

Survival of and In Situ Gene Expression by *Vibrio vulnificus* at Varying Salinities in Estuarine Environments[∇]

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The opportunistic human pathogen *Vibrio vulnificus* survives in a wide range of ecological environments, which demonstrates its ability to adapt to highly variable conditions. Survival and gene expression under various conditions have been extensively studied in vitro; however, little work has been done to evaluate this bacterium in its natural habitat. Therefore, this study monitored the long-term survival of *V. vulnificus* in situ and simultaneously evaluated the expression of stress (*rpoS*, *relA*, *hfq*, and *groEL*) and putative virulence (*vvpE*, *smcR*, *viuB*, and *trkA*) genes at estuarine sites of varying salinity. Additionally, the survival and gene expression of an *rpoS* and an *oxyR* mutant were examined under the same conditions. Differences between the sampling sites in the long-term survival of any strain were not seen. However, differences were seen in the expression of *viuB*, *trkA*, and *relA* but our findings differed from what has been previously shown in vitro. These results also routinely demonstrated that genes required for survival under in vitro stress or host conditions are not necessarily required for survival in the water column. Overall, this study highlights the need for further in situ evaluation of this bacterium in order to gain a true understanding of its ecology and how it relates to its natural habitat.

Vibrio vulnificus is an opportunistic human pathogen indigenous to estuarine environments and commonly isolated from fish, shellfish, water, and sediment samples worldwide (3, 11, 25, 43). Waters where this bacterium is found typically have salinities that range from 5 to 35 ppt and temperatures that range from 9 to 31°C (16, 22, 26, 28). However, the prevalence of *V. vulnificus* at certain temperatures has been shown to be related to the salinity of the environment. At lower salinities, the bacterium can be found over a wide range of temperatures, but as salinity increases, the bacterium is typically found only at higher temperatures (33). Other studies have reported a lack of culturability at temperatures of less than 10°C, at which point the bacterium enters the viable but nonculturable (VBNC) state (27). In addition to being able to survive in a wide range of temperatures and salinities, *V. vulnificus* can survive for extended periods of time in nutrient-deficient environments, such as the water column, but is also routinely isolated from such nutrient-rich environments as the oyster gut (25).

The wide variety of ecological niches that this bacterium inhabits demonstrates its ability to survive under highly variable conditions, and these physiological conditions have been extensively studied in vitro (10, 15, 29). However, such studies can only replicate a fraction of the biotic and abiotic factors that may be encountered in a natural habitat. Thus, in situ studies using cells constrained in membrane diffusion chambers, particularly involving the VBNC state, have also been conducted in order to examine the survival of *V. vulnificus* in the environment (27). While such studies are valuable in characterizing the survival of bacteria in natural environments, they

generally rely on simple plate counts and offer no insight into the gene expression occurring during growth and/or survival.

Of the few studies that have investigated gene expression in *V. vulnificus*, most have been conducted under select physiological conditions in an attempt to identify altered niche-related phenotypes on the basis of select gene inactivation (12, 31, 32, 34, 35). However, many traits contribute to ecological survival and such studies are likely to overlook subtle changes that would occur in the bacterium's natural environment. In fact, the wide variety of habitats in which this bacterium resides suggests that numerous variations in gene expression occur in response to changes in environmental conditions. To date, the only studies that have evaluated in situ gene expression by *V. vulnificus* have involved differences due to temperature, studying gene expression by cells in natural estuarine waters during the summer months (38) or on entry into and resuscitation from the VBNC state during the winter (37). The present study set out to evaluate the effects of both high and moderate salinities on in situ gene expression by *V. vulnificus* incubated in a natural estuarine environment. Examined were the expression of several stress and putative virulence genes over time and the relationship of gene expression to bacterial survival. Additionally, as the stress response proteins (namely, the alternative sigma factor *rpoS* and the oxidative stress protein *oxyR*) are thought to be involved in bacterial survival under stressful conditions (8, 39), in situ gene expression by an *rpoS* mutant and an *oxyR* mutant was examined simultaneously with that by the parent strain.

