Dissemination of cagA and cagE Virulence Genes Among H. Pylori Strains From Sudanese Patients With Gastric Discomfort

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Research note

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Abstract

Objectives

*Helicobacter pylori* cytotoxin-associated gene pathogenicity island (*cag*-PAI) is one of the strain-specific genes (they do not exist in all strains). *cag*-PAI is involved in inducing inflammation, ulceration, and carcinogenesis. This study aimed to detect and characterize *cag*A and *cag*E virulence genes among *H. pylori* strains from Sudanese patients with gastric discomfort.

Result

Out of 288 gastric biopsies screened for the presence of *H. pylori*, 34% (98/288) were positive, *cag*A gene was present in 41% (40/98) of specimens, mostly in patients with gastritis 72.5% (29/40), followed by duodenal ulcer 15% (6/40), esophagitis 5% (2/40), and 7.5% (3/40) in patients diagnosed normal by endoscopy. *cag*E gene was present in 39% (38/98) of specimens, the majority 73.7% (28/38) were from patients with gastritis, 10.5% (4/38) duodenal ulcer, 5.3% (2/38) esophagitis, 2.6% (1/38) gastric ulcer, and 7.9% (3/38) were diagnosed as normal. The *cag*A and *cag*E protein sequences have synonymous amino acid variations.

Introduction

*Helicobacter pylori* (*H. pylori*) is a gram-negative bacterium, grows in a microaerophilic environment at 37°C. It is one of the causative agents of gastric diseases [1], like chronic gastritis, ulcers (duodenal or gastric), gastric adenocarcinoma, and gastric mucosa-associated lymphoid tissue (MALT) lymphoma [2]. Previous studies showed that soil, water, animal manure, and human stool are the primary resources for *H. pylori* [3]. There is an opportunity for transmission from person to person [4]. Some factors affect the incidence of *H. pylori* infection; race, gender, social and socioeconomic status of the population [5].

The pathogenicity of *H. pylori* infection is mediated by environmental, bacterial virulence factors, and internal interaction with the host [6].

The virulence genes of *H. pylori* are classified into 3 categories; strain-specific genes, phase variable genes, and genes with variable genotypes or structures depending on the strain [7]. Cytotoxin-associated gene pathogenicity island (*cag*-PAI) and plasticity island genes are strain-specific genes, and they do not exist in all strains [7].

The *cag*-PAI is a DNA fragment that contains 31 genes, including; *cag*A and *cag*E in addition to other genes [8]. *cag*-PAI genes involved in inducing inflammation, ulceration, and carcinogenesis [3]. The presence of *cag*-PAI is strongly associated with the outcomes of *H. pylori* infection [9].

The *cag*A protein behaves as a bacterial oncoprotein, affects the expression of vital proteins or its function in oncogenic or tumor suppressor signaling pathways [10].
The \textit{cagE} gene encodes a secretory protein that induces secretion of IL-8 from infected epithelial cells; also this gene is important for translocation and phosphorylation of the \textit{cagA} protein, some studies have suggested that it is a more accurate marker of an intact pathogenicity island and can be used with \textit{cagA} as a marker for \textit{cag-PAI} [11, 12, 13].

To date there are limited published studies about the detection and characterization of \textit{cagA} and \textit{cagE} genes of \textit{H. pylori} in Sudan, this information is of great importance in determining the virulence and clinical outcomes of \textit{H. pylori} strains. Thus, this study was conducted to detect and characterize \textit{cagA} and \textit{cagE} genes of \textit{H. pylori} strains from gastric biopsies of patients with gastric discomfort in Khartoum State.

**Materials And Methods**

This was a descriptive cross-sectional study conducted during the period from August 2018 to October 2019. The sampling technique was convenience (Non-probability).

**Collection of clinical specimens**

A total of 288 gastric biopsies were collected by gastric endoscope by the gastroenterologist in different Khartoum hospitals: Omdurman Medical Military Hospital, Police Hospital, Al-Buqa'a Specialized Hospital, Ibn Sina Hospital, and Fedail Hospital. The specimens were collected from patients suffering from gastric discomfort and indicated for endoscopy, in which both antrum and corpus were sampled. The specimens were collected using endoscopes, using 1.0 ml brain heart infusion broth with 15\% glycerol as transport media.

