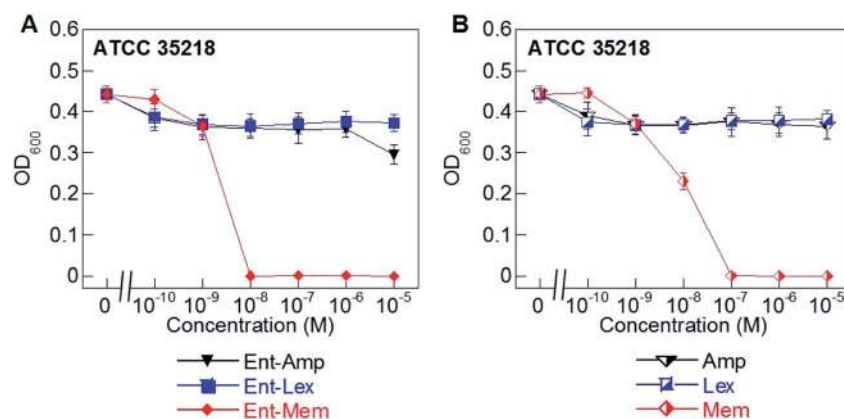


Fig. 3 Antibacterial activity of (A) Ent-Amp 1, -Lex 3, and -Mem 5 and (B) Amp, Lex, and Mem against the class A serine  $\beta$ -lactamase producer *E. coli* ATCC 35218. Assays were performed in modified M9 medium (20 h, 30 °C; mean  $\pm$  standard deviation,  $n = 3$ ). A summary of all MIC values, including those for the DGE- $\beta$ -lactam conjugates, are provided in Table S4.†



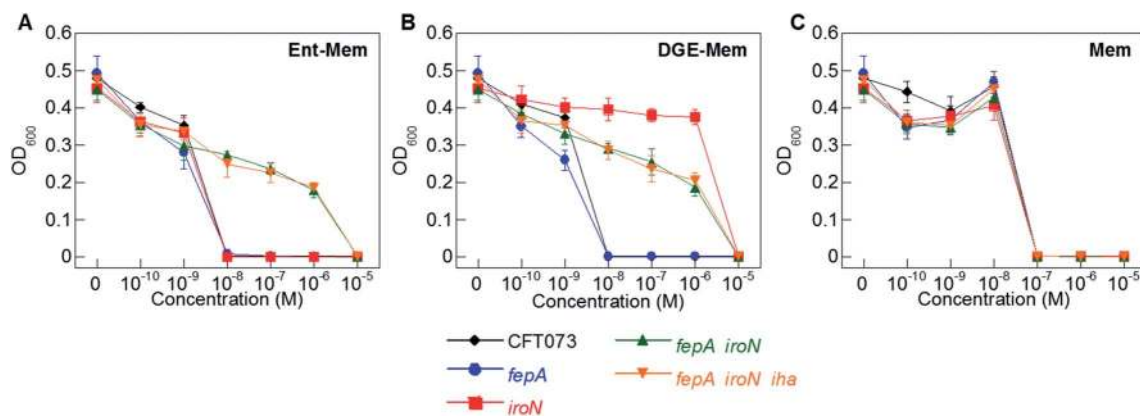


Fig. 4 Antibacterial activity of (A) Ent-Mem 5, (B) DGE-Mem 6 and (C) unmodified Mem against uropathogenic *E. coli* CFT073 wild-type and its isogenic OM receptor mutants in *fepA*, *iroN*, *fepA iroN*, and *fepA iroN iha*. All assays were performed in modified M9 medium (20 h, 30 °C; mean  $\pm$  standard deviation,  $n = 3$ ). The AMA plots for the OM receptor mutants treated with Ent/DGE-Amp/Lex 1–4 are shown in Fig. S3,† whereas the IM receptor mutants *fepC* and *fepDG* are shown in Fig. S4.†

transport the Ent/DGE conjugates across the membrane (e.g. IreA),<sup>88</sup> albeit less efficiently than FepA and IroN. Lastly, it is also possible that CFT073 secretes hydrolases that linearize Ent/DGE and subsequently imports the resulting iron-bound hydrolyzed conjugates into the periplasm *via* Cir or Fiu. These possibilities warrant further investigation, which could uncover yet-unknown pathways of Ent/DGE- $\beta$ -lactam uptake that explain the increased sensitivity of CFT073.

As expected, unmodified Mem exhibited potent AMA (MIC of  $10^{-7}$  M) against the six CFT073 OM mutants (Fig. 4C). This activity is consistent with porin-mediated uptake of unmodified  $\beta$ -lactams, which is unaffected by siderophore receptor expression.

Next, we investigated whether conjugate transport into the cytoplasm affects the antibacterial activity of Ent/DGE-Mem 5 and 6. Both the carrier and the cargo moieties of the Ent- and DGE- $\beta$ -lactam conjugates have binding partners in the periplasm. The  $\beta$ -lactam warheads target PBPs and form covalent acyl-enzyme species with an active site serine residue, whereas the siderophore moiety can be captured by FepB for delivery to the IM transporter FepCDG. To determine whether the presence

of FepCDG influences the antibacterial activity of the conjugates, we examined the susceptibility of a *fepC* mutant and a *fepDG* mutant. Neither deletion of *fepC* nor *fepDG* affected the antibacterial activity of the Ent- and DGE-Mem 5, 6 conjugates (Fig. S4†). These results indicate that, if any transport of the conjugates through FepCDG occurs, this process has negligible impact on AMA against *E. coli* CFT073. Therefore, we presume that the majority of the molecules are being trapped in the periplasm following covalent attachment to the PBPs. This conclusion agrees with our prior analysis of the *E. coli* K-12 *fepC* mutant, which showed comparable susceptibility to Ent-Amp/Amx as the parent strain.<sup>34</sup>

### Siderophore conjugation accelerates the killing of *E. coli* to a greater extent for Amp than for Mem

In previous studies, we found that conjugation of Amp/Amx to either Ent or DGE accelerated the rate of killing of *E. coli* CFT073 and UTI89 relative to the unmodified  $\beta$ -lactams.<sup>34,35</sup> We therefore sought to determine whether siderophore modification also enhances the time-kill kinetics of other  $\beta$ -lactams, and

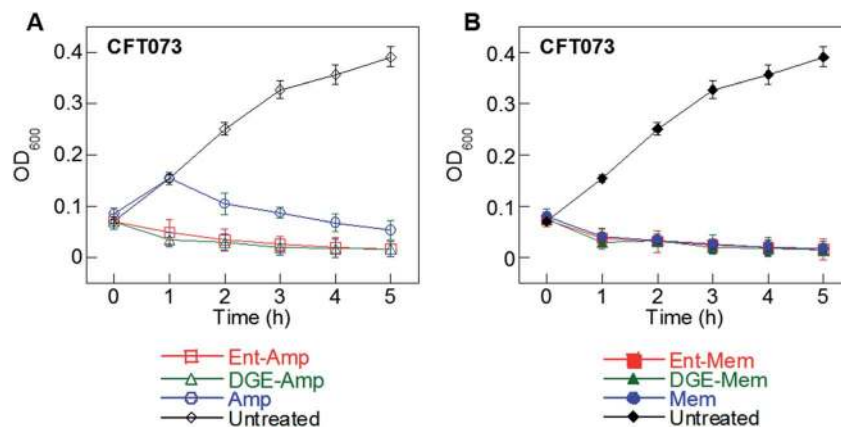


Fig. 5 Time-kill kinetics of (A) Ent/DGE-Amp 1, 2 and (B) Ent/DGE-Mem 5, 6 against *E. coli* CFT073 ( $\sim 10^8$  CFU mL<sup>-1</sup>) treated with 50  $\mu$ M Ent/DGE- $\beta$ -lactams and 100  $\mu$ M unmodified Amp/Mem. All assays were performed in modified M9 medium (37 °C; mean  $\pm$  standard deviation,  $n = 3$ ).



