Regulation of *Helicobacter pylori* Virulence Within the Context of Iron Deficiency

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*Helicobacter pylori* strains that harbor the oncoprotein CagA increase gastric cancer risk, and this risk is augmented under iron-deficient conditions. We demonstrate here that iron depletion induces coccoid morphology in strains lacking cagA. To evaluate the stability of augmented *H. pylori* virulence phenotypes stimulated by low-iron conditions, *H. pylori* isolated from iron-depleted conditions in vivo were serially passaged in vitro. Long-term passage decreased the ability of hypervirulent strains to translocate CagA or induce interleukin 8, indicating that hypervirulent phenotypes stimulated by low-level iron conditions are reversible. Therefore, rectifying iron deficiency may attenuate disease among *H. pylori*-infected persons with no response to antibiotics.

**Keywords.** *Helicobacter pylori*; iron deficiency; CagA; gastric cancer.

*Helicobacter pylori* is the strongest known risk factor for the development of gastric cancer, and *H. pylori* strains harboring the cag pathogenicity island further augment cancer risk [1]. The cag island encodes a bacterial type IV secretion system (T4SS), which translocates the bacterial protein CagA into host cells. Intracellular CagA aberrantly activates numerous signaling pathways that are mutated during gastric cancer, including β-catenin, resulting in cellular responses that lower the threshold for carcino-

ogenesis, such as hyperproliferation and pro-inflammatory cytokine production [2, 3]. Further, transgenic mice overexpressing CagA have been shown to develop gastric adenocarcinoma [4], thus validating this effector as an oncoprotein.

Iron deficiency is also associated with an increased risk for neoplasms that arise within the gastrointestinal tract [5], and *H. pylori* is linked with iron deficiency [6]. CagA facilitates *H. pylori* colonization by mediating bacterial iron acquisition [7], suggesting that iron deficiency may influence the virulence of this pathogen. We recently demonstrated that iron deficiency augments and accelerates the development of gastric carcinogenesis within the context of *H. pylori* infection and that this is mediated by increased assembly and function of the cag secretion system [8]. Whether this phenotype is due to a stable genetic modification or is reversible remains unknown, which likely influences therapeutic strategies. Therefore, the aim of this study was to define the dynamics of *H. pylori* virulence phenotypes that are augmented by iron deficiency.

**METHODS**

*H. pylori* Strains

Wild-type carcinogenic *H. pylori* strain 7.13 and its cagA- isogenic mutant were minimally passaged and used to infect Mongolian gerbils maintained on iron-replete or iron-depleted diets, as previously described [8]. Gerbils were euthanized 12 weeks after challenge, and in vivo–adapted strains were harvested from gerbils maintained on iron-depleted diets [8]. The Vanderbilt University Institutional Animal Care and Use Committee approved all procedures.

Parental strain 7.13 or in vivo–adapted strains harvested from 5 iron-depleted gerbils were serially passaged every 24–48 hours for 1, 5, 10, 15, 20, 25, and 30 passages. At each passage, strains were grown in *Brucella* broth supplemented with 10% fetal bovine serum (FBS) for 16 hours at 37°C with 5% CO₂ for coculture with gastric epithelial cells.

**Gastric Epithelial Cells**

Human AGS gastric epithelial cells were cocultured with *H. pylori* strains at a multiplicity of infection of 100:1 for 6 hours.

**Immunofluorescence and Confocal and Scanning Electron Microscopy**

Gastric tissue specimens from gerbils were processed for immunofluorescence as previously described [9]. Tissue samples were fixed in 2% paraformaldehyde and embedded in agar, and 100-µM sections were cut (Leica). Rabbit anti-*H. pylori*...
antibodies (Dako), Alexa Fluor 594 phalloidin (Invitrogen), and DAPI (Invitrogen) were used for visualization of *H. pylori*, actin, and nuclei, respectively. Samples were imaged with a Zeiss LSM 700 confocal microscope, and z-stacks were reconstructed using Volocity software (Improvision). For scanning electron microscopy, gastric tissue was fixed in paraformaldehyde phosphate buffer fixative and processed, as previously described [10].

