QStatin, a Selective Inhibitor of Quorum Sensing in Vibrio Species

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ABSTRACT Pathogenic Vibrio species cause diseases in diverse marine animals reared in aquaculture. Since their pathogenesis, persistence, and survival in marine environments are regulated by quorum sensing (QS), QS interference has attracted attention as a means to control these bacteria in aquatic settings. A few QS inhibitors of Vibrio species have been reported, but detailed molecular mechanisms are lacking. Here, we identified a novel, potent, and selective Vibrio QS inhibitor, named QStatin [1-(5-bromothiophene-2-sulfonyl)-1H-pyrazole], which affects Vibrio harveyi LuxR homologues, the well-conserved master transcriptional regulators for QS in Vibrio species. Crystallographic and biochemical analyses showed that QStatin binds tightly to a putative ligand-binding pocket in SmcR, the LuxR homologue in V. vulnificus, and changes the flexibility of the protein, thereby altering its transcription regulatory activity. Transcriptome analysis revealed that QStatin results in SmcR dysfunction, affecting the expression of SmcR regulon required for virulence, motility/chemotaxis, and biofilm dynamics. Notably, QStatin attenuated representative QS-regulated phenotypes in various Vibrio species, including virulence against the brine shrimp (Artemia franciscana). Together, these results provide molecular insights into the mechanism of action of an effective, sustainable QS inhibitor that is less susceptible to resistance than other antimicrobial agents and useful in controlling the virulence of Vibrio species in aquacultures.

IMPORTANCE Yields of aquaculture, such as penaeid shrimp hatcheries, are greatly affected by vibriosis, a disease caused by pathogenic Vibrio infections. Since bacterial cell-to-cell communication, known as quorum sensing (QS), regulates pathogenesis of Vibrio species in marine environments, QS inhibitors have attracted attention as alternatives to conventional antibiotics in aquatic settings. Here, we used target-based high-throughput screening to identify QStatin, a potent and selective inhibitor of V. harveyi LuxR homologues, which are well-conserved master QS regulators in Vibrio species. Structural and biochemical analyses revealed that QStatin binds tightly to a putative ligand-binding pocket on SmcR, the LuxR homologue in V. vulnificus, and affects expression of QS-regulated genes. Remarkably, QStatin attenuated diverse QS-regulated phenotypes in various Vibrio species, including pathogenesis against brine shrimp, with no impact on bacterial viability. Taken together, the results suggest that QStatin may be a sustainable antivibriosis agent useful in aquacultures.

KEYWORDS LuxR, Vibrio, quorum sensing, quorum-sensing inhibitor

Vibrio species are metabolically versatile and abundant in diverse marine environments. As commensals or mutualistic symbionts, they commonly live in or on a wide range of marine organisms, including corals, zooplankton, crustaceans, shellfish,
and fish (1–3). However, they also cause infectious diseases, especially in marine animals reared in aquaculture. For example, *Vibrio harveyi* causes luminescent vibriosis in shrimp and lobsters; *V. vulnificus*, *V. anguillarum*, and *V. alginolyticus* infect several fish species; and *V. crassostreae* infects oysters (4, 5). Therefore, control of pathogenic *Vibrio* species is critical for the aquaculture industry. Antibiotics have been extensively used for this purpose but have associated resistance problems. Consequently, more-sustainable alternatives that control bacterial virulence, without directly affecting bacterial viability, have attracted attention (6, 7).

Bacterial cell-to-cell communication (known as quorum sensing [QS]) makes individual cells enter “population mode” by synchronizing gene expression according to cell density. To monitor their population, each cell produces, secretes, and detects diffusible signaling molecules, called autoinducers (AIs) (8). One of the best-characterized QS systems is present in the squid symbiont *V. fischeri*, whose QS controls luminescence. In this bacterium, LuxI produces AIs, whereas LuxR (here referred to as *V. fischeri* LuxR [LuxR<sub>vfn</sub>]), a cytosolic transcriptional regulator protein, senses them directly (9). In addition to bioluminescence, diverse bacterial phenotypes requiring group cooperation such as sporulation, competence, biofilm formation, and resistance to bacteriophage are regulated by QS. Notably, pathogenesis-associated genes in many human pathogens causing chronic diseases, such as *Pseudomonas aeruginosa* and *Staphylococcus aureus*, are also controlled by QS (10). In pathogenic *Vibrio* species, QS regulates production of exoprotease/metalloprotease that causes severe diseases in marine animal hosts (11–16). Consistent with this, QS inhibition in *V. harveyi, V. campbellii, V. para-haemolyticus*, and *V. vulnificus* results in attenuated virulence against an aquatic model host, brine shrimp (17–19). Furthermore, QS provides grazing resistance and starvation-adaptation ability to *Vibrio* species and thus supports their persistence and survival in marine environments (20, 21). Accordingly, anti-QS strategies have been explored as a way to counteract the activity of pathogenic vibrios, as well as that of other chronic human pathogens. However, to our knowledge, the precise molecular mechanisms of *Vibrio* QS inhibitors remain unknown, which limits the application of this type of strategy in the field (18, 22, 23).

