

ORIGINAL ARTICLE

Members of native coral microbiota inhibit glycosidases and thwart colonization of coral mucus by an opportunistic pathogen

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The outcome of the interactions between native commensal microorganisms and opportunistic pathogens is crucial to the health of the coral holobiont. During the establishment within the coral surface mucus layer, opportunistic pathogens, including a white pox pathogen *Serratia marcescens* PDL100, compete with native bacteria for available nutrients. Both commensals and pathogens employ glycosidases and *N*-acetyl-glucosaminidase to utilize components of coral mucus. This study tested the hypothesis that specific glycosidases were critical for the growth of *S. marcescens* on mucus and that their inhibition by native coral microbiota reduces fitness of the pathogen. Consistent with this hypothesis, a *S. marcescens* transposon mutant with reduced glycosidase and *N*-acetyl-glucosaminidase activities was unable to compete with the wild type on the mucus of the host coral *Acropora palmata*, although it was at least as competitive as the wild type on a minimal medium with glycerol and casamino acids. Virulence of the mutant was modestly reduced in the *Aiptasia* model. A survey revealed that ~8% of culturable coral commensal bacteria have the ability to inhibit glycosidases in the pathogen. A small molecular weight, ethanol-soluble substance(s) produced by the coral commensal *Exiguobacterium* sp. was capable of the inhibition of the induction of catabolic enzymes in *S. marcescens*. This inhibition was in part responsible for the 10–100-fold reduction in the ability of the pathogen to grow on coral mucus. These results provide insight into potential mechanisms of commensal interference with early colonization and infection behaviors in opportunistic pathogens and highlight an important function for the native microbiota in coral health.

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Introduction

The foundation of reef ecosystems is the coral holobiont: a dynamic symbiosis comprised of the coral polyp, the dinoflagellate *Symbiodinium* spp. and their associated microbial communities. Interactions within this tripartite symbiosis determine the overall health of the coral colony and its resistance to opportunistic pathogens (Reshef *et al.*, 2006; Rosenberg *et al.*, 2007; Teplitski and Ritchie, 2009). The subject of this investigation is *Serratia marcescens* PDL100, one of the strains that causes white pox disease in *Acropora palmata* and is associated with anthropogenic sources (Patterson *et al.*, 2002; Sutherland *et al.*, 2011). To colonize and infect the

coral, this multi-host opportunistic pathogen must first establish within the coral surface mucus layer.

Coral mucus is a polymer made in specialized mucocytes of the polyp from the photosynthate produced by their endosymbiotic dinoflagellates and then excreted onto the coral surface (Brown and Bythell, 2005; Bythell and Wild, 2011). The chemical structure of coral mucus has been determined for only few species (Ducklow and Mitchell, 1979; Meikle *et al.*, 1987, 1988; Coddeville *et al.*, 2011). The mucus polymer of *Acroporid* corals is a glycoprotein, which consists of a serine and threonine-rich polypeptide backbone with sulfated oligosaccharide side chains attached to it through *O*-glycosidic linkages (Meikle *et al.*, 1987). The major sugar residues in the oligosaccharide side chains of *A. formosa* mucus are *D*-arabinose, *D*-mannose, *N*-acetyl-*D*-glucosamine and *D*-galactose (Meikle *et al.*, 1988). The patterns of induction of glycosidase activities in *S. marcescens* and coral commensals, when grown on coral mucus, are consistent with its chemical structure (Krediet *et al.*, 2009a, b).

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Coral mucus supports growth of bacteria up to 10^6 – 10^8 cfu ml⁻¹ (Sharon and Rosenberg, 2008; Krediet *et al.*, 2009b; Garren and Azam, 2010). Nutrients and chemicals from the host excreted with the mucus determine the composition and structure of the associated microbial communities (Ritchie, 2006; Garren and Azam, 2012). The ability to efficiently utilize carbon and nitrogen sources within mucus likely directly affects competitiveness of the microorganisms in this environment.

Aside from the need to be able to grow on available substrates, successful establishment within a niche requires that microorganisms efficiently spread to and colonize the sites where nutrients are available. Coordinated, multi-cellular spreading over semi-solid surfaces (known as 'swarming') is one such mechanism of expansion and surface colonization (Verstraeten *et al.*, 2008). Swarming is often coregulated with certain metabolic pathways (Toguchi *et al.*, 2000; Wang *et al.*, 2004). Global regulatory networks (GacS/GacA-Csr, FlhDC-FlhA and EnvZ/OmpR) contribute to the control of both the motility and catabolism of specific carbon sources (Park and Forst, 2006; Jones *et al.*, 2008a; Timmermans and Van Melderen, 2010). This coregulation may be instrumental for the regulatory switch that contributes to the efficient establishment within a particular niche.

A better understanding of the interactions between coral commensals and opportunistic pathogens led to the appreciation of the fact that commensals have evolved to protect their nutrient niche (and the coral hosts) from the invading opportunists (Reshef *et al.*, 2006). Coral commensals produce antibiotics, inhibitors of cell-to-cell communication, and currently unknown compounds capable of inhibiting swarming and biofilm formation (Ritchie, 2006; Nissimov *et al.*, 2009; Shnit-Orland and Kushmaro, 2009; Rypien *et al.*, 2010; Tait *et al.*, 2010; Alagely *et al.*, 2011). When coral commensals capable of interfering with swarming and biofilm formation in a pathogen were pre-colonized onto a model polyp, they completely inhibited the progression of a disease caused by *S. marcescens* (Alagely *et al.*, 2011). This is similar to the ability of the commensal microbiota of vertebrates to act as a barrier to colonization and infection by pathogens (Leatham *et al.*, 2009).

With this study we aimed to test whether the ability of the native commensal microbiota to inhibit specific catabolic enzymes in the coral pathogen *S. marcescens* disrupts the ability of the pathogen to establish within coral mucus. *S. marcescens* PDL100 possesses a broad suite of catabolic enzymes. Although many of the tested glycosidases are produced apparently constitutively, β -D-galactosidase, N-acetyl- β -glucosaminidase, and α -D-glucopyranosidase were differentially accumulated when the pathogen was grown on mucus (Krediet *et al.*, 2009a). The over-arching hypothesis was tested with a three-step experiment: first, a *Serratia* mutant

defective in mucus utilization was identified to test the role of specific catabolic enzymes in growth on mucus; second, coral commensals capable of inhibiting specific enzymatic activities were identified; third, interactions of the wild-type *S. marcescens* and its isogenic mutant defective in mucus utilization with the commensals were tested in mesocosm experiments.

Materials and Methods

Bacterial strains, media and growth conditions

Bacterial strains used in this study are listed in Table 1. Coral-associated bacteria were isolated from mucus of *A. palmata* by dilution plating onto Glycerol Artificial Sea Water medium as in Ritchie, 2006. The identities of the marine isolates were confirmed by PCR-amplifying and sequencing fragments of their 16S rRNA genes as previously described (Alagely *et al.*, 2011). Data are deposited in GenBank (Accession numbers JQ954975; JQ954976; JQ954977; JQ954978; JQ954979; JQ954980).

