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OhrRA functions as a redox-responsive system controlling toxinogenesis in Bacillus cereus

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ABSTRACT

Bacillus cereus OhrR is a member of the subgroup of the MarR (multiple antibiotic resistance) family of transcriptional regulators that use a cysteine-based redox sensing mechanism. OhrA is a thiol-dependent, peroxidase-like protein. The dual OhrRA system triggers B. cereus adaptation in response to redox changes, such as those encountered in the environment of the gastrointestinal tract. Here, we investigated the role of OhrRA in toxinogenesis. Comparative shotgun analysis of exoproteomes from ΔohrA, ΔohrR and wild-type cells revealed significant changes in the abundance levels of toxin-related proteins depending on the extracellular redox potential. We complemented these data by measuring the DNA binding activity of reduced and oxidized recombinant OhrR on toxin and putative toxin promoter regions. Furthermore, transcriptomic data and OhrRA-dependent, antiproliferative activity of the B. cereus exoproteome on Caco-2 human epithelial cells were recorded. The results indicate that OhrR controlled toxin gene expression directly or indirectly in a redox- and toxin-dependent manner, and may function as a repressor or an activator. Moreover, we found that OhrR restricts toxin-dependent antiproliferative activity of the B. cereus exoproteome whatever the growth conditions, while the restrictive impact of OhrA occurs only under low ORP anoxic conditions.

Biological significance

B. cereus is a notorious foodborne pathogen which causes gastroenteritis. Fatal and severe cases have been reported. The pathogenicity of B. cereus is intimately associated with the production of epithelial cell-destructive toxins in the small intestine. The small intestine poses several challenges for a pathogen because it is sliced into various niches with different oxygen concentrations and different redox potentials. We recently showed that the organic hydroperoxide resistance OhrRA system was crucial to the successful adaptation of B. cereus to extreme redox environments such as those encountered in the lumen (high reducing anoxic environment) and on the intestinal epithelium (transient oxic environment). Here we provide evidence that this bacterial system is a major virulence determinant in B. cereus in that it coordinates toxinogenesis in a redox dependent manner. Specifically, our comparative exoproteomic analyses reveal that OhrR strongly restricts B. cereus toxinogenesis under high reducing anoxic conditions while OhrA boosts toxinogenesis. Based on exoproteomic analyses, we further examined the role of OhrR and found that it functions as a...
1. Introduction

Bacillus cereus is a ubiquitous, globally distributed, foodborne pathogen. This human pathogen [1,2] is mainly associated with two types of gastrointestinal (GI) infection, both causing diarrhea and vomiting after the ingestion of contaminated food. The diarrheal syndrome is mainly attributed to toxins secreted by vegetative cells colonizing the human small intestine [3-5]. A total of 14 toxin-related proteins were identified in the culture supernatant of B. cereus cells grown in low oxygen reduction potential (ORP) anoxic conditions, which are considered to mimic those encountered in the human intestine [6]. These 14 toxin-related proteins include the four components (HblL1, HblL2, HblB and HblB’ ) of hemolysin BL (Hbl), the three components (NheA, NheB, NheC) of the nonhemolytic enterotoxin, Nhe, and the single-component protein, CytK [5,7-9]. Hbl, Nhe and CytK are secreted, pore-forming cytotoxins, which are currently considered to be the main causative agents of diarrheal syndrome [9-11]. However, several other toxin-related proteins may contribute to this syndrome, including hemolysin I (cereolysin O, HlyI), which is a thiol-activated, cholesteryl-binding cytolsin, hemolysin II (HlyII) and enterotoxin FM (EntFM) [12-17]. Enterotoxin-like EntA, EntB and EntC, which are not regulated by the pleiotropic virulence gene regulator PlcR, cannot be excluded as causative agents of diarrheal syndrome [6]. Although toxin-related proteins most probably play a prevalent role in B. cereus pathogenicity, other proteins present in the extracellular milieu may be required, such as components of flagella [18], cell-wall peptidases [13,19,20], metalloproteases [21], antioxidant enzymes [22], siderophores for iron acquisition from the host [23] and moonlighting proteins [24-26]. All of these exoproteins are classified as virulence factors and constitute the major part of the B. cereus exoproteome as observed by shotgun, high-throughput methodology [6,27].

The regulation of B. cereus toxin gene expression mobilizes a complex machinery [28] that includes several pleiotropic regulators, such as the PlcR-PapR quorum sensing system [29,30], the CodY repressor [31], the ferric uptake regulator Fur [32], the catabolite control protein A (CcpA) [33], the two-component regulatory system ResDE, and the redox regulator Fnr. ResDE and Fnr are also involved in maintaining redox homeostasis by regulating catabolism [34-38]. Like ResDE and Fnr, the OhrRA system is involved in redox homeostasis [39]. Unlike ResDE and Fnr, the involvement of OhrRA in toxigenesis has not been yet reported in the literature. The B. cereus OhrRA system comprises OhrA (organic hydroperoxide resistance protein A), an antioxidant protein, and OhrR, a potential pleiotropic regulator belonging to the MarR (multiple antibiotic resistance) family of winged, helix-turn-helix, DNA-binding proteins [40,41]. However, B. cereus OhrR appears to be an atypical member of this family because it contains four cysteine residues at its N-terminal domain while most members have two cysteines [39].