Although the “LuxI-LuxR<sub>vfn</sub> system” described above is found in a range of Gram-negative bacteria, this system is not the rule for the QS in species of the *Vibrio* genus. In fact, other types of AI synthases and membrane-bound AI receptors are used by most *Vibrio* species for which QS systems have been characterized. In this “*Vibrio* QS system,” a signaling cascade initiated from the membrane-bound receptors culminates in expression of the master transcriptional regulator of the QS regulon (24, 25). The *Vibrio* QS master regulator characterized first was *V. harveyi* LuxR (here referred to as LuxR<sub>vfn</sub>), which also controls the bioluminescence of this bacterium (26). However, LuxR<sub>vfn</sub>, and its homologues conserved in other *Vibrio* species are distinct from LuxR<sub>vpr</sub> in terms of structure and biochemical properties. LuxR<sub>vfn</sub> homologues include *V. vulnificus* SmcR, *V. para-haemolyticus* OpaR, *V. anguillarum* VanT, *V. cholerae* HapR, and *V. fischeri* LitR (25). In fact, expression of LuxR<sub>vfn</sub> in *V. fischeri* is regulated directly by LitR (27, 28), supporting the idea of the central role of LuxR<sub>vfn</sub> homologues as QS master regulators in *Vibrio* species.

Previously, we and others determined the crystal structures of SmcR and HapR, which reveal a putative ligand-binding pocket within the dimerization domain (29, 30). Since they belong to the TetR family of transcriptional regulators whose DNA-binding activity is controlled in a ligand-dependent manner (31), we hypothesized that it might be possible to identify a small molecule that would bind to the pocket and regulate the function of LuxR<sub>vfn</sub> homologues.

In this study, we performed high-throughput screening of 8,844 compounds and identified QStatin [1-{(S)-bromothiophene-2-sulfonyl}-1H-pyrazole] as a potent SmcR inhibitor in *V. vulnificus*. Structural analysis of SmcR complexed with QStatin revealed that the latter binds tightly to the putative ligand-binding pocket, thereby altering the DNA-binding properties of SmcR and leading to dysregulation of the QS regulon. QStatin showed pan-QS inhibitor activity in diverse *Vibrio* species that have LuxR<sub>vfn</sub>
RESULTS

Small molecules interfering with SmcR. To identify a selective inhibitor of Vibrio QS master regulators, we chose V. vulnificus SmcR as a representative target (29). A heterologous system, namely, that of Escherichia coli, was used to rule out the presence of false-positive molecules that inhibit QS signaling components other than SmcR. Accordingly, we cotransformed E. coli with the pBSS wild type (pBSS-WT) (carrying the arabinose-inducible smcR) and with pBS0918 (carrying the promoterless lux operon fused to the SmcR-repressible promoter P_{VVMO6_03194}). This screening strain remains nonluminescent unless a potential hit molecule inhibits either the function or expression of SmcR (Fig. 1a). Screening of a total of 8,844 compounds (concentration, 20 μM) identified four hit molecules (see Fig. S1a in the supplemental material). When these molecules were reexamined at the same time, compounds 357D10 and 377B6 induced expression at significantly higher relative luminescence unit (RLU) levels than were seen with the dimethyl sulfoxide (DMSO)-treated negative-control strain. The RLUs for these compound-treated strains were almost equal to those for the positive-control strain containing the empty vector (pBAD24) instead of pBSS-WT (Fig. 1b).

The results were further validated using V. vulnificus reporter strains that harbor pBB1, a LuxR_{Vh}-activated lux reporter plasmid. As previously reported for V. harveyi and V. cholerae (32, 33), the RLU level of the V. vulnificus WT (pBB1) decreased soon after inoculation into fresh LB supplemented with 2.0% (wt/vol) NaCl (LBS) but increased back to the initial level as the cells grew (Fig. 1c, DMSO). However, the increase of RLU homologues with high sequence conservation and attenuated their virulence in an aquatic host.
was halted when the strain was treated with hit molecules (Fig. 1c). Notably, the increase of RLU was also halted when *V. vulnificus* Δ*luxO* (pBB1) was treated with the compounds (Fig. 1d), indicating that the SmcR protein, constitutively expressed in this particular strain, was the target of these compounds. Consistent with this, the compounds did not reduce the RLU level for the *V. vulnificus* Δ*smcR* (pBB1) strain (Fig. S1b).

**Identification of QStatin as an inhibitor of SmcR activity.** Because 377B6 [1-(5-bromothiophene-2-sulfonyl)-1H-pyrazole] showed the strongest inhibition of SmcR activity without significant growth attenuation of *V. vulnificus* (Fig. S1c), it was selected as a specific SmcR inhibitor and named “QStatin” to designate its function (Fig. 2a). SmcR activity was assessed by measuring the RLU level for *V. vulnificus* WT (pBB1) in the presence of different concentrations of QStatin, which revealed that the half-maximal effective concentration (EC$_{50}$) of QStatin was 208.9 nM (Fig. 2b).

We furthermore examined the effect of QStatin on the expression of known SmcR-activated virulence factors in *V. vulnificus*, (e.g., protease and elastase) (13, 34). As shown in Fig. 2c, the activities of both virulence factors were reduced by QStatin in a dose-dependent manner. When the β-galactosidase gene was transcriptionally fused downstream of the elastase gene promoter, $\beta$Gal$_{Vv}$, β-galactosidase activity was also reduced by the presence of QStatin (Fig. S1d), indicating that the reduction in elastase activity by QStatin was due to its lower expression caused by SmcR dysfunction rather than to impairment of elastase catalytic activity itself. Most importantly, despite significant attenuation of SmcR function, QStatin had no effect on the cellular levels of SmcR (Fig. 2c), indicating that the compound did not affect the expression or stability of SmcR. Taken together, these results suggest that QStatin is a selective and potent inhibitor of SmcR activity.