MATERIALS AND METHODS

Strains and growth conditions. *V. vulnificus* strain C7184/K2 (wild-type) and mutant derivatives AH1 (*rpoS*; 12) and K853 (*oxyR*; 17) were grown in heart infusion (HI) broth (Difco, Detroit, MI). Antibiotics (1 μg/ml chloramphenicol [Sigma, St. Louis, MO] for AH1) or additives (80 μg/ml pyruvate [Sigma] for

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TABLE 1. Primers used for gene expression analysis

Gene	Primer sequence	
	Forward (5' to 3')	Reverse (5' to 3')
<i>rpoS</i>	GCAAGATGAAGACATGCGTG	CTTCAACCTGAATCTGGCG
<i>groEL</i>	CGCAAGCTATCGTAAATGTGGG	GAGTTCGCAGAGATGGTG CCTAC
<i>relA</i>	ACTGGGCGACGTGATGTG	CCAAAGGCCAGCCAATGT
<i>smcR</i>	TTCTGTGGCGACCGTCTTCAAC	CTTCACCACGCTCAAT GGCT
<i>trkA</i>	GTCAGGTGCGCTTACTCT	TTCGTTCTCGTTGGTGAG
<i>vvpE</i>	ACTGTTTGCCGTCCAATAC	CGGGTGAAGCGGCAGAGT
<i>viuB</i>	GGTTGGGCACTAAAGGCAGAT	TCGCTTCTCCGGGGCGG
<i>hfq</i>	AGGGGCAATCTCTACAAG	TCTACTGTGGTTCCTGCT

K853) were added in order to maintain the mutations or allow growth, respectively. Cells were grown overnight at room temperature with periodic inversion.

In situ experiments. Two preparations were created for each strain by inoculating 1 ml of an overnight culture into 50 ml of 1/2 ASW (artificial seawater) (42), generating a 1:50 dilution (final concentration, ca. 10^7 CFU/ml). Appropriate concentrations of chloramphenicol or pyruvate were added to AH1 and K853 preparations, respectively, to preserve the integrity of the mutations before the cells were placed into the environment. Membrane diffusion chambers (21) were inoculated with 15-ml aliquots of one of the bacterial preparations, with four chambers inoculated for each strain. Two chambers of each strain were placed at a high-salinity site (site 1; temperature and salinity readings of 24°C and 31 ppt), and the remaining two were at a lower-salinity site (site 2; temperature and salinity readings of 26.4°C and 21 ppt). Site 1 was a private-access portion in Banks Channel, NC, and site 2 was located near a public boat ramp at Topsail Sound, NC. At 0 and 30 min and 4, 12, 24, and 48 h, 1 ml was aseptically withdrawn from each chamber. Samples were immediately centrifuged to remove cells from the environmental matrix. The supernatant was discarded, and the pellets were resuspended in RNA Protect (Qiagen, Valencia, CA) in accordance with the manufacturer's instructions and stored at -20°C until RNA extraction was performed. An additional 1 ml was withdrawn at $t = 0, 24, 48$ h to determine CFU counts on HI agar (with additives as appropriate). Plates were incubated overnight at 30°C.

RNA extraction. RNA was purified from samples treated with RNA Protect with the TRIzol Max bacterial RNA isolation kit (Invitrogen, Carlsbad, CA) in accordance with the manufacturer's instructions, with slight modifications that have been shown to increase the pure-RNA yield from *V. vulnificus* cells (37).