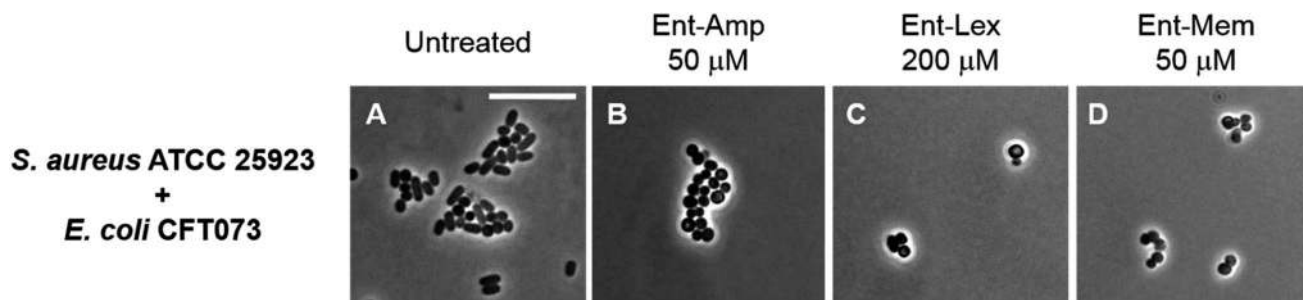


Fig. 6 Phase-contrast micrographs of mixed cultures of *E. coli* CFT073 (rods) with *S. aureus* ATCC 25923 (cocci) treated with MIC doses of Ent- $\beta$ -lactam conjugates. All cell cultures were incubated with the conjugates in 50% MHB in the presence of 100  $\mu$ M Bpy for 20 h at 30  $^{\circ}$ C prior to the microscopy imaging. Scale bar: 10  $\mu$ m. The micrographs of the mixed cultures treated with DGE- $\beta$ -lactam conjugates are provided in Fig. S7 $\dagger$

compared the time-kill kinetics of the conjugates with the lowest MIC values, Ent/DGE-Amp 1, 2 and Ent/DGE-Mem 5, 6, against *E. coli* CFT073.

Of the two  $\beta$ -lactams, Mem is more hydrophilic than Amp (*vide supra* for log  $P$  values) and is transported by OM porins more readily.<sup>89–92</sup> Indeed, the time-kill curves of Amp (Fig. 5A) and Mem (Fig. 5B) show that Mem reduces OD<sub>600</sub> of *E. coli* CFT073 almost to baseline within 1 h, whereas Amp requires 5 h to reach a similar OD<sub>600</sub> value. Conjugation to the Ent/DGE scaffolds changes the uptake pathway of the  $\beta$ -lactam warhead from passive diffusion through porins to TonB-powered active transport, which results in acceleration of killing for Ent/DGE-Amp 1, 2. Consequently, Ent/DGE-Amp/Mem 1, 2, 5, and 6 reduce the OD<sub>600</sub> of CFT073 to almost baseline within 1–2 h (Fig. 5A and B). Hence, these results suggest that the rate of bacterial cell killing will be accelerated to a greater extent for more lipophilic  $\beta$ -lactams such as Amp.

#### Ent/DGE- $\beta$ -lactam conjugates selectively kill *E. coli* in co-culture with *S. aureus*

To investigate species selectivity of the new Ent- $\beta$ -lactams, we performed mixed-species assays to ascertain if the conjugates selectively kill *E. coli* CFT073 in co-culture with another bacterial species. We employed phase-contrast microscopy and selected to co-culture *E. coli* CFT073 with the Gram-positive pathogen *Staphylococcus aureus* ATCC 25923. The cell shape of *E. coli* (rod) is distinct from the cell shape of *S. aureus* (coccus), which facilitates microscopic differentiation of these species. Since *S. aureus* did not grow in the modified M9 medium, we performed these assays in 50% MHB containing 100  $\mu$ M Bpy, which provides iron limitation and promotes expression of the siderophore transport machinery. The co-cultures were treated with MIC doses of the Ent/DGE- $\beta$ -lactams or parent antibiotics adjusted with respect to the cell density used in each imaging experiment. In the co-cultures treated with the unmodified antibiotics (Fig. S6 $\dagger$ ), both *E. coli* and *S. aureus* were killed. In contrast, the Ent/DGE- $\beta$ -lactam conjugates selectively killed *E. coli* as evidenced by the presence of cocci and absence of rod-shaped bacteria remaining in cultures following treatment (Fig. 6 and S7 $\dagger$ ). Together with LIVE/DEAD viability assays (*vide infra*, Fig. S8 $\dagger$ ), these results indicate that Ent/DGE conjugation narrows the AMA spectrum of Amp, Lex and Mem and

attenuates the AMA of these  $\beta$ -lactams against *S. aureus*. The latter outcome is in agreement with our previous studies on HardyCHROM™ UTI plates that showed both *E. coli* selective killing and attenuated AMA against *S. aureus* ATCC 25923 for Ent-Amp/Amx.<sup>34</sup>

#### Ent/DGE-Amp/Lex induce filamentous growth, whereas Ent/DGE-Mem cause formation of spheroplasts, in *E. coli*

The three  $\beta$ -lactam antibiotics used as cargos in this study target different class B high-molecular-weight PBPs when administered at sub-MIC doses, inducing different bacterial cell shapes as a result of inhibition of the corresponding PBPs.<sup>53,54</sup> Amp and Lex primarily bind to PBP3 that participates in cell division,<sup>93–96</sup> whereas Mem targets PBP2 that participates in cell elongation.<sup>93,97,98</sup> Consequently, at sub-MIC quantities, Amp and Lex induce bacterial cell filamentation, whereas Mem induces the formation of spheroplasts, which are CW-deficient cells. At quantities equal to or greater than the MIC, all three  $\beta$ -lactams induce cell lysis due to concomitant inhibition of multiple PBPs.<sup>54,99,100</sup> To further examine the consequences of Ent- and DGE- $\beta$ -lactam treatment on *E. coli* and thereby inform mechanism, we employed phase-contrast microscopy to visualize the cellular morphologies caused by these conjugates (Fig. 7).