S. marcescens PDL100 and coral commensals were grown at 30 °C on the Glycerol Artificial Sea Water medium or marine broth (Difco-Becton, Dickinson and Company, Franklin Lakes, NJ, USA); *Escherichia coli* strains were grown at 37 °C in Luria-Bertani broth (Fisher Scientific, Atlanta, GA, USA) and when necessary on 1.5% agar plates. Unless otherwise specified, *S. marcescens* cultures were supplemented with tetracycline 10 μ g ml⁻¹ (Tc10), to which *S. marcescens* PDL100 is spontaneously resistant.

Coral mucus was collected from apparently healthy colonies of *A. palmata* at Looe Key Reef, Florida (24° 32.764' N; 81° 24.304' W) in July 2011 and prepared as previously described (Ritchie, 2006; Krediet *et al.*, 2009a).

Mariner transposon mutagenesis of Serratia marcescens PDL100 and characterization of the mutants

Tri-parental conjugation was conducted using *E. coli* SM10- λ pir pBT20, *S. marcescens* PDL100 and the helper strain *E. coli* DH5 α pRK600 as previously described (Kalivoda *et al.*, 2008). One thousand colonies from each of the three independent matings were compiled into a library. This library was then screened for β -galactosidase activity (see below). To characterize mutants of interest, inverse PCR of DNA sequences flanking the transposon insertion was performed (Ochman *et al.*, 1988; Supplementary Material).

For the experiments with *E. coli*, individual *E. coli* BW25113 mutants were obtained through the Keio Collection (Baba *et al.*, 2006) and each of the mutants was transduced into strain W3110 (*lac*⁺) using phage P1 (Lennox, 1955; Ikeda and Tomizawa, 1965).

Table 1 Bacterial strains and plasmids

Isolate or strain	Relevant characteristic(s)	Source or reference
<i>Escherichia coli</i> strains		
DH5 α	F- ϕ 80 <i>lacZ</i> Δ M15 Δ (<i>lacZYA-argF</i>) U169 <i>recA1 endA1 hsdR17</i> (rk-, mk+) <i>phoA supE44 λ-thi-1 gyrA96 relA1</i>	Invitrogen
DH5 α λ -pir	DH5 α lysogenized with a bacteriophage λ_{434} derivative containing the <i>pir</i> gene	(Macinga <i>et al.</i> , 1995)
Top10	F <i>mcrA</i> (<i>mrr-hsdRMS-mcrBC</i>) 80 <i>lacZ</i> M15 <i>lacX74 deoR recA1 araD139</i> (<i>ara-leu</i>)7697 <i>galU galK</i>	Invitrogen
W3110	<i>rpsL endA1 nupG</i>	(Hayashi <i>et al.</i> , 2006)
BW25113	<i>rrnB3 ΔlacZ4787 hsdR514 Δ(araBAD)567 Δ(rhaBAD)568 rph-1</i>	(Baba <i>et al.</i> , 2006)
BW25113 Δ <i>malG15::kan</i>	Single knockout deletion mutant of <i>malG</i>	(Baba <i>et al.</i> , 2006)
BW25113 Δ <i>malF16::kan</i>	Single knockout deletion mutant of <i>malF</i>	(Baba <i>et al.</i> , 2006)
BW25113 Δ <i>malE17::kan</i>	Single knockout deletion mutant of <i>malE</i>	(Baba <i>et al.</i> , 2006)
W3110 Δ <i>malG15</i>	<i>malG</i> transduced into W3110 by P1	This study
W3110 Δ <i>malF16</i>	<i>malF</i> transduced into W3110 by P1	This study
W3110 Δ <i>malE17</i>	<i>malE</i> transduced into W3110 by P1	This study
<i>Serratia marcescens</i> strains		
<i>Serratia marcescens</i> PDL100	Coral pathogen isolated from white pox lesion on <i>Acropora palmata</i> , ATCC BAA-632, Tet ^R	ATCC, (Patterson <i>et al.</i> , 2002)
<i>Serratia marcescens</i> CK2A4	Mariner transposon mutant deficient in β -galactosidase activity	This study
<i>Serratia marcescens</i> MG1	Isolated from rotten cucumber	(Givskov <i>et al.</i> , 1997)
<i>Serratia marcescens</i> ATCC 43422	Human throat isolate, pigmented	ATCC, (Krediet <i>et al.</i> , 2009b)
Coral commensal strains		
<i>Photobacterium</i> sp. 33G2	Isolated from <i>A. palmata</i> mucus; identity confirmed by 16S rRNA gene sequencing; JQ954975	This study, (Ritchie, 2006)
<i>Photobacterium damselae</i> 33G4	Isolated from <i>A. palmata</i> mucus; identity confirmed by 16S rRNA gene sequencing; JQ954976	This study, (Ritchie, 2006)
<i>Photobacterium leiognathi</i> 33E3	Isolated from <i>A. palmata</i> mucus; identity confirmed by 16S rRNA gene sequencing; JQ954977	This study, (Ritchie, 2006)
<i>Photobacterium leiognathi</i> 33C4	Isolated from <i>A. palmata</i> mucus; identity confirmed by 16S rRNA gene sequencing; JQ954978	This study, (Ritchie, 2006)
<i>Exiguobacterium</i> sp. 33G8	Isolated from <i>A. palmata</i> mucus; identity confirmed by 16S rRNA gene sequencing; JQ954979	This study, (Ritchie, 2006)
<i>Vibrio harveyi</i> 34B3	Isolated from <i>A. palmata</i> mucus; identity confirmed by 16S rRNA gene sequencing; JQ954980	This study, (Ritchie, 2006)
Plasmids		
pCR2.1-TOPO	Cloning vector; Km ^R	Invitrogen
pRK600	ColE1 replicon with RK2 <i>tra</i> genes, conjugation helper plasmid	(Karunakaran <i>et al.</i> , 2005)
pBT20	Mariner transposon delivery vector, Gm ^R , R6K <i>ori</i>	(Kalivoda <i>et al.</i> , 2008)
pCP20	Temperature-sensitive replicon, thermal induction of FLP synthesis, Cm ^R , Ap ^R	(Datsenko and Wanner, 2000)
pBBR1-MCS2-kan	Cloning vector, Km ^R	(Kovach <i>et al.</i> , 1995)
pBAD18-kan	Cloning vector, Km ^R , under arabinose inducible <i>araBAD</i> promoter	(Guzman <i>et al.</i> , 1995)
pCJK16	<i>malEFG</i> from PDL100 PCR amplified and cloned into pCR2.1-TOPO, Ap ^R , Km ^R	This study
pCJK17	<i>malEFG</i> from pCJK16 subcloned into <i>EcoRI</i> site of pBAD18-kan, Km ^R	This study
pCJK18	<i>malEFG</i> from pCJK17 subcloned into <i>EcoRI</i> site of pBBR1-MCS2-kan, Km ^R	This study

Abbreviations: Ap^r, ampicillin resistance; Cm^r, chloramphenicol resistance; Gm^r, gentamicin resistance; Km^r, kanamycin resistance; Tet^r, tetracycline resistance.