**DNA extraction and polymerase chain reaction (PCR)**

DNA was extracted using the guanidine chloride extraction method [14]. The extracted DNA was stored at -70\°C until use.

The PCR was performed using a thermocycler (Convergys, Germany) and specific primer (Table 1) according to the following protocols:

For \textit{glmM} gene: initial activation at 94\°C for 3 minutes, followed by 35 cycles at 94\°C for 30 seconds, 58\°C for 30 seconds, and 72\°C for 30 seconds, then a final extension at 72\°C for 3 minutes [15].

For 16S rRNA gene: initial activation at 94\°C for 3 minutes, followed by 35 cycles at 94\°C for 30 seconds, 53\°C for 30 seconds, and 72\°C for 45 seconds, then a final extension at 72\°C for 5 minutes [16].

For \textit{cagA} gene: initial activation at 94\°C for 3 minutes, followed by 35 cycles at 94\°C for 30 seconds, 53\°C for 30 seconds, and 72\°C for 45 seconds, then a final extension at 72\°C for 5 minutes [15].

For \textit{cagE} genes: initial activation at 94\°C for 3 minutes, followed by 35 cycles at 94\°C for 30 seconds, 53\°C for 30 seconds, and 72\°C for 45 seconds, then a final extension at 72\°C for 5 minutes [17].
Table (1): Sequences of primers used for the detection of *H. pylori* 16S rRNA, *glmM*, *cagA*, and *cagE* genes.

| Gene   | Primer sequence (from 5’-3’) | Size of PCR product (base pairs) | References |
|--------|-----------------------------|---------------------------------|------------|
| 16S rRNA | F: GCTAAGAGATCAGCCTATGTCC R: TGGCAATCAGCGTCAGGTAAT G | 532bp | [16] |
| *glmM* | F: GGATAAGCTTTTAGGGGTGTTAGGGG R: GCTTACTTTCTAACAACAAGCGC | 294bp | [15] |
| *cagA* | F: ATAATGCTAATTAGACAACCTTGAGCGA R: AGAAACAAAAGCAATACGCATTC | 128bp | [15] |
| *cagE* | F: TTGAAAACCTTCAAGGATAGGATAGAGC R: GCCTAGCGTAATATCACCATTACC | 508bp | [17] |

**DNA sequencing**

A total of 21 *cagE* and 5 *cagA* amplified gene products were sequenced by BGI Company in China. The BLAST program was used to align the nucleotide sequences with the reference sequences from GenBank. The nucleotide sequences of *cagA* and *cagE* genes were translated into their corresponding amino acid using ExPASY available online at [https://web.expasy.org/translate/](https://web.expasy.org/translate/). BioEdit software was used to compare the translated amino acid sequences with four reference sequences (JF798705.1 and MK074991.1 for *cagA*, and AB191082.1 and AY153111.1 for *cagE*).

The *cagE* nucleotide sequences have been deposited in Genbank. The *cagA* sequences were shorter than 200bp so it was not accepted in Genbank.

**Statistical Analysis**

Data were analyzed using the Statistical Package for Social Science (SPSSversion11.5) computer program. Frequencies were calculated, the Chi-square test was used to test the association between variables, *p*-value ≤0.05 was considered significant.

**Results**

**Gender and age groups**
Out of 288 patients, there were 160 (56%) males and 128 (44%) females; they have been divided into two age groups; 10 adolescents (2 males and 4 females) and 278 adults (158 males and 124 females).

**Clinical diagnosis**

Based on the gastroscopy findings; 188 (65.3%) of the patients had gastritis, 25 (8.7%) had a duodenal ulcer, 29 (10.1%) had a gastric ulcer, 15 (5.2%) had esophagitis, 1 (0.3%) had MALTL, 1 (0.3%) had duodenitis, and 29 (10.1%) had normal gastric mucosa.