Analysis of gene expression. The following genes were examined for their expression at each time point sampled: *rpoS* (alternative sigma factor), *groEL* (protein chaperone), *smcR* (*V. harveyi luxR* homologue), *trkA* (potassium [K^+] uptake), *vvpE* (zinc metalloprotease), *hfq* (small protein involved in posttranscriptional regulation), *relA* (ppGpp synthase), and *viuB* (iron acquisition). Expression was evaluated by amplifying mRNA with the Access reverse transcription (RT)-PCR system (Promega). The primers used for each of these genes are listed in Table 1. The RT-PCR cycling conditions for *groEL*, *rpoS*, *smcR*, *relA*, and *viuB* were as follows: 45°C for 45 min, 94°C for 5 min, and then 30 cycles of 94°C for 30 s, 61°C for 30 s, and 72°C for 1 min, followed by a final extension at 72°C for 7 min. Cycling conditions were altered slightly for amplification of *hfq*, *trkA*, and *vvpE* as follows: 45°C for 45 min, 94°C for 3 min, and then 40 cycles of 94°C for 40 s, 57°C for 40 s, and 72°C for 1 min, followed by a final extension at 72°C for 7 min. Results presented represent data obtained from two individual chambers for each strain at each salinity site.

RESULTS AND DISCUSSION

Survival of *V. vulnificus*. *V. vulnificus* and two single-gene mutants were analyzed for the ability to survive in situ in the estuarine environment. One mutant (AH1) possesses a mutation in the *rpoS* gene, which encodes an alternative sigma factor typically involved in stress condition responses (8). The second mutant strain (K853) possesses a mutation in *oxyR*, which is part of an oxidative stress regulon for proteins involved in stress-induced protection against unrelated stresses (39). Insertion stability studies indicated that both mutations are stable, at least over a 72-h period (data not shown),

whether cells were growing in HI broth or incubated in 1/2 ASW. Culture concentrations for all strains were measured during incubation at high- and moderate-salinity sites at 0, 24, and 48 h. Results showed little variation in cell counts over the 48-h study period (Fig. 1). During in situ incubation, a slight decrease in culture concentration was observed in all strains between 0 and 24 h; however, these changes were not significant ($P \leq 0.4496$) for any strain at either sampling site. At the high-salinity site, where no recreational use of the water typically occurred, membrane diffusion chambers were left in the water column and sampled for cell concentration after 30 days. This final time point revealed that all strains remained culturable at day 30, although levels had dropped to ca. 10^5 CFU/ml (data not shown), demonstrating that *V. vulnificus* cells are able to survive at high concentrations for extended periods of time in natural environments. Indeed, we have cells of *V. vulnificus* strain C7184 that have been incubated at room temperature in ASW for more than 17 years and still retain a considerable culturable population (data not shown).

Interestingly, comparisons of cell concentrations between the mutants and the parent strain demonstrated no significant differences ($P \geq 0.6745$) in survivability at either high or moderate salinity. These results suggest that loss of either *rpoS* or *oxyR* function does not have an adverse affect on long-term survival in the water column. It was expected that *rpoS*, which is known to play a major role in the stress response in other bacteria, may play a role in survival, as it has been shown to regulate several genes that are induced upon incubation in ASW in *Escherichia coli* (36). However, our results suggest that incubation in estuarine water at warm temperatures (24 to 26°C) may not provide a considerable stress for *V. vulnificus*, and thus, *rpoS* may not be essential for survival under these conditions. Survivability studies also indicated that *oxyR* may not be required for long-term survival of *V. vulnificus* in these environments. However, while this gene has been shown to be essential for resuscitation of *V. vulnificus* out of the VBNC state (17), the present studies were conducted when water temperatures were well above the level (10 to 13°C) known to induce this dormancy state in this species. Therefore, differ-

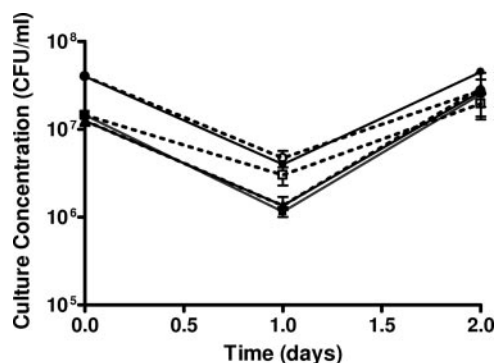


FIG. 1. Culture concentrations in chambers at moderate and high salinities. Shown are the concentrations of wild-type and mutant cultures of *V. vulnificus* incubated in membrane diffusion chambers with high-salinity (31‰) and moderate-salinity (24‰) environments over 48 h. Shown are C7184/K2 at 31 ppt (■) and 21 ppt (□), K853 at 31 ppt (▲) and 21 ppt (△), and AH1 at 31 ppt (●) and 21 ppt (○).