β -Galactosidase assays

Independent screens were conducted either with a library of transposon mutants or with coral commensals. The ability of the commensals to inhibit β -galactosidase in *S. marcescens* PDL100 was tested by coinoculating coral commensals with PDL100 in 0.3% marine agar in 96-well plates supplemented with 5-bromo-4-chloro-indolyl- β -D-galactopyranoside (X-gal) at 40 μ g ml⁻¹ (Gold Biotechnology, St Louis, MO, USA). Plates were incubated at 30 °C overnight, and were monitored for blue color. Candidates were

retested in cross-streaks on marine agar. Inhibitory capabilities of the coral commensals were tested against PDL100 in β -galactosidase assays through coincubation of *S. marcescens* in the spent cell-free culture supernatants of coral commensals. The ability of the cell-free supernatants to inhibit activity of the purified *E. coli* β -galactosidase enzyme (Sigma-Aldrich, St Louis, MO, USA) was also tested (Supplementary Material).

Individual transposon mutants were assayed in soft marine agar with X-gal and were monitored for

blue color development. Mutants deficient in β -galactosidase activity (compared to the wild type) were isolated to single colony on X-gal containing media and then assayed quantitatively against the wild type using ONPG (Miller, 1972).

Competitive fitness assays

Overnight cultures of *S. marcescens* PDL100 and CK2A4 were grown from glycerol stock in marine broth to an approximate OD_{600} of 1.7 and cultures were serially diluted and mixed in 1:1 ratio. Strains were inoculated individually or as a mix at starting concentrations of 10^2 cfu ml⁻¹ into filter sterilized coral mucus, high-molecular-weight fraction of coral mucus from *A. palmata*, and an artificial seawater-based casamino acids/glycerol media (in artificial seawater: 0.05% Casamino acids, 0.4% glycerol, 10 mM HEPES, pH 7.1). All cultures were grown in triplicates. For enumeration, cultures were dilution plated at 0, 12, 24, 48, 72 and 96 h on marine agar supplemented with Tc10. All assays on each medium were repeated at least twice. To distinguish between the wild type and the mutant, individual colonies were patched onto selective media with Tc10 and gentamycin (Gm) 50 μ g ml⁻¹.

For the three-strain competition assay, *S. marcescens* PDL100, CK2A4 and commensals with β -galactosidase inhibitory properties, strains were grown individually in marine broth overnight and diluted as above. *S. marcescens* PDL100 was inoculated at 10^2 cfu ml⁻¹ and the cocktail of the six inhibitory coral commensals was inoculated at 10^4 cfu ml⁻¹ into high-molecular-weight coral mucus from *A. palmata*. Competitive indices were calculated for each treatment using the formula $(M_{out}/WT_{out})/(M_{in}/WT_{in})$, where M is the proportion of mutant cells and WT is the proportion of wild-type cells in the inoculum (_{in}) or in the recovered samples (_{out}). Statistical significance of each competitive index was established by comparing log values of the competitive indices using a two-tailed *t*-test.

Preliminary characterization of inhibitory compounds in *Exiguobacterium* sp. 33G8

Potential antibiotic production by the coral commensal strains was tested by filling agar cores of the commensal strains into semi-solid marine agar (0.5%) seeded with PDL100. The coinoculated strains were incubated at 30 °C overnight and monitored for inhibition of growth in PDL100.

Inhibition of β -galactopyranosidase, *N*-acetyl- β -D-glucosaminidase and α -D-glucopyranosidase enzymatic activities in PDL100 by coral commensal bacteria were tested through coinoculation of the strains using dialysis pouches. Cellulose ester dialysis membrane (MWCO: 1000 000) stored in 0.05% sodium azide (Spectrum Laboratories, Inc., Rancho Dominguez, CA, USA) was cut into 10-cm strips and washed in sterile DI water (10 water

changes) at 4 °C. *S. marcescens* PDL100 was grown from glycerol stock in 5 ml marine broth and then subcultured into 100 ml marine broth. *Exiguobacterium* sp. 33G8 was grown from glycerol stock in 5 ml of marine broth overnight at 30 °C and subcultured into 100 ml of sterile marine broth. After overnight incubation, the OD_{600} of PDL100 was 0.7 and *Exiguobacterium* 33G8 was at 1.8. The PDL100 culture was concentrated to an OD_{600} of 1.2 and 5 ml of these cells were transferred into the dialysis pouches. The entire 100 ml culture of *Exiguobacterium* 33G8 was transferred into 500 ml of sterile marine broth. A dialysis tube containing PDL100 cells was placed into the 33G8 culture and into a flask of sterile marine broth as a control. The cocultures were incubated at 30 °C for 6 h and enzymatic activities of all three enzymes in PDL100 were measured at the beginning of the experiment and 6 h later using nitrophenyl substrates as in Miller, 1972.

To begin to determine the type of inhibitory activity produced by *Exiguobacterium* sp. 33G8 in coculture, cell-free supernatant from the *Exiguobacterium-Serratia* coinoculation experiment was subjected to flash ion-exchange chromatography (DOWEX 50-W, Sigma-Aldrich) and the flow-through was then subjected to reverse-phase Silica-C₁₈ chromatography (Alltech Associates, Inc., Deerfield, IL, USA) (Supplementary Material).

Each of the eluted fractions was dried, re-extracted as described in Materials and methods, serially diluted (threefold) and then subjected to a bioassay. The highest dilution contained ~1.5 ml of the 'culture equivalent'. The dilutions and solvent controls were evaporated in the chamber of the Centrivap Concentrator (Labconco, Kansas City, MO, USA) at 50 °C for 30 min. Six hundred and fifty microliters of *S. marcescens* PDL100 culture (OD_{600} =0.922) that had been washed in equal volume of sterile marine broth were added to each sample. Cells were incubated with the diluted fractions for 6 h at 30 °C and assayed for α -D-glucopyranosidase activity as described by Miller, 1972.

Swarming assays

Twenty microliters of cultures of individual commensal strains grown overnight in marine broth were spotted onto a sterile glass fiber GF/C filter disk and placed on a marine agar plate. Five microliters of *S. marcescens* cultures were spotted onto AB swarm agar plate ~2 cm from the disk containing commensals as described previously (Alagely *et al.*, 2011). For swimming motility, 5 μ l of each culture were stab inoculated into marine broth 0.3% agar plates 2 cm apart. Plates were incubated at 30 °C overnight and photographed with a Canon EOS Rebel Xsi digital camera (Canon USA Inc., San Jose, CA, USA). Images were corrected for auto levels in Adobe Photoshop.