*E. coli* CFT073 and its *fepA iron* and *fepA iron tha* mutants were incubated with the Ent-Amp/Lex/Mem conjugates 1, 3, 5 at a high cell density ( $\sim 10^8$  CFU mL<sup>-1</sup>) to facilitate microscopic visualization of the bacterial cells. To probe the effect of the conjugates at sub-MIC quantities, the bacteria were treated with 50  $\mu$ M of Ent-Amp/Mem 1 and 5 and 100  $\mu$ M of Ent-Lex 3. Significant cell elongation was observed for *E. coli* CFT073 treated with Ent-Amp 1 and Ent-Lex 3 (Fig. 7B and C) compared to the untreated control (Fig. 7A). In contrast, *E. coli* CFT073 cells treated with 50  $\mu$ M Ent-Mem 5 transformed into round or ovoid-shaped spheroplasts (Fig. 7D). These observations are consistent with the inhibition of the primary PBP targets of the three  $\beta$ -lactams. Treatment of the double and triple mutants *fepA iron* and *fepA iron tha* with sub-MIC concentrations of the Ent- $\beta$ -lactam conjugates showed that some of the cells exhibit slight elongation following exposure to Ent-Amp/Lex 1, 3 (Fig. 7F, G, J and K) and that some cells form spheroplasts following exposure to Ent-Mem 5 (Fig. 7H and L). The observation of these morphologies is consistent with the weak AMA



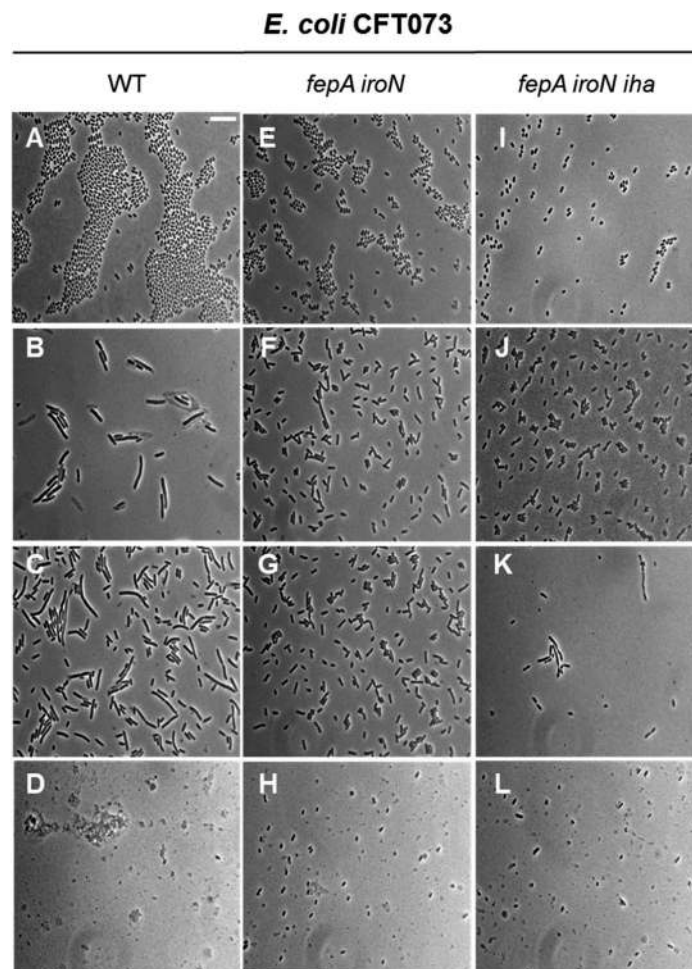


Fig. 7 Full-field view micrographs of *E. coli* CFT073 WT (A–D) and the indicated outer-membrane receptor mutants (E–L) treated with sub-MIC amounts of Ent- $\beta$ -lactams in modified M9 medium for 20 h at 30 °C. Scale bar: 10  $\mu$ m. The micrographs of *E. coli* and outer-membrane receptor mutants treated with sub-MIC amounts of DGE- $\beta$ -lactams are shown in Fig. S9.†

observed for these mutants in the AMA assays (Fig. 4), which might suggest the existence of alternative pathways of Ent- $\beta$ -lactam uptake in *E. coli* CFT073.

When the DGE- $\beta$ -lactam conjugates 2, 4, and 6 were added to bacteria at sub-MIC doses, similar morphologies were observed

as for the cells treated with Ent- $\beta$ -lactams 1, 3, and 5 (Fig. S9†). Thus, the  $\beta$ -lactam warhead induces specific morphological changes to the bacterial cell shape independent of the carrier to which it is attached. Curiously, the triple-mutant *fepA iroN iha* appears to be more sensitive to the action of DGE- $\beta$ -lactams (Fig. S9J–L†) than to the corresponding Ent- $\beta$ -lactams (Fig. 7J–L). Further investigation into this finding may uncover additional mechanisms of Ent/DGE uptake and better explain the enhanced sensitivity of *E. coli* CFT073 to the Ent/DGE conjugates.

Next, we treated *E. coli* CFT073 with an estimated MIC dose of Ent-Amp or DGE-Amp (100  $\mu$ M) determined by extrapolating to  $\sim 10^8$  CFU mL<sup>-1</sup> of the MIC values obtained in the AMA assays. This concentration of the conjugates resulted in cell lysis, indicated by the cell debris observed in the micrographs (Fig. 8B and C). Moreover, the presence of enlarged cells in the bacterial cultures treated with Ent/DGE-Amp 1, 2 (Fig. 8B and C) is consistent with the concomitant inhibition of multiple essential PBPs by Amp at doses that are equal or exceeding the MIC.<sup>54</sup> Similar cell shapes and debris were observed at MIC doses of Ent/DGE-Lex 3, 4 and Ent/DGE-Mem 5, 6 (Fig. S10†).

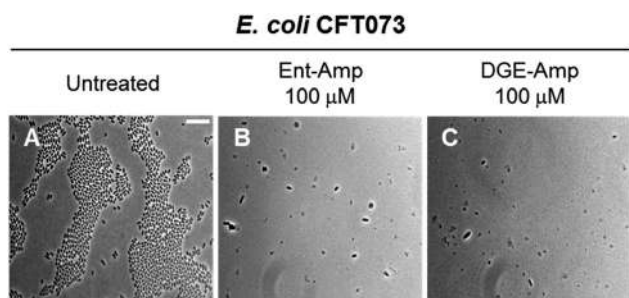
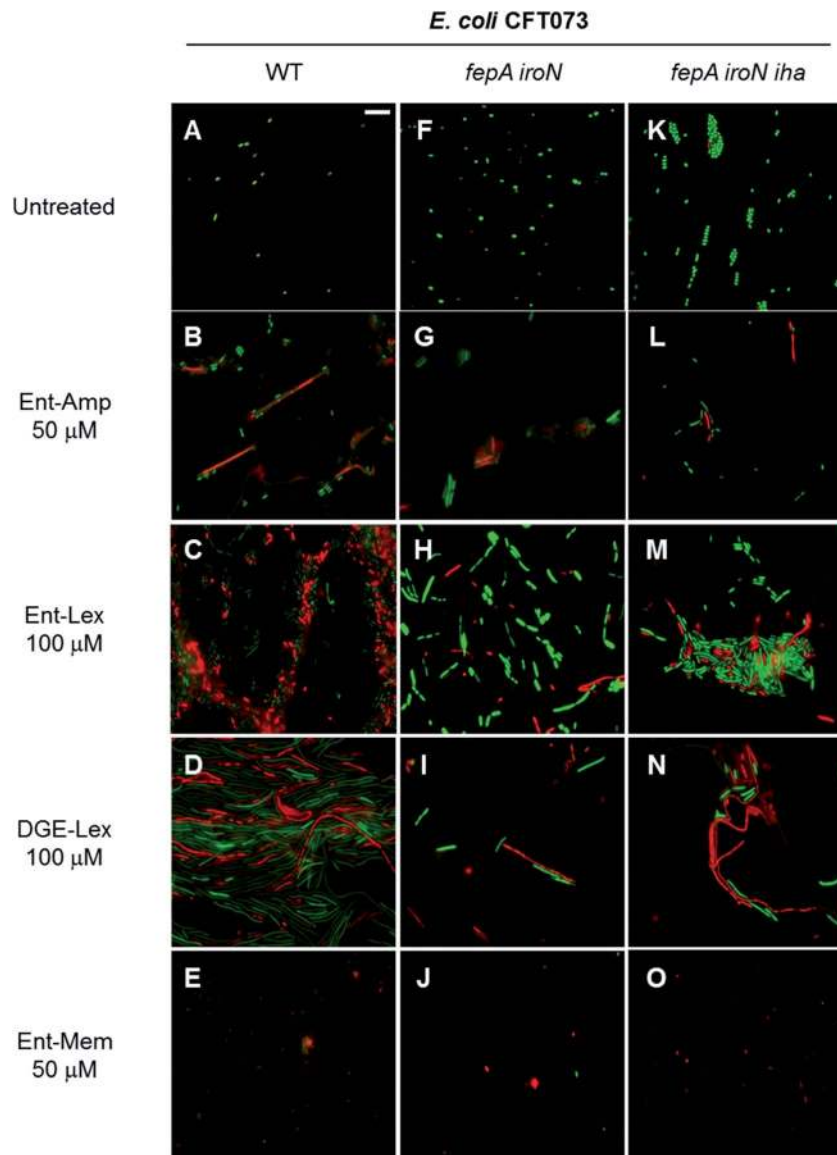