Here, we investigated the impact of B. cereus OhrR and OhrA on toxigenesis. We compared the exoproteomes of ΔohrR and ΔohrA mutants with their parental strains from cells grown under three different ORP— one under aerobicosis and two under anaerobiosis. We found that OhrRA controls the toxigenic profile of B. cereus. The effect of OhrR on the production of toxin-related proteins is mediated at both the transcriptional (mainly by OhrR) and post-transcriptional (mainly by OhrA) levels. OhrR restricts toxin-dependent, antiproliferative activity of the B. cereus exoproteome on Caco-2 cells whatever the growth conditions, while OhrA restricts this activity only under low-ORP anaerobiosis.

2. Materials and methods

2.1. Bacterial strains and growth conditions

Escherichia coli BL21 CodonPlus(DE3)-RIL (Stratagene) was used for the heterologous production of OhrR. B. cereus ATCC 15479 was used as the parental strain [42]. Strains lacking OhrA (ΔohrA) and OhrR (ΔohrR) were constructed previously [6]. Wild-type and mutant strains were grown in pH-regulated batch cultures on glucose-containing MOD medium [43,44] under low-ORP anaerobiosis (initial ORP = -410 mV, pO2 = 0%), high-ORP anaerobiosis (iORP = -10 mV, pO2 = 0%) and full aerobicosis (iORP = +140 mV, pO2 = 100%), as previously described [39]. Supernatants were collected at the log/exponential growth phase, i.e. 1.5 h after reaching the maximal growth rate (when μ = 80 ±10 % of μmax). After growth, the culture supernatants were first filtered through a 45-μm-pore-size, low-adsorption, cellulose-acetate membrane filter (Sartorius) and then through a 20-μm-pore-size filter. Filtered fractions (40 mL) were quickly frozen and kept at −20 °C until use.

2.2. Relative quantification of gene expression using real time RT-PCR

RT-PCR was performed using SYBR Green technology on a Lightcyt instrument (Roche applied Science) as described previously [34]. The primers used in this study are shown in Table S1.

2.3. Exoprotein sample preparation and nanoLC-MS/MS analysis

Three independent biological replicates were harvested for each of the three conditions (aerobiosis, low- and high-ORP anaerobiosis) and three strains (wild-type, and ΔohrA and ΔohrR mutant strains). Extracellular proteins were extracted from the 27 culture supernatants, subjected to SDS-PAGE, and then identified after trypsin proteolysis and nanoLC-MS/MS tandem mass spectrometry, as previously described [6]. The Mascot search engine (version 2.2.04) was used to search all MS/MS spectra against an in-house polypeptide sequence database containing the sequence of all annotated proteins.
encoded by the B. cereus ATCC 14579 chromosome (NC_004722) and plasmid pBClin15 (NC_004721). This database comprises 5255 polypeptide sequences, totaling 1,455,982 amino acids. Searches for tryptic peptides were performed with the following parameters: full-trypsin specificity, a mass tolerance of 10 ppm on the parent ion and 0.6 Da on the MS/MS, static modifications of carboxyamidomethylated Cys (+57.0215), and dynamic modifications of oxidized Met (+15.9949). The maximum number of missed cleavages was set at 2. All peptide matches with a peptide score below a p-value of 0.01 were filtered by the IRMA 1.21.0 parser [45]. A protein was considered to be validated when at least two different peptides were detected in the same experiment. The false-positive rate for protein identification was estimated using the appropriate decoy database to be below 0.1% with such stringent parameters. The MS/MS data were deposited in the PRIDE Proteomics Identifications database (http://www.ebi.ac.uk/pride/) under accession numbers 27209–27211 via the Proteomexchange platform (ProteomeXchange DOI: http://dx.doi.org/10.6019/PXD000024).

2.4. Spectral count-based protein quantification and statistical analysis

The number of MS/MS spectra per protein recorded by nanoLC-MS/MS was extracted for each biological triplicate for all three growth conditions assayed. Proteins were further considered for comparison only if peptides were detected in at least two of the three replicates. The resulting datasets were normalized taking into account the total protein concentration of the corresponding supernatant. Protein concentrations in B. cereus were determined using the Reducing Agent Compatible Detergent Compatible (RCDC) protein assay (Bio-Rad) following the supplier’s instructions. MS/MS spectral counts were compared with the TFold method using the PatternLab software program 2.0.6.13 [46]. Fold-change and p-value cutoffs were set at 2 and 0.05, respectively. Log2 (fold-change) was then calculated for comparison. Principal component analysis (PCA) was carried out using Tanagra software 1.4.44 [47].