Virulence in a sea anemone model

Individual polyp infections with PDL100 wild type and CK2A4 were performed as described previously (Krediet *et al.*, in review). The polyps were monitored for disease signs and Kaplan–Meier survivorship analyses were performed for each infection using JMP 9.0 Pro statistical software (SAS Institute Inc., Cary, NC, USA) and mortality across treatments was compared with a Cox proportional hazards χ^2 model (Hosmer and Lemeshow, 1999).

Results

Characterization of the *S. marcescens* mutant defective in glycosidase and *N*-acetyl-glucosaminidase activities
To test the importance of β -galactosidase, *N*-acetyl- β -D-glucosaminidase and α -D-glucopyranosidase for the ability of *S. marcescens* to utilize components of coral mucus, a transposon mutant, *S. marcescens* CK2A4, deficient in these activities was isolated. To characterize the mutant, inverse PCR was carried out, and a single PCR band was detected and then sequenced. The sites flanking the transposon insertion were identified as homologous to the *malF* gene (Spro_4473; base pairs 283–715) of *Serratia proteamaculans*. However, attempts to complement the transposon mutant with the entire predicted 4.8-kb '*malEFG*' operon amplified from *S. marcescens* PDL100 genomic DNA were not successful (data not shown).

Because the complementation of the CK2A4 mutant was unsuccessful, glycosidase and *N*-acetyl-glucosaminidase activities of the defined *malE*, *malF* and *malG* mutants were tested in *E. coli* W3110 to corroborate the identification of CK2A4 as a *malF* mutant to establish whether the phenotype of the *S. marcescens* mutant is consistent with the mutation in the *mal* operon. Enzymatic activities in the defined *E. coli* mutants and in CK2A4 were similarly decreased (Figure 1). Note, that *E. coli* W3110 does not produce *N*-acetyl- β -D-glucosaminidase (Figure 1). Therefore, even though the *malF* mutation in CK2A4 could not be complemented, its phenotype is consistent with the phenotypes of three defined *E. coli* mutants in the *malEFG* operon.

Competitive fitness of *S. marcescens* PDL100 and CK2A4

The ecological importance of β -galactosidase and other catabolic enzymes for utilization of coral mucus from *A. palmata* was tested by growing wild-type PDL100 and the CK2A4 mutant on total coral mucus and high-molecular-weight fraction of mucus both individually and in a 1:1 coculture. Individual growth curves for both strains showed the same growth trend on total mucus and high-molecular-weight coral mucus (Figures 2a and c). Importantly, however, within the coculture, CK2A4 was not competitively fit on either total coral mucus or on high-molecular-weight fraction of the mucus

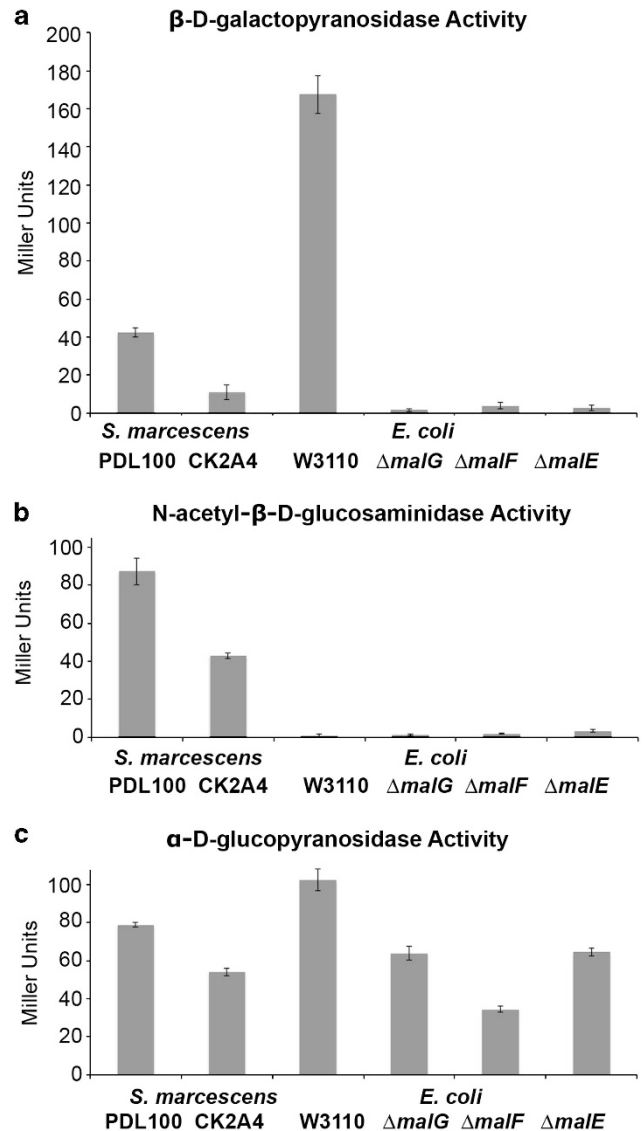


Figure 1 Enzymatic activities in mutants of *S. marcescens* and *E. coli*. Overnight cultures of wild-type *S. marcescens* PDL100 and the transposon mutant (CK2A4) were tested for activities of (a) β -D-galactopyranosidase, (b) *N*-acetyl- β -D-glucosaminidase and (c) α -D-glucopyranosidase using a Miller assay (Miller, 1972) with appropriate *p*-nitrophenyl-glycoside substrates. *E. coli* W3110 and individual in-frame single gene knockouts of *malE*, *malF* and *malG* were also tested to compare phenotypes of CK2A4 with those of the defined well-characterized *E. coli* mutants in the *malEFG* operon. All cultures were grown overnight before the assays; *S. marcescens* was cultured in Marine broth, *E. coli* was grown in LB. Averages of three biological and four technical replications are shown. Error bars are standard error. There was a significant reduction in the enzymatic activities between wild-type *S. marcescens* and CK2A4 (β -D-galactopyranosidase: $t = 13.55$, $dF = 6$, $P < 0.0001$; *N*-acetyl- β -D-glucosaminidase: $t = 12.22$, $dF = 3.3$, $P = 0.0008$; α -D-glucopyranosidase: $t = 20.86$, $dF = 4.87$, $P < 0.0001$). In *E. coli*, β -D-galactopyranosidase and α -D-glucopyranosidase activities were reduced in *malE*, *malF* and *malG* mutants.