TABLE 2. Gene expression by C7184/K2 over time

Gene	Expression at high/moderate salinity					
	0 min	30 min	4 h	12 h	24 h	48 h
<i>rpoS</i>	+/+	+/+	+/+	+/+	+/+	+/+
<i>groEL</i>	+/+	+/+	+/-	+/+	+/+	-/-
<i>smcR</i>	+/+	+/+	+/+	+/+	+/+	+/+
<i>relA</i>	+/+	+/+ ^a	+/+	+/+	-/+	-/+
<i>hfq</i>	+/+	+/+	+/+	+/+	+/+	+/+
<i>viuB</i>	+/+	+/-	+/-	-/-	-/-	-/-
<i>trkA</i>	+/+	+/+	+/-	+/-	+/-	+/-
<i>vvpE</i>	+/+	+/+	+/+	+/+	+/+	+/+

^a One replicate was positive for gene expression, while the other replicate was negative.

ences in long-term survival between this mutant strain and the wild type may only be apparent at colder temperatures.

Gene expression analysis. It is unknown what mechanisms allow *V. vulnificus* to survive for extended periods of time in the marine environment. A necessary step in gaining an understanding of the natural ecology of this bacterium is to evaluate gene expression during long-term survival. Thus, the expression of various stress and putative virulence genes was monitored in *V. vulnificus* over 48 h in high- and moderate-salinity environments. A variety of genes have been shown to play a role in the survival or pathogenesis of *V. vulnificus* under in vitro and/or in vivo conditions (6, 12, 14, 20), and several of these genes (*vvpE*, *smcR*, *viuB*, *rpoS*, and *trkA*) were selected for evaluation of their role in survival in situ. Additionally, because the estuarine environment has been assumed to be a stressful habitat for bacteria due to temperature and salinity fluctuations and the lack of available nutrients, the expression of genes involved in stress and stringent responses (*relA*, *hfq*, and *groEL*) was also examined.

Expression of all eight genes occurred in strain C7184/K2 for at least 4 h after introduction into high-salinity estuarine water, and five of the eight genes remained on through 48 h under these conditions (Table 2). However, the expression of several genes was markedly altered when this strain was incubated at moderate salinity (Table 2), with the most dramatic differences in expression being seen for *viuB*, *trkA*, and *relA*.

Expression of *viuB* (whose product is involved in siderophore-mediated iron acquisition) dropped below the limit of detection before all other genes at both salinities in the wild-type strain. However, the lack of detection was seen much sooner (30 min) at moderate salinity than at high salinity, where expression ended after 4 to 12 h of incubation (Table 2). These results suggest that *viuB* may not be required for extended survival in the water column and may instead be used primarily for survival once inside an oyster or human host. As iron levels in seawater are only $\sim 0.0001 \mu\text{mol/kg}$ (2), it might be expected that acquisition of iron would be necessary for environmental survival, and *viuB* expression might be expected to remain on for extended periods of time at both salinities. However, while the role of siderophores has been examined in the presence of serum and in host models (19, 24), we are aware of no research that has explored their role in iron uptake in natural marine environments. Indeed, Panicker et al. (30) demonstrated that only 24% of the environmental *V. vulnificus* isolates tested possessed *viuB*, suggesting that alternative iron

acquisition systems may be present and necessary for environmental survival.

