SHORT COMMUNICATION

*Helicobacter pylori* protein oxidation influences the colonization process

Renata Godlewskaa, Artur Dzwonekb, Michał Mikula b, Jerzy Ostrowski b, Marcin Pawłowskic, Janusz M. Bujnickic, Elżbieta K. Jagusztyn-Krynickaa,⁎

aDepartment of Bacterial Genetics, Institute of Microbiology, Warsaw University, Miecznikowa 1, PL-02-096 Warsaw, Poland
bDepartment of Gastroenterology, Medical Center for Postgraduate Education, Maria Skłodowska-Curie Memorial Cancer Center, and the Institute of Oncology, PL-02-781 Warsaw, Poland
cLaboratory of Bioinformatics and Protein Engineering, International Institute of Cell and Molecular Biology, Trojdena 4, PL-02-109 Warsaw, Poland

Received 15 March 2005; received in revised form 20 October 2005; accepted 8 November 2005

Abstract

Dsb proteins control the formation and rearrangement of disulfide bonds during the folding of membrane and exported proteins. Here we examined the role of DsbI protein in *Helicobacter pylori* pathogenesis and demonstrated that a *dsbI* mutant impaired in disulfide bond formation revealed a greatly reduced ability to colonize mice gastric mucosa.

© 2006 Elsevier GmbH. All rights reserved.

Keywords: *Helicobacter pylori*; Colonization; Protein oxidation

Introduction

*Helicobacter pylori* is a Gram-negative spiral-shaped *e*-proteobacterium, which colonizes the gastric mucosa of humans. Infections caused by *H. pylori* are associated with different symptoms ranging from mild gastritis to ulcers. In addition, some chronic infections, acquired early in life, confer an increased risk of gastric cancer (Nardone and Morgner, 2003; Isakov and Malfertheiner, 2003). Although more than 50% of the human population is infected with *H. pylori*, only a subset develops a disease. Moreover, it has been documented that disease symptoms are correlated with the genotype of the two interacting organisms, the pathogen and the host. To date, numerous virulence factors responsible for the diversity of aspects of pathogen–host cells interaction have been described and characterized. The majority are extracytoplasmic proteins, mainly soluble periplasmic, membrane or exported proteins (Lamarque and Peek, 2003). Some essential virulence factors such as VacA or CagA are translocated into target cells via different secretion mechanisms (Hatakeyama, 2004; Montecucco and de Bernard, 2003). It was documented that the proper folding of a three-dimensional active structure of many extracytoplasmic virulence factors of Gram-negative pathogens require the complex action of Dsb proteins involved in disulfide bond formation and isomerization (Kadokura, 2003).

Materials and methods

Bacterial strains and culture conditions

*H. pylori* strain SS1 was grown under microaerobic conditions at 37 °C on blood agar base No. 2 containing 10% horse blood. The medium for the *H. pylori* mutant

⁎Corresponding author. Tel.: +48 22 554 1216; fax: +48 22 554 1402.
E-mail address: kjkryn@biol.uw.edu.pl (E.K. Jagusztyn-Krynicka).
contained kanamycin (25 µg/ml). A preculture was used to inoculate liquid tryptophane soy broth supplemented with 5% fetal bovine serum. The flasks were kept under microaerobic conditions at 37 °C with constant shaking (150 rpm), and bacterial cells were used for mice inoculation.

**Recombinant DNA techniques**

Gene replacement was used to construct the *H. pylori dsbI* isogenic mutant. The construction of the plasmid containing the *dsbI* gene disrupted by a kanamycin-resistant cassette (*Campylobacter jejuni aph(3′)-III* gene) was described by Raczko et al. (2005). This mutation does not have any polar effect on downstream genes. The Km cassette DNA fragment lacking a transcriptional termination signal was inserted in the orientation corresponding to HP595 transcription. Its orientation was checked by PCR. Additionally, we also investigated HP596 transcription and found that HP596 (located downstream) is transcribed from its own promoter. The *H. pylori* SS1 strain, naturally competent for genetic transformation, was used for generation of isogenic *H. pylori dsbI* mutants. The obtained kanamycin-resistant transformants were screened by hybridization using cloning vector DNA as a probe to confirm that no vector DNA was integrated into the chromosome through single cross-over and analyzed by PCR to prove that the predicted double crossover of the mutated allele had occurred. Further experiments showed that the lack of *dsbI* gene product is not lethal for bacterial cells. The mutant showed typical *Helicobacter* morphology and no alterations in growth rate.