2.5. Recombinant overexpression of ohrR and protein purification

B. cereus OhrR was produced as a C-terminal His-tagged protein using E. coli BL21(pET101ohrR). For this, the coding sequence for ohrR was PCR amplified from B. cereus ATCC 14579 genomic DNA using the primers, ohrE1 and ohrE2 (Table S1). The ampiclon was cloned as a blunt-end PCR product into pET101/D-TOPO, yielding pET101ohrR. E. coli BL21 CodonPlus(DE3)-RIL(pET101ohrR) recombinant cells were grown until the OD600 reached ~0.7. Protein overproduction was then triggered with 1 mM isopropyl-β-D-thiogalactopyranoside. After incubation for 4 h, cells were harvested by centrifugation (10,000 g for 10 min at 4 °C) and suspended in 50 mM Tris-HCl buffer (pH 7.9) containing 200 mM NaCl. The purity of the protein was estimated as 98% by SD200 gel filtration chromatography and UV signal integration. The protein concentration (1.2 mg/mL) was determined using a Bradford assay (Intergen) with bovine serum albumin as the reference. Protein was stored at –80 °C until analysis.

2.6. Dynamic light scattering (DLS)

To evaluate the quaternary structure of OhrR in solution, purified protein sample was concentrated on an Ultracell YM10 microcon (Millipore) device to 2.4 mg/mL. An 80 μL sample of OhrR was loaded onto a 24 mL Superdex 200 10/300 GL column (GE Healthcare) previously equilibrated with 50 mM KH2PO4/Na2HPO4 buffered at pH 7.4 and containing 50 mM NaCl. The column was operated at a flow rate of 0.5 mL/min with the same buffer on an Agilent 1100 series, reverse-phase HPLC system equipped with a G1322A degasser, G1311A quaternary pump, and G1313A autosampler. The elution profile was monitored with a G1315B diode array detector (Agilent), a miniDawn Tristar multilange laser static light scattering detector (three angles: 45°, 90°, and 135°) coupled to a DynaPro Titan light-scattering instrument (Wyatt Technology) placed at 90°, and an Optilab rEX differential refractometer (Wyatt Technology). The 90° multilange light-scattering detector was calibrated with pure toluene, and bovine serum albumin was then used to normalize the other detectors (45° and 135°) in the corresponding buffer. Light scattering data were analyzed using ASTRA software (Wyatt technology).

2.7. Nonreducing SDS-PAGE

The presence and formation of disulfide bonds were monitored by nonreducing SDS-PAGE. Reduced protein samples were prepared by incubating OhrR (20 μg/L) with 1 to 100 μM dithiothreitol (DTT) for 30 min at 37 °C. Excess DTT was then removed by ultrafiltration. Oxidized OhrR was obtained by incubating reduced protein samples for 30 min at 37 °C with either 5 mM or 500 μM cumene hydroperoxide (CHP) or 1 μM H2O2. To test the reducing effect of LMW thiols and dihydroxylic acid, purified OhrR was first treated with 50 μM CHP as described above, then incubated for 30 min with increasing concentrations (0.05 mM to 25 mM) of either cysteine (Cys, Sigma), bacillithiol (BSH, JEMA Biosciences), coenzyme A (CoASH, Sigma) or dihydroxylic acid (DHLA, Sigma). Protein samples were then subjected to electrophoresis on nonreducing novex NuPAGE® SDS-PAGE 4–12% polyacrylamide gels (Invitrogen) and stained with SimplyBlue™ SafeStain (Invitrogen).

2.8. Electromobility shift assays (EMSAs)

Nucleic acid fragments containing the promoter regions of ohrRA, nhe, hbl, hblB, entA, entB, entC, hlyI, hlyII, entFM and cytK were PCR-amplified using biotinylated forward primers and nonbiotinylated reverse primers (listed in Table S6). The DNA
used as a negative control was a fragment of the ssuRNA BC0007 sequence (NC_004722) that was amplified with the biotinylated ssuBioF and ssuR primer pair. The 5’-labeled amplicons were purified using the High Pure PCR Product Purification Kit (Roche). Binding reactions were performed for 30 min at 37 °C by incubating biotin-labeled DNA fragments (2 nM per reaction) with either 0.05, 0.1, 0.2, 0.4, 0.8, 1 or 2 μM reduced OhrR (with 100 μM DTT) in 10 mM Tris-HCl buffered at pH 7.5 and containing 50 mM KCl, 2.5% glycerol, 5 mM MgCl₂ and 5 mg/L poly(dl-dc). The samples were resolved by electrophoresis on a 6% nondenaturing polyacrylamide gel and electrotransferred onto Hybond N + Nylon membranes (Amersham). Biotin-labeled DNA was detected using the LightShift Chemiluminescent EMSA Kit (Pierce). Oxidized OhrR was obtained by adding 50 μM CHP to the binding buffer. The DNA binding activity of oxidized OhrR was assayed in the presence of LMW thiol with 5 mM cysteine added to the binding buffer.

2.9. Growth inhibition tests with human Caco-2 cells

Growth inhibition of human Caco-2 cells with filtrate supernatants from aerobic, high and low-ORP anaerobic B. cereus cultures was assayed as previously described [6]. For each of the nine biological samples, growth inhibition tests were assayed twice.