Fig. 8 Full-field view micrographs of *E. coli* CFT073 WT (A–C) treated with MIC amounts of Ent/DGE-Amp 1, 2 in modified M9 medium for 20 h at 30 °C. Scale bar: 10  $\mu$ m. The micrographs of *E. coli* CFT073 treated with MIC amounts of Ent/DGE-Lex/Mem 3–6 are shown in Fig. S10.†







**Fig. 9** Fluorescence micrographs acquired after incubating *E. coli* CFT073 and the *fepA iroN* and *fepA iroN iha* mutants with the LIVE/DEAD dyes at 30 °C for 15 min. The cell cultures were incubated with the Ent/DGE conjugates in modified M9 for 20 h at 30 °C prior to staining. Scale bar: 10  $\mu$ m. The micrographs acquired after incubation of DGE-Amp/Mem are shown in Fig. S11.†

Overall, phase-contrast microscopy imaging corroborated the AMA assays and illuminated the differences in cell morphologies due to variable affinities of the  $\beta$ -lactam warheads towards PBP2 and PBP3. It should be noted, however, that sub-MIC doses of  $\beta$ -lactam antibiotics can also inhibit nonessential PBPs (e.g. PBP4)<sup>101</sup> in Gram-positive and Gram-negative bacterial strains.<sup>54,101–105</sup> Inhibition of nonessential PBPs leads to more subtle morphological effects than inhibition of PBP1, 2 and 3;<sup>106–109</sup> thus, further studies are needed to elucidate whether the Ent/DGE- $\beta$ -lactam conjugates also inhibit nonessential PBPs similar to the parent  $\beta$ -lactams.

#### Inhibition of PBP2 and PBP3 by the Ent/DGE- $\beta$ -lactam conjugates concludes with cell lysis

Next, we employed the LIVE/DEAD viability assay to examine the fate of the bacterial cell filaments and spheroplasts formed as

result of Ent/DGE- $\beta$ -lactam treatment. This assay utilizes fluorescent dyes to distinguish between cells with intact OM (stain green with SYTO 9) and cells with compromised OM (stain red with propidium iodide, PI).<sup>110</sup>

As expected, elongation/filamentation was detected in all strains treated with sub-MIC quantities of the PBP3 inhibitors Ent-Amp/Lex 1, 3 and DGE-Amp/Lex 2, 4 (Fig. 9B–D and S11B†), whereas the PBP2 inhibitors Ent/DGE-Mem 5, 6 led to spheroplast formation (Fig. 9E and S11C†). Similarly to the phase-contrast microscopy and the AMA assays, filamentation (Fig. 9G–I, L–N and S11E, H†) and spheroplasts formation (Fig. 9J, O and S11F, I†) were also observed, to different degrees, in the *fepA iroN* and *fepA iroN iha* mutants.

All untreated cells stained green with SYTO 9, indicating viability (Fig. 9A, F and K). In contrast, addition of sub-MIC quantities of the conjugates lead to a mixture of green and



red cells (Fig. 9B–E, G–J, L–O and S11†). The latter indicated that cell lysis occurred for a subset of the cells under these conditions. Thus, after undergoing morphological change as determined by inhibition of either PBP3 (Ent-Amp/Lex 1, 3 and Ent/DGE-Lex 2, 4) (Fig. 9B–D, G–I, L–N and S11B†) or PBP2 (Ent/DGE-Mem 5, 6) (Fig. 9E, J, O and S11C†), the bacterial cells lose OM integrity and can be considered dead based on the PI staining. Lastly, the partial red staining of the *fepA iroN* (Fig. 9G–J) and *fepA iroN iha* (Fig. 9L–O) cells is another indication that the conjugates exert some AMA against these cells and are likely transported into the periplasm.

Overall, the LIVE/DEAD assay showed that the treatment of *E. coli* CFT073 and its mutants with sub-MIC quantities of conjugates leads to the death of both filamentous and CW-deficient cells as result of OM rupture. Therefore, the  $\beta$ -lactam warhead is responsible for the bactericidal effect of the Ent/DGE- $\beta$ -lactam conjugates 1–6 as it inhibits the target PBPs causing characteristic phenotypes and compromises OM integrity conducive to cell lysis.

### Ent-Mem induces explosive cell lysis in *E. coli*

To obtain a visual depiction of the course of cell killing by the Ent/DGE conjugates, we added MIC amounts of Ent/DGE-Amp 1, 2 and Ent-Mem 5 to *E. coli* CFT073 and acquired time-lapse images of the process at 37 °C. The Ent/DGE-Lex 3, 4 conjugates were not studied due to their relatively high MIC values. The images of CFT073 cells treated with 100  $\mu$ M Ent/DGE-Amp 1, 2 show that the cells ceased to grow and divide. Moreover, these cells did not elongate or undergo explosive lysis over the course of 4 h. These observations indicate that non-explosive rupture of the cell membranes occurs and results in cell lysis without bursting (Fig. 10) similar to the red-stained cells in the LIVE/DEAD assay (Fig. 9).

In contrast to Ent/DGE-Amp 1, 2, the treatment of *E. coli* CFT073 with 100  $\mu$ M Ent-Mem 3 (Fig. 10) is accompanied by explosive lysis that starts within the first 60 min and is virtually complete by 4 h. Thus, targeting of different primary PBPs by the Ent/DGE- $\beta$ -lactam conjugates determines not only the morphologies of the cells treated with sub-MIC doses, but also the character of lysis when MIC amounts are reached.