Another possible explanation for the lack of *viuB* expression seen throughout most of our study may be related to the expression of the iron regulator Fur. It was recently shown in *V. vulnificus* that Fur is found in increasing amounts under iron-limiting conditions (18). Additionally, it appears that the activity of Fur in *V. vulnificus* may differ from that in other organisms. In *E. coli*, Fur forms a complex with Fe^{2+} , which then binds to consensus sequences upstream of genes encoding siderophores and other proteins involved in iron acquisition, leading to the repression of these genes. However, regulation by Fur in *V. vulnificus* does not appear to require Fe^{2+} to bind to these consensus sequences (18). The role of Fur in the regulation of these genes in *V. vulnificus* is still unknown, but recent research opens the possibility that overexpression of Fur under iron-limiting conditions, such as those present in the marine environment, may lead to the differential expression of genes like *viuB*. Furthermore, it has been shown that Fur can bind other divalent cations, including zinc (1). Zinc occurs in seawater at approximately 10 times the concentration of iron (2). This potential for competitive binding to Fur may represent an additional or alternative aspect of iron, and thus *viuB*, regulation.

A third possible explanation for the early cessation of *viuB* expression may be related to quorum sensing. Recently, new theories as to the role of autoinducers in the environmental survival of bacteria have been proposed. One such theory is "efficiency sensing," which suggests that autoinducers are used by bacteria as indicators of their external environment (9). Autoinducers are less energetically expensive than other macromolecules, are detected with high sensitivity, and therefore can be used to determine if the environment will allow the return of secreted macromolecules to the bacterium. The ultimate value of siderophores to *V. vulnificus* lies in the ability of the bacterium to use the iron that has been scavenged by the proteins. Thus, in an open system (such as the water column), the likelihood of a siderophore returning to the bacterium is very low. Therefore, in the efficiency-sensing model, siderophore production would be turned off. This hypothesis gains additional support from our data in that *smcR* (a *V. harveyi luxR* homologue) remained on throughout our study at both salinities and in all strains (Tables 2 to 4), suggesting that the bacterium may be constantly testing the surrounding environment.

Differences in *viuB* expression were also noted between the parent strain and mutant strains, but only at moderate salinity. In the mutant strains, expression of this gene mimicked the high-salinity results, becoming undetectable between 4 and 12 h at both salinities (Tables 2 to 4). These differences may indicate that *rpoS* and *oxyR* are involved in *viuB* expression at some salinities. However, given the possibility that *viuB* is not involved in environmental iron acquisition, it may be that the observed differences in gene expression between the mutants and the parent strain are related not to salinity but rather to the overall effect of losing a stress response system.

Differential expression of *trkA* (a regulator of potassium uptake) by C7184/K2 was also observed between the moderate- and high-salinity environments. At moderate salinity, *trkA* expression dropped below the limit of detection within 4 h

TABLE 3. Gene expression by AH1 over time

Gene	Expression at high/moderate salinity					
	0 min	30 min	4 h	12 h	24 h	48 h
<i>groEL</i>	+/+	+/+	+/+	+/+	++	-/-
<i>smcR</i>	+/+	+/+	+/+	+/+	+/+	+/+
<i>relA</i>	+/+	+/+ ^a	+/+	+/+ ^a	+/+	-/+
<i>hfq</i>	+/+	+/+	+/+	+/+	+/+	+/+
<i>viuB</i>	+/+	+/+	+/+	-/-	-/-	-/-
<i>trkA</i>	+/+	+/+	+/-	+/-	+/-	+/-
<i>vvpE</i>	-/-	+/+	+/+	+/+	+/+	+/+

^a One replicate was positive for gene expression, while the other replicate was negative.

(Table 2). This result falls in line with previous research that indicates that bacteria respond immediately to osmotic upshifts by inducing a massive uptake of K⁺, which is quickly followed by downregulation of Trk and efflux of potassium (4). However, this effect on *trkA* expression was not observed at high salinity as *trkA* remained on throughout the study (Table 2) and was, in fact, one of only two genes that remained on at day 30 (data not shown).