3. Results

3.1. Exoproteome analysis of B. cereus ΔohrR and ΔohrA cells

Toxins are extracellular proteins and are the main components of the B. cereus exoproteome [6]. To determine whether Ohra contributes to toxin production, we identified changes in the exoproteomes of ΔohrR and ΔohrA mutants in comparison with the wild-type strain [6,27]. For this, cells were grown in pH-regulated batch cultures on glucose-containing medium under low-ORP anaerobiosis, high-ORP anaerobiosis and full aerobiosis. Three independent biological replicates were harvested for each of the three conditions and three strains. Extracellular proteins were extracted from the 27 culture supernatants and then identified and quantified by tandem mass spectrometry with a high-resolution LTQ-Orbitrap XL mass spectrometer. A dataset comprising 171,776 MS/MS spectra was recorded for the 27 samples. A total of 65,579 different spectra were confidently assigned to B. cereus peptides, resulting in the identification of 181 proteins (Table S2). The number of MS/MS spectra assigned to each protein was normalized for comparison using the protein concentration of the samples (Table S3). We reported the fold-change observed for each detected protein, taking into account spectral counts as input (Table S4). We systematically performed blast analyses to annotate wrong-gene functional assignments and to retrieve missing functions. The detected proteins were catalogued into five functional groups: 1) toxins, 2) degradative enzymes, 3) flagella components, 4) metabolism, and 5) others. The latter, which comprised less than 5% of the detected proteins, includes proteins associated with defense mechanisms, transport and membrane biogenesis, translation and transcription, and proteins without functional assignment. Fig. 1 and Table S4 show that the global

![Fig. 1 - Distribution of exoproteins in terms of function. The diagrams indicate the mean ratio of MS/MS spectra assigned to proteins classified into functional categories for the exoproteomes from wild-type (green), ΔohrA (blue) and ΔohrR (red) cells grown under low-ORP anaerobiosis, high-ORP anaerobiosis, and aerobic conditions. See Table S3 for details on the mean ratio estimated from triplicate independent experiments. Statistical differences between wild-type and mutant strains were evaluated with a Student’s t-test. * = P < 0.05.](image-url)
distribution of proteins among these five functional groups was significantly different in the mutant strains compared with the wild-type strain. This was observed whatever the growth condition. Specifically, the ΔohrR exoproteome exhibited a higher toxin content but lower flagella component content in the three conditions tested. The disruption of ohrA significantly decreases the toxin content of low-ORP anaerobically grown cells and increases the flagella component content. These results indicate that OhrR and OhrA modulate the B. cereus exoproteome in a redox-dependent manner.

3.2. Impact of OhrR and OhrA on exoprotein-specific changes

3.2.1. Evidence for novel exoproteins

Detailed analysis of the ΔahrA or ΔohrR exoproteomes revealed the presence of 40 additional proteins compared with the parental strain exoproteome (Table S3). A total of 8 and 20 proteins were found exclusively in the ΔahrA and ΔohrR exoproteomes, respectively. Among these, one and three proteins of unknown functions – annotated as NP_832962.1 and NP_831667.1, NP_834698.1 and NP_831845.1 – were shown to be abundant components of the ΔahrA and ΔohrR exoproteomes, respectively. NP_832962.1 encompasses an LPXTG-motif and shows similarities to a cell wall anchor domain. NP_831667.1 is annotated as a phage protein. NP_834698.1 exhibits similarities with the ErfK/YukD LD-transpeptidase catalytic domain (COG1376) and could be involved in peptidoglycan crosslinking. NP_831845.1 belongs to the uncharacterized COG4849 orthologous group and shows some similarities to the small ESAT-6 protein that has been characterized as central to virulence and protective immunity in Mycobacterium tuberculosis [50].

3.2.2. Focus on toxin-related proteins

Table 1 lists the abundance level changes of toxin-related proteins in ΔahrA and ΔohrA mutants. We noted that (i) the abundance levels of Hbl and Nhe components significantly decreased in ΔahrA cells and increased in ΔohrR cells when grown under low-ORP anaerobiosis, (ii) the HlyI abundance level was significantly decreased in the ΔahrA exoproteome, whatever the conditions, (iii) HlyI was significantly more abundant in the ΔohrR exoproteome, respectively. NP_832962.1 encompasses an LPXTG-motif and shows similarities to a cell wall anchor domain. NP_831667.1 is annotated as a phage protein. NP_834698.1 exhibits similarities with the ErfK/YukD LD-transpeptidase catalytic domain (COG1376) and could be involved in peptidoglycan crosslinking. NP_831845.1 belongs to the uncharacterized COG4849 orthologous group and shows some similarities to the small ESAT-6 protein that has been characterized as central to virulence and protective immunity in Mycobacterium tuberculosis [50].

3.3. Dual role of B. cereus OhrR as a repressor and an activator of toxin gene expression

Based on (i) the functions of its orthologs [51], (ii) our previous proteomic data [39], and (iii) our exoproteomic results, we believe that OhrR, unlike OhrA, controls the toxin content of the B. cereus exoproteome at the transcriptional level. We thus quantified the expression of operons or genes encoding toxin-related proteins in wild-type and ΔohrR and ΔohrA cells using RT-PCR. Fig. 2 associates the mRNA level changes with the protein level changes in mutant strains relative to the wild-type strain. As expected, there is no correlation between protein and mRNA level variations of the toxin genes in the ΔohrA mutant, suggesting that OhrA-dependent toxin secretion may occur mainly at the post-transcriptional level, whatever the redox conditions. However, the variations of mRNA levels are consistent with proteomic data in the ΔohrR mutant, with some exceptions: (i) HlyI and CytK in the three culture conditions tested, (ii) HlyI under both high-ORP anaerobiosis and aerobiosis, and (iii) NheC under aerobiosis. This characteristic distinguishes these four toxins from the others and suggests that OhrR-dependent regulation of the HlyI, HlyI, CytK and NheC abundance level may occur mainly at the post-transcriptional level in these growth conditions. Moreover, the results indicate that OhrR may act as a transcriptional activator of entB and a repressor of hblABC, nheA, nheB, hblB, entFM, entA and entC, whatever the conditions.

