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Functional Characterization of Crp/Fnr-Type Global Transcriptional Regulators in *Desulfovibrio vulgaris* Hildenborough

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Crp/Fnr-type global transcriptional regulators regulate various metabolic pathways in bacteria and typically function in response to environmental changes. However, little is known about the function of four annotated Crp/Fnr homologs (DVU0379, DVU2097, DVU2547, and DVU3111) in *Desulfovibrio vulgaris* Hildenborough. A systematic study using bioinformatic, transcriptomic, genetic, and physiological approaches was conducted to characterize their roles in stress responses. Similar growth phenotypes were observed for the *crp/fnr* deletion mutants under multiple stress conditions. Nevertheless, the idea of distinct functions of Crp/Fnr-type regulators in stress responses was supported by phylogeny, gene transcription changes, fitness changes, and physiological differences. The four *D. vulgaris* Crp/Fnr homologs are localized in three subfamilies (HcpR, CooA, and cc). The *crp/fnr* knockout mutants were well separated by transcriptional profiling using detrended correspondence analysis (DCA), and more genes significantly changed in expression in a Δ DVU3111 mutant (JW9013) than in the other three paralogs. In fitness studies, strain JW9013 showed the lowest fitness under standard growth conditions (i.e., sulfate reduction) and the highest fitness under NaCl or chromate stress conditions; better fitness was observed for a Δ DVU2547 mutant (JW9011) under nitrite stress conditions and a Δ DVU2097 mutant (JW9009) under air stress conditions. A higher Cr(VI) reduction rate was observed for strain JW9013 in experiments with washed cells. These results suggested that the four Crp/Fnr-type global regulators play distinct roles in stress responses of *D. vulgaris*. DVU3111 is implicated in responses to NaCl and chromate stresses, DVU2547 in nitrite stress responses, and DVU2097 in air stress responses.

Transcription factors, promoter sequences, and regulatory circuit architecture are often referred as “the regulatory genome” (27, 33), and transcription factors are central to the function of regulatory networks (5). Organisms rely on regulatory networks to orchestrate the transcription of genes in response to internal or external environmental changes in order to optimize metabolism and enhance survival. Crp/Fnr regulators are global transcriptional regulators widely distributed in bacteria. The characteristic structure of Crp/Fnr is a C-terminal helix-turn-helix (HTH) motif that fits the DNA major groove and an N-terminal nucleotide binding domain (34). This class of transcriptional factors is named after the first two representatives discovered in *Escherichia coli*, i.e., the Crp (cyclic AMP [cAMP] receptor protein) and Fnr (fumarate and nitrate reductase regulator protein) (4, 21, 24, 27). Crp/Fnr regulators are generally classified into different subfamilies (e.g., CooA, Crp, Dnr, FixK, Fnr, HbaR, NnrR, etc.) according to phylogenetic affiliations (50). The increasing number of sequenced whole genomes has added to a growing list of Crp/Fnr family regulators identified in bacteria (9, 26); however, functions for most are not well defined.

Members of the Crp/Fnr family of global transcription regulators typically function as transcriptional activators. Previous studies on several bacterial species suggest that Crp/Fnr regulators are involved in responses to a variety of intracellular or extracellular signals, such as anoxia, carbon monoxide, temperature, nitric oxide, and oxidative and nitrosative stress (1, 13, 35, 42) and control metabolic pathways such as photosynthesis, nitrogen fixation, aromatic compound degradation, and respiration (11, 25, 31, 39).

The *E. coli* Crp, also known as catabolite activator protein (CAP), regulates expression of over 100 genes, the majority of which are involved in energy metabolism (5). Fnr, the other Crp/Fnr-type transcriptional regulator in *E. coli*, positively or negatively regulates the expression of approximately 40 genes under anaerobic conditions, including the fumarate reductase, nitrate and nitrite reductase, and cytochrome *d* oxidase genes (29). It does so by its direct interactions with molecular oxygen to regulate the metabolic transition between aerobic and anaerobic growth (23, 41). In contrast, *Rhodospseudomonas palustris* HbaR, another Crp/Fnr-type transcriptional factor, regulates gene expression of the 4-hydroxybenzoate coenzyme A ligase involved in the initial enzymatic step of aromatic compound degradation (8, 11, 14). CooA, the only known heme protein of the Crp/Fnr-type regulators, functions as a CO-sensing transcriptional regulator and positively regulates the *coo* operon encoding enzymes for the oxidation of CO to CO₂ coupled with proton reduction in *Rhodospira*

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TABLE 1 Bacterial strains and plasmids used in this study

Strain or plasmid	Description and relevant feature(s) ^a	Source or reference
Strains		
<i>D. vulgaris</i> Hildenborough	Wild type; NCIMB 8303	19
<i>D. vulgaris</i> Hildenborough JW9007	ΔDVU0379, Km ^r	This study
<i>D. vulgaris</i> Hildenborough JW9009	ΔDVU2097, Km ^r	This study
<i>D. vulgaris</i> Hildenborough JW9011	ΔDVU2547, Km ^r	This study
<i>D. vulgaris</i> Hildenborough JW9013	ΔDVU3111, Km ^r	This study
Plasmids		
pMO719	<i>Desulfovibrio</i> shuttle vector; SRB replicon pBG1 (EcoRI-EcoRI from pSC27) ligated to the EcoRI-bounded pCR8/GW/TOPO fragment; Spec ^r	22
pMO719-DVU0379	Complementation plasmid for JW9007; DNA fragment 425654-429479 (3,826 bp) cloned into the EcoRV site in pMO719; chromosome position of DVU0379, 428780-429457; Spec ^r	This study
pMO719-DVU2097	Complementation plasmid for JW9009; DNA fragment 2185067-2186208 (1,142 bp) cloned into the EcoRV site in pMO719; chromosome position of DVU2097, 2185336-2186007; Spec ^r	This study
pMO719-DVU2547	Complementation plasmid for JW9011; DNA fragment 2659090-2660860 (1,771 bp) cloned into the EcoRV site in pMO719; chromosome position of DVU2547, 2659978-2660664; Spec ^r	This study
pMO719-DVU3111	Complementation plasmid for JW9013; DNA fragment 3260525-3261583 (1,059 bp) cloned into the EcoRV site in pMO719; chromosome position of DVU3111, 3260648-3261382; Spec ^r	This study

^a Km^r, kanamycin resistance; Spec^r, spectinomycin resistance.

rillum rubrum (17, 38), highlighting the functional diversity of the Crp/Fnr-type regulators.

