

# A *Vibrio cholerae* LysR Homolog, AphB, Cooperates with AphA at the *tcpPH* Promoter To Activate Expression of the ToxR Virulence Cascade

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**We describe here a new member of the LysR family of transcriptional regulators, AphB, which is required for activation of the *Vibrio cholerae* ToxR virulence cascade. AphB activates the transcription of the *tcpPH* operon in response to environmental stimuli, and this process requires cooperation with a second protein, AphA. The expression of neither *aphA* or *aphB* is strongly regulated by environmental stimuli, raising the possibility that the activities of the proteins themselves may be influenced under various conditions. Strains of the El Tor biotype of *V. cholerae* typically exhibit lower expression of ToxR-regulated virulence genes in vitro than classical strains and require specialized culture conditions (AKI medium) to induce high-level expression. We show here that expression of *aphB* from the *tac* promoter in El Tor biotype strains dramatically increases virulence gene expression to levels similar to those observed in classical strains under all growth conditions examined. These results suggest that AphB plays a role in the differential regulation of virulence genes between the two disease-causing biotypes.**

Cholera is a life-threatening diarrheal disease caused by the gram-negative bacterium *Vibrio cholerae*. The organism colonizes the upper intestine, and the toxin-coregulated pilus (TCP) is the primary factor involved in this process (36). The severe diarrhea associated with the disease results from the action of the secreted cholera toxin (CT) on intestinal epithelial cells (reviewed in reference 17). The genes required for the biogenesis of TCP are located in an operon on a large pathogenicity island termed the TCP-ACF element, or vibrio pathogenicity island (18, 19). The subunits of CT are encoded by the *ctxA* and *ctxB* genes on a separate genetic element which comprises the genome of the lysogenic filamentous bacteriophage CTX $\phi$  (38).

Many of the genes involved in the pathogenesis of *V. cholerae* comprise what is known as the ToxR virulence regulon, since they are coordinately expressed and dependent upon the transcriptional activator ToxR (23, 26). ToxR is a transmembrane DNA binding protein whose activity is enhanced by a second transmembrane protein, ToxS (5, 21, 23). The *toxR* and *toxS* genes, which are expressed as an operon, are not associated with either the TCP-ACF or CTX elements but appear to be part of the “ancestral chromosome” and have other important regulatory roles (22). TcpP is a transcriptional activator encoded on the TCP-ACF element which has recently been shown to share significant homology with ToxR and which cooperates with it to initiate gene expression (13, 25). The *tcpP* gene is coexpressed with a second gene, *tcpH*, which encodes a protein that enhances the activity of TcpP (2). TcpP and TcpH appear to have a similar membrane topology to ToxR and ToxS.

ToxRS and TcpPH control the expression of the ToxR virulence regulon by their ability to activate the expression of a third transcriptional activator, ToxT, which is also encoded on the TCP-ACF element (7, 13). ToxT is a cytoplasmic protein

that is a member of the AraC family of transcriptional activators (15). Once its expression is activated by ToxRS and TcpPH, ToxT then directly activates various genes within the regulon, such as the *tcp* and *ctx* operons (3, 7). The *toxT* gene is located within the *tcp* operon, and its expression is dependent upon a promoter located immediately upstream of the gene (14) as well as by one located at the beginning of the *tcp* operon which may function in an autoregulatory capacity (1).

The expression of the *tcp* and *ctx* operons are strongly influenced by specific environmental cues such as pH and temperature. Since the expression of *tcpPH* is also influenced by both of these parameters (2, 32), the mechanisms that regulate the expression of this operon are likely to be of central importance in the control of virulence gene expression by environmental stimuli. AphA is a 20-kDa *V. cholerae* protein which has recently been shown to be required for expression of the *tcpPH* operon and for its response to environmental stimuli (32). Since the basal level of *tcpPH* expression in a  $\Delta$ *aphA* mutant still appeared to be influenced by pH and temperature, it was hypothesized that factors in addition to AphA might also play a role in the expression of *tcpPH*. We describe here a new member of the LysR family of transcriptional regulators, AphB, which is required for transcriptional activation of *tcpPH* as well as its response to environmental stimuli. AphB functions synergistically with AphA to activate the expression of *tcpPH*, and it also appears to contribute to the differences in virulence gene expression between the two major disease-causing biotypes, classical and El Tor. Since neither AphA nor AphB is encoded within the TCP-ACF element, these proteins may have other regulatory roles in *V. cholerae*, and the expression of the *tcpPH* operon may have evolved to come under their control.

## MATERIALS AND METHODS

**Bacterial strains and media.** The *V. cholerae* and *Escherichia coli* strains and plasmids used in this study are listed in Table 1. Bacteria were maintained at  $-70^{\circ}\text{C}$  in Luria-Bertani (LB) medium (20) containing 30% (vol/vol) glycerol. Antibiotics were used at the following concentrations in LB medium or AKI medium (16): ampicillin, 100  $\mu\text{g/ml}$ ; kanamycin, 45  $\mu\text{g/ml}$ ; tetracycline, 7.5  $\mu\text{g/ml}$  for *V. cholerae* and 15  $\mu\text{g/ml}$  for *E. coli*; and streptomycin, 100  $\mu\text{g/ml}$ , except

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TABLE 1. Bacterial strains and plasmids used in this study