**Molecular interaction between SmcR and QStatin.** Next, we analyzed the interaction between SmcR and QStatin by performing isothermal titration calorimetry (ITC) experiments. Exothermic reactions were observed (Fig. S2a to c) and a best fit was achieved using a two-sequential binding model. Although two identical putative ligand-binding pockets are present in the SmcR dimer (29), the sequential binding model revealed a significant difference in free energy changes; the first binding ($K_d$ [dissociation constant] = 0.47 μM) was about 10-fold tighter than the second binding ($K_d$ = 5.00 μM) (Fig. 3a).

To gain further insight into the interaction between SmcR and QStatin, we determined the crystal structure of the SmcR-QStatin complex. Consistent with the results of the ITC experiments, we found that two QStatin molecules bound to the putative...
ligand-binding pockets of the SmcR dimer (Fig. 3b). The hydrophilic sulfonyl group and pyrazole ring of QStatin were orientated toward the pocket entrance, which was predominantly surrounded by polar residues (Fig. 3c). Specifically, the oxygen atom of the sulfonyl group and a nitrogen atom in the pyrazole ring of QStatin formed hydrogen bonds with the side chains of Asn133 and Gln137 residues, respectively. The pyrazole ring also engaged in hydrophobic interactions with the Trp114 and Phe129 residues, as well as in stacking interactions with the imidazole ring of His167 (Fig. 3c). On the other hand, the bromothiophene ring of QStatin was located within the inner part of the hydrophobic ligand-binding pocket formed by Phe75, Phe78, Leu79, Ile96, Met100, Val140, Ala163, Phe166, and Cys170 (Fig. 3c). To verify the biological relevance of the interaction between SmcR and QStatin, we substituted the His167 residue involved in the interaction with QStatin with alanine. When we analyzed this substitution mutant SmcR using the E. coli screening strain, we found that the repressive activity of the mutant SmcR on the P

FIG 3 QStatin binds to the putative ligand-binding pocket of SmcR. (a) ITC analysis showing that QStatin binds directly to SmcR with high affinity. Data are representative of two experiments with similar results. Kₐ, absorption rate constant. (b) Structure of the SmcR-QStatin complex. QStatin is represented by a ball-and-stick model. (c) Electron density difference map showing QStatin bound within the putative ligand-binding pocket of SmcR. The FO-FC map was calculated before the inclusion of QStatin and is contoured at 3.0σ. The SmcR residues involved in the interaction with QStatin are represented by white sticks. (d) Superimposition of the apo-SmcR and SmcR-QStatin complexes. (e) QStatin reduces the flexibility of SmcR. The structures of apo-SmcR and SmcR-QStatin were compared according to their B-factor values. High and low B-factors are represented by dark and light colors, respectively. QStatin is represented by magenta sticks.
by QStatin (20 μM), whereas that of the WT SmcR was (Fig. S2d). This indicates that the His167 residue is critical for the interaction with QStatin.

The overall structure of the SmcR-QStatin complex is quite similar to that of apo-SmcR (Fig. 3d) (29). However, the average B-factor (Å²) for QStatin-SmcR was 26.7, whereas that for apo-SmcR was 36.1 (Fig. 3e). The B-factor of a protein structure reflects the fluctuation of atoms around an average position and provides important information about protein dynamics (35); thus, it indicates the degree of thermal motion and static disorder of atoms in a protein crystal structure (36). Since a higher B-factor (reflected by a dark color) implies that an atom/residue is more flexible, the results indicate that QStatin binding led to a significant reduction in the flexibility of SmcR ligand-binding and DNA-binding domains, which comprised the most flexible glycine-rich hinge region (red) (Fig. 3e). These changes in flexibility may affect the interaction between SmcR and its target promoter DNAs in V. vulnificus.

QStatin alters the interaction between SmcR and its target promoter DNAs. To examine whether QStatin affects the DNA-binding activity of SmcR, we performed an electrophoretic mobility shift assay (EMSA) with P_vvpE DNA (13). Although SmcR binding to the DNA was reduced specifically by QStatin in a dose-dependent manner, the interaction was not completely abolished, even in the presence of large amounts (100 μM) of the molecule (Fig. 4a and b).

Thus, we used ITC experiments to examine other biochemical properties of the interactions between target promoter DNAs and either apo-SmcR or QStatin-SmcR. Consistent with the previous results, the binding affinity of apo-SmcR for the P_vvpE DNA (K_d = 3.7 μM) was only slightly higher than that of QStatin-SmcR (K_d = 5.0 μM). However, the characteristics of the interaction were significantly different; the former interaction was exothermic, whereas the latter was endothermic (Fig. 4c; see also Fig. S3a). Thus, the enthalpic (ΔH) and entropic (TΔS) components of the free binding energy of SmcR-DNA interaction were markedly changed by QStatin, indicating