3.4. B. cereus OhrR is mainly a noncovalent dimer in its reduced form and a covalent dimer in its oxidized form

Orthologs of B. cereus OhrR regulate their target genes by directly binding to the operator site located in the vicinity of its target promoters [52]. To investigate whether B. cereus OhrR may bind to toxin-related protein gene promoters in in vitro experiments, ohrR was expressed in E. coli and purified as a recombinant, His-tagged protein. The Mr of the purified polypeptide, as estimated by SDS-PAGE under reducing conditions, was ~20 kDa as expected (Fig. S1). Its native quaternary structure was monitored by gel filtration coupled to a Multi-Angle laser Light Scattering (MALS) detector. The protein behaved as a relaxed homodimer in solution with a molecular weight estimated at ~62 (±1) kDa and a hydrodynamic radius of 3.2 (±0.3) nm (Fig. S2). Nonreducing SDS-PAGE revealed an abundant protein band corresponding to OhrR monomer and a doublet corresponding to OhrR covalently linked dimer (Fig. 3A). Taken together, these data indicate that the dimeric OhrR population detected by Dynamic Light Scattering (DLS) is a mixed population comprising mainly noncovalently associated subunits, but also covalent dimers corresponding to oxidized forms of OhrR. Moreover, covalent dimers could adopt two conformations differing in terms of rigidity, with the more relaxed structure migrating more slowly than the rigid structure. To test the ability of noncovalent OhrR dimer to convert into covalent dimer, purified protein (20 μM) was treated with the oxidizing agent, cumene hydroperoxide (CHP). Fig. 3A shows the CHP dose-dependent formation of rigid, covalently linked OhrR dimers as analyzed by SDS-PAGE in nonreducing conditions. Increasing the concentration of CHP from 5 μM to 50 mM led to the disappearance of the protein band corresponding to OhrR monomer and a concomitant increase in the amount of the faster-migrating band corresponding to a covalent dimer. Moreover, at the highest CHP concentrations, we observed faint bands corresponding to tetrameric and octomeric species. Interestingly, only the slower-migrating band corresponding to covalently linked OhrR dimer in a relaxed form was detected after treatment with 500 mM CHP. In addition, the tetrameric
Table 1 – Fold-changes in abundance levels (log2 ratio) of exoproteins classified as toxins.

<table>
<thead>
<tr>
<th>Protein</th>
<th>Functional annotation</th>
<th>Accession</th>
<th>ΔohrA versus wild-type</th>
<th>ΔohrR versus wild-type</th>
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</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>Low ORP anaerobiosis</td>
<td></td>
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<tr>
<td></td>
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<td>High ORP anaerobiosis</td>
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<td>Low ORP anaerobiosis</td>
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<td>High ORP anaerobiosis</td>
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<td>Hbl2 Enterotoxin HBL component L2</td>
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<td>-0.286 0.240</td>
<td>0.324 0.259</td>
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<tr>
<td>HblB Enterotoxin HBL component B</td>
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<td>0.918 0.048</td>
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<tr>
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<td>-0.902 0.087</td>
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* Each log2 ratio value represents the mean protein level of the mutant in relation to the wild-type sample. The relative amount of each protein was determined using PatternLab software. Plus and minus symbols indicate up- and down-regulation of the protein, respectively. Numbers in bold indicate data that satisfied statistical criteria (p-value < 0.05)
and octameric species showed higher apparent molecular weights at the highest CHP concentrations than those detected with lower CHP concentrations.

The reversibility of CHP oxidation was assayed by incubating OhrR with either DTT (E₀ = −0.33 V) or free dihydrolipoate (DHLA, E₀ = −0.32 V) as a dithiol/disulfide redox component, or the three major low-molecular-weight (LMW) monothiols found in B. cereus [55], i.e. cysteine (Cys, E₀ = −0.21 V), bacillithiol (BSH, standard redox potential is unknown) and Coenzyme A (CoASH, E₀ = −0.33 V). Fig. 3B shows representative nonreducing SDS-PAGE gels where dose-dependent samples were resolved. Whatever the reductant, OhrR reduction led to appearance of monomeric OhrR concomitantly with i) first, the disappearance of the faster-migrating form of the OhrR covalent dimer, and ii) second, bleaching of the slower band. These data indicated that the rigid and relaxed conformations of the covalent OhrR dimers differ in terms of the presence of disulfide bridges. The more rigid form may result from a reversible intramolecular disulfide bridge that is easily accessible to the reductant. The relaxed form may result from a less-accessible intermolecular disulfide bridge. As shown in Fig. 3, we found that the dihthiol, DHLA, is a stronger reductant for the OhrR protein than DTT and the monothiols, Cys, BSH and CoASH. Of the three LMW monothiols, Cys appeared to be the strongest reductant. The relaxed form may result from a disulfide bridge that is easily accessible to OhrR dimers in its reduced form and a covalent dimer in its oxidized form. We also showed that DHLA and LMW thiols may participate, in vivo, in the recovery of reduced OhrR from the oxidized form.

