The Lactic Acid-Induced Acid Tolerance Response in Salmonella enterica Serovar Typhimurium Induces Sensitivity to Hydrogen Peroxide†

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Transcriptome analyses of Salmonella enterica serovar Typhimurium revealed that 15 genes were significantly up-regulated after 2 h of adaptation with lactic acid. cadB was the most highly up-regulated gene and was shown to be an essential component. Lactic acid-adapted cells exhibited sensitivity to hydrogen peroxide, likely due to down-regulation of the OxyR regulon.

Salmonella enterica serovar Typhimurium employs both a logarithmic-phase acid tolerance response (ATR) and a stationary-phase ATR to cope with acid stress (23). The logarithmic-phase HCl-induced ATR requires both pre- and post-acid-shock adaptation phases to be fully tolerant to a pHo of ≤3.0 at 37°C (13). The pre-acid-shock adaptation phase occurs after exposure to pH 5.8 and involves the production of at least 12 acid shock proteins (5). The gene products of the cad operon (1) and the cspB genes (6, 7) all appear to be important in this preshock adaptation phase. Salmonella commonly encounters stress from organic acids, either as by-products of its own metabolism (25), as food preservatives, or as volatile fatty acids in the intestine (12). Limited studies have shown that these organic acids protect against subsequent HCl stress (e.g., gastric acid) (9, 10).

We have recently shown that lactic acid induces a transient ATR in Salmonella enterica Serovar Typhimurium at 20°C, and the maximum ATR occurs at 2 h (9). We used a transcriptome approach to identify the genes involved in the lactic acid-induced ATR (L-ATR). Mid-logarithmic cultures of Salmonella serovar Typhimurium SL1344 were produced and adapted for 2 h with 0.015 M lactic acid in tryptone soy broth (TSB) (Difco) at pH5.8 as previously described (9). After adaptation, RNA was stabilized by addition of a phenol-ethanol solution (5:95) and was immediately transferred to ice prior to extraction. RNA samples were prepared using a Promega SV total RNA purification kit, and the quality was checked using an RNA nanochip (with an Agilent 2100 bioanalyzer). The microarray experiment and analyses of transcriptome data were performed as previously described (4). We used an indirect “type 2” experimental design (26) with genomic DNA as a common reference. We used two technical replicates for each of two biological replicates. The relevant protocols are described at http://www.ifr.bbsrc.ac.uk/safety/microarrays/protocols.html. The 4,114-feature microarray based on the serovar Typhimurium LT2 genome sequence (18) has been described previously (4).

Following statistical filtering (false discovery rate, 0.05), changes in gene expression of >twofold were considered to be significant (4). A total of 76 genes changed during the L-ATR, and 15 of them were significantly up-regulated (see Table S1 in the supplemental material). Only three of these up-regulated genes, cadA, cadB, and cspB, exhibited more than a fivefold increase in the level of expression (Table 1). Expression of cadBA can be induced in Escherichia coli by lowering the external pH in the presence of lysine (20). This raises the possibility that the high level of cadBA induction observed in the L-ATR may not have been caused by lactic acid per se but could have reflected the decrease in the pHo in the presence of lysine in the medium.

The cadB gene is a component of the lysine decarboxylase system and has a known role in the HCl-induced ATR of Salmonella serovar Typhimurium (21). As cadB was the most highly up-regulated gene in the L-ATR, its role was investigated further. Using the λ Red method (3), we replaced the cadB structural gene in SL1344 with a kanamycin resistance marker to generate strain JH2917. The mutation was P22 transduced to a wild-type SL1344 background, and the genotype of the mutant strain was verified by PCR. The following primers were used for PCR amplification of the DNA fragment that was recombined with the chromosome from the pKD4 plasmid (3): CadB-KO-F (5′-TAA GCCCGGTTCCTTAAATAATCAGCTACGGAGAAATGAA CGTGTAAGGCTGGAGCTGCTTC) and CadB-KO-R (5′-GC AGGTGACGAAAGGGGCTTTGAGAAAAAGGAGTTA GCCGCATATGAAATCCCTCCTTA). It is possible that the cadB:kan mutation could have a polar effect on the downstream cadA gene. Even if this were the case, the mutant phenotype would remain as a defect in lysine decarboxylase. To test whether the lysine decarboxylase system was required for the L-ATR, cultures of Salmonella serovar Typhimurium SL1344 and JH2917 were grown to the mid-logarithmic phase and either adapted with lactic acid prior to acid shock or immediately acid shocked (unadapted cultures) as described previously (9). JH2917 was not able to mount an ATR (Fig. 1), showing that the lysine decarboxylase system was an essential component of the L-ATR. CadA has been shown to be important in the ATR of Vibrio cholerae (19), and CadB has been shown to be important but not essential for the HCl-induced ATR in Salmonella serovar Typhimurium (8, 21), in which limitation that the high level of cadBA induction observed in the L-ATR may not have been caused by lactic acid per se but could have reflected the decrease in the pHo in the presence of lysine in the medium.