FIG 4 QStatin affects the interaction between SmcR and target promoter DNAs. (a) For EMSA, a 200-bp P_vvpE DNA fragment was radioactively labeled and then used as a DNA probe. Purified SmcR protein was added to the probe (15 ng) along with either QStatin (100 μM) or DMSO (2%). As a control, a random molecule (100 μM) from the library that showed no SmcR-inhibiting activity during initial screening was added instead of QStatin. B, bound DNA; F, free DNA. (b) EMSA was performed as described for panel a, except that SmcR was mixed with DMSO (2%) or increasing amounts of QStatin as indicated. (c to e) QStatin alters the interaction between SmcR and its target promoter DNAs. The promoter DNA of vvpE (c), flhF (d), or VVMO6_03194 (e) was titrated with the apo-SmcR complex (left) or the SmcR-QStatin complex (right). The thermodynamic values calculated from the reactions are shown. Data are representative of two experiments with similar results. (f) The in vivo interaction between SmcR and the promoter DNAs in the presence (20 μM) or absence (0.02% DMSO) of QStatin was analyzed by ChIP. V. vulnificus ΔsmcR was included as a control. A representative image from two independent experiments is shown.
that QStatin affects the characteristics of the interaction between SmcR and P_vvpE DNA. Similarly, SmcR interactions with the promoter DNAs of the flhF (P_{flhF}) and VVMO6_03194 (P_{VVMO6_03194}) genes, both of which are directly repressed by SmcR (37), were affected by QStatin. In these cases, the effect of QStatin was more noticeable, as the compound decreased the binding affinities of SmcR for the DNAs about 5-fold (for P_{flhF}) or 8-fold (for P_{VVMO6_03194}) (Fig. 4d and e; see also Fig. S3b and c).

Next, we used chromatin immunoprecipitation (ChiP) assays to assess whether changes in the DNA-binding affinity of SmcR upon exposure to QStatin reflect QStatin-mediated dysregulation of SmcR binding to its target promoter DNAs in vivo. To this end, a FLAG-tagged SmcR protein which was expressed and that functions and responds to QStatin in a manner identical to that of native SmcR (Fig. S3d and e) was immunoprecipitated with anti-FLAG magnetic beads from cross-linked V. vulnificus grown in the presence or absence of QStatin (20 \mu M). When the coprecipitated chromosomal DNAs were reverse cross-linked and amplified by PCR using primers specific for each promoter (P_vvpE, P_{flhF}, and P_{VVMO6_03194}), all three promoters were amplified (even from the sample treated with QStatin); however, the intensities of the promoter bands were slightly weaker than intensities of those from the DMSO-treated sample (Fig. 4f). Thus, the results support the finding that QStatin-induced changes in the structural flexibility of SmcR have marked effects on the interaction between SmcR and its various target promoter DNAs, although mild changes in the affinity of the interaction with target DNAs do occur which may lead to SmcR dysfunction.

QStatin globally affects the expression of the SmcR regulon in vivo. We examined whether QStatin (20 \mu M) affects the expression of SmcR regulon in vivo. Both quantitative reverse transcription-PCR (qRT-PCR) and RNA sequencing analyses revealed that QStatin-treated WT V. vulnificus (WT+QStatin) and DMSO-treated ΔsmcR mutant V. vulnificus (ΔsmcR+DMSO) had expression profiles similar to those seen with the SmcR regulon (Fig. 5a and b; see also Fig. S4), indicating that QStatin switches SmcR to a dysfunctional state and globally affects the expression of the SmcR regulon in vivo. Also, QStatin does not seem to inhibit transcriptional regulators other than SmcR, because gene expression profiles in the ΔsmcR mutant were not affected by QStatin treatment (Fig. 5b); thus, the ΔsmcR+QStatin sample clustered with the ΔsmcR+DMSO and WT+QStatin samples in a principal-component analysis (Fig. 5c). Taken together, these transcriptome-level in vivo analyses further indicate that QStatin affects the expression of the whole SmcR regulon by directly and selectively inhibiting SmcR.

QStatin is a pan-QS inhibitor attenuating the virulence of pathogenic Vibrio species. One of the known SmcR-mediated QS phenotypes in V. vulnificus is biofilm dispersion, which helps bacteria colonize new sites (38). Consistent with this, QStatin (20 \mu M) significantly impaired the biofilm dispersion of V. vulnificus WT, which resembled that of the DMSO-treated ΔsmcR mutant (Fig. 6a, WT+QStatin and ΔsmcR+DMSO). Because QStatin did not increase the mass of ΔsmcR mutant biofilms, QStatin did not directly enhance biofilm formation but impaired SmcR-mediated biofilm dispersion (Fig. 6a, ΔsmcR+QStatin). RNA sequencing results also showed that the expression levels of various biofilm formation/dispersion-related genes were altered similarly by QStatin treatment and the ΔsmcR mutation (Fig. 5b).

We hypothesized that QStatin may also affect the function of other Vibrio LuxR_vv homologues because the residues forming the ligand-binding pocket of SmcR were highly conserved across different LuxR_vv homologues (Fig. 3c; see also Fig. S5a). Thus, we examined the effects of QStatin on the representative QS phenotypes of each Vibrio species, namely, bioluminescence of V. harveyi and V. fischeri, colony opacity of V. para-haemolyticus, and protease production of V. anguillarum (28, 39–41). All phenotypes were markedly affected by QStatin (Fig. 6b and c; see also Fig. S5b and c), indicating that this compound is a bona fide pan-QS inhibitor of Vibrio species.