### 3.5. The redox state of OhrR is critical for its binding to toxin gene promoters

In order to determine if reduced OhrR binds directly to toxin gene promoter regions to mediate transcriptional repression or activation, electrophoretic mobility shift assays (EMSA) were performed using DTT-reduced, purified OhrR and DNA fragments corresponding to promoter regions of the toxin-related operons or genes. Because of its very large size, the promoter region of hbl was divided into two overlapping fragments of 636 bp (hbl1) and 610 bp (hbl2). Binding specificity was shown by the absence of a mobility shift with the negative control performed with the ssu promoter region (Fig. 4A), the appearance of protein–nucleic acid complexes upon addition of an excess (50-fold) of unlabeled heterologous promoter regions, and the maintenance of protein–nucleic acid complexes in competition assays using a ten-fold excess of unlabeled heterologous promoter regions assayed except entFM. Reduced OhrR showed markedly different behavior in gel-shift titration assays depending on the promoter region: (i) OhrR bound to the entFM promoter region in an ordered manner, producing a first protein–nucleic acid species faintly visible from 0.05 μM to clearly visible at 0.4 μM OhrR, and then a second, slowly moving species at concentrations above 0.4 μM; (ii) increasing amounts of OhrR

<table>
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<th>Low-ORP anaerobiosis</th>
<th>High-ORP anaerobiosis</th>
<th>Aerobiosis</th>
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<tbody>
<tr>
<td>ΔohrA</td>
<td>ΔohrA</td>
<td>ΔohrA</td>
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<tr>
<td>log₂(fold-change) mRNA</td>
<td>log₂(fold-change) proteins</td>
<td>log₂(fold-change) proteins</td>
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Fig. 2 – Correlation of mRNA level changes with protein abundance changes in ΔohrA and ΔohrR cells relative to wild-type cells when grown under low-ORP anaerobiosis, high-ORP aerobiosis, and aerobiosis. Relative transcript abundance data are plotted against relative exoprotein abundance data.
3.6. OhrR and OhrA modulate the inhibitory effect of the B. cereus exoproteome on the growth of Caco-2 cells

We showed previously that the B. cereus exoproteome had inhibitory effects on the cellular growth of human colonic epithelial Caco-2 cells, which are recognized as a model of the intestinal mucosa [56]. To evaluate the contribution of OhrR and OhrA to the anti-proliferative activity of the B. cereus exoproteome, we compared the inhibitory effects of ΔohrR, ΔohrR and wild-type culture filtrate supernatant on Caco-2 cell growth. Two exoprotein concentrations were assayed: 1.6 ± 0.1 mg/L and 0.3 ± 0.1 mg/L (Table 2). At the higher exoprotein concentration, the ΔohrR culture filtrate supernatant abolished Caco-2 cell growth in all cases while the ΔohrR culture filtrate supernatant abolished Caco-2 cell growth only when prepared from low-ORP, anaerobically grown cells. At the lower exoprotein concentration the inhibitory effects were less pronounced, resulting in a poor difference between the ΔohrA and wild-type culture filtrate supernatants, whatever the conditions. However, the ΔohrR culture filtrate supernatant retained its high inhibitory activity when prepared from low-ORP, anaerobically grown cells, and exhibited a higher inhibitory activity than the high concentration culture filtrate supernatant of the wild-type strain obtained from high-ORP, anaerobically or aerobically grown cells. Taken together, these data indicate that OhrR restricts the Caco-2-anti-proliferative activity of the B. cereus exoproteome, whatever the growth conditions — markedly so when the B. cereus cells are grown under low-ORP anaerobiosis. Similarly to OhrR, but to a lesser extent, OhrA restricts the Caco-2-anti-proliferative activity of the B. cereus exoproteome when bacteria are grown under low-ORP anaerobiosis.

3.7. Changes in toxin abundance levels explain the OhrR- and OhrA-dependent inhibitory effect of the B. cereus exoproteome on the growth of Caco-2 cells