Analysis of the genome of the model sulfate-reducing bacterium (SRB) *Desulfovibrio vulgaris* Hildenborough revealed four annotated Crp/Fnr-type regulators (DVU0379, DVU2097, DVU2547, and DVU3111), suggesting potential functional differentiations among the multiple Crp/Fnr-type regulators, which exhibit a remarkable ability to survive and adapt to environmental perturbations and stress conditions (32, 40, 46, 52). Indeed, functional genomics studies of the response of *D. vulgaris* to various stress conditions have shown that the expression of the four Crp/Fnr-type transcriptional regulator genes significantly changed when *D. vulgaris* cells were exposed to nitrate (15), nitrite (16), oxygen (37), air (37), H₂O₂ (51), ethanol (unpublished data [<http://www.microbesonline.org>]), heat (7), or acetone (unpublished [<http://www.microbesonline.org>]) stress, suggesting that Crp/Fnr-type global regulators may play important roles in regulating a variety of stress responses in *D. vulgaris*. However, the functions and specificity of the Crp/Fnr-type regulators in *D. vulgaris* remain unclear.

Therefore, the objectives of this study were to (i) examine the activities of the four *fnr/crp*-type regulatory genes in the control of stress responses and (ii) identify the potential specificity of each Crp/Fnr-type regulator for various stress conditions in *D. vulgaris*. To address these objectives, knockout mutants for each Crp/Fnr regulator were generated and functionally characterized by their response to various environmental stress conditions that included air, NaCl, nitrite, and Cr(VI) by growth phenotype, transcriptional profiling, and competition analysis. Our results indicate that these four Crp/Fnr-type global transcriptional regulators have distinct functions in *D. vulgaris* in response to various environmental conditions.

MATERIALS AND METHODS

Phylogenetic analysis of Crp/Fnr superfamily members. An initial data set of all annotated Crp/Fnr genes in all currently available SRB genomes was compiled by extracting the genes from the Joint Genome Institute

Integrated Microbial Genomes database (33). Additional reference sequences were also extracted based on homology to SRB sequences to obtain a representative data set of the major Crp/Fnr subfamilies. The final data set containing 1,588 Crp/Fnr sequences was aligned using MUSCLE (10). The evolutionary history was inferred by the neighbor-joining method (44). The percentages of replicate trees in which the associated taxa clustered together were analyzed with the bootstrap test (500 replicates) (12). The evolutionary distances were computed by the Poisson correction method (53), and units represent the number of amino acid substitutions per site. The rate of variation among sites was modeled with a gamma distribution (shape parameter = 0.7241). All positions containing alignment gaps and missing data were eliminated only in pairwise sequence comparisons. Phylogenetic analyses were conducted in MEGA4 (47). The final tree was prepared for publication with Interactive Tree of Life (IToL) software (28).

Bacterial strains, growth conditions, and biomass production. *D. vulgaris* was cultured in defined lactate/sulfate medium (LS4D) (36) at 37°C. A knockout mutant for each *crp/fnr* pair was generated by a marker replacement approach (Table 1) (2). For mutant complementation, the entire operons of DVU2097, DVU2547, and DVU3111 or a partial operon of DVU0379 as well as the upstream and downstream sequences were included in the complementation plasmids (Table 1). The DNA fragments were amplified from wild-type (WT) *D. vulgaris* genomic DNA (gDNA) by PCR with Phusion high-fidelity DNA polymerase (Finnzymes Oy, Finland) and the primers listed in Table S1 in the supplemental material and were cloned into pMO719, which is stably replicated in *D. vulgaris* (22). The plasmids were propagated in *Escherichia coli* DH5α, and the correctness of the construct was confirmed with Sanger sequencing. The complementation plasmids carrying the right inserts were electroporated (1,750 V, 1-mm gap) into competent cells of each *crp/fnr* knockout mutant. The transformants were confirmed by plasmid isolation and digestion of plasmid DNA.

To test the effects of nitrite, NaCl, and chromate on *D. vulgaris* growth, the wild-type (WT) strain and four knockout mutants were inoculated into LS4D medium containing different concentrations of nitrite (0 mM, 0.5 mM, 1.0 mM, and 1.5 mM), NaCl (0 mM, 200 mM, 300 mM, and 400 mM), or chromate (0 mM, 0.5 mM, 0.7 mM, and 0.9 mM). An anaerobic titanium solution [titanium (III) chloride (20%) (wt/vol), sodium citrate (0.2 M), and sodium carbonate (8% wt/vol)] was used as a reductant for the LS4D medium before inoculation. To avoid an abiotic reaction, cysteine HCl (0.025%) prepared in NaHCO₃ (8 mM) was used as a reductant

in the chromate-related stress response test. For air stress conditions, 10 ml of air was aseptically injected into the mid-exponential-phase cell culture (10 ml of culture with 20 ml of headspace with N₂ gas) and 10 ml of the headspace gas was removed before air injection. Cell growth was monitored at 600 nm with a spectrophotometer.

For the transcriptomic analysis, mid-exponential-phase (optical density at 600 nm [OD₆₀₀] of about 0.4) cell cultures of WT *D. vulgaris* and the four Crp/Fnr mutants were harvested and flash-frozen in liquid nitrogen. Three biological replicates were prepared for each sample.

Microarray database search. A survey of the gene transcription level changes of four *crp/fnr* genes as well as *recA* in *D. vulgaris* under different growth conditions or stress conditions was carried out by searching the microarray database (<http://www.microbesonline.org/cgi-bin/microarray/viewExp.cgi>). Most of the data sets have been published and deposited in the NCBI GEO database as follows: nitrate stress (accession no. GSE20079), nitrite stress (16), air stress (GSE8196), H₂O₂ stress (GSE21441), ethanol stress (GSE7489), heat shock stress (7), and salt adaptation stress (GSE14343).

Reverse transcription-PCR. Expression patterns of four *crp/fnr* genes in cultures of cells at the mid-exponential-growth phase and the stationary-growth phase and of cells in the mid-exponential-growth phase harvested at 60 min after the application of different stressors such as 300 mM NaCl, 2.0 mM NaNO₂, 0.5 mM potassium chromate [Cr(VI)], and 20 ml of air were determined. The application of air stress was performed as described above. Isolation of total RNA and on-column DNase digestion were performed as described previously (7). Total RNAs (1.5 μg) were reverse transcribed with SuperScript reverse transcriptase (Qiagen, Valencia, CA) in a final reaction volume of 20 μl following the manufacturer's manual. A 1-μl volume of the reverse transcription product was used for PCR to amplify the full-length open reading frame (ORF) of *crp/fnr* genes (28 cycles) and the *recA* control gene (25 cycles) with primers listed in Table S1 in the supplemental material.

Transcriptional profiling assay. Isolation, fluorophore dye labeling, and handling of total RNA or gDNA were performed as described previously (48). DNA microarrays used in this study covered 3,482 of the 3,531 annotated protein-coding sequences of the *D. vulgaris* genome (16). Each array was hybridized with Cy3-labeled gDNA (as a control for normalization) and Cy5-labeled cDNA on a Tecan HS4800 hybridization station (Tecan Group, Durham, NC). Hybridization conditions as well as washing conditions and performance of image scanning were as described previously (7, 16, 36, 51). The microarray raw data processing was performed as described previously (36).