**Genotyping of *H. pylori***

Out of 288 samples, 98 (34%) were positive for *H. pylori* (*glmM* and/or 16S rRNA positive), 88 (31%) were 16S rRNA gene-positive and 42 (15%) were *glmM* gene-positive, 32 (33%) were positive for both genes (Figure 1). 50/98 (51%) of the *H. pylori*-positive patients were females and 48/98 (49%) were males. The majority of the *H. pylori*-positive patients were adults 95/98 (97%), and 3/98 (3%) were adolescents.

**Cytotoxin associated genes (*cagA* and *cagE*)**

*cagA* gene was present in 40/98 (41%) specimens, their clinical diagnosis as follows: 29 (72.5%) gastritis, 6 (15%) duodenal ulcer, 2 (5%) esophagitis, and 3 (7.5%) were diagnosed as normal by endoscopy. *cagE* gene was present in 38/98 (39%) of the specimens, their clinical diagnosis as follows: 28 (73.7%) gastritis, 1 (2.6%) gastric ulcer, 4 (10.5%) duodenal ulcer, 2 (5.3%) esophagitis and 3 (7.9%) were diagnosed as normal (Figure 2).

From *H. pylori*-positive specimens (98); 27/98 (28%) were positive for both *cagA* and *cagE* genes, 13 specimens (13%) were positive for *cagA* gene only, 11/98 (11%) were positive for *cagE* gene only, 47/98 (48%) were negative for both genes.

There was no statistically significant association between the presence of *cagA* and *cagE* virulence genes and clinical diagnosis (*p*-value= 0.305).

**DNA sequencing**

The comparison of 5 *cagA* protein sequences with the JF798705.1 reference from GenBank revealed substitution of proline (P) with alanine (A), with 100% identity to MK074991.1 reference sequence (Figure 3).

The comparison of 21 *cagE* protein sequences with the AY153111.1 reference sequence from GenBank, revealed a substitution mutation in one sequence, the sequence has valine (V) instead of isoleucine (I), with 100% identity to AB191082.1 reference sequence (Figure S1 and Figure S2).

**Discussion**
In the present study, out of 288 gastric biopsies, 98 (34%) were positive for *H. pylori*, using PCR targeting 16S rRNA and *glmM* genes. These genes are more sensitive than other genes in detecting *H. pylori* infection from gastric biopsies [18]. So, we have used them for the detection of *H. pylori* in biopsy samples.

*H. pylori* infection prevalence was 79.1% in Africa, 54.7% in Asia, and 37.1% in Northern America [19]. The geographic variation in the prevalence may be due to the uneven distribution of *H. pylori* in the stomach in the different clinical settings, gender, race, social and socioeconomic status of the population [5, 20].

In Sudan, there is scanty information about the prevalence of *H. pylori* infection [14]. The prevalence in this study may indicate a low rate of *H. pylori* infection. Most of the previous studies investigating *H. pylori* seroprevalence using tests for the detection of *H. pylori* IgM and IgG antibodies. They reported a seroprevalence ranging from 20% up to 70% [21]. While PCR was used in this study which has significantly higher accuracy than stool antigen test and antibody-based tests used in the mentioned Sudanese studies.

In this study 50/98 (51%) of the infected patients were females, this may attributed to the possibility of transmission from animal and their products, as women may deal with foods with animal origins when preparing food. This result in contrasts with other reports, noting a high existence of infection among males [22]. The majority of the infected patients were adults 96/98 (98%), the infected adolescents were 2/98 (2%), this result is in harmony with El-Shenawy [23] and Salih [24] studies, they mentioned that; the prevalence of *H. pylori* infection increases with age. 

*cagA* in this study was detected in 41% (40/98) of specimens, with high frequency in patients with gastritis (72.5%). The *cagA* prevalence in the present study is similar to that obtained in Nigeria (40%), Cyprus (42.5%), Bahrain and Jordan (25–60%), and Iran (44–94%), and it is lower than results obtained from other countries including; Egypt (65.2%), Palestine (65.9%), Tunisia (61.6%), Russia (85%), and India (60% to 80%) [15, 27]. These data indicate that various parts of the world have different *cagA* prevalence. 