**Infection of mice and DNA preparation**

C57BL/6, outbred specific pathogen-free mice (6–8-week-old) were experimentally infected with either *H. pylori* SS1 or isogenic *dsbI*-mutated *H. pylori* SS1. DNA from gastric mucosa was obtained as described previously (Dzwonek et al., 2004; Mikula et al., 2003). Non-infected 6–8-week-old C57BL/6 mice were used as a control.

**Real-time PCR**

The infection with *H. pylori* was quantified 3 weeks post-challenge by real-time PCR of the 26-kDa species-specific antigen gene, as described previously (Mikula et al., 2003). The calculated amounts of total *H. pylori* DNA and the number of bacterial cells present in each sample were normalized to murine *GAPDH* DNA. Student’s *t*-test was used to calculate the significance of the effects of the *dsbI* mutant on the colonization process. *p* < 0.05 was considered to be significant.

**Bioinformatics analysis**

The homology search and multiple alignment were performed using Psi-blast program (Altschul et al., 1997) and ClustalW (Thompson et al., 1997). The predictions of extracytoplasmic proteins of *H. pylori* were performed using the ConPred II server (Masafumi et al., 2004) and the LipoP server (Juncker et al., 2003).

**Results and discussion**

The most extensively studied disulfide oxidoreductases are those of *Escherichia coli* which generate two distinct pathways functioning in the periplasmic space: the oxidative pathway (*DsbA, DsbB*) and the isomerization pathway (*DsbC, DsbD, DsbE*) (Kadokura, 2003). We have previously identified and characterized a new *H. pylori* DsbB-like thiol-oxidoreductase encoded by the JHP0542 gene of J99 and HP0595 of 26695 (Raczko et al., 2005). HP0595 belongs to the DsbI protein family, which is paralogous to the DsbB family. DsbB accepts electrons from the periplasmic protein DsbA, while the substrates of DsbI still remain unknown. We used PSI-BLAST to search the genomes of *e*-Proteobacteria for the orthologs of *dsbA, dsbB*, and *dsbI* genes, using their protein family members from *H. pylori* and *C. jejuni* (Thompson et al., 1997; Altschul et al., 1997). As expected, the results of our analysis (Table 1) show that the distribution of DsbA and DsbB is strongly correlated. The only notable exception is the lack of a DsbB ortholog in *C. lari*. Interestingly, the genome of this bacterium contains an additional copy of a DsbI ortholog. We hypothesize that one of the copies of DsbI in *C. lari* has taken over the function of the lost DsbB protein and acquired the ability to interact with DsbA. We have previously shown that in *C. jejuni*, DsbB can complement

<table>
<thead>
<tr>
<th>Organism</th>
<th>DsbI</th>
<th>DsbB</th>
<th>DsbA</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Campylobacter coli</em> RM2228</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td><em>Campylobacter jejuni</em> RM1221</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td><em>Campylobacter jejuni</em> subsp. jejuni NCTC 11168</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td><em>Campylobacter lari</em> RM2100</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td><em>Campylobacter upsaliensis</em> RM3195</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td><em>Helicobacter hepaticus</em> ATCC 51449</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td><em>Helicobacter pylori</em> 26695</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td><em>Helicobacter pylori</em> 399</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td><em>Wolinella succinogenes</em> DSM 1740</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
</tbody>
</table>

+ or – indicate the presence or absence of a member of the respective family. + + indicates the presence of two members.

*Unfinished genome, whole genome shotgun sequence.*
for the lack of DsbI, but not vice versa. Thus, it will be interesting to carry out complementation tests with both DsbI proteins from C. lari.

The analysis of both H. pylori strains (26695, J99) revealed the presence of DsbI proteins in the absence of DsbA, which suggests that “orthodox” DsbI proteins most likely act on other substrates. In this case, the oxygenation of a pair of cysteines in target proteins must be carried out by some other, yet unknown mechanism. This observation points to the differences in the interactions between Dsb proteins acting in two closely related microorganisms – Campylobacter and Helicobacter – and strongly implies a leading role for DsbI in oxidative folding of Helicobacter proteins.