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other amino acid decarboxylase systems can be invoked to restore the ATR phenotype (21). However, our study demonstrated that JH2917 was unable to mount an ATR even though the organism was provided with a rich source of alternate amino acids. Recently, it has been shown that the inhibition of porins by excreted cadaverine may be part of the adaptive mechanism that helps E. coli survive in acidic environments (24). Our results indicate that a similar adaptive mechanism may be essential for the Salmonella serovar Typhimurium L-ATR, as it requires cadaverine transport mediated by the CadB transport protein.

The transcriptome data also showed that five members of the OxyR regulon were down-regulated during the L-ATR (Table 1); this regulon is critical for the survival of Salmonella under oxidative stress conditions (2). Such down-regulation during the L-ATR raised the possibility that although adaptation with lactic acid protected against acidic environments, it was incompatible with the response to oxidative stress. To investigate this, lactic acid-adapted and unadapted cultures of Salmonella serovar Typhimurium SL1344 were exposed to an oxidative stress. Cells were removed from TSB by filtration through a 0.22-μm membrane filter (Millipore) and were resuspended by vortexing in fresh TSB, and enough of a 30% chlorous acid (16). Our study demonstrated that the L-ATR of Salmonella serovar Typhimurium causes sensitivity to hydrogen peroxide; the L-ATR appears to inactivate the oxidative stress response via down-regulation of the OxyR regulon.

In this study, we identified genes involved in the L-ATR of Salmonella serovar Typhimurium and related transcriptional

**TABLE 1. Salmonella serovar Typhimurium genes exhibiting more than a fivefold increase in the level of expression during the lactic acid-induced acid tolerance response and members of the OxyR regulon that are down-regulated during the response**

<table>
<thead>
<tr>
<th>Common name</th>
<th>Systematic name</th>
<th>Expression level after exposure to lactic acid at pH 5.8 compared to expression at pH 7</th>
<th>Function or comment</th>
</tr>
</thead>
<tbody>
<tr>
<td>cadA</td>
<td>STM2559</td>
<td>5.9</td>
<td>Lytic decarboxylase 1</td>
</tr>
<tr>
<td>cadB</td>
<td>STM2558</td>
<td>50.94</td>
<td>Lysine/cadaverine transport protein</td>
</tr>
<tr>
<td>cspB</td>
<td>STM1996</td>
<td>8.97</td>
<td>Putative cold shock protein</td>
</tr>
<tr>
<td>ahpC</td>
<td>STM0608</td>
<td>0.2</td>
<td>Detoxification of hydroperoxides</td>
</tr>
<tr>
<td>ahpF</td>
<td>STM0609</td>
<td>0.06</td>
<td>Detoxification of hydroperoxides</td>
</tr>
<tr>
<td>dps</td>
<td>STM0831</td>
<td>0.14</td>
<td>Starvation-induced resistance to H2O2 catalase: hydroperoxidase HP1(1)</td>
</tr>
<tr>
<td>katG</td>
<td>STM4106</td>
<td>0.09</td>
<td>Catalase:hydroperoxidase HP1(1)</td>
</tr>
<tr>
<td>oxyR</td>
<td>STM4125</td>
<td>0.57</td>
<td>Regulatory protein sensor for oxidative stress</td>
</tr>
</tbody>
</table>

*The complete set of transcriptome data has been submitted to the ArrayExpress database (www.ebi.ac.uk/arrayexpress).*

![FIG. 1. Survival of lactic acid-adapted ΔcadB mutant JH2917 and wild-type Salmonella serovar Typhimurium strain SL1344 during acid shock at an external pH of 3.0 poised with HCl. Cells were adapted at pH5.8 for 2 h. □, unadapted wild-type cells; ■, adapted wild-type cells; ▲, unadapted JH2917 cells; ●, adapted JH2917 cells. The values are means for two biological replicates. The coefficient of variation was <20%.

![FIG. 2. Survival of lactic acid-adapted and unadapted cells of Salmonella serovar Typhimurium SL1344 during exposure to hydrogen peroxide. Cells were adapted with lactic acid at pH5.8 for 2 h. The bacterial cells were subsequently challenged with 100 mM H2O2. ▲, unadapted cells; □, lactic acid-adapted cells. The values are means for two biological replicates. The coefficient of variation was <20%.*
changes in bacterial phenotypes, including resistance to acidity and sensitivity to oxidative stress. This type of approach highlights the value of transcriptome analysis with DNA microarrays for the food industry, since the protective responses and any linked vulnerabilities associated with particular environmental stresses can be identified and exploited to enhance food preservation regiments.

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REFERENCES