Strain or plasmid	Relevant genotype	Source or reference
<b>Strains</b>		
<i>V. cholerae</i>		
CG842	O395 (classical Ogawa Sm <sup>r</sup> ) $\Delta lacZ$	4
KSK218	CG842 <i>ctx-lacZ</i> Sm <sup>r</sup> Cm <sup>r</sup>	30
KSK404	KSK218 <i>aphB::TnphoA</i>	This work
GK91	KSK218 <i>aphB::pGKK17</i>	This work
GK122	KSK218 $\Delta aphB1$	This work
KSK618	CG842 <i>tcpP-lacZ</i>	32
GK121	KSK618 $\Delta aphB1$	This work
KSK647	KSK618 $\Delta aphA1$	32
KSK805	KSK618 $\Delta aphA1 \Delta aphB1$	This work
KSK666	CG842 <i>aphA-lacZ</i>	32
GK130	CG842 <i>aphB-lacZ</i>	This work
C6706str2	El Tor Inaba Sm <sup>r</sup>	37
KSK262	C6706str2 $\Delta lacZ3$	This work
KSK725	KSK262 <i>tcpP-lacZ</i>	This work
GK138	KSK725 $\Delta aphB1$	This work
GK161	KSK725 $\Delta aphA1$	This work
GK142	KSK262 <i>aphB-lacZ</i>	This work
<i>E. coli</i>		
MC1061	$\Delta (ara-leu)7697 \Delta (lac)X74$	Laboratory collection
KSK782	MC1061 $\lambda$ KSPL1 ( <i>tcpP-lacZ</i> )	This work
<b>Plasmids</b>		
pKAS64	pKAS32 $\Delta rpsL$ , Ap <sup>r</sup>	31
pKAS110	pKAS64 $\Delta SmaI$	This work
pGKK17	pKAS110, 200-bp <i>aphB</i> fragment	This work
pGKK18	pGKK17 chromosomal capture plasmid	This work
pGKK25	pKAS46, $\Delta aphB1$ classical	This work
pGKK26	pGKK25, <i>aphB1-lacZ</i>	This work
pGKK28	pKAS46, $\Delta aphB1$ El Tor	This work
pGKK29	pGKK28, <i>aphB1-lacZ</i>	This work
pGKK35	pKAS46, $\Delta aphA1$ El Tor	This work
pKAS48	pKAS46, $\Delta lacZ3$	29
pKAS113	pKAS46, <i>tcpP-lacZ</i> El Tor	This work
pLAFR3	Tc <sup>r</sup> expression plasmid	33
pKAS116	pLAFR3 <i>aphB</i> (classical), Tc <sup>r</sup> Gm <sup>r</sup>	This work
pMMB66EH	Ap <sup>r</sup> expression plasmid	9
pKAS107	pMMB66EH <i>aphA</i> (classical), Ap <sup>r</sup>	32
pKAS117	pMMB66EH <i>aphB</i> (classical), Ap <sup>r</sup>	This work
pBAD22	Ap <sup>r</sup> expression plasmid	11
pKAS118	pBAD-TOPO <i>aphB</i> (classical), Ap <sup>r</sup>	This work
pKAS119	pBAD-TOPO <i>aphA</i> (classical), Ap <sup>r</sup>	This work
pKAS120	pBAD-TOPO <i>aphB</i> (El Tor), Ap <sup>r</sup>	This work

when selecting for loss of integrated plasmids in *V. cholerae*, where it was used at 1 mg/ml. 5-Bromo-4-chloro-3-indolyl- $\beta$ -D-galactopyranoside (X-Gal) was used in LB agar at 40  $\mu$ g/ml.

**Identification of *aphB*.** Random insertion of *TnphoA* into the chromosome of strain KSK218 was as previously described (30, 35, 36). Chromosomal DNA from *V. cholerae* transposon mutant KSK404 was digested with *SphI* and ligated into an *oriR6K* plasmid lacking *rpsL* (pKAS64). The ligated DNA was subjected to two rounds of PCR amplification, the first using a plasmid-specific primer, *ORIR6K* (5'-GGTTTAAACGGTTGTGGACAAC), and a transposon-specific primer, *TNPHOA-1* (30); and the second using *ORIR6K* with a nested transposon-specific primer, *TNPHOA-2* (5'-AGCAGCCGGTTTCCAGAAC). The resulting 200-bp fragment, which contained a portion of the *aphB* open reading frame, was ligated into another *oriR6K* plasmid lacking *rpsL* (pKAS110), generating pGKK17. Plasmid pGKK17 was integrated into the *aphB* gene of KSK218, generating strain GK91. Chromosomal DNA was isolated from GK91, digested with *SphI*, ligated, and transformed into *E. coli*. The resulting plasmid, pGKK18, was then used to obtain the complete *aphB* nucleotide sequence with the ABI PRISM Dye System (Perkin-Elmer).