The similarities of the transcription profiles of the *crp* knockout mutants were analyzed with detrended correspondence analysis (DCA). The transcript level of each ORF was calculated as the ratio of signal intensity of Cy5 to that of Cy3, and the genes detected in at least two out of three replicates were kept for statistical analysis. The ratio of (Cy5/Cy3_{mutant})/(Cy5/Cy3_{wildtype}) with three biological replicates for each mutant was used for DCA with PC-ORD (version 5; MjM Software Design). The heat map of gene expression data was rendered as vector graphics in Encapsulated PostScript format with JColorGrid (20).

Cr(VI) reduction assay with washed cell. Experiments with washed cells were conducted to determine the chromate reduction capability of the WT strain and the four *crp/fnr* knockout mutants. Mid-exponential-phase cells (100 ml each; OD₆₀₀ of approximately 0.4) were anoxically harvested by centrifugation. After being washed twice with anoxic bicarbonate buffer (30 mM NaHCO₃, 62 mM sodium lactate, 20 mM Na₂SO₄, 0.025% cysteine HCl), the cells were resuspended in 1 ml of anoxic bicarbonate buffer. Then, 0.2-ml cells were transferred to 9.8 ml of anoxic bicarbonate buffer containing 0.5 mM K₂CrO₄ and the culture was incubated at 37°C. Samples taken at different time points were centrifuged, and the supernatants were used for quantification of Cr(VI) (30). Cr(VI) concentrations were determined colorimetrically (OD₅₄₀) using the diphenylcarbazide method with a Hach ChromaVer 3 chromium reagent powder pillow (Hach Company, Loveland, CO) (49).

Fitness assay. WT *D. vulgaris* and the four *crp/fnr* knockout mutants were subjected to competition under standard growth conditions (LS4D) or stress conditions with extra NaCl (200 mM), nitrite (0.1 mM), or chromate (0.2 mM) in LS4D. Low concentrations of stressors were chosen to obtain enough biomass at the sampling time point. As mentioned above, cysteine HCl (0.025%) was used as a reductant for chromate stress. For competition under air stress conditions, 10 ml of air was injected into the medium about 1 h after the reduction of the medium. All strains were revived, and approximately equal numbers of cells were mixed and inoculated into the competition media and allowed to compete for 2 days. Samples were taken at day 0 and day 2. Genomic DNA was isolated from each sample and quantified with PicoGreen (Quant-iT PicoGreen dsDNA assay kit; Invitrogen). Equal amounts of template DNA (20 ng) were used for real-time PCR with strain-specific PCR primers (see Table S1 in the supplemental material). Copy numbers were used to represent the cell numbers of each strain in the population. Fitness (*w*) was calculated as follows:

$$w = \frac{\ln(\text{mutant cell number at day 2} / \text{mutant cell number at day 0})}{\ln(\text{wild-type cell number at day 2} / \text{wild-type cell number at day 0})}$$

Microarray data accession number. Microarray data for this study have been deposited in the NCBI GEO database under accession number GSE29380.

RESULTS

Phylogeny of *D. vulgaris* Crp/Fnr homologs. A total of 1,588 protein sequences, including all annotated Crp/Fnr proteins from sequenced SRB genomes, were used for phylogenetic analysis. As shown in Fig. 1, the four *D. vulgaris* Crp/Fnr homologs are localized in three subfamilies. DVU0379 and DVU3111 are both located in the *cc* subfamily, which is constituted entirely of SRB sequences. DVU2547 is located in the HcpR branch and is highly conserved in most SRB species analyzed. In addition, DVU2097 is located in the *cooA* subfamily with other SRB genes and its localization in the genome with the *coo* operon suggests that it is the likely regulator of carbon monoxide dehydrogenase operon expression. While it contains no *D. vulgaris* genes, the phylogeny also revealed a novel clade (related to subfamily G) containing numerous SRB genes. The phylogenetic positioning of the SRB genes suggests that the Crp/Fnr homologs in *D. vulgaris* and other SRB species may have different functions in gene regulation and that the diversity of *crp/fnr* genes in SRB species is much greater than originally envisioned.

Expression changes of *crp/fnr* under different stress or growth conditions. To examine whether these four *crp/fnr* homologs are actively expressed, gene expression changes of the four genes were examined with a currently available microarray database. As shown in Fig. 2, the four genes were differentially transcribed in stress responses or growth conditions with different electron donors or acceptors. For instance, the DVU2547 expression increased in the nitrite stress response and DVU3111 in the chromate stress response; the expression of both DVU2547 and DVU3111 significantly increased in the heat shock response; DVU2097, DVU2547, and DVU3111 altered their expression under conditions of salt (NaCl) adaptation; expression of DVU2097 and DVU0379 increased when *D. vulgaris* was cultured with different electron donors or acceptors; and the expression of DVU0379, DVU2097, and DVU3111 increased under conditions of oxygen stress, but expression of DVU2097 decreased in the H₂O₂ stress response. Similar expression changes of four *crp/fnr* genes under stress conditions, in addition to their expression at both the mid-exponential phase and stationary phase, were de-