(Figures 2b and d; log competitive indices -1.64 and -1.23 , respectively), consistent with the predicted role of the β -galactosidase, *N*-acetyl- β -D-glucosaminidase and α -D-glucosidase enzymes in

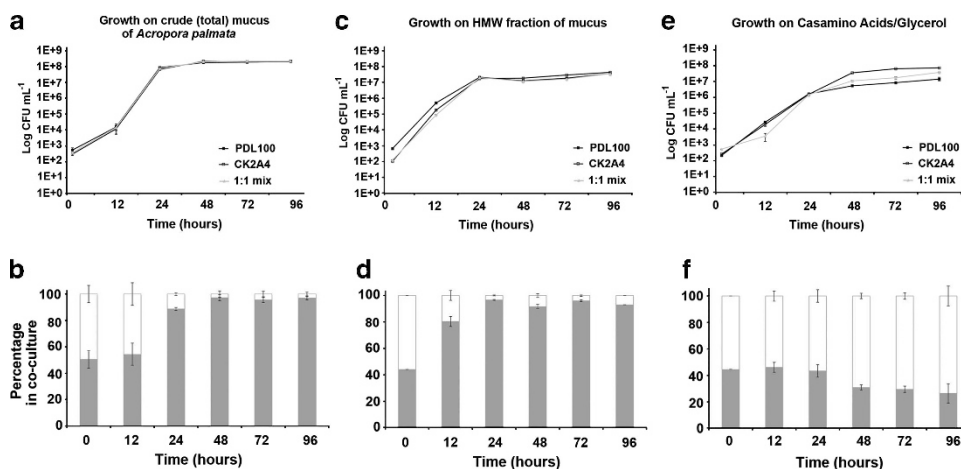


Figure 2 Competitive fitness of *S. marcescens* PDL100 and CK2A4. To test the importance of catabolic enzymes for growth of *S. marcescens* on coral mucus, both wild type and CK2A4 were grown both individually and in coculture on mucus of *A. palmata* and in the HEPES-buffered (10 mM, pH 7.1) artificial seawater with casamino acids (0.5 g l⁻¹) and glycerol (0.4% vol/vol). For competition experiments, overnight inocula were serially diluted and mixed (1:1 ratio). Cultures in (a) crude (total) mucus of *Acropora palmata*, (c) high-molecular-weight (>5 kDa) fraction of *A. palmata* mucus or (e) casamino acids/glycerol were inoculated at 10² cfu ml⁻¹. Cultures were incubated at 30 °C with shaking. To enumerate cells, at 12, 24, 48, 72 and 96 h cultures were dilution plated onto marine agar supplemented with Tc10. Averages of three biological replications of the experiment (three independent cultures) are shown, error bars are standard error. In monocultures, all strains grew similarly and reached the same final population densities. To estimate the percentage of each strain within each coculture (competitive fitness), colonies were patched onto marine agar supplemented with Tc10 and Gm50 to differentiate between the wild type and the transposon mutant. The relative proportion of the wild type is shown as the gray portion of the stacked column. Competitive fitness experiments were repeated at least twice, and averages are shown (error bars are standard errors). On total mucus (b) and high-molecular-weight fraction of the mucus (d), the mutant was not competitive against the wild type. CK2A4 had a fitness advantage on the casamino acid/glycerol defined medium and grew to higher population densities than wild type and the 1:1 coculture (f).

mucus degradation (Krediet *et al.*, 2009a, b). After 12–24 h, the wild type dominated the culture and after 48 h the wild type represented ~95% of the culture. However, in the absence of the substrates for glycosidases or *N*-acetyl-glucosaminidases (that is, when grown on a minimal medium with casamino acids and glycerol), the CK2A4 mutant had a modest competitive advantage (Figure 2e; log competitive index 0.480). This indicates that the reduced fitness of the mutant on coral mucus is not due to a general reduction in the metabolism, but rather is due to a disruption of the specific catabolic activities required for the utilization of coral mucus.

Virulence of *S. marcescens* PDL100 and CK2A4 in a polyp model

Sixty-nine *Aiptasia pallida* polyps were used to test the virulence of *S. marcescens* PDL100 and CK2A4 over a 7-day period. After 24 h of incubation at the highest concentration (10⁸ cfu ml⁻¹) the polyps compressed their stalk length, retracted their tentacles and exhibited darkening of the tissue. These responses were seen to a lesser degree at the 10⁷ cfu ml⁻¹ treatment and rarely at the 10⁶ cfu ml⁻¹ concentration in the wild type but no mortality was observed at the lower concentrations in CK2A4. The survivorship of the infected polyps is plotted in Figure 3. The differences in survivorship between the different treatments were statistically significant

(PDL100: $\chi^2 = 38.60$; dF = 4; $P < 0.0001$; CK2A4: $\chi^2 = 28.25$; dF = 3; $P < 0.0001$).

Coral commensal bacteria capable of inhibiting enzymatic activities limit growth of *S. marcescens* in coculture

Because the ability to cleave sugar and amino-sugar residues off coral mucus appears important for the successful colonization of the coral mucus surface layer, we next tested whether the native commensals have evolved the mechanisms to inhibit glycosidases in the invading opportunists. Initial screens of a library of commensal isolates collected from the mucus surface layer of a healthy *A. palmata* (which is the only coral known to be susceptible to *S. marcescens* infections) revealed that ~8% of isolates are capable of inhibiting β -galactosidase activity in *S. marcescens* (Supplementary Figure 1). The competitive fitness of PDL100 and CK2A4 on high-molecular-weight mucus was tested in the presence of these inhibitory coral commensal strains (*Photobacterium* sp. 33G2, *P. damsela* 33G4, *Exiguobacterium* sp. 33G8, *P. leiognathi* 33C4, and *P. leiognathi* 33E3, and *Vibrio harveyi* 34B3). We tested the hypothesis that in the presence of the commensals that inhibit glycosidase and *N*-acetyl-glucosaminidase activities, the CK2A4 mutant will be as fit as the wild type due to the ability of the commensals to inhibit these enzymatic activities in the wild type. In the coral mucus coculture of *Serratia* with the