**Construction of in-frame deletions and *lacZ* fusions.** The in-frame  $\Delta aphB1$  mutations in both classical and El Tor biotypes were constructed by PCR amplifying two 200-bp fragments encompassing the regions upstream and downstream of the *aphB* gene, respectively, from either O395 or C6706str2 (37) by using primer pair CO2-3 (5'-GATCGTCTAGAAATGGTTTTCAATAAATCA

TC) and CO2-4 (5'-GATCGGCGGCCGCATGTTCATTGAAGCGAGACGC TC) and primer pair CO2-5 (5'-GATCGGCGGCCGCCTGTATAACCACAA AGATCAC) and CO2-6 (5'-GATCGGAATTCAGCCATGCAAATGGCGG CC). The resulting fragments were ligated into pKAS46 (29), generating pGKK25 and pGKK28, respectively, and the deletions were introduced into *V. cholerae* by allelic exchange. To construct the *aphB-lacZ* fusions, a promoterless *E. coli lacZ* gene was inserted into the plasmids described above, generating pGKK26 and pGKK29, prior to allelic exchange. The classical  $\Delta aphA1$  deletion was previously described (32). The El Tor  $\Delta aphA1$  deletion was constructed in a similar manner, except that primer YF-7 (5'-GATCGGAATTCACCATGTCA TTACCACACGTTATCC) was used in place of YF-1 and the fragments were ligated into pKAS46, generating pGKK35, prior to allelic exchange.

**Construction of chromosomal *tcpP-lacZ* fusions.** Plasmid pKAS48 (29) was used to construct the  $\Delta lacZ3$  deletion in El Tor strain C6706str2 (37) by allelic exchange, generating strain KSK262. The El Tor *tcpP-lacZ* operon fusion in KSK262 was constructed in a manner similar to that of the classical *tcpP-lacZ* fusion (32), except that primers TP-BAME (5'-GATCGGGATCCAGTAATG CCGGTAATTCATG) and TP-SEE (5'-GATCGGTGCAGCAATTCAGCC GTTAGCAGCTTGTAAAG) were used in place of TP-BAM and TP-SE for amplification from C6706str2. The resulting fusion in plasmid pKAS113 was introduced into *V. cholerae* by allelic exchange. The *tcpP-lacZ* fusion on  $\lambda$  KSPL1 was previously described (32).

**Construction and mobilization of expression plasmids.** The expression plasmids constructed in this study are listed in Table 1. The *aphB* gene was amplified from either the classical (O395) or El Tor (C6706str2) biotypes by using primers CO2-7 (5'-GATCGGAATTCATAAATTAGCGATAGTTGC) and CO2-8 (5'-GATCGAAGCTTGAAGGCGCGAAGCC). The *aphA* gene was amplified from O395 by using primers YF-5 (5'-GATCGGAATTCATAAATGCGT TGATATGCGTGCC) and YF-6 (32). Plasmids derived from pLAFR3 (33), pMMB66EH (9), and pBAD-TOPO (Invitrogen), respectively, were introduced into *V. cholerae* by mating with *E. coli* SM10 (28), triparental mating with *E. coli* MM294 carrying pRK2013 (8), and electroporation.

**$\beta$ -Galactosidase assays.**  $\beta$ -Galactosidase assays (20) were carried out with *tcpP-lacZ*, *aphA-lacZ*, or *aphB-lacZ* fusion strains during mid-logarithmic growth and with *ctx-lacZ* fusion strains after overnight growth. In AKI medium, cultures were assayed after 4 h without rotation. The bicinchoninic acid procedure (Pierce) was used to determine the total amount of protein in each reaction from the overnight cultures. The data are averaged results from at least two experiments.

**Immunoblot analysis.** Cell extracts from overnight cultures were subjected to sodium dodecyl sulfate–12.5% polyacrylamide gel electrophoresis, transferred to nitrocellulose, and probed with anti-TcpA antibody (34) by using the ECL (enhanced chemiluminescence) detection system (Amersham).

**Nucleotide sequence accession number.** The accession number for the nucleotide sequence of *aphB* in GenBank is AF148502.

## RESULTS

**The *aphB* gene is required for virulence gene expression.** *V. cholerae* TnphoA mutant KSK404 was identified as a derivative of the *ctx-lacZ* fusion strain KSK218, which showed reduced  $\beta$ -galactosidase production under environmental conditions normally optimal for its expression (i.e., LB medium [pH 6.5] at 30°C) and failed to produce TCP. A 200-bp DNA fragment encompassing the region adjacent to the transposon in KSK404 was obtained by ligating restriction-digested chromosomal DNA from the mutant into a plasmid and performing two rounds of PCR with primers specific for the plasmid and for the transposon. The DNA fragment, which contained a portion of the *aphB* open reading frame, was then inserted into an oriR6K plasmid and used to disrupt the wild-type *aphB* gene in KSK218. After confirming that the *aphB* disruption in the resulting strain, GK91, caused a defect in virulence gene expression similar to that of the original transposon mutant, the entire *aphB* gene was isolated from this strain by using chromosomal capture (30, 31) and sequenced.

To verify that the disruption of *aphB* was solely responsible for the defect in virulence gene expression in strain GK91, an in-frame deletion of *aphB* was constructed in KSK218 (*ctx-lacZ*), strain GK122, and this defect was complemented by inducing a wild-type *aphB* gene expressed from the *tac* promoter of plasmid pKAS117. As shown in Fig. 1, the  $\Delta$ *aphB* mutation in GK122 significantly reduced the production of  $\beta$ -galactosidase under inducing conditions (LB medium [pH 6.5] at 30°C) and expression of *aphB* from pKAS117 restored its production to wild-type levels under these conditions. The mutation had only a small effect on the already low levels of  $\beta$ -galactosidase under repressing conditions (LB medium [pH 8.5] at 30 or at 37°C). However, expression of *aphB* from pKAS117 increased  $\beta$ -galactosidase production at pH 8.5 at 30°C in both the parental strain and the  $\Delta$ *aphB* mutant to close to the levels observed under inducing conditions. These results indicate that *aphB* plays a role in activating *ctx* expression and that it is also involved in its regulation by environmental stimuli such as pH. Induction of *aphB* from pKAS117 also increased  $\beta$ -galactosidase production at 37°C, but to a smaller extent than at pH 8.5 at 30°C (Fig. 1).