Recently, Chen et al. (6) examined the role of *trkA* in *V. vulnificus* growth under a variety of K⁺ concentrations. The use of a *trkA* mutant demonstrated reduced growth compared to the wild type in media with moderate K⁺ concentrations (1 to 20 mM), suggesting that *trkA* is required for normal growth under these conditions. It has been estimated that the concentration of K⁺ in 35 ppt seawater is around 10.2 mmol/kg (2). Therefore, the K⁺ levels at the high- and moderate-salinity sites would fall into the "moderate" range, where *trkA* has been shown to influence growth, indicating that there should be no differences in expression between the two sites. However, these studies were conducted in vitro and it is known that the environment persistence of *V. vulnificus* at various salinities is dependent upon temperature (33). Examination of the two in situ sampling sites used in this study revealed that the temperature (24.1°C) and salinity (31 ppt) conditions of the high-salinity site would lead to lower levels of *V. vulnificus*, and these differences in survival are likely correlated with differences in gene expression. Therefore, it is possible that the observed differences in *trkA* expression between the two sampling sites may be related to the normal survival cycles of the bacterium. Furthermore, it is known that many bacteria, including *V. vulnificus*, have multiple K⁺ uptake systems (6, 7). Therefore, as redundant systems are present, the observed differences in the in situ expression of *trkA* in *V. vulnificus* may also be linked to the function of other K⁺ uptake systems, and not necessarily to *trkA*.

The third gene whose expression differed between the high- and moderate-salinity sites in C7184/K2 was *relA*, which encodes a synthase for the alarmone ppGpp. At moderate salinity, the expression of *relA* remained detectable throughout the entire study but dropped below the limit of detection somewhat earlier (by 24 h) at high salinity (Table 2). There is no evidence that the expression of *relA* is influenced by changing salinities but rather is activated by amino acid and carbon or energy limitation (13). Therefore, the observed differences in *relA* expression in *V. vulnificus* may have to do with other

TABLE 4. Gene expression by K853 over time

Gene	Expression at high/moderate salinity					
	0 min	30 min	4 h	12 h	24 h	48 h
<i>rpoS</i>	-/-	+/+	+/+	+/+	+/+	+/+
<i>groEL</i>	+/+	+/+	+/-	+/+	+/+	-/-
<i>smcR</i>	+/+	+/+	+/+	+/+	+/+	+/+
<i>relA</i>	+/+	+/+	+/+	+/+	+/+ ^a	-/+
<i>hfq</i>	+/+ ^a	+/+	+/+	+/+	+/+	-/+
<i>viuB</i>	+/+	+/+	+/+	-/-	-/-	-/-
<i>trkA</i>	+/+	+/+	+/+	+/-	+/+	+/-
<i>vvpE</i>	+/+	+/+	+/+	-/+	-/+	-/-

^a One replicate was positive for gene expression, while the other replicate was negative.

physical factors that differed between the sites (i.e., quantities and/or types of nutrients, etc.) rather than salinity.

Differences in *relA* expression were also observed between the wild-type and mutant strains, but only at high salinity. As with *viuB* expression, this gene dropped below the limit of detection earlier in the parent (after 12 h) than in the mutant strains (through 24 h; (Tables 2 to 4)). It is difficult to make a direct link between differences in expression and the individual mutations; however, it is possible that having all of the stress response systems intact in the parent strain allows the bacterium to respond to environmental stresses more rapidly than strains that have lost one of those systems.

Overall, the *relA* results suggest that estuarine waters with moderate to high salinities may not be as stressful for *V. vulnificus* as might be expected given the involvement of *relA* in stress responses. This hypothesis is supported by the survival data for the *rpoS* mutant (discussed above), as well as the expression data for *groEL*. GroEL is another major molecular chaperone that is activated during stress (i.e., heat, osmotic, or pH stress), when misfolding of proteins can occur (41). Its potential importance in *V. vulnificus* is evidenced by the two complete copies found in both sequenced strains of the bacterium (at VV1_1259-1260 and VV2_1134-1135 in CMCP6 and at VV3106-3107 and VVA1659-1660 in YJ016). However, in situ expression of this gene dropped below the limit of detection after 24 h at both salinities in all of the strains (Tables 2 to 4), again suggesting that these stress-related factors are not required for the long-term survival of this bacterium in the marine environment. Unfortunately, this assertion is not supported by the *rpoS* expression data. It was found that this gene remained on throughout the entire study (Table 2), but this does not necessarily negate the hypothesis that the environment is less stressful than once assumed given that *rpoS* is known to be constitutively expressed in *V. vulnificus* in vivo (S. Limthammahisorn, C. Arias, and Y. Brady, presented at the 107th General Meeting of the American Society for Microbiology, 2007). Studies are currently under way in our laboratory to quantitatively measure *rpoS* expression in *V. vulnificus* under a variety of conditions over time in order to confirm the constitutive expression of this gene.