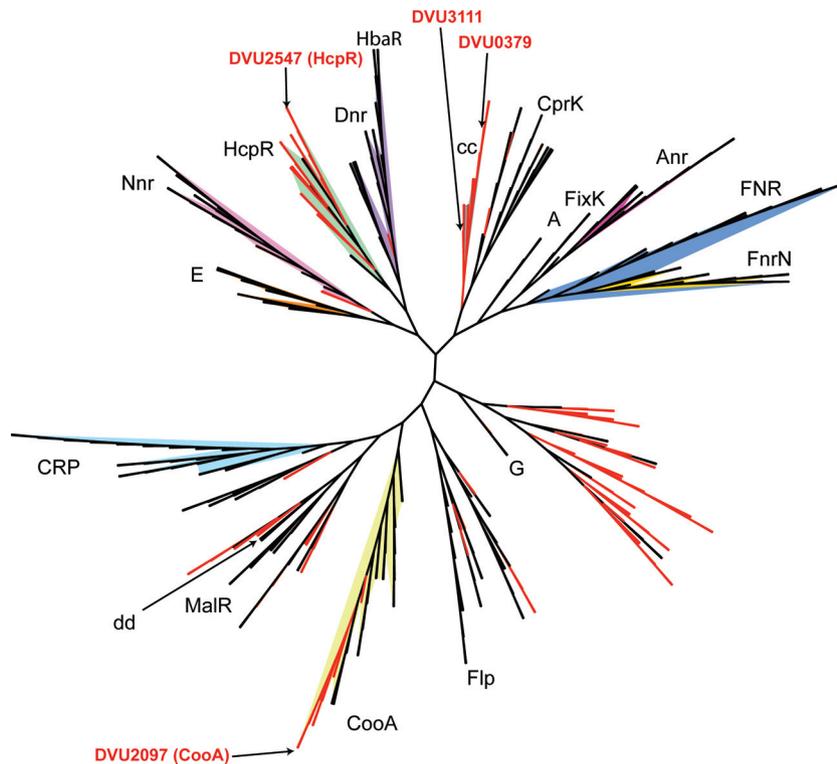


FIG 1 Phylogenetic analysis of Crp/Fnr superfamily enzymes (1,588 proteins) with an emphasis on SRB sequences. Red indicates genes from SRB species. Established subfamily names are marked for each clade. Multiple sequence alignment was conducted using MUSCLE. The evolutionary history was inferred using the neighbor-joining method. The optimal tree with the sum of branch lengths = 394.5 is shown. The percentage of replicate trees in which the associated taxa clustered together was assessed using the bootstrap test (500 replicates). Branch lengths are ignored for clarity. The evolutionary distances were computed using the Poisson correction method, and units represent the number of amino acid substitutions per site. The rate variation among sites was modeled with a gamma distribution (shape parameter = 0.7241). All positions containing alignment gaps and missing data were eliminated only in pairwise sequence comparisons. Phylogenetic analyses were conducted in MEGA4. The final tree was prepared using the Interactive Tree of Life (IToL) software.

tected with reverse transcription-PCR (see Fig. S2 in the supplemental material). It is likely that these regulators are involved in various stress responses but play different roles to some extent.

Characterization of knockout mutants for *crp/fnr*. To further investigate the function of the four Crp/Fnr-type regulator genes, single knockout mutants for each *crp/fnr* gene were generated (Table 1) and confirmed by PCR amplification of the full-length open reading frames (see Fig. S1 in the supplemental material). The growth phenotypes of *crp/fnr* knockout mutants under different environmental stress conditions, including air, NaCl, nitrite, and chromate stress, were examined.

(i) Air stress. To find out whether these *crp/fnr* ortholog regulators are involved in defense against increased oxygen levels, the growth phenotypes of each *crp/fnr* knockout mutant in response to air shock were examined. The growth of WT *D. vulgaris* and the growth of *crp/fnr* knockout mutants under standard growth conditions were quite similar except for a higher final optical density in the WT culture (Fig. 3A). Although the growth of WT and knockout mutants was inhibited by air shock, the four knockout mutants were more resistant to air stress conditions, with quicker recovery from the stress and higher biomass yield (Fig. 3B). The results suggest the involvement of these *crp/fnr* regulators in response to altered oxygen levels in *D. vulgaris*.

(ii) Salt stress. Growth inhibition by extra NaCl (200 mM or 300 mM) in the medium was observed in this study. Very limited growth was observed with the medium supplemented with 400

mM NaCl. Interestingly, all *crp/fnr* mutants had increased resistance to NaCl at all concentrations tested. As an example, growth phenotypes of the WT and mutants with extra 300 mM NaCl in the medium are shown in Fig. 3C. Higher growth rates, shorter lag phases, and higher biomass yields were observed for mutants. These data indicate that *crp/fnr* regulators are involved in the response of *D. vulgaris* to NaCl stress.

(iii) Nitrite stress. Growth of *D. vulgaris* was inhibited by nitrite in the medium. With 0.5 mM nitrite in the medium, longer lag phase was observed for both WT and all *crp/fnr* knockout mutants. A lower final biomass yield was obtained from mutants, although they grew faster than WT *D. vulgaris* (Fig. 3D). Growth of all strains was further delayed with 1.0 mM nitrite in the medium, and complete growth arrest was observed for all strains with 1.5 mM nitrite in the medium (data not shown). These results provide evidence for the involvement of *crp/fnr* genes in nitrite stress response.

(iv) Chromate stress. Growth was slightly stimulated for both WT and four mutants with a low concentration (0.5 mM) of Cr(VI) in the medium (data not shown). However, growth was inhibited by an increase in the Cr(VI) concentration (e.g., 0.7 mM) in the medium, with strain JW9013 containing a deletion of DVU3111 showing the highest growth rate (Fig. 3E). With 0.9 mM chromate in the medium, the growth of all *crp/fnr* mutants was completely inhibited; WT *D. vulgaris* was able to survive but produced only half of the final biomass seen with WT in the absence of

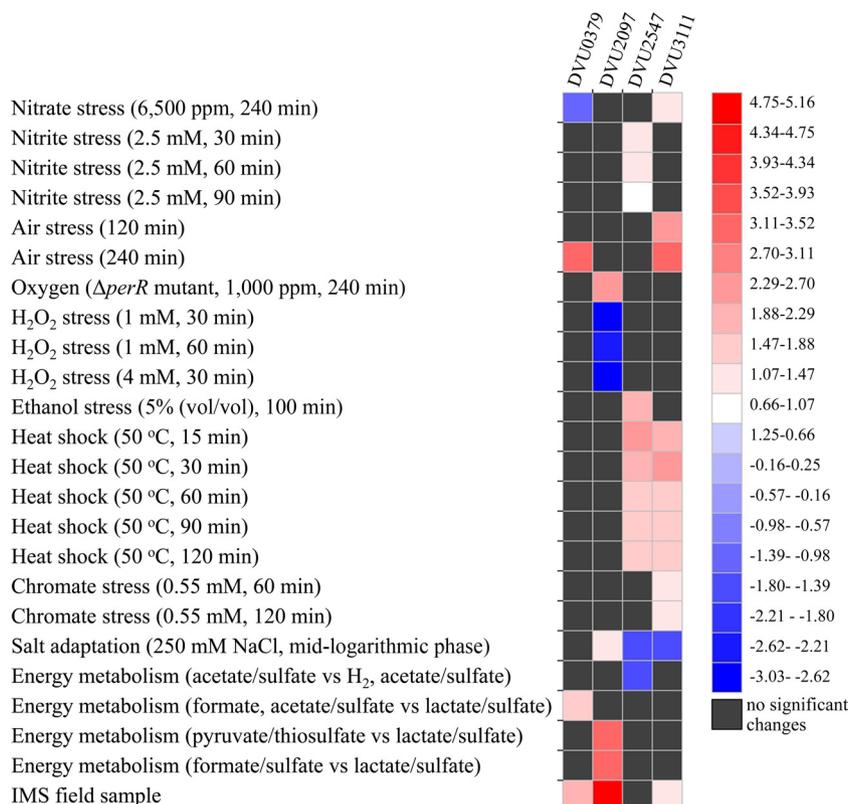


FIG 2 Expression changes of the four *crp/fnr* genes under different stress conditions or with different electron donors or acceptors. JColorgrid was used to generate the heat map.

stress (data not shown). These results indicate that these four regulators are involved in chromate response and that DVU3111 may play more important roles in the response of *D. vulgaris* to chromate.