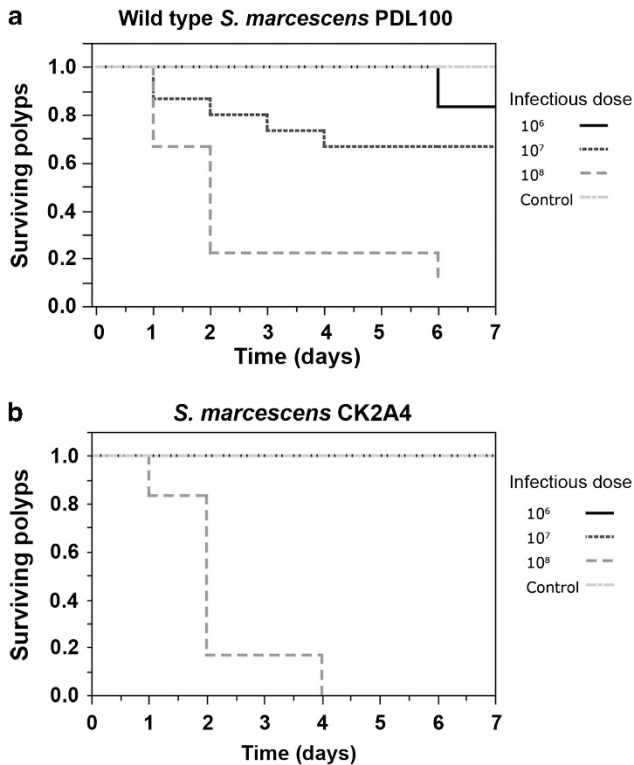


Figure 3 Survivorship of *Aiptasia pallida* polyps infected with *S. marcescens*. Ten-fold dilutions of overnight cultures of *S. marcescens* PDL100 and CK2A4 were inoculated into microtiter plate wells with *A. pallida*. Infections were maintained at room temperature (22–24 °C) with illumination. Polyps were scored daily. Each line represents the proportion of polyps surviving at each concentration (10^6 cfu ml $^{-1}$, 10^7 cfu ml $^{-1}$, 10^8 cfu ml $^{-1}$, or control as shown in the key). Survivorship decreased significantly with increasing pathogenic concentration for polyps infected with (a) *S. marcescens* PDL100 ($\chi^2 = 38.604$; dF = 4; $P < 0.0001$) and (b) *S. marcescens* CK2A4 ($\chi^2 = 28.249$; dF = 3; $P < 0.0001$).

commensals, growth of the pathogen was reduced by 10–100-fold (compared with the monoculture of *Serratia*; Figure 4a). Furthermore, the CK2A4 mutant (defective in the enzymatic activities targeted by the commensals) was more competitive against the wild type compared with the mesocosms without the commensals at 12 and 24 h (Figures 2d vs 4b; log competitive index -0.729 vs -0.129 , $t_{3,82} = 4.769$, $P = 0.0099$). These observations suggest that the ability of commensals to specifically inhibit catabolic enzymes (glycosidases and *N*-acetyl-glucosaminidase) is at least in part responsible for the reduction in growth of *S. marcescens* on coral mucus in the presence of these commensals. Our subsequent experiments focused on the characterization of the activities produced by the commensals.

Preliminary characterization of inhibitory compounds in *Exiguobacterium* sp. 33G8

Exiguobacterium sp. 33G8 strongly inhibited β -galactosidase activity in *S. marcescens* strains

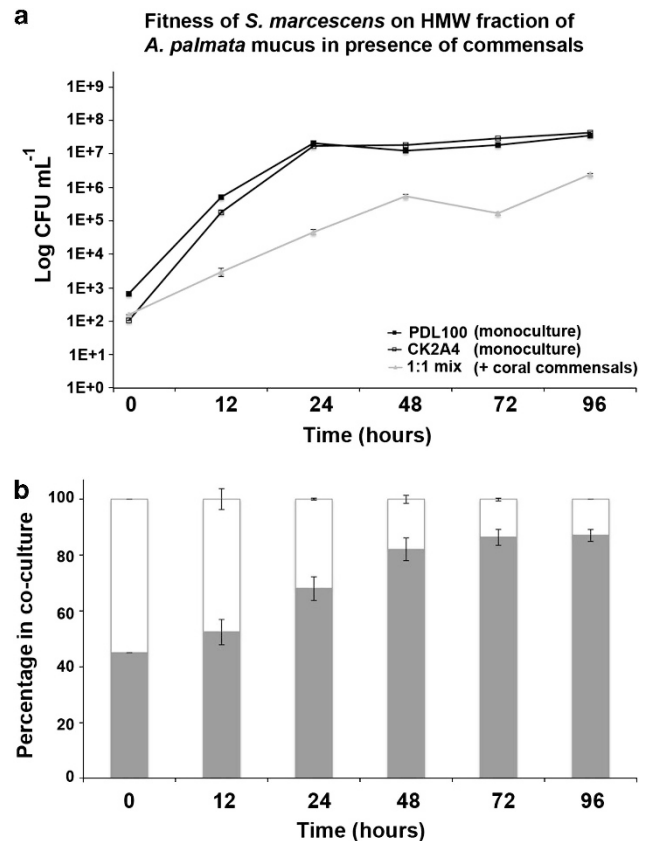


Figure 4 Competitive fitness of *S. marcescens* with coral commensals. Coral commensal strains shown to inhibit β -D-galactopyranosidase activity in *S. marcescens* PDL100 were grown in coculture on high-molecular-weight fraction of *A. palmata* mucus with wild-type PDL100 and transposon mutant CK2A4. To prepare inoculum of *S. marcescens* strains, overnight cultures were serially diluted and wild type and CK2A4 were mixed in 1:1 ratio and inoculated at 10^2 cfu ml $^{-1}$. The coral commensals (*Photobacterium* sp. 33G2, *P. damsela* 33G4, *Exiguobacterium* sp. 33G8, *P. leiognathi* 33C4, *P. leiognathi* 33E3 and *Vibrio harveyi* 34B3) were grown separately, mixed as a cocktail and inoculated at 10^4 cfu ml $^{-1}$ concurrently with *S. marcescens*. (a) In monocultures (black lines) of *S. marcescens* without coral commensals, the wild type (filled square) and CK2A4 mutant (empty square) reached the same population densities. In the coculture on high-molecular-weight fraction of *A. palmata* mucus in the presence of the coral commensals, growth of *S. marcescens* was significantly reduced (gray line). Averages of three biological replicates (three independent cultures) are shown. Error bars are standard errors. (b) Fitness of the mutant versus the wild type in the presence of the commensals was estimated by patching of *Serratia* colonies onto media with the appropriate antibiotics. The relative proportion of the wild type is shown as the gray portion of the stacked column.

PDL100 and MG1 (Supplementary Figure 1) and was chosen for further investigation. Our experiments aimed to (a) test mode of action of the compound(s) and (b) characterize chemical properties of the activity.

Reduction in enzymatic activity in PDL100 in the coculture with *Exiguobacterium* sp. was first assayed by growing PDL100 in a dialysis pouch in the presence of *Exiguobacterium* sp. At the start of the experiment, the enzymatic activities were not

significantly different between the control culture and PDL100 inoculated with *Exiguobacterium* sp. (Figure 5). After the 6 h coculture, the enzymatic activities of PDL100 in coculture were significantly reduced for each enzyme (β -galactosidase: $t = 9.340$, $dF = 4.2$, $P = 0.0005$; *N*-acetyl- β -D-glucosaminidase: $t = 4.410$, $dF = 3.27$, $P = 0.0180$; α -D-glucopyranosidase: $t = 22.191$; $dF = 4.34$; $P < 0.0001$; Figure 5). We then tested whether the substance(s) produced by the commensal directly inhibit the enzymatic activity or regulation of the enzyme synthesis. Incubation of *Serratia* in the cell-free supernatant from the monoculture of *Exiguobacterium* neither reduced the wild-type β -galactosidase activity, nor did the coral commensal cultures reduce activity of a purified β -galactosidase from *E. coli* (data not shown). This indicates that the substance is not a compound that inhibits the function of the β -galactosidase enzyme. The inhibitory activity was extracellular, but was produced by *Exiguobacterium* sp. only when it was cultured with *S. marcescens*.