Finally, we evaluated the efficacy of QStatin against the virulence of pathogenic Vibrio species using their model aquatic host, Artemia franciscana (17–19, 42). QStatin (20 \mu M) markedly increased the survival rate of shrimp nauplii challenged with V. vul-
FIG 5 Effects of QStatin on SmcR regulon expression. (a) The fold changes in expression of each gene in either the QStatin-treated WT or DMSO-treated ΔsmcR mutant strain are presented relative to those in the DMSO-treated WT strain. Error bars represent the SD of results from three independent experiments. (b) The fold changes of whole transcriptome expression in the DMSO-treated ΔsmcR mutant, QStatin-treated ΔsmcR mutant, and QStatin-treated WT biofilms relative to that in the DMSO-treated WT biofilm were examined by RNA sequencing. Among the genes differentially expressed in DMSO-treated ΔsmcR mutant biofilm relative to DMSO-treated WT biofilm (P = 0.05; fold change, ≥2), 19 genes potentially involved in virulence, motility, and biofilm formation/dispersion were selected. Fold changes of the expression of 19 genes in the indicated samples relative to those in the DMSO-treated WT biofilm are shown in the heatmap with colors representing the log2 RPKM ratio. Locus tags of genes in the V. vulnificus MO6-24/O genome (GenBank accession numbers CP002469.1 and CP002470.1) and their gene products are shown. Please refer to Fig. S4 and Data Set S1 for expression changes in other genes. (c) Principal-component analysis of the whole-gene expression profiles of the samples. Each symbol represents the transcriptome of a single sample from two biological replicates per sample group.
nificus, V. harveyi, or V. parahaemolyticus, without affecting the viability of bacteria (Fig. 6d to f; see also Fig. S1c and S5d to f). No such virulence attenuation was observed with the V. vulnificus ΔsmcR mutant, demonstrating that virulence attenuation by QStatin is SmcR mediated (Fig. 6d). Remarkably, V. harveyi, a known shrimp pathogen, showed the most significant virulence attenuation; the survival rate of the nauplii increased from 35.7% ± 6.8% (DMSO) to 85.5% ± 8.1% (QStatin).

**DISCUSSION**

Diverse strategies have been explored to control QS, including inhibition of Ai synthesis, degradation of Ai, and interference with Ai detection (43). However, certain Vibrio species produce different kinds of AIs and sense them using specific cognate receptors (32, 44). Thus, simultaneous inhibition of all Ai-specific pathways is necessary to block Vibrio QS. However, LuxRv, homologues function as master QS regulators at the center of the Vibrio QS pathway (25). Moreover, these homologues show high sequence similarity and may have structural similarity, making them the most attractive targets for Vibrio QS inhibition. QStatin inhibited SmcR activity with an EC50 in the range of hundreds of nanomolars and markedly affected QS in all Vibrio species examined. Its potent and broad-spectrum activity would be particularly important in practical settings, since multiple Vibrio species can cause vibriosis in aquaculture. Actually, the unit of Vibrio pathogenesis in naturally infected oysters is the Vibrio population and not the clone (45).

Some Vibrio species opportunistically infect human and cause acute diseases using virulence factors such as cholera toxin, toxin coregulated pili (Tcp), and hemolysin (1). Because QS represses the expression of such virulence factors (33, 46), pro-QS strategies
have been proposed to treat patients (47, 48). However, this does not seem to be the case for aquatic animals, the more relevant natural hosts for *Vibrio* species (17). Indeed, virulence of *V. vulnificus* against the shrimp was considerably attenuated by QStatin (Fig. 6d) despite an increase in the expression of the hemolysin gene (Fig. 5b). This apparently conflicting result is not unprecedented, because a *V. harveyi* QS mutant also exhibits reduced virulence to brine shrimp despite expressing more type 3 secretion system (T3SS) components (17, 49). QStatin decreased the expression of many other virulence factors (Fig. 5b), including exoprotease/metalloprotease (VvpE) (19), a spike protein of T6SS apparatus (VgrG) (50), and a phenazine biosynthesis protein (PhzF) (51). Thus, these virulence factors seem to be more critical than hemolysin or T3SS in *Vibrio* pathogenesis in aquatic environments.

Furthermore, the genes governing motility (FlaE), chemotaxis (methyl-accepting chemotaxis proteins), and biofilm formation/dispersion (CabBC- and c-diGMP-regulating enzymes), which can affect both virulence and environmental adaptation of bacteria (52–54), were also significantly dysregulated by QStatin (Fig. 5b). In fact, QS has been reported to contribute to bacterial persistence and survival in the presence of grazing predators and bacteriophages in natural environments (20, 21, 55–57). Therefore, it is less likely that QStatin causes the pathogenic vibrios to bloom in the relevant environments, although the shrimp were persistently infected by vibrios under our gnotobiotic experimental conditions.

To the best of our knowledge, QStatin is the first ligand to have been shown to bind to the putative ligand-binding pocket of LuxR<sub>v</sub> homologues. Furthermore, our results provide new insights into the *Vibrio* QS. First, the tight binding of QStatin to the conserved binding pocket suggests the presence of an authentic natural ligand regulating LuxR<sub>v</sub> homologues. If it exists, such a ligand might have pharmacophore properties similar to those of QStatin. In this regard, it is fascinating that halogenated furanones, which bind to LuxR<sub>v</sub> and affect its DNA-binding activity, are produced by the marine alga *Delisea pulchra* (58). Future examination of furanone binding to the ligand-binding pocket of LuxR<sub>v</sub> homologues would reveal the relationship between *Vibrio* species and their ecologic neighborhoods at the molecular level.

