

RESEARCH ARTICLE

Open Access

Matrix metalloproteinase-3 promoter polymorphisms but not *dupA*-*H. pylori* correlate to duodenal ulcers in *H. pylori*-infected females

Yi-Chun Yeh¹, Hsiu-Chi Cheng^{2,3}, Wei-Lun Chang^{2,3}, Hsiao-Bai Yang^{4,5}, Bor-Shyang Sheu^{2,3*}

Abstract

Background: This study investigated if the *H. pylori dupA* genotype and certain host single nucleotide polymorphisms (SNPs) of matrix metalloproteinases (MMPs) and their inhibitors (TIMPs), including MMP-3, MMP-7, MMP-9, TIMP-1 and TIMP-2, might correlate with ulcer risk of *H. pylori*-infected Taiwanese patients.

Results: Of the 549 *H. pylori*-infected patients enrolled, 470 patients (265 with gastritis, 118 with duodenal ulcer, and 87 with gastric ulcer) received SNPs analysis of MMP-3₋₁₆₁₂ 6A > 5A, MMP-7₋₁₈₁ A > G, MMP-9_{exon 6} A > G, TIMP-1₃₇₂ T > C and TIMP-2₄₁₈ G > C by PCR-RFLP. The 181 collected *H. pylori* isolates were detected for the *dupA* genotype by PCR. The rates of *dupA*-positive *H. pylori* infection were similar among patients with duodenal ulcer (22.8%), gastric ulcer (20.0%), and gastritis (25.5%) ($p > 0.05$). Males had higher rates of duodenal ulcer and gastric ulcer than females ($p < 0.01$). Of *H. pylori*-infected patients, the MMP-3 6A6A genotype were more common in patients with duodenal ulcers than in those with gastritis (87.7% vs. 74.9%, $p < 0.05$) in females. This genotype had a 2.4-fold (95% CI: 1.02-5.66) increased risk of duodenal ulcer, compared to those with the 5A carrier. Combining the MMP-3/TIMP-1 genotype as 6A6A/CC, the risk of duodenal ulcer increased up to 3.6 fold ($p < 0.05$) in *H. pylori*-infected females.

Conclusions: The MMP-3 promoter polymorphism, but not the *dupA*-status, may correlate with susceptibility to duodenal ulcer after *H. pylori* infection in Taiwanese females.

Background

Helicobacter pylori infection leads to chronic gastritis and in some individuals, to peptic ulcer disease or even gastric carcinoma [1]. Diverse outcomes may depend on complex interactions among bacterial virulence factors, host genetics, and environmental factors [2,3]. In Taiwan, despite the nearly 100% prevalence of the so-called triple-genopositive *cagA-vacA-babA2* virulent *H. pylori* infections, there is a lack of correlation to different disease outcomes [4,5]. It will be useful for Taiwan to validate new virulence factors or any host genomic predisposition in relation to severe *H. pylori*-infected clinical outcomes.

Recently, a duodenal ulcer-promoting gene A (*dupA*) encompassing *jhp0917* and *jhp0918* has been suggested to lead into higher IL-8 production of epithelial cells

and thus, triggering dense neutrophil infiltration and increased risk of duodenal ulcers [6]. However, even in such large-scale validation, those with duodenal ulcer have a nearly 55% *dupA*-positive infection [6]. Moreover, prevalence of *dupA* and relationships between *dupA*-positive *H. pylori* and clinical outcomes are different in distinct populations [7-11]. It may indicate that *dupA* serves a promoting role leading to duodenal ulcer after *H. pylori* infection. Alternatively, it is necessary to validate host factors that predispose patients to gastro-duodenal ulcer, especially with *dupA*-negative infection.

H. pylori infection stimulates the production of pro-inflammatory cytokines, such as IL-1, which play important roles in gastric inflammation and physiology. However, IL-1 beta or IL-1RN polymorphisms are not associated with gastric ulcer in the Taiwanese population [12].

Matrix metalloproteinases (MMPs) are a family of enzymes that degrade most extracellular matrix and

* Correspondence: sheubs@mail.ncku.edu.tw

²Institute of Clinical Medicine, National Cheng Kung University Hospital, Medical College, Tainan

Full list of author information is available at the end of the article

correlate with ulcer formation or repairs [13]. *H. pylori* infection can up-regulate MMP-3, MMP-7, and MMP-9 in the gastric mucosa and even sera [14-16]. A large-scale German survey has further validated that the single-nucleotide polymorphisms (SNP) genotype as MMP-7₋₁₈₁ G allele and MMP-9_{exon 6} A allele increase the risk of gastric ulcer after *H. pylori* infection [17]. A deletion at MMP-3 promoter -1612, and A to G substitution at MMP-7 promoter -181 may affect transcriptional activity, leading to alterations in gene expression [18,19]. Moreover, A to G substitution at MMP-9 exon 6 causes the amino acid change required for binding to its substrate and affects its binding ability [20].