To further confirm whether the phenotypes observed in knockout mutants resulted from the deletion of each specific *crp/fnr* gene, the phenotypes of the complemented knockout mutants carrying the WT copy of *crp/fnr* and the flanking upstream and downstream sequences were analyzed under conditions of NaCl stress, which resulted in the most conspicuous changes in growth between the knockout mutants and WT strain. As expected, WT-like phenotypes were observed for all complemented strains (Fig. 3F), indicating that these *crp/fnr* genes were indeed responsible for the observed phenotypes under these conditions.

Altered global gene transcription profiling in *crp/fnr* knockout mutants. In order to better understand the roles of Crp/Fnr-type regulators in *D. vulgaris*, the global transcriptional profiling of the four knockout mutants at the mid-exponential phase was analyzed. Compared with the WT strain, a small percentage of *D. vulgaris* genes with a significant change in expression ($|\log_2 R| > 1.5$; Z score > 1.5) was detected in each knockout mutant, with 136 in JW9013, 33 in JW9011, 31 in JW9009, and 23 in JW9007. Nevertheless, gene transcription profiling showed that the four *crp/fnr* mutants were well separated in a DCA plot (Fig. 4A). The gene transcription profiling of JW9011 was quite different from that of other three mutants, and the profile was separated by from the others by axis 1, which explains 42.93% of the total variance among the four mutants. The transcriptional profile of JW9013

was separated from those of JW9007 and JW9011 by axis 1 and that of JW9009 by axis 2, which explains 8.47% of the total variance. In addition, the transcriptional profile of JW9009 was separated from that of JW9007 by axis 1 (Fig. 4A). The greater number of transcriptionally changed genes detected in JW9013 suggested that DVU3111 might be involved in more cellular pathways than other *crp/fnr* gene products in *D. vulgaris*. A further analysis of transcription profiling demonstrated that the four *crp/fnr* products regulate both common and distinct genes (Table 2). A few categories of clusters of orthologous groups (COG) consisting of genes involved in stress responses, energy metabolism, and signal transduction are described below, further supporting the distinct functions of the four *crp/fnr* regulators in stress responses of *D. vulgaris*.

(i) Amino acid transport and metabolism. *glnH* (DVU0386) encoding an amino acid ABC transporter was found significantly upregulated in three mutants, JW9009, JW9011, and JW9013 (Table 2). In contrast, *trp* operon genes (DVU0465 to DVU0471 for tryptophan synthase) were upregulated in JW9007 only (Table 2). Increased expression of *trp* operon genes was observed when mid-exponential-phase *D. vulgaris* cells encountered NaCl shock (36), and glutamine was accumulated in salt-adapted *D. vulgaris* (18). Therefore, these gene expression changes may contribute to the enhanced salt resistance phenotype of the four *crp/fnr* mutants.

(ii) Inorganic ion transport and metabolism. Changes in the expression of the genes involved in inorganic ion transport, particularly iron homeostasis, were found in all *crp/fnr* mutants. The expression of the primary iron uptake system genes, *feoAB*

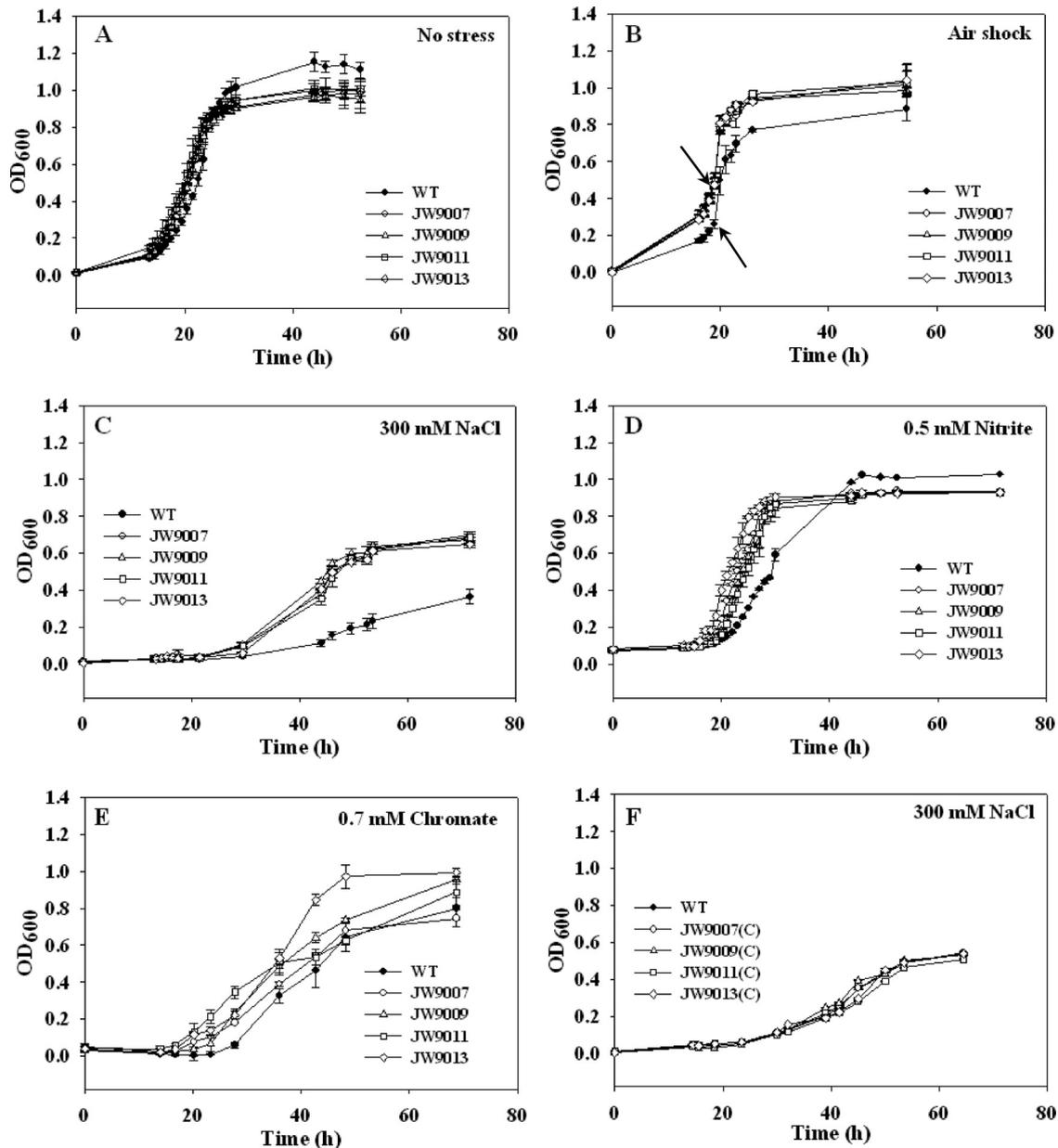


FIG 3 Growth phenotypes of the four *crp/fnr* knockout mutants. (A) no stress; (B) air shock; (C) 300 mM NaCl; (D) 0.5 mM nitrite; (E) 0.7 mM chromate (VI); (F) growth phenotype of complemented mutants with extra 300 mM NaCl in the medium. The standard deviations of the results from four biological replicate experiments are shown as error bars. The arrows in panel B indicate the addition of air (10 ml).