**Transmission Electron Microscopy**

*H. pylori* were grown in *Brucella* broth supplemented with 10% FBS alone or with 100 µM FeCl₃ (iron-replete), 100 µM dipyridyl (iron restricted), or 100 µM dipyridyl plus 100 µM FeCl₃ (iron restricted with iron supplementation) for 16 hours at 37°C with 5% CO₂. *H. pylori* were harvested in 0.05-M sodium cacodylate buffer, spotted onto Formvar-coated grids, and negatively stained with 1% ammonium molybdate. Samples were viewed with a Philips C-12 transmission electron microscope, as previously described [10].

**Western Blot Analysis**

*H. pylori*:AGS coculture protein lysates were harvested, separated by sodium dodecyl sulfate polyacrylamide gel electrophoresis, and transferred to polyvinylidene fluoride (PVDF) membranes. Levels of total CagA (anti-CagA antibody; Austral Biologicals), phosphorylated CagA (anti-pY99 antibody; Santa Cruz) as a measure of translocated CagA, and β-actin (anti-β-actin antibody; Sigma) were determined. Protein intensities were quantified using ImageJ (National Institutes of Health) and standardized to levels of β-actin.

**Quantitative Reverse Transcription Polymerase Chain Reaction (RT-PCR)**

RNA was extracted (Qiagen) from *H. pylori*:AGS cocultures, and quantitative RT-PCR was performed according to the manufacturer’s instructions. IL-8 messenger RNA (mRNA) expression was normalized to levels of GAPDH mRNA expression (TaqMan; Applied Biosystems).

**Interleukin 8 (IL-8) Enzyme-Linked Immunosorbent Assay (ELISA)**

*H. pylori*:AGS coculture supernatants were collected, and levels of IL-8 were determined by the Quantikine IL-8 ELISA (R&D Systems) and analyzed with Gen5 software (Synergy4; BioTek).

**Statistical Analysis**

Mean values with standard errors are shown from experiments performed on at least 3 independent occasions. Mann–Whitney *U* tests and 1-way analysis of variance were used for comparisons. A *P* value of ≤ .05 was considered statistically significant.

**RESULTS**

We previously demonstrated that (1) iron deficiency augments gastric carcinogenesis within the context of *H. pylori* infection in Mongolian gerbils [8], and (2) loss of cagA significantly decreases *H. pylori* colonization under iron-deficient conditions but not normal iron conditions [7], suggesting that CagA provides a survival advantage in vivo that is revealed under conditions of iron deficiency. To define mechanisms that may account for these findings, we assessed the morphology of the cagA⁻ isogenic mutant in gastric tissue under iron-replete and iron-depleted conditions, using confocal microscopy (Figure 1A) and scanning electron microscopy (Figure 1B). Under iron-replete conditions, the cagA⁻ isogenic mutant exhibited an elongated and spiral morphology; however, iron depletion led to the development of coccoid phenotypes. This phenotype was not observed with wild-type *H. pylori*, which maintained its spiral morphology under both iron-replete and iron-depleted conditions (data not shown). To determine whether the transition to coccoid morphology under conditions of iron depletion could be replicated in vitro, we next assessed morphologic phenotypes of wild-type strain 7.13 and its cagA⁻ mutant under control, iron-replete, iron-restricted, or iron-restricted conditions with iron supplementation, by transmission electron microscopy (Figure 1C). Loss of cagA led to coccoid morphology but only under conditions of iron depletion, confirming our ex vivo results (Figure 1A and 1B). Of note, this phenotype was prevented by the addition of exogenous iron (Figure 1C).

*H. pylori* strains grown under iron-limiting conditions developed an increased capacity to assemble the cag type IV secretion system, which is accompanied by increased expression and translocation of CagA, and increased production of IL-8 by infected gastric epithelial cells [8]. Our data (Figure 1C) demonstrated that altered morphology of the cagA⁻ mutant under conditions of iron deficiency was reversible. Therefore, we next sought to determine whether augmented cagA-dependent virulence phenotypes were stable or reversible.

The parental strain 7.13 or in vivo–adapted strains harvested from gerbils maintained on iron-depleted diets were serially passaged in vitro and then cocultured with AGS gastric epithelial cells at each passage number. Compared with minimally passaged *H. pylori* isolates, long-term in vitro passage resulted in significantly decreased levels of CagA expression and translocation (Figure 2A and 2B). This occurred rapidly, after only 10 passages in vitro, indicating that heightened CagA expression and translocation is reversible in *H. pylori* strains that have been exposed to iron-deficient conditions.