### Perspectives on siderophore- $\beta$ -lactam conjugates

The growing problem of antibiotic resistance and the insufficient amount of new antibiotics in the drug pipeline pose a serious threat to public health. Development of narrow-spectrum antibiotics that kill pathogenic species and spare commensal microbes is considered a promising solution in the face of infections caused by multidrug-resistant Gram-negative bacteria. Leveraging the essential nutrient uptake machineries of such bacteria for a targeted delivery of antibiotics has been reported for a number of different compounds.<sup>111–113</sup> Recently, FDA approved the first antibiotic that is reported to hijack Fe<sup>3+</sup> uptake machinery to deliver a cephalosporin cargo to the periplasm of Gram-negative bacteria in the treatment of complicated urinary tract infections.<sup>114–116</sup>

Our approach consists of using the native siderophore scaffolds Ent and DGE as antibiotic carriers.<sup>34–36</sup> The Ent/DGE-Lex/Mem conjugates synthesized and studied in this work expand the arsenal of clinically-approved broad-spectrum antibiotics that can be endowed with selectivity towards pathogenic strains that express Iron. This selectivity is achieved by targeting the enterobactin/salmochelin uptake machinery of *E. coli* and is not affected by the  $\beta$ -lactam structure. Moreover, conjugation to Ent/DGE further enhances the MIC of the parent  $\beta$ -lactams by 10–1000-fold. The time-kill kinetics of the conjugated  $\beta$ -lactams is enhanced to a greater extent for the more lipophilic antibiotics

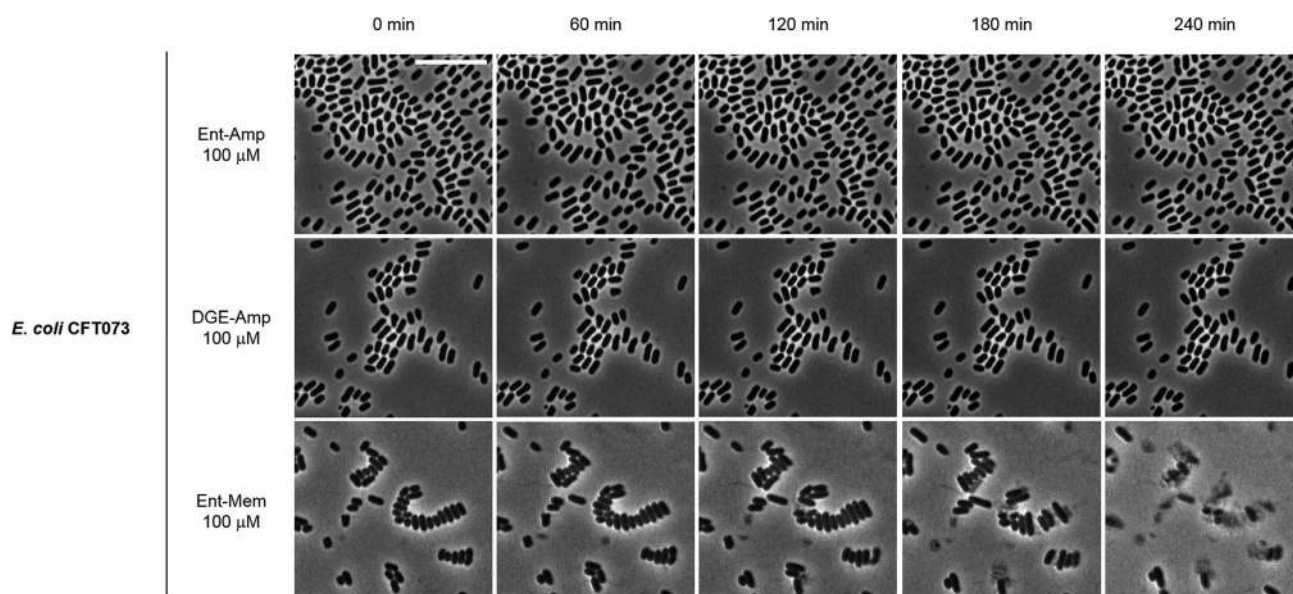


Fig. 10 Time-lapse phase-contrast micrographs of *E. coli* CFT073 incubated with 100  $\mu$ M Ent/DGE-Amp 1, 2 and Ent-Mem 5 in modified M9 medium (37 °C). Scale bar: 10  $\mu$ m.



such as Amp due to the change of the uptake route from porins (Amp) to TonB-powered FepA or IroN (Ent-Amp and DGE-Amp). These results indicate that the Ent/DGE scaffold could serve as a versatile tool for pharmaceutical development of antibacterial therapeutics that target IroN-expressing strains (e.g. DGE conjugates), enhancing the MIC values and decreasing the time needed to exert its bactericidal effect.

The three  $\beta$ -lactams, Amp, Lex, and Mem, conjugated to the Ent/DGE scaffolds in this work, exert AMA by inhibiting class B high-molecular-weight PBP2 and PBP3 as indicated by the phase-contrast microscopy and LIVE/DEAD staining assays. These results along with the AMA assays of the OM receptor mutants demonstrated that the  $\beta$ -lactam cargo is indeed responsible for the killing of bacteria while the uptake of the Ent/DGE conjugates with different  $\beta$ -lactam cargos takes place via FepA and IroN. In prior work, we hypothesized that Iha may contribute to the sensitivity of *E. coli* CFT073 to the Ent/DGE- $\beta$ -lactams. However, the AMA assays and microscopy study of the *fepA iroN* and *fepA iroN iha* mutants indicated that Iha does not play an important role in the transport of either Ent- or DGE- $\beta$ -lactams. Nevertheless, we have observed weak AMA and signs of uptake of the conjugates by the *fepA iroN* and the *fepA iroN iha* mutants, which suggests that CFT073 possesses other yet-unknown routes of Ent/DGE uptake.

Although uptake through the OM siderophore receptors allows the  $\beta$ -lactam cargos of the Ent/DGE conjugates to exert antibacterial activity, there may be additional mechanisms in place that explain the enhancement of the MIC values of conjugates relative to parent antibiotics. For instance, it has been shown that accumulation of apo Ent in the periplasm of *E. coli* leads to deleterious effects for the bacterial cells.<sup>117</sup> However, under the assay conditions employed in this work, we did not observe any of the abnormal morphologies that were attributed to the accumulation of Ent in the bacterial periplasm.<sup>117</sup>

Of the four  $\beta$ -lactam cargos we have thus far conjugated to Ent/DGE – Amp, Amx,<sup>34,35</sup> Lex, and Mem – Mem seems to be the most promising for further evaluation. The exceptional stability of the carbapenem ring to the action of extended spectrum  $\beta$ -lactamases, its enhanced MIC upon conjugation to Ent/DGE, and induction of explosive cell lysis at nanomolar concentrations make Mem an outstanding warhead for siderophore conjugation. However, use of Mem as warhead could be limited by the spread of carbapenemase-producing strains, which render Mem inactive.<sup>80,82</sup> Likewise, the selectivity afforded by conjugation of antibiotics to Ent/DGE could be compromised by the existence of commensals that can cross-feed using Ent/DGE as xenosiderophores.<sup>118</sup> Therefore, further studies are needed to better understand the host microbiome and evaluate the potential of Ent/DGE antibiotic conjugates as pathogen-selective therapeutics.