The influence of AphB on the production of TCP, shown in Fig. 2, is similar to the above results observed with *ctx*. The  $\Delta$ *aphB* mutation in strain GK122 prevented the production of the 20.5-kDa major pilin protein TcpA under inducing conditions (Fig. 2, lane 3). Induction of *aphB* expression from

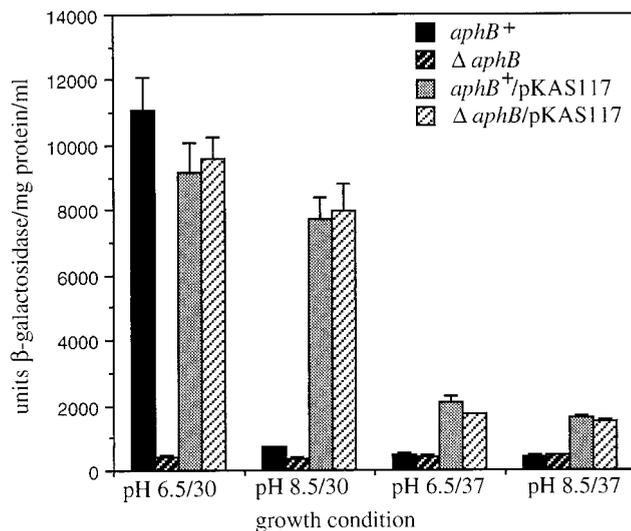


FIG. 1. Influence of AphB on the expression of a *ctx-lacZ* fusion. Cultures were grown in LB medium at pH 6.5 or 8.5 at 30 or 37°C. Those with pKAS117 also contained 1 mM isopropyl- $\beta$ -D-thiogalactopyranoside. Black bars, KSK218; striped black bars, GK122 ( $\Delta$ *aphB*); gray bars, KSK218 with pKAS117 (AphB); striped gray bars, GK122 with pKAS117 (AphB).

pKAS117 in this mutant restored TcpA production (Fig. 2, lane 4) and permitted the cells to autoagglutinate in culture, a property associated with wild-type levels of TCP. Thus, AphB influences the expression of both the *ctx* and *tcp* operons in *V. cholerae*.

**AphB activates the expression of the *tcpPH* promoter.** The significant impact of *aphB* on the expression of the *ctx* and *tcp* genes prompted us to investigate whether genes required earlier in the virulence cascade, *toxRS*, *tcpPH*, or *aphA*, were also influenced by AphB. The  $\Delta$ *aphB* mutant GK122 did not produce the outer membrane protein OmpT in place of OmpU (data not shown), suggesting that *toxR* expression was not altered in the strain (22). To assess its effects on the expression of *tcpPH* and *aphA*, the  $\Delta$ *aphB* mutation was introduced into the classical *tcpP-lacZ* fusion strain KSK618 and the classical *aphA-lacZ* fusion strain KSK666 (32). Table 2 shows that the expression of the *tcpP-lacZ* fusion in the  $\Delta$ *aphB* strain, GK121, was significantly reduced under each environmental condition examined relative to the parental strain. Furthermore, the basal level of *tcpPH* expression in the absence of *aphB* did not significantly respond to environmental stimuli. When *aphB* was induced from the *tac* promoter of pKAS117, the expression of *tcpP-lacZ* in the  $\Delta$ *aphB* mutant GK121 was increased under all environmental conditions examined (Table 2). This increase was most dramatic under the strongest repressive condition,

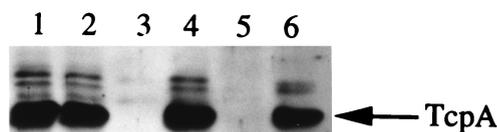


FIG. 2. AphB influences TcpA production in both classical and El Tor biotypes. Samples were prepared from KSK218 (classical [lane 1]), KSK218 with pKAS117 (AphB [lane 2]), GK122 (classical  $\Delta$ *aphB* [lane 3]), GK122 with pKAS117 (AphB [lane 4]), KSK262 (El Tor [lane 5]), and KSK262 with pKAS117 (AphB [lane 6]). Cultures were grown overnight in LB medium (pH 6.5) at 30°C. Those with pKAS117 also contained 1 mM isopropyl- $\beta$ -D-thiogalactopyranoside. Samples were analyzed by Western blotting with anti-TCP antiserum (34). TcpA is indicated by the arrow to the right.