(DVU2571 to DVU2574), decreased significantly in all *crp/fnr* mutants (Table 2). The decreased expression of the *feo* system in the knockout mutants provided evidence for the involvement of these *crp/fnr* regulators as activators for the acquisition of Fe(III).

(iii) **Energy production and conversion.** The expression of a hybrid cluster protein-encoding gene, *hcp2* (DVU2543), a predicted target gene for DVU2547 (43), was significantly downregulated in JW9011 (Table 2). Also, gene DVU2544, encoding an iron-sulfur cluster binding protein in the same operon, was concurrently downregulated in JW9011 (Table 2). Downregulation of these two genes was observed in JW9013 as well (Table 2). Another gene, *fld* (DVU2680), predicted to be an iron-repressed flavodoxin, was significantly downregulated in all *crp/fnr* mutants (Ta-

ble 2). Moderately increased expression of DVU3171 (a cytochrome *c*₃-encoding gene) was detected in JW9013 and JW9009 (Table 2). Cytochrome *c*₃ was previously reported to be the major electron carrier for metal ion reduction (19), and the upregulation of this gene may be related to an increased chromate reduction capacity in *crp/fnr* mutants. The alkyl hydroperoxide reductase C gene *ahpC* (DVU2247), known for reducing hydrogen peroxide (H₂O₂) into H₂O (6), was downregulated in JW9007 (Table 2), suggesting the possible role of DVU0379 in oxidative stress response.

(iv) **Transcription.** The same trend of changes was found in all *crp/fnr* mutants. For example, increased expression of genes for a GntR family transcription regulator, *pdhR* (DVU2785), and a

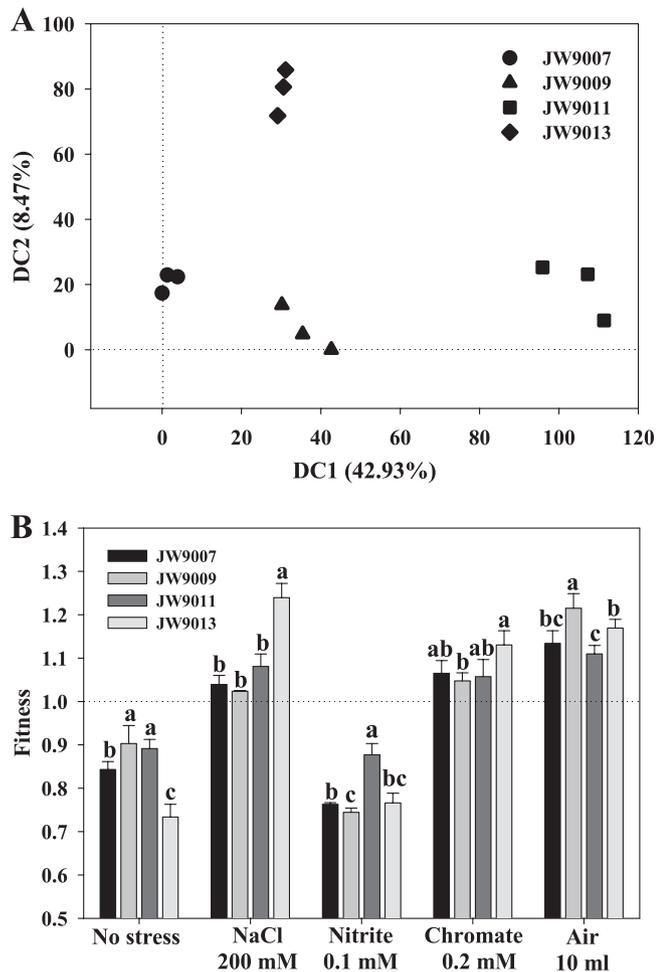


FIG 4 Transcriptional profiling of *crp/fnr* mutants analyzed by DCA (A) and the fitness changes of the *crp/fnr* mutants compared to the WT under the standard growth and stress conditions (B). Each symbol in panel A indicates microarray data from one biological replicate experiment. Fitness data represent averages of the results of three biological replicate experiments, with the standard deviations shown as error bars. A pairwise *t* test was performed to assess the significance of fitness among four mutants, with a, b, and c indicating significant differences at the $P < 0.05$ level.

LuxR family transcriptional regulator (DVU2577) was detected in all mutants (Table 2). Also, expression of DVU2827, a sigma-54-dependent transcriptional regulator, decreased in all mutants. Future work is needed to reveal the relationship between Crp/Fnr and other transcription regulators, which may form complex networks in *D. vulgaris*.

(v) **Signal transduction mechanisms.** Many genes involved in signal transduction mechanisms were found to be shared by some or all *crp/fnr* mutants (Table 2). Expression of two “carbon starvation protein A” genes in the operon DVU0598-DVU0599 was significantly downregulated in all *crp/fnr* mutants. Also, expression of a predicted methyl-accepting chemotaxis protein (MCP) encoded by DVU1169 increased significantly in three *crp/fnr* mutants (but not the JW9007 mutant), and expression of chemotaxis protein-encoding gene *cheA-1* (DVU1594) and chemotaxis protein methyltransferase-encoding gene *cheR-1* (DVU1595) decreased more than 2-fold in three *crp/fnr* mutants (but not the JW9007 mutant). Decreased expression of MCP gene *dcrA*

(DVU3182) in JW9009 and JW9013 and *dcrH* (DVU3155) in all mutants was detected. *dcrA* and *dcrH* were previously reported to be involved in locating SRB in the gradients of oxic and anoxic interfaces in pipelines or sewers (3). Therefore, these results suggest the roles of the four *crp/fnr* regulators in regulation of alternative energy pathways during carbon starvation and DVU2097, DVU2547, and DVU3111 in cell chemotaxis.