We showed that OhrR, and to a lesser extent OhrA, impact both the toxin abundance level (Table 1) and the inhibitory effect of the B. cereus exoproteome on Caco-2 cell growth (Table 2). To determine whether changes in toxin levels could explain the differences between the antiproliferative activities of ΔohrR, ΔohrA and the wild-type exoproteome, we performed a principal component analysis (PCA). Variables were organized as active and supplementary. The active variable group included the content of each toxin-related protein as evaluated by our shotgun proteomic approach and the supplemental variable group included the percentage of growth inhibition on Caco-2 cells (noted as cytotoxicity in Fig. 5). The corresponding data and analytical details are given in Table S5. Projection of the variables on the PC1–PC2 plane synthesized the PCA (Fig. S) and shows that (i) by restricting the exoproteome to toxin-related protein we reconstructed the variability of "cytotoxicity", (ii) the variability of a subset of toxin components, namely EntA, EntB, EntC, EntFM, HblL1, HblL2 and HblB, resulted in a gradual decrease of hbl1, hbl2 and nhe DNA mobility above 0.8, 0.2, and 0.8 μM, respectively, and (iii) higher concentrations of reduced OhrR were needed to form DNA complexes with hbl1 and hbl2, hblB, nhe, hlyI, hlyII, entA and entC promoter regions than with its own promoter region. Adding 50 μM CHP to the EMSA reaction mixtures abolished the binding of 1 μM reduced OhrR to its own promoter and to all the toxin gene promoters except entB. Only the faster-migrating protein-entB species was observed with CHP-oxidized OhrR (Fig. 4A). The initial binding behavior was recovered after addition of 100 μM DTT, whatever the promoter region (data not shown).
NheA, NheB and NheC, and (iv) the variability in “cytotoxicity” was highly negatively correlated with EntB variability, whatever the redox growth condition, and highly positively correlated with HlyII variability under low-ORP anaerobiosis. From these results, and those reported in Table 2, we concluded that EntB is the toxin which contributes the most to the overall difference between ΔohrR and wild-type antiproliferative activities on Caco-2 cells.

Table 2 – Mean percentage of Caco-2 cell growth inhibition by B. cereus exoproteomes.

<table>
<thead>
<tr>
<th>Exoprotein concentration</th>
<th>Low-ORP anaerobiosis</th>
<th>High-ORP anaerobiosis</th>
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<tbody>
<tr>
<td></td>
<td>WT</td>
<td>ΔohrR</td>
<td>ΔohrA</td>
</tr>
<tr>
<td>1.6 mg/L</td>
<td>28&lt;sup&gt;a&lt;/sup&gt;</td>
<td>100</td>
<td>100</td>
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<tr>
<td>0.3 mg/L</td>
<td>13</td>
<td>100</td>
<td>20</td>
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<sup>a</sup> Percentage of Caco-2 cell growth inhibition was measured after 96 h-incubation with exoproteomes containing 1.6 and 0.3 mg/L of proteins. Exoproteomes were prepared from cells grown under low-ORP anaerobiosis, high-ORP anaerobiosis, and aerobiosis. Data are the means of 6 values. For clarity, standard deviations (below 10%) are not shown.
Fig. 5 – Correlation circles for principal component analysis (PCA) on toxin-related proteins of wild-type and \( \Delta ohrR \) and \( \Delta ohrR \) mutant strains. Vectors are the loadings on PC1 (x-axis) and PC2 (y-axis). Vector length indicates the strength of the relationship and the angle between two vectors gives the degree of correlation (adjacent = highly correlated, orthogonal (90°) = uncorrelated, and opposite (180°) = negatively correlated). For details, see Table S5.

4. Discussion

Similarly to its orthologs in other bacteria, B. cereus OhrR functions as a transcriptional regulator that binds to its own promoter in its dimeric, reduced form [52,57-60]. Unlike most of its orthologs, B. cereus OhrR may regulate genes other than ohrRA. Notably, we found that OhrR may regulate, directly or indirectly, the transcription of all operons and genes encoding toxin-related proteins that have been identified in the B. cereus exoproteome. Based on our EMSA data (Fig. 4), OhrR does not directly control the transcription of the entFM locus. However, entFM expression was increased in the \( \Delta ohrR \) mutant (Table 2). This indicates that the OhrR-dependent regulation of entFM transcription is indirect. Transcriptomic analysis indicated little, if any, change in cytK mRNA in the absence of OhrR, suggesting a weak transcriptional activity of OhrR on cytK in our growth conditions. We also demonstrated a striking dissimilarity between OhrR-dependent regulation of entB and of other toxin-related genes, in that: (i) OhrR was an activator of entB and a repressor of other toxin- and putative toxin-encoding genes, and (ii) oxidized OhrR is capable of binding entB but did not form a 2× dimers complex, as observed with the reduced form. Previous studies have defined the consensus binding site of OhrR from other bacteria as an AT-rich, palindromic sequence of approximate-2 × 7 nucleotides separated by a variable-length spacer of 1 to 17 nucleotides [58]. Inspection of the nucleotide sequences for the promoter regions tested here indicated the recurrent presence of AT-rich sequences. A perfect inverted-repeat motif (TAAAATT-N4-AATT) is present in the entB promoter region and could be a good candidate as the binding site for OhrR (Fig. 4B). Another AT-rich dual motif showing only one single nucleotide deviation (TAAAATT-N6-AATT) flanks the first binding site. The presence of tandem binding sites on entB DNA and the presence of two migrating species in the corresponding EMSA (Fig. 4A) suggest occupation of two binding sites at a moderately low concentration of reduced OhrR and occupation of both DNA binding sites at a higher concentration of reduced OhrR. This indicates that one entB DNA molecule could bind two noncovalent dimers of OhrR (reduced form) and only one covalent-linked dimer of OhrR (oxidized form), as only the faster-migrating protein-DNA complex was observed with the oxidized form. For ohrR, we found two seven-nucleotide motifs that are almost perfect inverted repeats (Fig. 4B) but differ in sequence compared with the other motifs identified. This could explain why B. cereus OhrR showed lower DNA binding affinity to its own promoter than did its orthologs [52,61]. There are two other, major possible explanations. First, B. cereus OhrR differs structurally from its orthologs, and second, we chose a longer target DNA to sample the entire ohrRA promoter region. Four putative binding sites were found in the hbl promoter region (two in hbl11 and two in hbl2), three were found in the nhe promoter region, two in entA, and one in the hblB′, cytK, hlyI, hlyII, and entC promoter regions, while no motif could be detected within the entFM promoter region. We noted that putative binding sites in cytK, hlyI and hlyII differed significantly from the consensus described in other bacteria, because the two binding motifs were either separated by more than 17 nucleotides or did...
not involve the core tetranucleotide AATT sequence. Accordingly, EMSA data showed that reduced OhrR bound to cytK, hlyI and hlyII promoter regions with a lower affinity than to the other toxin gene promoter regions. Thus, direct binding of OhrR to promoter regions for toxin-related proteins may depend on both its redox state and its AT-rich target binding site. Globally, OhrR binding to toxin promoter regions required a high protein concentration in our in vitro analysis. Given that toxin gene expression is multi-regulated, it is possible that the stability of the toxin promoter-OhrR complex requires several additional protein partners in B. cereus [62].