In this report, considering the potentially essential physiological role of DsbI, we decided to analyze the influence of a dsbI gene knockout on the colonization process. To achieve this, an isogenic dsbI-mutated H. pylori SS1 strain was used for experiments on mice and the level of colonization was monitored by RT-PCR. H. pylori SS1 is a mouse-adapted strain, isolated by Lee et al. (1997). Although some pathological symptoms differ from those provoked by human infection, the murine model has been widely used to evaluate different aspects of Helicobacter pathogenesis. Recently, it has been shown that the immunoproteome of infected mice resembles that of infected humans, which further validated this experimental model (Bumann et al., 2002).

The mutation in dsbI almost eliminated the ability of H. pylori to colonize the stomach (p < 0.01). These results support the importance of the disulfide bond formation process in the infection. In one out of seven mice, real-time PCR confirmed a low level of residual infection, but the remaining animals remained uncolonized. The extrapolated amount of bacterial DNA quantified by real-time PCR corresponded to 2230–18975 bacterial cells/100ng of mouse DNA in gastric samples from seven control mice (infected by the wild-type strain SS1) and 0–183 bacterial cells/100ng of mouse DNA in samples from seven mice infected with the isogenic mutant in dsbI (Fig. 1). Mucosa specimens were also checked for H. pylori colonization using the qualitative rapid urease test (CLO test) which was read routinely 1 h after gastric mucosa specimen collection and, if negative, 3 and 24 h later. Development of a pink color was indicative of a positive test. In each mouse infected by SS1 wild-type strain the CLO test was positive, whereas it was negative in the case of all mice infected by the mutated strain. The complementation experiment (dsbI mutant transformed with dsbI gene on a plasmid) was not performed due to the fact that H. pylori strain SS1 is poorly tolerant to non-self DNA (Gorrell et al., 2005). A complementation test was conducted using H. pylori J99 as described in the previously published paper (Raczko et al., 2005) but this strain does not colonize mice.

Proteins, which are affected by dsbI mutations are still unknown, but as in many pathogenic bacteria, the dsbI defect may have an adverse effect on the conformation of many virulence determinants resulting in their misfolding and instability. We carried out a prediction of extracytoplasmic proteins in H. pylori. The criteria used were: identification of putative transmembrane helices (Masafumi et al., 2004) or identification of an Spl- or SpII-type signal sequence (Juncker et al., 2003). We found 366 extracytoplasmic proteins in H. pylori, 211 of which contain at least two cysteines. This number represents the upper estimation of a set of possible targets of DsbI.

Analysis of experimental data from two-hybrid assays of H. pylori available in the http://dpi.nhri.org.tw/hp/database (Lin et al., 2005) reveals that HP0595 interacts with: HP0025 (Omp2), HP0060, HP0870 (FlgE), HP1052 (EnvA), HP1173, and HP1430. Among these proteins, perhaps the most interesting in terms of pathogenesis is HP1173. It bears a resemblance to a periplasmic protein LipoP, predicted to be anchored with its N-terminus in the inner membrane. HP1173 has two Cys residues, which are most probably located on the surface of the protein (according to our preliminary structure prediction), suggesting that they could be oxygenated by HP0595. It is noteworthy that HP1173 has an ortholog in the J99 strain, but not in any other bacterial genomes analyzed. HP1173 itself interacts with four proteins, which may be involved in pathogenicity of Helicobacter.

Another interesting protein interacting with HP0595 is Omp2 (HP0025). Forty paralogs of hp0025 were found in the H. pylori genome, 35 of these paralogs do not have a homolog in other hitherto sequenced genomes. Proteins from the HP0025 family possess characteristic Cys residues, which again make them possible substrates of HP0595. Moreover, 27 of these proteins are, like HP0595, extracytoplasmic proteins (according to LipoP analysis). However, no experimental data are available, to suggest that other members of the HP0025 family interact with HP0595.
Acknowledgments

This work was supported by the Ministry of Scientific Research and Information Technology (Grants no. 4 P05A 078 17 and 6 P06 K 043 21).

References