Second, the results provide new perspectives into how LuxR<sub>v</sub> homologues directly regulate many different target genes. In fact, the members of the TetR family of transcriptional regulators are known to bind to one or two promoters containing a symmetrical palindrome sequence (31). In contrast, LuxR<sub>v</sub> homologues bind to hundreds of promoters harboring imperfect, asymmetrical consensus sequences in which one half is more conserved than the other half (37, 59–61). Thus, LuxR<sub>v</sub> homologues are speculated to have evolved structural flexibility, allowing it to bind to less-conserved, diverse sequences (61). In the present study, we showed that QStatin reduced the structural flexibility of SmcR (Fig. 3e), altering its DNA-binding properties in vitro (Fig. 4a to e) and thereby dysregulating gene expression in vivo (Fig. 5; see also Fig. S4 in the supplemental material). Notably, QStatin affects the flexibility of the glycine-rich hinge region of apo-SmcR (Fig. 3e). Consistent with this, a previous study revealed that a natural variant of HapR with a mutation (G39D) in the glycine-rich hinge region is defective with respect to target promoter regulation (62). Thus, our results provide direct evidence that flexibility is an essential molecular feature of LuxR<sub>v</sub> homologues, permitting them to function as global transcriptional regulators. If QStatin were to bind to LuxR<sub>v</sub> homologues, they would become less flexible, resulting in nonfunctional binding to target promoter DNAs (Fig. 4c to f and 7). One possible explanation for this nonfunctionality is that the rigid LuxR<sub>v</sub> homologues could not interact properly with other transcriptional regulators required for regulation of target promoters. Indeed, markedly rigid residues within QStatin-bound SmcR include Leu139 and Asn142, which are predicted to be essential for LuxR<sub>v</sub>-RNA polymerase interactions (61). Since one monomer of the SmcR dimer is more flexible than the other (Fig. 3e), we propose a model in which the less flexible monomer binds to the more conserved half and the other, more flexible monomer is “induced-fitted” into the less conserved half of the consensus sequence for functional binding (Fig. 7).
In conclusion, we identified QStatin as a potent pan-QS inhibitor that selectively inhibits the activity of LuxR homologues in *Vibrio* species. Since QStatin showed a marked antivirulence effect with no direct bactericidal or bacteriostatic activity, it could be used to control vibriosis in aquacultures while avoiding the resistance associated with other antimicrobial agents. The data revealing the structure of the SmcR-QStatin complex should help us to design a more effective *Vibrio* QS inhibitor in the future. Importantly, the effect of QStatin on the persistence and survival of *Vibrio* species in real marine environments needs to be investigated, as well as any eventual mechanisms of QStatin resistance.

MATERIALS AND METHODS

**Bacterial strains, plasmids, and culture media.** The strains and plasmids used in this study are listed in Table S1 in the supplemental material. *E. coli* and *Vibrio* strains were grown in Luria-Bertani medium (LB) at 37°C and in LB supplemented with 2.0% (wt/vol) NaCl (LBS) at 30°C, respectively, with appropriate antibiotics. The small-molecule library was generously provided by the Korea Chemical Bank ([http://eng.chembank.org/](http://eng.chembank.org/)), and the molecules were dissolved in DMSO. Hit molecules were either purchased from ChemDiv (San Diego, CA) or synthesized as described below. Other chemicals were purchased from Sigma-Aldrich (St. Louis, MO).

**High-throughput screening.** *E. coli* DH5α was cotransformed with pBSS-WT, carrying the arabinose-inducible smcR gene, and with pBS0918, a reporter plasmid carrying the SmcR-repressed promoter of *VVMO6* _03194_ (P <sub>VVMO6_03194</sub>) (37) fused to the lux operon. The resulting strain was cultured to an A<sub>600</sub> of 0.5 in fresh LB containing 0.0002% (wt/vol) l-arabinose, and then 100 μl of culture was transferred to each well of a 96-well microtiter plate (Optilux; BD Falcon, Bedford, MA) containing a 20 μM concentration of each molecule or 2% DMSO. The plates were incubated at 37°C with shaking, and luminescence and growth (A<sub>600</sub>) were measured three times at 1.5-h intervals using an Infinite M200 microplate reader (Tecan, Männedorf, Switzerland). RLU was calculated by dividing the luminescence value by the A<sub>600</sub> value (13). Information related to the high-throughput screening is summarized in Table S2.

**Verification and determination of the EC<sub>50</sub> of hit molecules.** The plasmid pBB1 carrying the LuxR<sub>smcR</sub> homologue-activated lux operon (39) was conjugally transferred into the *V. vulnificus* wild-type (WT) strain, the ΔsmcR mutant, or the ΔluxO mutant (63). These *V. vulnificus* reporter strains were grown overnight, diluted 1:1,000 in fresh LBS, and treated with hit molecules as described above. RLU values were calculated every hour. To determine the EC<sub>50</sub> QStatin (10<sup>-12</sup> to 10<sup>-4</sup> M) or 2% DMSO as a control was added to the *V. vulnificus* WT reporter strain and RLU was measured after 5 h. The percentage of
SmcR activity of the sample at a given concentration of QStatin was determined using the following equation: percent SmcR activity = sample RLU/control RLU × 100. The EC<sub>50</sub> of QStatin (the concentration reducing the SmcR activity to 50%) was calculated from a plot of the percentages of SmcR activity versus QStatin concentrations using GraphPad Prism 6.0 (GraphPad Software, Inc., San Diego, CA).