The remaining genes examined in this study (*hfq*, *vvpE*, *groEL*, and *smcR*) did not show differences in expression between the sampling sites in the parent strain, and only two of these genes (*hfq* and *vvpE*) demonstrated differences in expression between the parent and one or both of the mutant strains.

The *hfq* gene encodes an RNA binding protein that acts as a chaperone for stress-related genes and stimulates *rpoS* production (8). In situ expression of *hfq* ceased earlier in both mutants (after 24 h) than in the wild type (on through 48 h). The wide role that *hfq* plays in the physiology and virulence of bacteria makes it difficult to pinpoint the exact reasons for differences in gene expression (Tables 2 to 4); however, results do suggest that loss of either RpoS or OxyR can lead to differences in gene regulation compared to the parent strain. Studies have shown that *hfq* regulates *rpoS*, but there are no data to suggest that changes in *rpoS* expression influence the expression of *hfq*. However, the observation that *hfq* expression is lost in the *oxyR* mutant while *rpoS* expression remains on is not unusual, as it has been shown that *rpoS* continues to be expressed in the absence of *hfq*, albeit at low levels (5, 23). The use of quantitative RT-PCR will be beneficial in further establishing this *rpoS*-*hfq* connection in *V. vulnificus*.

When examining the expression of *vvpE*, differences were only found between the parent and one of the mutant strains. *vvpE* expression in C7184/K2 remained on through 48 h at both salinities (Table 2). However, in the *oxyR* mutant, *vvpE* expression stopped between 4 and 12 h at high salinity and between 24 and 48 h at moderate salinity (Table 4). In *V. vulnificus*, *vvpE* encodes a protease that is thought to be involved in virulence but is not a definitive virulence factor of this bacterium (14). Research examining the expression of *vvpE* in vitro showed that its expression is reduced in iron-limited media (reported as $\leq 1.0 \mu\text{g}/\text{dl}$) and suggests that iron is required for the efficient transcription of *vvpE* (40). However, these results are not supported by our in situ expression data, which show that, even at the extremely low iron concentrations that are found in the marine environment, *vvpE* expression remains on for extended periods. These results once again highlight the marked differences in gene expression between in vitro and in situ studies.

Overall, our study highlights the obvious need for in situ examination of gene expression in *V. vulnificus*. The environmental expression and regulation of genes are undoubtedly complex, and while in vitro studies have significant value for indicating genes and pathways that are important for *V. vulnificus* survival, they are unable to adequately mimic what actually occurs in the natural environment. This is evidenced by the differences found between this in situ study of gene expression and previously conducted studies examining the involvement of singular genes under in vitro conditions (6, 40). Furthermore, this study demonstrated that previous assumptions about the stressful nature of the estuarine environment may also be inaccurate, at least to some degree. This was particularly evident when evaluating the survival of the *rpoS* mutant strain, where results showed that loss of this stress system did not adversely affect the long-term (30 days) survival of *V. vulnificus*. It is likely, however, that the conditions under which survival was tested in this study did not stress the bacterium but that other changes in the natural environment (higher or lower salinity, pH, temperature, etc.) may provide sufficient stresses for which *rpoS* would be required. For this reason, we are continuing such studies in this laboratory in an attempt to better understand how *V. vulnificus* relates to its natural habitat.

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