Functions of *crp/fnr* regulators confirmed by competition assay. The assay of competition between the WT strain and the four knockout mutants provided further evidence of the specific contributions of each *crp/fnr* gene in response to different stresses (Fig. 4B). A significant ($P \leq 0.05$) decrease of fitness was observed for all *crp/fnr* mutants compared to the WT strain under the standard growth conditions, and significant ($P \leq 0.05$) differences were observed among mutants, with the lowest fitness exhibited by JW9013, medium fitness exhibited by JW9007, and the best fitness exhibited by JW9009 and JW9011 (but with no significant difference between these two mutants). Consistent with the growth phenotype, a fitness decrease was also observed for all knockout mutants under nitrite stress conditions, and significant ($P \leq 0.05$) differences occurred among four mutants, with the highest fitness for JW9011, the lowest for JW9009, and medium fitness for JW9007, but no significant difference between JW9009 and JW9013 or between JW9007 and JW9013. In contrast, all four mutants generally showed improved fitness under air, NaCl, or chromate stress conditions compared to the WT. Under NaCl stress conditions, the highest fitness was seen in JW9013, but no significant difference was observed for the three other mutants. Under chromate stress conditions, the highest fitness was found in JW9013, and significant differences appeared only between JW9009 and JW9013. Under air shock conditions, the highest fitness was observed in JW9009, and significant differences were observed between JW9009 and other three mutants and also between JW9011 and JW9013 (Fig. 4B). These results indicate that each *crp/fnr* gene product performs a different role(s) in response to different stresses. For example, more important roles were seen with DVU3111 in response to NaCl or chromate stresses, DVU2547 in response to nitrite stress, and DVU2097 in response to air stress.

Increased chromate reduction in *crp/fnr* knockout mutants. To further confirm the function of Crp/Fnr regulators in hexavalent chromate Cr(VI) reduction, experiments with washed cells were conducted to examine the Cr(VI) reduction capability of each strain. Increased Cr(VI) reduction was observed in four *crp/fnr* mutants with 0.5 mM Cr(VI) present in the medium (Fig. 5). Consistent with the fitness data, the highest chromate reduction rate was observed in JW9013.

DISCUSSION

Regulation of gene expression is critical for bacterial response and survival in an ever-changing environment. Crp/Fnr regulators are global transcriptional factors present in many bacteria and serve key roles in response to environmental stresses. In this study, taking advantage of bioinformatics, mutagenesis, high-throughput transcriptomics, and competition analysis, functional characterization of four annotated Crp/Fnr-type regulators in *D. vulgaris* was performed to better understand the physiological responses associated with the regulators.

Each of the *crp/fnr* regulators may have distinct functions in response to different environmental stresses in *D. vulgaris*. First,

TABLE 2 Selected genes with significant expression level changes in *crp/fnr* knockout mutants

Regulator and category	Gene name	Annotation	Log ₂ R (mutant/wild type) ^a			
			JW9007	JW9009	JW9011	JW9013
Amino acid transport and metabolism						
DVU0465	<i>trpE</i>	Anthranilate synthase, component I	1.33 (1.78)	0.23 (0.32)	N/A	N/A
DVU0466	<i>trpG</i>	Anthranilate synthase, glutamine amidotransferase	1.52 (2.35)	0.08 (0.12)	-1.33 (-1.66)	-0.81 (-0.60)
DVU0467	<i>trpD</i>	Anthranilate phosphoribosyltransferase	1.44 (2.24)	-0.17 (-0.18)	-0.32 (-0.49)	-0.08 (-0.12)
DVU0468	<i>trpC</i>	Indole-3-glycerol phosphate synthase	1.23 (2.08)	0.12 (0.21)	-0.90 (-1.41)	-0.19 (-0.28)
DVU0469	<i>trpF-1</i>	N-(5-phosphoribosyl)anthranilate isomerase	0.88 (1.30)	-0.17 (-0.24)	-1.00 (-1.45)	-0.13 (-0.16)
DVU0470	<i>trpB-2</i>	Tryptophan synthase, beta subunit	1.43 (2.52)	0.34 (0.56)	-0.56 (-0.95)	0.15 (0.25)
DVU0471	<i>trpA</i>	Tryptophan synthase, alpha subunit	1.48 (2.66)	0.42 (0.70)	-0.44 (-0.74)	0.32 (0.54)
DVU0386	<i>glmH</i>	Amino acid ABC transporter, periplasmic amino acid binding protein	0.55 (1.01)	1.83 (3.32)	1.64 (2.89)	1.51 (2.76)
Inorganic ion transport and metabolism						
DVU2571	<i>feoB</i>	Ferrous iron transport protein B	-2.11 (-2.53)	-1.81 (-2.16)	-1.32 (-2.14)	-1.78 (-1.96)
DVU2572	<i>feoA</i>	Ferrous iron transport protein A	-1.69 (-2.51)	-1.73 (-2.17)	-1.50 (-2.25)	-1.75 (-1.97)
DVU2573	N/A	Hypothetical protein	-0.62 (-0.96)	-0.52 (-0.85)	-0.64 (-1.13)	-0.46 (-0.62)
DVU2574	<i>feoA</i>	Ferrous iron transporter component <i>feoA</i>	-2.17 (-2.79)	-1.34 (-1.30)	-1.70 (-2.51)	-2.00 (-2.98)
Energy production and conversion						
DVU3171	N/A	Cytochrome <i>c</i> ₃	0.63 (0.64)	1.26 (1.30)	0.88 (0.93)	1.45 (1.52)
DVU2784	<i>lldD</i>	Dehydrogenase, flavin mononucleotide - dependent family	0.63 (0.97)	1.36 (2.32)	1.07 (1.90)	1.51 (2.15)
DVU3143	N/A	Iron-sulfur cluster-binding protein	1.30 (1.68)	0.97 (1.32)	1.21 (1.72)	1.74 (2.08)
DVU1165	<i>ndh</i>	NADH respiratory dehydrogenase	-0.54 (-0.55)	-0.80 (-0.66)	-0.75 (-1.06)	-1.42 (-2.27)
DVU2543	<i>b0873</i>	Hybrid cluster protein	-0.72 (-1.07)	-1.22 (-1.81)	-2.79 (-3.61)	-1.46 (-1.96)
DVU2544	N/A	Iron-sulfur cluster binding protein	-0.45 (-0.59)	-0.82 (-0.95)	-1.26 (-1.32)	-1.06 (-1.33)
DVU2680	<i>fld</i>	Flavodoxin, iron repressed	-2.66 (-2.64)	-2.64 (-2.44)	-2.91 (-3.67)	-2.78 (-3.26)
DVU2247	<i>ahpC</i>	Alkyl hydroperoxide reductase C	-1.41 (-2.47)	-0.29 (-0.47)	-0.34 (-0.61)	-0.74 (-1.33)
Transcription						
DVU2577	N/A	DNA-binding response regulator, LuxR family	1.60 (2.53)	1.15 (1.76)	0.71 (1.09)	1.01 (1.57)
DVU2785	<i>pdhR</i>	Transcriptional regulator, GntR family	1.28 (2.00)	1.46 (2.14)	1.16 (1.82)	1.94 (2.15)
DVU2827	N/A	Sigma-54 dependent transcriptional regulator	-0.76 (-0.91)	-2.20 (-1.75)	-0.63 (-1.12)	-1.24 (-1.64)
Signal transduction mechanisms						
DVU0878	N/A	<i>dnaK</i> suppressor protein, putative	2.13 (3.75)	0.68 (1.13)	0.19 (0.31)	1.00 (1.83)
DVU0888	N/A	Response regulator	0.62 (0.67)	0.72 (0.82)	0.21 (0.24)	1.81 (1.57)
DVU1169	N/A	Methyl-accepting chemotaxis protein	0.72 (0.77)	1.57 (2.01)	1.97 (2.83)	1.94 (2.86)
DVU1594	<i>cheA-1</i>	Chemotaxis protein CheA	-0.98 (-1.33)	-1.07 (-1.27)	-1.40 (-2.25)	-1.71 (-2.56)
DVU1595	<i>cheR-1</i>	Chemotaxis protein methyltransferase	-0.54 (-0.66)	-1.33 (-1.04)	-1.55 (-2.03)	-1.76 (-2.47)
DVU3182	<i>dcrA</i>	Methyl-accepting chemotaxis protein DcrA	-0.38 (-0.49)	-1.15 (-0.75)	-0.21 (-0.35)	-1.69 (-2.14)
DVU3155	<i>dcrH</i>	Methyl-accepting chemotaxis protein DcrH	-1.28 (-1.36)	-1.83 (-1.62)	-1.29 (-1.53)	-1.75 (-2.20)
DVU2577	N/A	DNA-binding response regulator, LuxR family	1.60 (2.53)	1.15 (1.76)	0.71 (1.09)	1.01 (1.57)
DVU0039	N/A	C4-type zinc finger protein, DksA/TraR family	0.58 (0.62)	0.78 (0.77)	0.82 (0.92)	2.14 (1.80)
DVU0598	N/A	Carbon starvation protein A, putative	-2.76 (-2.15)	-1.79 (-2.04)	-1.69 (-2.91)	-3.07 (-4.02)
DVU0599	N/A	Carbon starvation protein A, putative	-1.38 (-1.23)	-1.44 (-1.32)	-1.05 (-1.09)	-1.69 (-1.87)
DVU3023	<i>atoC</i>	Sigma-54-dependent DNA-binding response	-1.49 (-1.33)	-1.50 (-1.02)	-0.08 (-0.11)	-2.01 (-1.65)