*cagE* in this study was present in 39% (38/98) of specimens, with high frequency in patients with gastritis (73.7%). The presence of this gene was also varied geographically; in Turkey (28%), Brazil (53.2%), Malaysia (70%), Thailand (88%), India (77%), the United States of America (64%), and the United Kingdom (71%) [27].

In the present study, the *cagA* gene was more frequent than *cagE*, and this result in agreement with Lima [28] study in which 28% of specimens were positive for both genes (*cagA* and *cagE*), which might be attributed to the importance of *cagE* for *cagA* translocation, as mention in Yong [10] study.

There was no statistically significant association between the presence of these virulence genes and the clinical outcomes in this study. This result is similar to Moaddeb [29] study, which found that; there is no significant association between *cagA*-positive strains and gastroduodenal diseases. In contrast with
Yong [10] and Skoog [9] studies, they mentioned that the presence of cag-PAI is strongly associated with the outcomes of *H. pylori* infection.

The differences in *H. pylori* virulence may be attributed to geographic variations in the strain-specific (non-conserved) genes and reduced virulence of African *H. pylori* strains, as in Bullock [30] study, who also mentioned that cagA proteins produced by East Asian strains cause more extensive alterations in gastric epithelial cells than that produced by strains from other parts of the world.

The analysis of 5 cagA protein sequences in this study revealed a substitution mutation in two cagA protein sequences. This variation may be due to the high level of geographic variability in *H. pylori* protein sequences, which may lead to differences in *H. pylori* virulence. The comparison of 21 cagE protein sequences in this study revealed a substitution mutation in one cagE protein sequence. This variation was synonymous according to Hamad's [27] study. Bullock [30] study mentioned that there is a high level of geographic variation in sequences of *H. pylori* proteins.

**Conclusion**

*H. pylori* low prevalence in this study (34%) indicates that Sudan is in the low-risk region for infection. There is high dissemination of cagA and cagE genes in *H. pylori* from Sudanese patients. Still, they may have reduced virulence. The cagA and cagE protein sequences have synonymous amino acid variations.

**Limitations**

- DNA sequencing is not done to all specimens.
- Using of short cagA primers (128bp) for DNA sequencing.

**Abbreviations**

GC: gastric cancer.

cag-PAI: cytotoxin-associated gene pathogenicity island.

cagA: cytotoxin associated gene A.

cagE: cytotoxin associated gene E.

T4SS: type IV secretion system.

SPSS: Statistical Package for Social Science.

**Declarations**

Ethical approval and consent to participate
The research was approved by the Khartoum state Ministry of health research department on 1/3/2018. Written consent was obtained from patients.

Availability of data and materials

The cagE DNA sequences data were uploaded to the NCBI database under accession numbers: (MN418976, MN418977, MN418978, MN418979, MN418980, MN418981, MN418982, MN418983, MN418984, MN418985, MN418986, MN418987, MN418988, MN418989, MN418990, MN418991, MN418992, MN418993, MN418994, MN418995, and MN418996). Other data are available at: http://doi.org/10.6084/m9.figshare.13620422.

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Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

MMM, HNA, EHO, AMA, and LOM designed the study. EHO, MMM, and AMA performed the experience. EHO, HNA, and LOM analyzed the data. EHO wrote the manuscript. All authors read and approved the final manuscript.

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**Figures**

![Figure A](image1)

![Figure B](image2)

**Figure 1**
Amplified DNA of the glmM gene (A) (Lanes 7, 8& 9 contain gene does not included in this manuscript) and 16S rRNA gene (B) on 1.5% agarose gel electrophoresis.

Figure 2
Amplified DNA of the cagA gene (A) and cagE gene (B) on 1.5% agarose gel electrophoresis.
Figure 3

Alignment of cagA protein sequences with amino acid sequences of JF798705.1 reference sequence (A) and MK074991.1 reference sequence (B).

Supplementary Files

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