In addition to toxins, OhrR regulates the abundance level of flagellins (Table S4). Inspection of the promoter region of the fla operon [42], which encodes flagellins A, B, and C, led to the identification of an inverted repeat sequence (TACAATT-N8-AATTGAT) highly resembling the OhrR-like, AT-rich box found in toxin promoter regions. OhrR may, thus, regulate motility by directly modulating fla gene expression levels. Our previous analysis of the B. cereus intracellular proteome showed significant changes in the abundance levels of the pleiotropic regulators, SinR, AbrB, CcpA and CodY [39]. These regulators were recognized as modulating toxin production and/or motility [31,33,63,64]. Inspection of the promoter regions of their genes indicated a putative OhrR-like box, except for CcpA (Fig. S3). Thus, OhrR could orchestrate the expression of virulence determinants in B. cereus, both directly, by modulating toxin and flagella gene expression, and indirectly, by impacting key regulatory circuits. Several studies converge to suggest the existence of a connection between motility and cytotoxicity in B. cereus, as is the case for several other pathogenic bacteria [65–69]. Our findings suggest that OhrR is a key player in this connection.

As for OhrR, OhrA controls the abundance level of toxin-related proteins in the B. cereus exoproteome in a redox- and toxin-dependent manner (Tables 1 and S4). Our data suggest that OhrA may act as a post-transcriptional regulator by controlling the stability and/or correct localization of toxin-related proteins in a protein- and redox-dependent manner. Given that NheC (NP_831584), unlike NheA (NP_831582) and NheB (NP_831583), contains two cysteines and thus features thiols vulnerable to oxidative modifications, OhrA could be needed to fully protect NheC against ROS damage and to sustain high-level secretion of NheC via the Sec pathway under aerobiosis. Unlike other toxins, HlyI has been recognized as a thiol-activated cytolsin [70]. This could offer two different targets for OhrA control compared with other toxins and could also explain why HlyI was positively controlled by OhrA, whatever the conditions. Why does HlyII escape redox-dependent OhrA control? The answer is probably linked to the existence of a very different control of HlyII export than for the other toxins [32]. Clearly the redox-dependent OhrA control of toxin level is a complex process and needs further investigation. Our exoproteomic analyses have, however, revealed a critical requirement for OhrA activity when OhrR is present, to maintain similar levels of toxin (except HlyII) whatever the conditions. Finally, as for OhrR activity, the activity of OhrA appeared to be higher under low-ORP anaerobiosis than under high-ORP anaerobiosis and aerobiosis.

Earlier studies [reviewed in [5]] have demonstrated that the level of toxin gene expression plays a major role in determining the cytotoxic potential of a bacterial strain, thus relegating the presence of a specific toxin to a less important role. Our study suggests that EntB is a crucial determinant of the OhrR-dependent inhibitory effect of the B. cereus exoproteome on model intestinal cell lines. EntB is structurally similar to both EntA and EntC [6] and we observed here an OhrR-dependent negative correlation between the abundance level of EntB and those of EntA and EntC and Nhe, Hbl and EntFM. Evidently, further intensive work is now required to elucidate the function of EntB, which could be crucial in triggering EntA and EntC- and Nhe, Hbl and EntFM-dependent cytotoxicity of the B. cereus exoproteome.

5. Conclusion

In our previous study, we proposed that B. cereus restricts its fermentative growth under low-ORP anaerobiosis and sustains efficient aerobic respiratory metabolism, motility, and stress response via OhrRA-dependent intracellular proteome remodeling [39]. Here, we found that OhrRA plays a major role in toxinogenesis, especially under low-ORP anaerobiosis. From this aspect, B. cereus OhrRA is unique compared with other OhrRA systems [39,51,52]. In the natural evolutionary context, the B. cereus OhrRA system appears to have been shaped to facilitate B. cereus survival and toxinogenesis under various redox growth conditions, such as those encountered in human intestine.

Supplementary data to this article can be found online at http://dx.doi.org/10.1016/j.jprot.2013.10.024.

Acknowledgments

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