TABLE 2. Activation of a classical biotype *tcpP-lacZ* fusion by AphB and AphA

Strain <sup>a</sup>	$\beta$ -Galactosidase activity <sup>b</sup> at pH and temp:			
	6.5, 30°C	8.5, 30°C	6.5, 37°C	8.5, 37°C
KSK618 ( <i>tcpP-lacZ</i> )	3,940 $\pm$ 24	931 $\pm$ 8	2,091 $\pm$ 31	502 $\pm$ 11
GK121 ( $\Delta$ <i>aphB</i> )	190 $\pm$ 22	145 $\pm$ 3	133 $\pm$ 8	98 $\pm$ 15
KSK647 ( $\Delta$ <i>aphA</i> )	350 $\pm$ 75	104 $\pm$ 4	201 $\pm$ 6	85 $\pm$ 13
KSK805 ( $\Delta$ <i>aphA</i> $\Delta$ <i>aphB</i> )	117 $\pm$ 7	97 $\pm$ 1	94 $\pm$ 7	74 $\pm$ 9
GK121(pKAS117) (AphB)	6,271 $\pm$ 680	5,428 $\pm$ 968	3,415 $\pm$ 74	2,616 $\pm$ 139
KSK647(pKAS117) (AphB)	3,283 $\pm$ 201	3,076 $\pm$ 408	1,669 $\pm$ 110	1,022 $\pm$ 61
GK121(pKAS107) (AphA)	502 $\pm$ 30	382 $\pm$ 88	288 $\pm$ 7	249 $\pm$ 1
KSK647(pKAS107) (AphA)	5,732 $\pm$ 723	2,589 $\pm$ 142	3,660 $\pm$ 846	1,816 $\pm$ 214

<sup>a</sup> All plasmids were induced with 1 mM isopropyl- $\beta$ -D-thiogalactopyranoside.

<sup>b</sup> Units per optical density at 600 nm of culture.

pH 8.5 at 37°C. These results indicate that AphB is required for the activation of the *tcpPH* operon in *V. cholerae* and for its response to environmental stimuli. The  $\Delta$ *aphB* mutation had no effect on the expression of the *aphA-lacZ* fusion in *V. cholerae* (data not shown), indicating that *aphB* is not influencing *tcpPH* expression indirectly through AphA.

#### AphB cooperates with AphA to activate *tcpPH* expression.

AphA has previously been shown to be required for activation of the *tcpPH* operon (32). The expression of *tcpPH* in a  $\Delta$ *aphA* mutant, strain KSK647 (32), is similar to that of the  $\Delta$ *aphB* mutant, except that the basal level of expression is still somewhat responsive to environmental stimuli (Table 2). Thus, loss of either AphA or AphB results in a dramatic decrease in the expression of the *tcpPH* operon. To determine if increased amounts of either protein could compensate for loss of the other, the *aphB* expression plasmid pKAS117 was introduced into the  $\Delta$ *aphA* mutant KSK647 and the *aphA* expression plasmid pKAS107 (32) was introduced into the  $\Delta$ *aphB* mutant GK121. Interestingly, high levels of AphB in the  $\Delta$ *aphA* mutant restored *tcpPH* expression to close to wild-type levels at pH 6.5 (and to greater than wild-type levels at pH 8.5) (Table 2), whereas high levels of AphA in the  $\Delta$ *aphB* mutant increased *tcpPH* expression somewhat, but did not restore it to wild-type levels. Thus, when present in sufficient amounts, either protein is capable of activating *tcpPH* transcription in the absence of the other, but AphA still requires AphB to achieve wild-type expression levels.

To further address whether AphA and AphB function sequentially or in separate pathways to activate *tcpPH* expression, the  $\Delta$ *aphA* mutation was introduced into the  $\Delta$ *aphB* mutant GK121, generating strain KSK805. The finding that the expression of *tcpPH* is lower in the double mutant under all environmental conditions than in either single mutant (Table 2) suggests that AphA and AphB function cooperatively to activate *tcpPH* transcription rather than sequentially. This notion is further supported by the results in Fig. 3 which show that in the  $\Delta$ *aphA*  $\Delta$ *aphB* double mutant, KSK805, and in an *E. coli* *tcpP-lacZ* fusion strain, KSK782, the presence of AphA and AphB together from plasmids pKAS119 and pKAS116 results in higher levels of  $\beta$ -galactosidase production than with either protein alone. Thus, it appears that AphA and AphB function synergistically to activate transcription at the *tcpPH* promoter.

**AphB is a LysR homolog.** The LysR family represents one of the most common types of prokaryotic transcriptional regulators. These proteins typically interact with small specific signal molecules known as coinducers to activate the expression of divergent or unlinked target genes which function in many diverse processes (for a review, see reference 27). Mem-

bers of this family show strong homology in their amino-terminal domains, much of which derives from conservation of a helix-turn-helix DNA-binding motif. AphB exhibits significant amino-terminal homology with a large number of these proteins, and an alignment of this region of AphB with several LysR family members is shown in Fig. 4. The *aphB* gene encodes a protein of 291 amino acids with a predicted molecular mass of 33.3 kDa. Two of the proteins with the strongest overall homology to AphB (27%) are PtxR, a positive regulator of exotoxin A production in *Pseudomonas aeruginosa* (12); and IrgB from *V. cholerae*, which positively regulates the expression of *irgA* in response to iron limitation (10).