## Conclusion and outlook

In closing, this work further demonstrates that conjugation of  $\beta$ -lactam antibiotics to the Ent/DGE scaffolds is an effective strategy to convert broad-spectrum antibiotics into compounds that selectively target bacteria expressing the catecholate

siderophore receptors FepA and IroN. Furthermore, the siderophore- $\beta$ -lactam conjugates presented in this work efficiently kill pathogenic *E. coli* regardless of the structure and polarity of the  $\beta$ -lactam cargo. Targeting pathogenesis with such conjugates could reduce the onset of secondary infections and slow the evolution of antibiotic resistance. In future work, studies using mouse models will aid the understanding of the stability and efficacy of such siderophore- $\beta$ -lactam conjugates *in vivo*. Lastly, it will be important to determine whether this strategy is applicable to other bacterial pathogens such as *Salmonella enterica* that cause human disease and utilize both Ent and DGE for  $\text{Fe}^{3+}$  acquisition.

## Experimental section

### Synthetic reagents

Anhydrous *N,N*-dimethylformamide (DMF) and dichloromethane (DCM) were purchased from VWR. Anhydrous dimethyl sulfoxide (DMSO) was purchased from Millipore Sigma and used as received. All other chemicals were purchased from Millipore Sigma or VWR and used as received. The syntheses of Ent-PEG<sub>3</sub>-N<sub>3</sub> **10**,<sup>34,35,119</sup> DGE-PEG<sub>3</sub>-N<sub>3</sub> **11**,<sup>35</sup> Amp-alkyne **7**,<sup>34,35</sup> Ent-Amp **1**,<sup>34,35</sup> and DGE-Amp **2** (ref. 35) are reported elsewhere.

### General methods and instrumentation

EMD TLC silica gel 60 F<sub>254</sub> plates were used for analytical thin layer chromatography. EMD PLC silica gel 60 F<sub>254</sub> plates of 2 mm thickness were used for preparative TLC. Sigma-Aldrich silica gel (70–230 mesh, 60 Å) was used for flash column chromatography.

<sup>1</sup>H NMR spectra were collected on a Bruker Avance III DPX 400 MHz spectrometer operated at ambient probe temperature (293 K). Q-LC/MS and Q-ToF MS analyses were carried out on an Agilent 6125B mass spectrometer attached to an Agilent 1260 Infinity LC and a high-resolution Agilent 6545 mass spectrometer, respectively. For all LC/MS analyses, solvent A was 0.1% formic acid/H<sub>2</sub>O and solvent B was 0.1% formic acid/MeCN (LC/MS grade, Millipore Sigma). The samples were analyzed using a solvent gradient of 5–95% B over 6 min with a flow rate of 0.4 mL min<sup>-1</sup>.

Analytical and semi-preparative HPLC were performed using an Agilent 1200 series HPLC system outfitted with a Cliepus reversed-phase C18 column (5  $\mu$ m pore size, 4.6  $\times$  250 mm, Higgins Analytical) at a flow rate of 1 mL min<sup>-1</sup> and an Agilent Zorbax reversed-phase C18 column (5  $\mu$ m pore size, 9.4  $\times$  250 mm) at a flow rate of 4 mL min<sup>-1</sup>. The multi-wavelength detector was set to detect absorbance at 220, 280, and 316 nm. HPLC-grade acetonitrile (MeCN) and trifluoroacetic acid (TFA) were purchased from Millipore Sigma. For HPLC analysis, solvent A was 0.1% v/v TFA/H<sub>2</sub>O and solvent B was 0.1% v/v TFA/MeCN. The percentage of TFA in the HPLC solvents used for antibiotic conjugate purification was 0.005% to avoid  $\beta$ -lactam ring hydrolysis. The HPLC solvents were prepared with HPLC grade MeCN and TFA, with Milli-Q water (18.2 M $\Omega$  cm), and filtered through a 0.2  $\mu$ m filter before use. To evaluate conjugate purity by analytical HPLC, the entire portion



of each HPLC-purified compound was dissolved in a 1 : 1 mixture of MeCN/H<sub>2</sub>O and an aliquot was taken for HPLC analysis. The remaining solution was subsequently lyophilized.

Optical absorption spectra were recorded on a Beckman Coulter DU800 spectrophotometer (1 cm quartz cuvettes, Starna). A BioTek Synergy HT plate reader was used to record absorbance at 600 nm (OD<sub>600</sub>) for antimicrobial activity assays and time-kill kinetics.

### Synthesis of Lex-alkyne, 8

Hex-5-ynoyl chloride (0.352 g, 2.70 mmol) was added dropwise to a 0 °C solution of Lex sodium salt (0.500 g, 1.35 mmol) and NaHCO<sub>3</sub> (0.567 g, 6.75 mmol) in 6.75 mL of water under vigorous stirring. The reaction mixture was warmed to room temperature and left to react for 1 h under stirring. Then, 5 mL of water was added to the solution and the resulting mixture was washed with EtOAc (2 × 10 mL). The aqueous layer was acidified to pH 2 using 1 M HCl and the product extracted with EtOAc (3 × 10 mL). The organic layer was dried over anhydrous Na<sub>2</sub>SO<sub>4</sub> and the solvent was removed under reduced pressure to yield a yellow solid. The crude product was triturated with hexane, then purified by semi-preparative HPLC (0–100% B over 30 min, 4 mL min<sup>-1</sup>; 0.005% TFA). Lyophilization of the HPLC fractions afforded 0.390 g (65%) of Lex-alkyne, 8, as an off-white powder. Analytical HPLC *R*<sub>t</sub> = 17.5 min (0–100% B over 30 min, 1 mL min<sup>-1</sup>; 0.005% TFA). <sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>, 400 MHz): δ, ppm 13.20 (br, 1H), 9.25 (d, *J* = 8.5 Hz, 1H), 8.56 (d, *J* = 8.1 Hz, 1H), 7.44 (d, *J* = 7.6 Hz, 2H), 7.25–7.36 (m, 3H), 5.68 (d, *J* = 8.2 Hz, 1H), 5.62 (dd, *J* = 4.7, 8.2 Hz, 1H), 4.96 (d, *J* = 4.7 Hz, 1H), 3.47 (d, *J* = 18.9 Hz, 1H), 3.28 (d, *J* = 18.2 Hz, 1H), 2.78 (s, 1H), 2.31 (t, *J* = 7.3 Hz, 2H), 2.13–2.22 (m, 2H), 1.99 (s, 3H), 1.67 (m, 2H). QToF-MS (*m/z*): mass calcd for (C<sub>22</sub>H<sub>23</sub>N<sub>3</sub>O<sub>5</sub>S + Na<sup>+</sup>) = 464.1256; found 464.1253.