^a N/A, no gene name has been assigned to the locus yet or no data for the gene. Z scores are shown in parentheses.

the overall similarities of the transcriptional profiles of each mutant are low (Fig. 4A). The differences were also obvious in terms of significantly changed gene numbers as well as the genes involved in different functional processes (Table 2). Second, fitness analysis of four *crp/fnr* mutants indicated that each *crp/fnr* mutant had different contributions in response to different stresses. The highest fitness was observed for JW9013 under NaCl stress or chromate stress conditions. JW9011 had the highest fitness under

conditions of nitrite stress, and highest fitness was observed for JW9009 under air stress conditions. Third, an analysis of chromate reduction provided further evidence for more important functions of DVU3111 in chromate reduction. Besides this experimental evidence, the phylogenetic position of the four Crp/Fnr homologs in three different branches and subfamilies also suggests different functions for the *crp/fnr* genes in *D. vulgaris*.

Results from this study suggest that four *crp/fnr* regulators *D.*

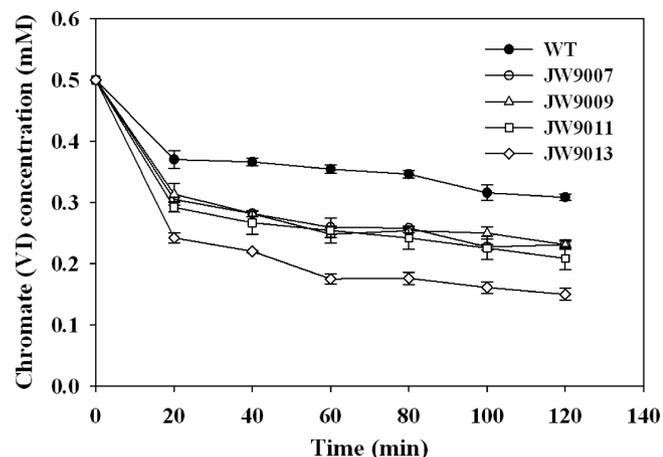


FIG 5 Cr(VI) reduction in experiments with washed cells. The error bars represent the standard deviations.

vulgaris may function differently from *E. coli fnr* but similarly to *E. coli crp* to some extent. In *E. coli*, nitrate and nitrite reductases have been shown to be regulated by Fnr (29); however, no significant changes in expression of nitrite reduction genes (e.g., the *nrfHA* operon encoding cytochrome *c* nitrite reductase) were observed in *crp/fnr* mutants of *D. vulgaris*. Increased resistance to air stress in *crp/fnr* knockout mutants suggested that *crp/fnr* regulators may act as transcriptional repressors in *D. vulgaris*, which is different from *E. coli* Fnr functioning as a global regulator under anaerobic growth conditions (21, 41). Decreased expression of two carbon starvation protein A genes (DVU0598 and DVU0599) in all *crp/fnr* mutants suggested the possible similarity between *crp/fnr* regulators in *D. vulgaris* and *E. coli crp*, which positively regulates about two-thirds of carbon starvation genes induced at the onset of carbon starvation (45).

Cooperation among *crp/fnr* and other transcriptional regulators in stress responses was also suggested by results from this study. Fur regulon genes such as the iron uptake system *feoAB* genes are repressed by Fur, and *trp* operon is repressed by iron-free Fur (2). A significant decrease in expression of *feoAB* and the iron-repressed flavodoxin gene *fld* (DVU2680) in the four *crp/fnr* mutants and upregulation of the *trp* operon in JW9007 (Table 2) suggested the possibility of cross-talk between Crp/Fnr and Fur. On the other hand, expression of DVU2543, one of the predicted target genes of DVU2547 (43), was found to be significantly downregulated in JW9011. However, no significant expression changes of other predicted target genes such as *apsAB*, *sat*, DVU1080-DVU1081, and genes in the *coo* operon (43) were found in this study. It is possible that additional cooperating factors contribute to the regulation of these genes. If such cooperation exists in *D. vulgaris*, complicated gene regulatory networks are expected.

In summary, the systematic study provided evidence for distinct functions of the *crp/fnr* regulators in *D. vulgaris*: for example, the roles of DVU3111 in responses to NaCl or chromate stresses, DVU2547 in nitrite stress response, and DVU2097 in air stress response. Future in-depth studies with double- or triple-knockout mutants and chromatin immunoprecipitation-microarray technology (ChIP-chip) should provide more and higher-resolution information about the functional redundancy and gene regulatory networks of Crp/Fnr regulators in *D. vulgaris*.

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