**AphB activates *tcpPH* expression in the El Tor biotype.** The expression of the ToxR virulence regulon in classical biotype strains is maximal in LB medium (pH 6.5) at 30°C. Strains of the El Tor biotype show reduced expression of the regulon under these conditions and require a bicarbonate-containing

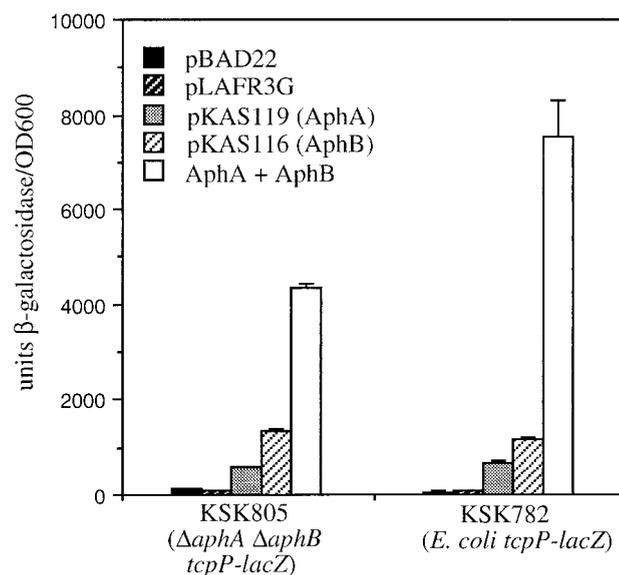


FIG. 3. Cooperation between AphA and AphB enhances *tcpP-lacZ* expression. *V. cholerae* KSK805 ( $\Delta$ *aphA*  $\Delta$ *aphB*) (left) was grown in LB medium (pH 6.5) at 30°C, and *E. coli* KSK782 (*tcpP-lacZ*) (right) was grown in LB medium (pH 7.0) at 37°C. Black bars, pBAD22 plus 0.2% arabinose; striped black bars, pLAFR3G plus 1 mM isopropyl- $\beta$ -D-thiogalactopyranoside (IPTG); gray bars, pKAS119 (AphA) plus 0.2% arabinose; striped gray bars, pKAS116 (AphB) plus 1 mM IPTG; open bars, pKAS119 plus pKAS116 (AphA plus AphB) plus 0.2% arabinose plus 1 mM IPTG. OD600, optical density at 600 nm.

M-LRHLRYFVAVAEH-S FTAAAE-LHVSQPT-SRO-R-LEDDLGV-LLE consensus  
 MKLDDLNLPRLLVVENGSTSTSKKTIIPVATITRRITQALEDLNLRLLN AphB  
 QDLSAVKAFHALCQHKSLTAAARALEQPKSTLSRRLAQLEEDLGQSLLM IrgB  
 LNLNHLIYAFVAVAEHNSTAAAEALGLSKSLLEQQLRRLEADLGIQLLT PtxR  
 VNLRLHIEIFHAVMTAGSLTEAAHLLHTSQPTVSRELARFEKVIIGLKLFE LysR

FIG. 4. Alignment of the amino-terminal region of *V. cholerae* AphB with those of several other members of the LysR family. The helix-turn-helix domain is underlined.

medium (AKI medium) at 37°C for high-level expression (16). Table 3 shows that the expression of an El Tor *tcpP-lacZ* fusion, strain KSK725, is significantly reduced in LB medium relative to the classical *tcpP-lacZ* fusion, strain KSK618 (Table 2), under all of the conditions examined. Although growth of the El Tor strain in AKI medium improved the expression of *tcpPH*, it was still significantly lower than that of the classical strain in LB medium (pH 6.5) at 30°C. To determine if AphB also activates *tcpPH* expression in the El Tor biotype, a  $\Delta$ *aphB* mutation was introduced into KSK725, generating strain GK138. The  $\Delta$ *aphB* mutation in this strain significantly decreased *tcpPH* expression under AKI conditions (Table 3), but had a smaller effect on the already low levels of expression in LB medium. A similar result was observed with a  $\Delta$ *aphA* mutation in this background, strain GK161 (Table 3). Thus, although the response of *tcpPH* in El Tor strains to environmental stimuli is different from that in classical strains, *aphA* and *aphB* play a role in its expression in both biotypes.

The mechanisms responsible for the differential expression of *tcpPH* in classical and El Tor biotype strains are not yet understood. Since expression of either *aphA* or *aphB* from the *tac* promoter significantly increased *tcpPH* expression under normally nonpermissive expression conditions in the classical biotype, it was of interest to determine whether either of these genes could increase *tcpPH* expression in the El Tor biotype as well. Table 3 shows that induction of *aphB* from pKAS117 in the El Tor fusion strain KSK725 increased *tcpPH* expression in both AKI and LB media. In LB medium, the levels of expression of *tcpPH* in the presence of pKAS117 were virtually identical to those of the classical *tcpP-lacZ* fusion strain (Table 2). El Tor strain KSK262 does not produce TCP detectable even by Western blotting when grown in LB medium (pH 6.5) at 30°C (Fig. 2, lane 5). However, induction of *aphB* expression from pKAS117 in KSK262 increased TCP production in LB medium (pH 6.5) at 30°C to a level similar to that of classical strains (Fig. 2, lane 6) and permitted the cells to autoagglutinate. Expression of *aphA* from pKAS107 also increased the expression of the El Tor *tcpP-lacZ* fusion, but to a lesser extent than *aphB* (Table 3), and did not permit strain KSK262 to produce TCP by Western blotting (data not shown). These findings indicate that the *tcpPH* promoter can be activated by

AphB and, to a lesser extent, AphA, in the El Tor biotype under conditions not normally permissive for its expression.