### Synthesis of Mem-alkyne, 9

Hex-5-ynoyl chloride (0.101 g, 0.78 mmol) was added dropwise to a 0 °C solution of Mem (0.150 g, 0.39 mmol) and Et<sub>3</sub>N (1.522 mL, 1.105 g, 10.92 mmol) in 4 mL of MeOH under vigorous stirring. The reaction mixture was warmed to room temperature and left to react for 1 h under stirring. Then, the solvent was removed under reduced pressure and the crude was purified by semi-preparative HPLC (0–100% B over 30 min, 4 mL min<sup>-1</sup>; 0.005% TFA). Lyophilization of the HPLC fractions containing product afforded 0.093 g (50%) of Mem-alkyne, 9, as a yellow powder. Analytical HPLC *R*<sub>t</sub> = 14.0 min (0–100% B over 30 min, 1 mL min<sup>-1</sup>; 0.005% TFA). The <sup>1</sup>H NMR (CD<sub>3</sub>OD, 400 MHz): δ, ppm 4.24 (t, *J* = 9.1 Hz, 1H), 4.07–4.17 (m, 1H), 3.82–3.92 (m, 1H), 3.53–3.64 (m, 1H), 3.48 (t, *J* = 10.2 Hz, 1H), 3.32 (dd, *J* = 2.7, 7.1 Hz, 2H), 3.19 (s, 3H), 3.05 (s, 1H), 2.98 (s, 3H), 2.73–2.84 (m, 1H), 2.33–2.60 (m, 2H), 2.24–2.31 (m, 4H), 1.77–1.87 (m, 2H), 1.31 (d, *J* = 6.7 Hz, 3H), 1.28 (d, *J* = 7.2 Hz, 3H). QToF-MS (*m/z*): mass calcd for (C<sub>23</sub>H<sub>31</sub>N<sub>3</sub>O<sub>6</sub>S + H<sup>+</sup>) = 478.2012; found 478.2010.

### General synthesis of Ent/DGE-Amp, Ent/DGE-Lex, Ent/DGE-Mem (1–6)

β-lactam alkyne 7–9 (50 μL of a 50 mM solution in DMSO, 2.5 μmol) and Ent/DGE-PEG<sub>3</sub>-N<sub>3</sub> 10, 11 (73 μL of an 11.3 mM

solution in DMSO, 0.825 μmol) were combined and 100 μL of DMSO was added. An aliquot of aqueous CuSO<sub>4</sub> (50 μL of a 90 mM solution, 4.5 μmol) and TBTA (100 μL of a 50 mM solution in DMSO, 5 μmol) were combined to give a blue solution, to which sodium ascorbate (NaAsc, 100 μL of a 180 mM solution in water, 18.0 μmol) was added. The color of the solution immediately changed from blue to pale yellow (reduction of Cu<sup>2+</sup> to Cu<sup>+</sup>) and the mixture was added to the alkyne/azide solution. The reaction was gently mixed on a bench-top rotator for 2.5 h at 25 °C (150 rpm). Then, the reaction mixture was diluted with 1 : 1 MeCN/water (3× the volume of solution) and purified by semi-preparative HPLC (0–100% B over 30 min, 4 mL min<sup>-1</sup>; 0.005% TFA). Lyophilization of the HPLC fractions afforded the corresponding conjugates (1–6) as white or off-white powders. Analytical data and the yields of the Ent/DGE conjugates are provided in Table S3.†

### Storage and handling of siderophores and siderophore-β-lactam antibiotic conjugates

All precursors and Ent/DGE conjugates were stored as either powders or DMSO stock solutions at –20 °C. The stock solution concentrations for the Ent/DGE conjugates ranged between 1–2 mM. These values were determined by diluting the DMSO stocks in MeOH and using the reported extinction coefficient for enterobactin in MeOH ( $\epsilon_{316} = 9500 \text{ M}^{-1} \text{ cm}^{-1}$ ).<sup>39</sup> To minimize multiple freeze-thaw cycles, the resulting solutions were divided into 50 μL aliquots and stored at –20 °C. The β-lactam antibiotic cargos as well as Ent/DGE are prone to hydrolysis, and aliquots were routinely analyzed by HPLC to confirm the integrity of the samples.

### General microbiology materials and methods

Information pertaining to all bacterial strains used in this study is listed in Table S1.† Freezer stocks of all *E. coli* strains were prepared from single colonies in 25% glycerol/lysogeny broth (LB) medium. Lysogeny broth (tryptone 10 g L<sup>-1</sup>, yeast extract 5 g L<sup>-1</sup>, NaCl 10 g L<sup>-1</sup>), modified M9 (Na<sub>2</sub>HPO<sub>4</sub> 6.8 g L<sup>-1</sup>, KH<sub>2</sub>PO<sub>4</sub> 3 g L<sup>-1</sup>, NaCl 0.5 g L<sup>-1</sup>, NH<sub>4</sub>Cl 1 g L<sup>-1</sup>, 0.4% glucose, 0.2% casein amino acids, 2 mM MgSO<sub>4</sub>, 0.1 mM CaCl<sub>2</sub>, 16.5 μg mL<sup>-1</sup> thiamine hydrochloride)<sup>36</sup> and agar were obtained from Becton Dickinson (BD). Mueller–Hinton broth (MHB; beef extract powder 2 g L<sup>-1</sup>, casein hydrolysate 17.5 g L<sup>-1</sup>, and soluble starch 1.5 g L<sup>-1</sup>) was purchased from Sigma-Aldrich. The iron chelator 2,2'-bipyridine (Bpy) was purchased from Sigma-Aldrich. All growth media and Milli-Q water used for bacterial cultures or for preparing stock solutions were sterilized in an autoclave. The modified M9 medium was filter-sterilized through a 0.22 μm filter. A 200 mM Bpy stock solution was prepared in anhydrous DMSO and utilized in bacterial growth inhibition assays. Working solutions of Ent and Ent/DGE-antibiotic conjugates were prepared by 10-fold serial dilutions in 10% DMSO/H<sub>2</sub>O. For all microbiology assays, the final cultures contained 1% v/v DMSO. Sterile polypropylene culture tubes and sterile polystyrene 96-well plates used for culturing were purchased from VWR and Corning Inc., respectively. The optical density at 600 nm (OD<sub>600</sub>) was recorded on



a Beckmann Coulter DU800 spectrophotometer or by using a BioTek Synergy HT plate reader.

### General procedure for antimicrobial activity (AMA) assays

Overnight bacterial cultures were prepared in 15 mL polypropylene tubes by inoculating 5 mL of growth medium (modified M9) with the appropriate freezer stock. The overnight cultures were then incubated at 37 °C for 16–18 h on a rotating wheel set at 150 rpm. Each overnight culture was diluted in a v/v ratio of 1 : 100 into 5 mL of fresh modified M9 medium and incubated at 37 °C and 150 rpm until OD<sub>600</sub> reached ~0.6 (mid-log phase). Each culture was subsequently diluted into fresh modified M9 to achieve a final OD<sub>600</sub> of 0.001. A 90 µL aliquot of the diluted culture was combined with a 10 µL aliquot of a 10× solution of the antibiotic or antibiotic conjugate in a 96-well plate. The covered plate was wrapped in a wet paper towel and Saran wrap, then incubated at 30 °C with shaking at 150 rpm for 20 h in a tabletop incubator. Bacterial growth was determined by measuring OD<sub>600</sub> (end point analysis) on a BioTek Synergy HT plate reader. Each well condition was prepared in duplicate (technical replicate) and at least three independent replicates (biological replicate) utilizing at least two synthetic batches of each conjugate were conducted on different days. The resulting mean OD<sub>600</sub> values are reported, and the error bars are the standard deviation of the mean (SDM) obtained from the biological replicates.