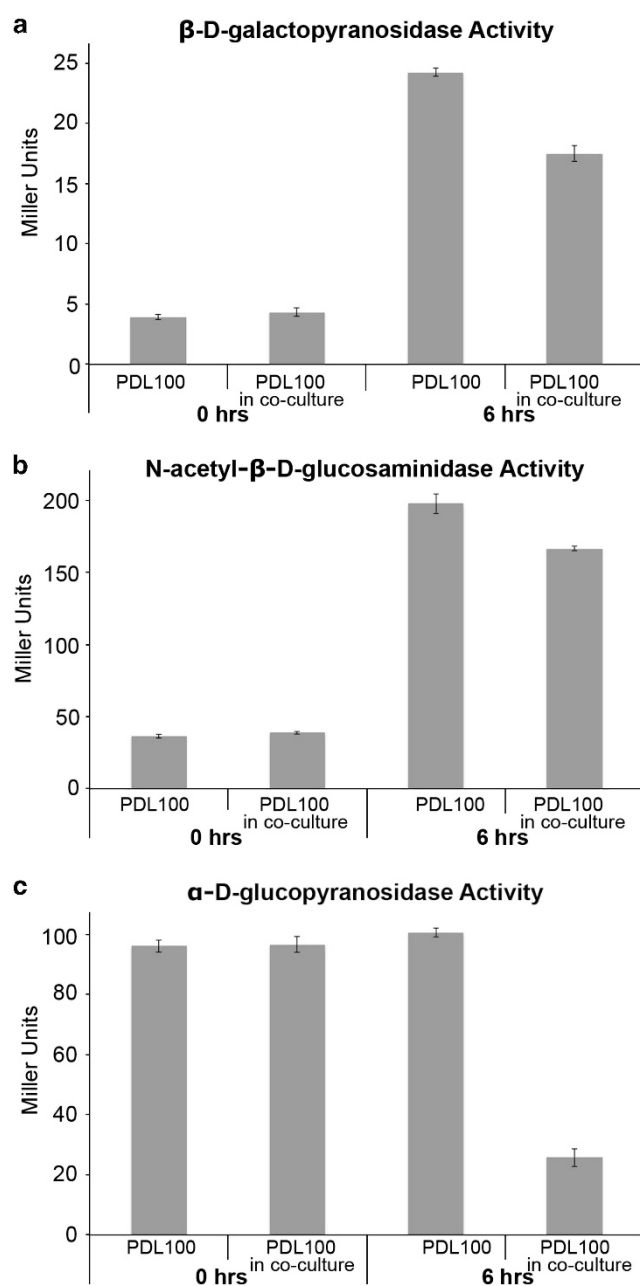
Preliminary characterization of the chemical produced by *Exiguobacterium* sp. 33G8 was performed by extraction of cell-free supernatant from the 33G8 culture used for the dialysis pouch coculture enzyme assay. Cell-free supernatant was passed through an ion-exchange column and then the flow-through from the ion-exchange column was subjected to reverse-phase Si C₁₈ chromatography. Eluent from both columns was dried, re-extracted with two different solvents and the resulting samples were assayed for their ability to reduce of α -D-glucopyranosidase activity in PDL100. This enzyme was chosen out of the three initially tested because of the strong inhibition seen in the coculture experiment (Figure 5). Of the fractions assayed, the ethanol-soluble extract of the eluent from the ion-exchange column showed the greatest reduction in activity (nearly 1.5-fold reduction),

Figure 5 Inhibition of enzymatic activities in *S. marcescens* PDL100 by *Exiguobacterium* sp. 33G8. *Exiguobacterium* sp. 33G8 was grown in coculture with PDL100 to test for inhibition of the enzymatic activities that are normally induced in *S. marcescens* grown on mucus of *A. palmata*. As detailed in Materials and methods, within the coculture, bacteria were separated with a dialysis membrane (MWCO: 1 000 000) for the ease of recovering cells for enzymatic assays. The dialysis pouch pore size was enough to retain cells but allow compounds produced by both organisms to pass through. Enzymatic assays for PDL100 (a) β -D-galactopyranosidase, (b) *N*-acetyl- β -D-glucosaminidase and (c) α -D-glucopyranosidase activities were conducted at the time of inoculation and after 6 h of coculture using nitrophenyl substrates as in Miller, 1972. Enzymatic activities of *S. marcescens* PDL100 in a monoculture or in the coculture with *Exiguobacterium* sp. 33G8 are shown. Experiments were repeated twice, each time with four technical replicates. Data from one replicated experiment are shown. There was no difference in initial activity between the treatments for each enzyme assayed. After coincubation for 6 h, *S. marcescens* PDL100 showed strongly significantly reduced enzymatic activities as a result of 33G8 for β -D-galactopyranosidase ($t_{4.2} = 9.340$, $P = 0.0005$) and α -D-glucopyranosidase ($t_{4.34} = 22.191$, $P < 0.0001$), and modestly significantly reduced activity of *N*-acetyl- β -D-glucosaminidase ($t_{3.27} = 4.410$, $P = 0.0180$).

as compared with the solvent alone control (Supplementary Figure 2A).

Swarming motility inhibition in *S. marcescens* PDL100 by coral bacteria

Even though the inhibition of enzymatic activities by the commensals was at least in part responsible for the reduced fitness of *S. marcescens* within coral mucus, it did not fully account for the observed phenotype. The conclusion that the activities produced by commensals target induction of enzymatic activities, rather than a function of the enzyme, suggests that the compounds may affect a higher-level regulator. Therefore, we explored the possibility that the commensals inhibit behaviors