**The AphB protein and its expression appear similar in both biotypes.** The significant effect of inducing *aphB* expression from pKAS117 on the activation of the El Tor *tcpPH* promoter in LB medium (pH 6.5) at 30°C suggested that, in this biotype, the AphB protein or its expression might be different from that in the classical biotype. The deduced amino acid sequences of the classical and El Tor AphB proteins, however, were found to be identical. In addition, when either the classical or El Tor *aphB* gene was induced from an arabinose promoter in plasmid pKAS118 or pKAS120, respectively, both activated an *E. coli tcpP-lacZ* fusion approximately 30-fold, suggesting that they are equally functional. To assess the expression of *aphB* in classical and El Tor strains, an *aphB-lacZ* fusion was constructed in each biotype. Table 4 shows that the levels of expression of *aphB* in the classical fusion strain GK130 and the El Tor fusion strain GK142 are similar. Since the AphB proteins from the classical and El Tor strains appear to be equally capable of activating *tcpPH* transcription and the levels of expression of the gene in both biotypes are similar, some other aspect of AphB function may be different in the two biotypes.

It has previously been shown that the expression of *aphA* is not strongly influenced by either pH or temperature (32). The results in Table 4 indicate that the expression of *aphB* is also not strongly influenced by these stimuli, nor does it completely reflect the pattern of expression that is observed with *tcpPH* under the different conditions. For example, expression of the classical *aphB-lacZ* fusion is not higher at pH 6.5 at 37°C than it is at pH 8.5 at 30°C, and expression of the El Tor *aphB-lacZ* fusion is not significantly higher in AKI medium than it is in LB medium at pH 6.5 at 30°C. It is also noteworthy that induction of either *aphA* or *aphB* from the *tac* promoter had no effect on the expression of the *aphB-lacZ* fusion or its response to environmental stimuli (data not shown). Since the expression of *tcpPH* in response to pH or temperature does not appear to solely depend upon the expression of either *aphA* or *aphB* in response to these stimuli, it is possible that the activities of the proteins themselves might be influenced under various conditions.

## DISCUSSION

Activation of the ToxR virulence cascade requires multiple factors encoded both within the "ancestral" *V. cholerae* chromosome and on discrete elements involved in pathogenicity. As shown in Fig. 5, ToxR and ToxS, a chromosomally encoded protein pair, cooperate with the TCP-ACF pathogenicity element-encoded TcpP and TcpH protein pair to positively regulate the expression of the TCP-ACF-encoded regulator, ToxT. ToxT, in turn, activates expression of the *ctx* and *tcp*

TABLE 3. Activation of an El Tor biotype *tcpP-lacZ* fusion by AphB and AphA

Strain <sup>a</sup>	$\beta$ -Galactosidase activity <sup>b</sup> at pH and temp:				
	AKI medium	LB medium			
		6.5, 30°C	8.5, 30°C	6.5, 37°C	8.5, 37°C
KSK725 ( <i>tcpP-lacZ</i> )	1,060 $\pm$ 27	362 $\pm$ 8	172 $\pm$ 0	199 $\pm$ 8	101 $\pm$ 2
GK138 ( $\Delta$ <i>aphB</i> )	129 $\pm$ 8	222 $\pm$ 6	162 $\pm$ 8	122 $\pm$ 8	94 $\pm$ 1
GK161 ( $\Delta$ <i>aphA</i> )	122 $\pm$ 3	148 $\pm$ 1	104 $\pm$ 1	90 $\pm$ 3	70 $\pm$ 1
KSK725(pKAS117) (AphB)	2,129 $\pm$ 37	3,785 $\pm$ 309	1,270 $\pm$ 173	2,053 $\pm$ 168	753 $\pm$ 197
KSK725(pKAS107) (AphA)	1,060 $\pm$ 26	1,689 $\pm$ 17	730 $\pm$ 21	621 $\pm$ 41	284 $\pm$ 11

<sup>a</sup> All plasmids were induced with 1 mM isopropyl- $\beta$ -D-thiogalactopyranoside.

<sup>b</sup> Units per optical density at 600 nm of culture.

TABLE 4. Comparison of *aphB* expression in classical and El Tor biotypes

Strain	$\beta$ -Galactosidase activity <sup>a</sup> at pH and temp:				
	AKI medium	LB medium			
		6.5, 30°C	8.5, 30°C	6.5, 37°C	8.5, 37°C
GK130 (classical <i>aphB-lacZ</i> )	201 $\pm$ 6	176 $\pm$ 3	129 $\pm$ 1	100 $\pm$ 3	78 $\pm$ 1
GK142 (El Tor <i>aphB-lacZ</i> )	223 $\pm$ 2	207 $\pm$ 2	137 $\pm$ 1	98 $\pm$ 6	83 $\pm$ 7

<sup>a</sup> Units per optical density at 600 nm of culture.

operons as part of a virulence gene regulatory cascade. In this report, we describe another chromosomally encoded protein pair required for the activation of the ToxR virulence cascade. AphB is a new member of the LysR family of transcriptional regulators which cooperates with the recently identified AphA protein (32) to activate the expression of the *tcpPH* operon.