**Determination of total protease, elastase, and β-galactosidase activities controlled by SmcR.** Total protease and elastase activities were determined as previously described (64), except that the *V. vulnificus* WT and ΔsmcR mutant strains were treated with QStatin (5, 20, or 50 μM) or DMSO (0.02%) as described above. To confirm the reduced expression of elastase gene, *V. vulnificus* DH0602 containing SmcR-expressing plasmid pBSH-WT (29) was treated with 20 μM QStatin for 16 h and then its β-galactosidase activity was measured as described previously (29). The amounts of cellular SmcR and DnaK were determined by immunoblotting using rat anti-SmcR polyclonal antiserum and mouse anti-E. coli DnaK monoclonal antibody (Enzo Lifesciences, Farmingdale, NY), respectively, with alkaline phosphatase or horseradish peroxidase (HRP)-conjugated anti-rat or anti-mouse IgG antibody, as described previously (13).

**Protein purification, crystallization, data collection, and structural analysis.** Native or selenomethionine (SeMet)-substituted SmcR was expressed and purified as described previously (29). High-quality SeMet-substituted SmcR crystals were produced under the following optimized conditions: 0.2 M Li<sub>2</sub>SO<sub>4</sub>, 7% polyethylene glycol (PEG) 3000, and 0.1 M imidazole (pH 8.0). Crystals appeared within 2 days and grew for a further 5 days. To obtain SmcR crystals complexed with QStatin, SmcR crystals were soaked for 30 min in a solution containing 2.5 mM QStatin, 0.2 M Li<sub>2</sub>SO<sub>4</sub>, 7% PEG 3000, 0.1 M imidazole (pH 8.0), and 10% glycerol. The crystals were then placed under a nitrogen gas stream (at −173°C). Diffraction data were collected at a resolution of 2.1 Å at beamline 7A (Pohang Accelerator Laboratory, Pohang, South Korea) and processed using the HKL2000 program suite (65). The structure of the SmcR-QStatin complex was solved using molecular replacement method and the MOLREP program (66), with the SmcR structure (PDB identifier [ID]: 3K29) used as a template. The structure was then refined using COOT (67) and refined using REFMACs (68). The refinement process included the translation-liberation-screw procedure. The crystallographic data are summarized in Table S3.

**EMSA and ChiP analysis.** EMSA of SmcR binding to the vvpE promoter region was performed as described previously (29), except that QStatin or a random molecule from the library was added to the reaction sample. ChiP analysis was performed as described elsewhere (61), with some modifications. Briefly, the ΔsmcR strain or a mutant strain expressing FLAG-smcR was grown for 16 h in the presence of either QStatin (20 μM) or DMSO (0.02%). After cross-linking with formaldehyde occurred, cells were lysed and sonicated to shear the genomic DNA. Clarified lysates were incubated for 6 h at 4°C with anti-FLAG M2 magnetic beads (Sigma-Aldrich). After washing was performed, the immunoprecipitated complexes were eluted and DNAs were reverse cross-linked. The presence of target promoter DNAs was analyzed by PCR.

**ITC analysis.** For ITC analysis of the SmcR-QStatin interaction, purified SmcR was dialyzed extensively against buffer (50 mM Tris [pH 7.0], 300 mM NaCl, 0.5% DMSO), and QStatin was diluted in the same buffer. The samples were degassed by vacuum aspiration for 15 min prior to titration at 25°C. SmcR (0.48 mM [in dimer]) in the syringe was titrated against QStatin (0.025 mM) in the reaction cell of VP-ITC (Microcal Inc., Northampton, MA). To evaluate how QStatin affects the SmcR-DNA interaction, the duplex DNAs of P<sub>vvpE</sub> (sequences are in Table S4) were synthesized and dialyzed against buffer. SmcR (0.42 mM [in dimer]) was incubated with QStatin (molar ratio, 1:4) prior to titration against each duplex DNA (0.02 mM). The mixture was stirred at 300 rpm, and the thermal power was recorded every 10 s. The thermograms were then analyzed using the Origin package (version 7) supplied with the instrument.

**qRT-PCR.** The *V. vulnificus* strains grown to an A<sub>600</sub> of 0.25 were treated with 20 μM QStatin or 2% DMSO and further incubated to the stationary phase (A<sub>600</sub> = 5). Total RNA was then isolated using RNPprotect bacterial reagent and an RNeasy minikit (Qiagen, Valencia, CA). Synthesis of cDNA and amplification of target genes were done using an iScript cDNA synthesis kit, iQ SYBR Green Supermix, and an iCycler iQ qRT-PCR system (Bio-Rad Laboratories, Hercules, CA). The sequences of the primers used are listed in Table S4. The relative expression levels of the genes were normalized to the expression of the 16S rRNA gene (internal reference), as described previously (69).

**Analysis of biofilm formation/dispersion.** Overnight-cultured *V. vulnificus* strains were diluted with *E. coli* minimal medium (70) containing 32.6 mM glycerol and either 20 μM QStatin or 0.04% DMSO. The diluted cultures (200 μl) were transferred to polystyrene microtiter plates (Nunc, Roskilde, Denmark) and incubated for the indicated times at 30°C to form biofilms, which were then quantitated as described previously (38).