### Time-kill kinetics

Overnight cultures were prepared by inoculating 5 mL of modified M9 medium with bacterial freezer stocks. The overnight cultures were diluted 1 : 100 into 5 mL of fresh modified M9 medium and incubated at 37 °C with shaking at 150 rpm until OD<sub>600</sub> reached ~0.3. The culture was centrifuged (3000 rpm × 10 min) and the resulting pellet was re-suspended in modified M9 and centrifuged (3000 rpm × 10 min). The resulting pellet was re-suspended again in fresh modified M9 and the OD<sub>600</sub> adjusted to 0.3. A 90 µL aliquot of the resulting culture was combined with a 10 µL aliquot of a 10× solution of parent antibiotic or Ent/DGE conjugate in a 96 well plate, which was wrapped in Saran film and incubated at 37 °C with shaking at 150 rpm. The OD<sub>600</sub> values were recorded at 1 h intervals. Each well condition was prepared in duplicate (technical replicate) and at least three independent replicates (biological replicate) utilizing at least two synthetic batches of each conjugate were conducted on different days. The resulting mean OD<sub>600</sub> values are reported, and the error bars are the standard deviation of the mean (SDM) obtained from the biological replicates.

### Calculation of Ent/DGE-β-lactam MIC for microscopy assays

The MIC values (Table S4†) of the Ent/DGE-Amp/Lex/Mem conjugates 1–6 were determined in modified M9 medium for mid-log phase (OD<sub>600</sub> ~ 0.6) cell cultures diluted to OD<sub>600</sub> = 0.001 (*vide supra*). Plating of the diluted cultures onto LB agar plates gave an average count of  $5 \times 10^4$  CFU mL<sup>-1</sup> ( $n > 3$ ). Thus, at  $5 \times 10^4$  CFU mL<sup>-1</sup>, the MIC of Ent/DGE-Amp/Mem 1, 2, 5,

and 6 for *E. coli* CFT073 is equal to  $1 \times 10^{-8}$  M (Table S4†). For microscopy experiments,  $5 \times 10^8$  CFU mL<sup>-1</sup> of mid-log phase CFT073 cultures were incubated with the Ent/DGE-β-lactams 1, 2, 5, and 6. Hence, the MIC of Ent/DGE-Amp/Mem 1, 2, 5, and 6 per  $5 \times 10^8$  CFU mL<sup>-1</sup> was estimated to be 100 µM. No cells were detected after incubating  $5 \times 10^8$  CFU mL<sup>-1</sup> of *E. coli* CFT073 with >100 µM of the Ent/DGE-Amp/Mem 1, 2, 5, and 6. For Ent/DGE-Lex 3 and 4, MIC =  $1 \times 10^{-5}$  M per  $5 \times 10^4$  CFU mL<sup>-1</sup>. To observe filamentous growth, 100 µM of Ent/DGE-Lex 3, 4 was incubated per  $5 \times 10^6$  CFU mL<sup>-1</sup> (sub-MIC).

### Phase-contrast microscopy

Phase-contrast microscopy imaging was carried out at the W. M. Keck Biological Imaging Facility of the Whitehead Institute (Cambridge, Massachusetts). The incubation of cell cultures with the Ent/DGE-β-lactams was performed on 96-well plates (100 µL sample per well) using  $5 \times 10^8$  CFU mL<sup>-1</sup> and 50 or 100 µM of Ent/DGE conjugates. For the Ent/DGE-Lex conjugates, the initial cell density was  $5 \times 10^6$  CFU mL<sup>-1</sup> to account for the higher MIC values of the Lex conjugates and use less compound. Following a 20 h incubation (30 °C, 150 rpm), a 5 µL sample from each culture was pipetted on an agarose (Bio-Rad, PCR grade) pad (1% w/w agarose/Milli-Q water) positioned on a microscope slide. The sample was then covered with a coverslip and imaged using a Zeiss Axioplan2 upright microscope equipped with a 100× oil-immersion objective lens. For each type of microscopy experiment, each condition was repeated in at least three independent replicates on different days. Representative micrographs for each condition are shown in the figures.

### Mixed-species microscopy assays

All bacterial strains were cultured in 50% MHB in the presence of 100 µM of 2,2'-bipyridine to an OD<sub>600</sub> ~ 0.6 (mid-log phase) (*S. aureus* ATCC 25923 does not grow in the modified M9 medium). Bacterial suspensions were then adjusted to  $1 \times 10^7$  CFU mL<sup>-1</sup>, mixed in a 1 : 1 ratio ( $10^7$  :  $10^7$  CFU mL<sup>-1</sup>) and incubated with 50 µM Ent/DGE-Amp/Mem per 100 µL sample in 96-well plates (20 h, 30 °C, 150 rpm). For Ent/DGE-Lex, bacterial suspensions were adjusted to  $1 \times 10^6$  CFU mL<sup>-1</sup> initial cell density and incubated with 200 µM Ent/DGE-Lex. A 5 µL sample from each culture was pipetted on an agarose pad (1% w/w agarose/Milli-Q water) positioned on a microscope slide. The specimen was then covered with a coverslip and imaged using a Zeiss Axioplan2 upright microscope equipped with a 100× oil-immersion objective lens.

### Time-lapse microscopy

The time-lapse studies were carried out using poly-D-lysine coated MatTek glass-bottom Petri dishes of 35 mm, with a 14 mm microwell and a No. 1.5 cover glass. The untreated bacteria ( $5 \times 10^8$  CFU mL<sup>-1</sup>) were mixed on 96-well plates (100 µL sample per well) to give a final concentration of 100 µM Ent/DGE conjugate. Once the conjugate was added to the culture, a 5 µL sample from each suspension was pipetted out onto the 14 mm microwell and covered with an agarose pad (1% w/w agarose/Milli-Q water). The Petri dish with the sample was



then immediately covered with the No. 1.5 coverslip and placed on the microscope stage for imaging at 37 °C. Image acquisition was carried out at 5 min intervals over 4.5–5 h. The time-lapse images were collected on a Nikon-TE 2000 U wide-field inverted microscope equipped with a 100× oil-immersion objective lens.

### LIVE/DEAD assays

All bacterial strains were cultured in the modified M9 growth medium up to an OD<sub>600</sub> ~ 0.6 (mid-log phase). Bacterial suspensions were then adjusted to 5 × 10<sup>8</sup> CFU mL<sup>-1</sup> and incubated with 50 or 100 μM Ent/DGE conjugates for 20 h (30 °C, 150 rpm). Subsequently, the cells were pelleted and re-suspended in 0.85% NaCl (centrifugation at 10 000g for 10 min). These NaCl-suspended cells were further incubated in 96-well plates wrapped in aluminum foil with the SYTO 9-propidium iodide (PI) dye mixture (1 : 5, 12 μM SYTO 9 : 60 μM PI) for 15 min (30 °C, 150 rpm). Following incubation, 5 μL of cell cultures were pipetted on agarose pads (1% w/w agarose/Milli-Q water) placed on microslides and covered with glass coverslip. Fluorescence microscopy images were recorded on the Zeiss Axioplan2 microscope. The Texas Red (λ<sub>Ex</sub> = 532–587 nm; λ<sub>Em</sub> = 608–683 nm) and GFP (λ<sub>Ex</sub> = 457–487 nm; λ<sub>Em</sub> = 502–538 nm) channels were utilized to obtain the images of the DEAD and LIVE cells, respectively.

### Image analysis

The microscopy images were processed using the FIJI software. For fluorescence images, fluorescence background subtraction was performed using a rolling ball method with a radius of 150 pixels. 8-bit image types were analyzed and the brightness intensity was set between 0 and 350.

## Conflicts of interest

E. M. N. and M. R. hold patents related to this work.

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