Consistent with the pattern observed for CagA expression and translocation, low-passage in vivo–adapted strains isolated from gerbils maintained on iron-depleted diets induced significantly higher levels of IL-8 mRNA and protein expression in gastric epithelial cells (Figure 2C and 2D), when compared to the parental strain 7.13. However, following in vitro passage of these strains (>5 passages), increased levels of IL-8 mRNA and protein expression decreased to levels observed with the parental strain 7.13 (Figure 2C and 2D). Collectively, these data
Figure 1. Loss of cagA alters Helicobacter pylori morphology in an iron-dependent manner. Immunofluorescence and confocal microscopy (A) and scanning electron microscopy (B) were performed on gastric tissue specimens recovered from gerbils infected with the cagA− isogenic mutant under iron-replete or iron-depleted conditions. A, H. pylori (green), actin (red), and nuclei (blue). Bars = 10 µm. C, Transmission electron microscopy was performed on wild-type H. pylori strain 7.13 (WT) and the cagA− isogenic mutant (cagA−) under control, iron-replete, iron-restricted, or iron-restricted conditions with iron supplementation. Bars = 1 µm.

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demonstrate that the hypervirulent phenotypes stimulated by iron depletion are reversible.

**DISCUSSION**

In the setting of iron deficiency, *H. pylori* accelerates the development of inflammation, dysplasia, and adenocarcinoma in an animal model of gastric cancer that closely resembles human disease [8]. This phenotype is mediated by enhanced deployment of the *cag* T4SS, increased CagA translocation into host cells, and increased proinflammatory immune responses. In addition to inducing pathogenic responses, CagA also facilitates *H. pylori* colonization through iron acquisition, indicating that CagA provides a survival advantage in vivo under iron-limiting...
conditions [7]. Our current results demonstrate that loss of cagA under conditions of iron depletion led to a conversion from a spiral morphology to a coccoid morphology, a phenotype that is prevented by iron supplementation. H. pylori typically exists in a helical form but can convert to a coccoid morphology during environmental stress. The functional status exerted by coccoid H. pylori, however, remains controversial. Data have shown that this morphological manifestation leads to nonviable H. pylori [12], while other results have demonstrated that coccoid forms are viable and maintain cell structure and virulence expression profiles [13, 14]. Coccoid forms of H. pylori have also been implicated in relapses of infection after antimicrobial treatment [15]. Our data have provided fresh insights into these questions as we have demonstrated that CagA, an oncoprotein, is required for normal growth and spiral morphology under iron-restricted conditions. Further, the ability of CagA to regulate bacterial fitness in the absence of host cell contact is a novel finding, thus providing new insights into how pathogenic factors may regulate microbial homeostasis.

In addition to defining the dynamics of bacterial morphology under changing conditions, we have now also demonstrated that H. pylori hypervirulent phenotypes mediated by iron deficiency are reversible. One potential explanation is that in vivo, there is positive selection for enhanced T4SS activity, whereas in vitro growth results in negative selection for such activity. However, we have previously demonstrated that when hypervirulent strains isolated from iron-depleted gerbils are reintroduced into iron-replete gerbils, they no longer induce increased frequency of disease [10], indicating that this is not simply due to negative selection in vitro.

Global and indiscriminate test-and-treat strategies for H. pylori have not been fully embraced, primarily because of the relatively low incidence of cancer among infected individuals and studies reporting that carriage of H. pylori is inversely related to the development of esophageal and atopic diseases [1]. These data underscore the importance of carefully identifying colonized persons who are highly predisposed to pathologic outcomes. Our earlier studies provided an important initial step in this direction by indicating that iron deficiency increases the virulence of H. pylori, thereby identifying a high-risk group of persons who may warrant therapy directed against H. pylori. Our new findings have extended these observations and indicate that hypervirulent H. pylori phenotypes stimulated by iron deficiency are reversible. Because many studies have reported that the efficacy of first-line anti–H. pylori therapy is around 80%–85%, a clinical strategy that focuses on treatment of iron deficiency among H. pylori–colonized individuals who do not respond therapy may ultimately attenuate disease. Collectively, these studies should enable clinicians to more appropriately focus diagnostic testing and eradication therapies on high-risk populations to effectively prevent the development of gastric cancer.

Notes

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