**RNA sequencing and analysis.** Total RNA was isolated from the biofilm developed as described above, except it was incubated for 13 h in polystyrene 6-well plates (SPL, Seoul, South Korea). The RNAs were further purified by removing DNA using TURBO DNase (Ambion, Austin, TX), and mRNA was selectively enriched by depleting rRNA using a Ribos-Zero rRNA removal kit (Epigenetix, Madison, WI). Then, the cDNA library was constructed using a TrueSeq Stranded mRNA Sample Prep kit (Illumina, San Diego, CA). The quality of the cDNA libraries was evaluated using an Agilent 2100 Bioanalyzer and Agilent DNA 1000 reagents (Agilent Technologies, Santa Clara, CA). Strand-specific single-ended 50-nucleotide sequences were read from each cDNA library using HiSeq 2500 (Illumina). The raw sequencing reads were analyzed using CLC Genomics workbench 5.5.1 (CLC Bio, Aarhus, Denmark) and mapped onto the *V. vulnificus* MO6-24/O reference genome (GenBank accession numbers CP002469.1 and CP002470.1), allowing up to two mismatches per read. The expression level of each gene was defined using a value corresponding to the number of reads per kilobase of transcript per million mapped reads (RPKM), as
described previously (71). Quantile-normalized RPKM values were then statistically analyzed by t tests to identify the genes that were differentially expressed (greater than 2-fold change with a P value of ≤0.05) from the DMSO-treated ΔsmcR mutant cells relative to the DMSO-treated WT cells. Genes with an RPKM value of <3 were considered not to be expressed and were excluded from the analysis. Heat maps were generated by the CIMminer program (72) using the RPKM-fold change for each gene in the test samples. The mapping statistics for the sequencing reads and the RPKM values, fold change values, and P values for entire genes under different conditions are shown in Data Set S1 in the supplemental material. CLC Genomics workbench 5.5.1 software was used for a principal-component analysis of the whole-gene expression profiles of the samples.

**Brine shrimp challenge test.** Cysts of *A. franciscana* (INVE Aquaculture, Salt Lake City, UT) were axenically hatched and challenged as described elsewhere (17), with the following modifications. Hatched nauplii were fed with autoclaved *Aeromonas hydrophila* strain KCTC 2358 at a concentration of 107 cells ml−1 of filtered and autoclaved artificial sea salt solution (Sigma-Aldrich) (40 g liter−1). The nauplii were challenged with 1 × 104 CFU of *V. vulnificus* and 1 × 105 CFU of *V. harveyi* or *V. parahaemolyticus* in the presence of 20 μM QStatin or 0.04% DMSO. In each experiment, at least four groups of 5 to 10 nauplii in 1 ml of the solution were transferred into each well of a 24-well plate and incubated at 28°C with gentle shaking until observation under a light microscope (Leica MZ125, Leica Microsystems, Inc., Switzerland) was performed.

**Chemical synthesis of QStatin.** Briefly, 4-bromothiophene-2-sulfonyl chloride (382 mg; 1.47 mM) and triethylamine (202 μl; 1.47 mM) were added to a solution of pyrazole (50 mg; 0.74 mM)–ethanol (10 ml). The reaction mixture was then refluxed for 6 h under a nitrogen atmosphere and cooled to room temperature. After the mixture was concentrated under reduced pressure, the crude product was extracted with methylene chloride (50 ml). The organic layer was then washed with brine, dried over Mg2SO4, and concentrated. The crude material was purified by silica gel chromatography using 10% to 30% (vol/vol) ethyl acetate in hexane as the eluent to yield QStatin as a solid (215 mg; 75% yield). The characteristics of the molecule were as follows: melting point, 112°C; 1H nuclear magnetic resonance (1H-NMR) (500 MHz, CDCl3) δ (ppm), 8.08 (dd, J = 2.8, 0.3 Hz, 1H), 7.79 (d, J = 1.2 Hz, 1H), 7.61 (d, J = 4.1 Hz, 1H), 7.09 (d, J = 4.1 Hz, 1H), and 6.44 (dd, J = 2.8, 1.6 Hz, 1H); 13C-NMR (126 MHz, CDCl3) δ (ppm), 145.8, 141.4, 135.1, 134.5, 131.2, 127.3, and 109.4.

**Statistical analysis.** Statistical analyses were performed as indicated in the figure legends using GraphPad Prism 6.0 software.

**Accession number(s).** The atomic coordinates and structure factors have been deposited in the Protein Data Bank (http://www.pdb.org) under PDB ID code 5X3R. All raw transcriptome data have been deposited in the NCBI BioProject database (https://www.ncbi.nlm.nih.gov/bioproject) under accession number PRJNA271541.

## SUPPLEMENTAL MATERIAL

Supplemental material for this article may be found at [https://doi.org/10.1128/mBio.02262-17](https://doi.org/10.1128/mBio.02262-17).

**FIG S1**, PDF file, 0.5 MB.

**FIG S2**, PDF file, 0.5 MB.

**FIG S3**, PDF file, 0.8 MB.

**FIG S4**, PDF file, 0.3 MB.

**FIG S5**, PDF file, 3.7 MB.

**TABLE S1**, DOCX file, 0.03 MB.

**TABLE S2**, DOCX file, 0.01 MB.

**TABLE S3**, DO CX file, 0.01 MB.

**TABLE S4**, DO CX file, 0.02 MB.

**DATA SET S1**, XLSX file, 1.3 MB.

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B.S.K., S.Y.J., M.H.K., and S.H.C. designed the research; B.S.K., S.Y.J., Y.-J.B., J.H., Y.K., and K.K.J. performed the research; B.S.K., S.Y.J., Y.-J.B., J.H., D.L., M.H.K., and S.H.C. analyzed the data; and B.S.K., M.H.K., and S.H.C. wrote the paper.
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