*V. cholerae* strains deficient in either *aphA* or *aphB* show reduced expression of the *tcpPH* operon and as a result do not produce virulence factors such as CT and TCP. That an *aphA aphB* double mutant shows lower expression of *tcpPH* than either single mutant suggests that AphA and AphB are not functioning sequentially in the same pathway but that they cooperate to activate *tcpPH* transcription. When expressed from their natural promoters in *V. cholerae*, neither protein significantly activates transcription in the absence of the other. When expressed from inducible promoters on plasmids in *V. cholerae* or *E. coli*, either protein is capable of activating the transcription of *tcpPH* in the absence of the other, with AphB showing a stronger effect than AphA and the former even compensating for the latter in *V. cholerae*. However, the expression of *tcpPH* is significantly greater with the two proteins together than with either one alone. It is possible that the presence of AphA enhances the ability of AphB to activate transcription.

The ToxR virulence regulon is strongly influenced by environmental cues such as pH and temperature. Although the mechanisms responsible for this regulation are not yet understood, the effect of environmental stimuli on the expression of the regulon may largely be the result of their influence over the expression of *tcpPH* (24). How pH and temperature control the expression of *tcpPH* is not yet understood, but AphA and AphB appear to play a role in this process. *V. cholerae* strains containing plasmids expressing either *aphA* or *aphB* show increased *tcpPH* transcription under both permissive and non-permissive environmental conditions. Supplying high levels of either of these two proteins in the presence of the other appears to be sufficient to almost completely override environmental regulation by pH and temperature. Since the expression of neither *aphA* (32) nor *aphB* is strongly regulated by environmental conditions, it is possible that their activities are influenced by them. Many LysR regulators activate gene expression only in the presence of specific coinducer molecules (27). Interaction of such a molecule with AphB only under certain environmental conditions might render it able to activate *tcpPH* transcription if AphA is present. High levels of either AphA or AphB might be sufficient to at least partly overcome the need for a coinducer to facilitate transcriptional activation. Alternatively, when present in high levels, AphA or AphB may effectively compete with other proteins that normally function to downregulate *tcpPH* expression under certain environmental conditions. Additional experiments are necessary in order to distinguish between these possibilities.

It is well established that *V. cholerae* strains of the El Tor biotype exhibit lower expression of the ToxR virulence regulon

in vitro than classical biotype strains. This appears to be the result of reduced expression of *toxT* and *tcpPH* in the El Tor biotype relative to the classical biotype (6, 24) (Table 3). Despite the fact that the expression of *tcpPH* is differentially regulated in classical and El Tor biotypes, *aphA* and *aphB* are involved in the activation of *tcpPH* in both. The observation that expression of *aphB* from the *tac* promoter increased *tcpPH* transcription in the El Tor biotype to classical levels in LB medium and permitted TCP production suggests that AphB might in some respect be different in the two biotypes. However, El Tor biotype strains encode a functional AphB protein and the expression of the gene is similar to that of classical strains. AphA alone does not appear to be responsible for the biotype-specific difference in expression, since induction of the *aphA* gene in the El Tor biotype did not increase *tcpPH* transcription to classical levels and it did not permit TCP production. These results raise the possibility that some other aspect of AphB function may be different in the two biotypes, such as the ability of the protein to assume a conformation that allows it to activate transcription at the *tcpPH* promoter. Experiments to address this issue are currently in progress.

It is not yet known whether AphA and AphB function alone or together in any other regulatory capacity in *V. cholerae*. It is common for LysR transcriptional regulators to be divergently transcribed from a promoter that is close to or that overlaps a regulated target gene (27). For example, the gene encoding the *V. cholerae* IrgB protein is divergently transcribed from the gene which it activates, *irgA* (10). The gene upstream of *aphB*, which is divergently transcribed, encodes a protein which shows a high degree of homology to response regulators of a number of bacterial two-component systems. Two-component systems frequently regulate gene expression in prokaryotes in response to environmental stimuli. It is tempting to speculate that AphB may also activate the expression of this gene, and experiments to determine this are currently under way.

This study describes a new chromosomal gene, *aphB*, which encodes a LysR homolog that functions in both biotypes of *V. cholerae* in concert with a second chromosomally encoded protein, AphA, to activate the expression of a virulence operon

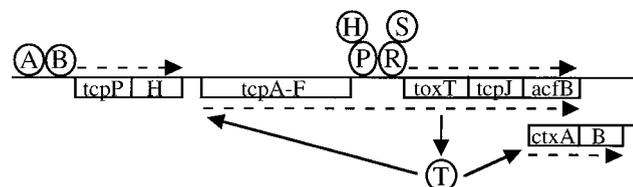


FIG. 5. Model of activation of the ToxR virulence cascade. In response to the appropriate environmental conditions, AphA and AphB activate transcription of the *tcpPH* operon. TcpPH, together with ToxRS, activate transcription of *toxT*. ToxT, in turn, activates expression of the *ctxAB* operon as well as expression of the entire *tcp* operon, including the *toxT* gene itself. The precise locations of the protein binding sites at the individual promoters have not yet been determined.

within a pathogenicity element. Further understanding of the mechanisms by which AphA and AphB activate gene expression may shed light on a number of questions regarding the pathogenesis of *V. cholerae*.

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