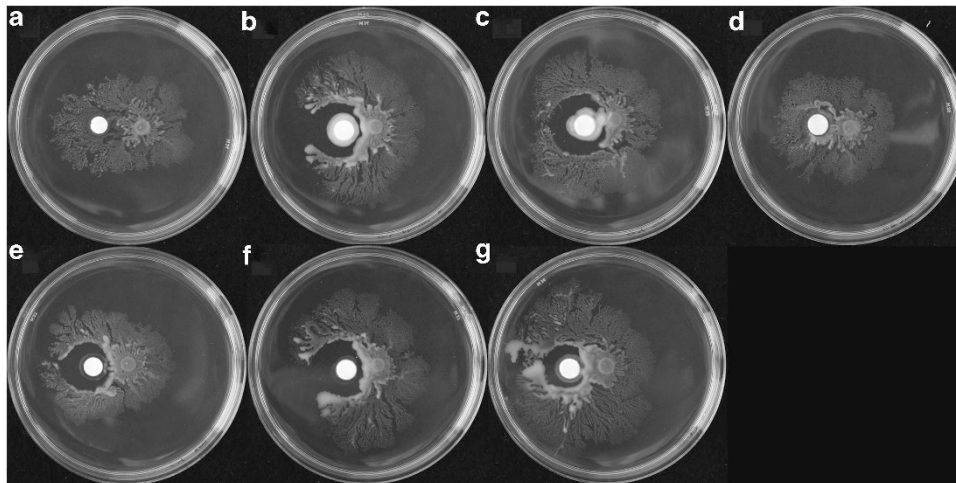


Figure 6 Inhibition of swarming of *S. marcescens* PDL100 by coral commensals. Swarming of *S. marcescens* PDL100 on AB 0.4% swarm agar next to a sterile glass fiber GF/C disk is shown in (a). Individual coral commensal strains (*Photobacterium* sp. 33G2 (b), *P. damselae* 33G4 (c), *Exiguobacterium* sp. 33G8 (d), *P. leiognathi* 33C4 (e), *P. leiognathi* 33E3 (f) and *Vibrio harveyi* 34B3 (g)) were grown in marine broth and 20 μ l were spotted onto a glass fiber GF/C disk on solid marine agar and disks were incubated at 30 °C overnight. Filter disks were placed next to PDL100 (5 μ l of overnight culture diluted 1:100 and grown for 2 h). Plates were incubated at 30 °C overnight and all strains, except 33G8, inhibited swarming motility in the pathogen. The experiments were repeated three times, data from one representative experiment are shown.

associated with niche colonization and coregulated with metabolism.

To further characterize the potential coregulation of metabolic enzymes and motility, swarming motility of PDL100 was assayed with and without the commensals as in Alagely *et al.*, 2011. When spotted onto AB swarm agar plate, PDL100 swarms (Figure 6a). Five of the six strains that were capable of inhibiting glycosidases, also inhibited swarming of *S. marcescens*. *Exiguobacterium* sp. 33G8 did not inhibit swarming although this strain showed the greatest inhibition of enzyme induction during growth in coculture (Figure 6d). We then tested three hypotheses that explain these observations: commensals produce antibiotics, motility inhibitors or specific inhibitors of swarming.

No growth inhibition of *S. marcescens* was observed in diffusion assays to test for the antibiotic production by the coral commensals (data not shown), indicating that neither the limitation of growth seen in the competitive fitness coculture test, nor inhibition of swarming is due to antibiotic production by commensals. Five of the six of the commensal strains tested reduced swarming motility in *S. marcescens*, however, none inhibited swimming within soft agar. These data suggest that the commensal strains may produce inhibitory compounds specific to the coregulation of swarming motility (not flagellar motility in general) and metabolism, perhaps by targeting a higher-level regulator that controls both pathways.

Discussion

Glycosidases and *N*-acetyl-glucosidase, which correspond to the available substrates in coral mucus,

are induced during growth of *Serratia marcescens* PDL100 on coral mucus (Krediet *et al.*, 2009a, b). This study sought to test the consequences of inhibition of these enzymatic activities in *S. marcescens* by coral-associated bacteria. A transposon mutant deficient in these activities was identified and tentatively mapped to the operon homologous to the *E. coli* *maleFG* maltose/malto-dextrin transporter. This high-affinity ABC transporter system consists of a periplasmic maltose-binding protein encoded by *malE* and a transport complex encoded by *malF*, *malG* and *malK* (Jones *et al.*, 2008b). On the basis of the transport functions of the operon, it was hypothesized that *maleFG* is necessary for proper colonization and utilization of coral mucus by the pathogen. Although the *S. marcescens* *maleFG* mutant showed a threefold decrease in β -galactosidase activity and a twofold decrease in *N*-acetyl- β -D-glucosaminidase activities compared with the wild type, the mutation did not fully eliminate the ability of the pathogen to utilize and grow on coral mucus. In a monoculture, the mutant grew to the same population density as the wild type, however, was not competitive against the wild type on coral mucus in coculture. It is plausible that the reduced affinity for the substrate resulting from the disruption of the transporters is responsible for the reduced competitive fitness of the strain. The reduction, but not a complete elimination, of the ability of the mutant to utilize mucus is likely due to the fact that bacteria rely on transporters with redundant functions for the uptake of carbon sources. The decrease in the virulence of the mutant in a surrogate host is also likely due to its reduced fitness within the surface mucus layer and its inability to establish population densities that are high enough to effectively invade the host.

Coral pathogens compete with the native coral-associated bacteria to establish within a niche and utilize preferred carbon sources of coral mucus. These coral-associated bacteria interact with and can interfere with the invading pathogen through antibiotic production, inhibition of quorum sensing and secondary metabolite production (Shnit-Orland and Kushmaro, 2009; Teplitski and Ritchie, 2009; Rypien *et al.*, 2010). This study characterized six coral-associated strains capable of interfering with metabolic enzymatic activities and decreasing the overall growth of *S. marcescens* on coral mucus. Interference of enzymatic activities was cell-associated and dependent on coculturing of commensals and the pathogen.

In addition to inhibiting enzymatic activities in *S. marcescens* that are involved in coral mucus utilization, the coral-associated bacteria also inhibited swarming motility in the pathogen. We do not yet know whether the same compound is involved in the inhibition of both swarming and catabolism of mucus components. However, if this were the case, it is possible to hypothesize that a higher-level regulatory system (GacS/GacA-Csr or FlhDC) (Romeo, 1998; Pruss *et al.*, 2003; Park and Forst, 2006; Timmermans and Van Melderen, 2010) may be the target. The GacS/GacA-Csr system is not likely to be affected by the activities of these commensals: first, the phenotype of the *gacA* mutant (Krediet *et al.*, in review) is distinct from what is observed when the wild type is exposed to the commensals tested here, and, second, the commensals did not affect the expression of the *gacA::lacZ* reporter (Krediet, unpublished data). It is also unlikely that FlhDC is affected by the commensals: even though FlhDC in *Enterobacteriaceae* is known to contribute to the regulation of both motility and metabolism (Pruss *et al.*, 2003), the fact that none of the commensals inhibited swimming motility suggests that FlhDC is also an unlikely target of these activities. Therefore, these observations suggest that either the compounds target another regulatory cascade, or that different compounds disrupt swarming and the induction of enzymatic activities separately.

The interactions between coral-associated mutualistic bacteria and invading pathogens are numerous, complex and still poorly understood. The results of this study demonstrate that coral-associated bacteria disrupt the coregulated metabolic activities and swarming in the invading opportunists, and thus the ability of the pathogens to compete with commensals within the coral mucus environment. Further research will elucidate the exact level of these regulatory pathways at which coral commensal strains interfere. It will be also important to establish whether the bacteria with the 'beneficial' properties are actively recruited by host (which would be consistent with the Hologenome Theory of Evolution (Rosenberg and Zilber-Rosenberg, 2011)), or whether there are other

hypotheses that explain their functions within the coral surface mucus layer.

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