Although MMP activity is in general counteracted by endogenous tissue inhibitors (TIMPs) [21], there remains no data to check whether TIMP-1 and TIMP-2 SNP genotypes relate to the risk of gastroduodenal ulcer after *H. pylori*-infection. As such, this study surveyed if the *H. pylori dupA* genotype and certain SNP genotypes of MMP-3, MMP-7, MMP-9, TIMP-1, and TIMP-2 predispose *H. pylori*-infected Taiwanese patients to ulcer risks.

Methods

Patients and study design

Five hundred and forty-nine consecutive *H. pylori*-infected patients documented by upper gastrointestinal endoscopy at National Cheng Kung University Medical Center, Tainan, Taiwan were enrolled. All were genetically unrelated ethnic Han Chinese from Tainan City and the surrounding regions. None had been treated with NSAIDs, proton pump inhibitor, or any antibiotics within two weeks prior to panendoscopy on enrollment, or a past history of anti-*H. pylori* treatment and peptic ulcer. The hospital Ethics Committee approved the study.

After obtaining informed consent, 470 patients had provided enough blood samplings for SNPs analysis of MMP-3₋₁₆₁₂ 6A > 5A, MMP-7₋₁₈₁ A > G, MMP-9_{exon 6} A > G, TIMP-1₃₇₂ T > C and TIMP-2₄₁₈ G > C by PCR-RFLP. Aside from endoscopic diagnosis for clinical diseases, at least three topographic gastric biopsies were sampled for histology or *H. pylori* culture, one each from the antrum, corpus, and cardia. These were stained with haematoxylin and eosin and reviewed for the *H. pylori*-related histology by the updated Sydney's system [4,22,23].

In addition, the study collected 181 *H. pylori* isolates for the detection of *dupA* genotype by PCR. One hundred and three isolates were collected from randomly selected patients who had agreed to undergo SNP analysis, while 78 isolates were from patients without SNP analysis. The *H. pylori* culture were conducted from the two additional gastric biopsies collected during the same endoscopy and processed with the method applied in previous publications [4,22].

For those with positive *H. pylori* culture, the isolates were extracted for genomic DNA to be analyzed for the *dupA* genotypes by PCR. The extraction of DNA was done with the same method as described previously [4,22]. Positive *H. pylori* infection was defined by positive histology or culture.

Genotypes of SNPs in MMPs and TIMPs

Peripheral blood 8 ml was obtained from each subject for genomic DNA, which was extracted from peripheral blood mononuclear cells according to the manufacturer's instructions (Viogene, Taipei, Taiwan). Five SNPs in MMP-3₋₁₆₁₂ 5A/6A, MMP-7₋₁₈₁ A/G, MMP-9_{exon 6} A/G, TIMP-1₃₇₂ C/T, and TIMP-2₄₁₈ G/C polymorphisms were determined by PCR-RFLP assays [18,24-26].

Using the extracted DNA as template, the regions of each MMP and TIMP were amplified by PCR using commercially available kits (GoTaq[®] Green Master Mix, Promega, Madison, WI, USA) following the manufacturer's instructions. The sequences of primers, PCR conditions, and restriction enzymes (obtained from New England Biosciences, U.S.) used were summarized in Table 1. After digestion, the products were separated by electrophoresis on a 4% agarose gel. The MMP and TIMP genotypes were shown as different gel examples (Figure 1).

Quality control for genotyping was achieved by including in each amplification a negative PCR control sample and three positive control samples for each SNP analyzed (homozygous for allele 1, heterozygous, and homozygous for allele 2). At least 10% of the samples were run twice in separate assays to reveal 100% concordance of the genotype designation for all of the polymorphisms. For the positive controls, the genopositive products were confirmed by direct sequencing.

Detection of *dupA* gene by PCR

The *dupA*-positive *H. pylori* was determined by positive PCR amplifications of at least 2 regions (*jhp0917* and *jhp0918*) of the gene using two specific primer pairs (Figure 2) for strains J99 and 26695 as templates (Table 1) [6]. DNA 2 µl were added to 50-µl reaction mixture, containing 1 × PCR buffer, 1.5 mM MgCl₂, 0.2 mM dNTP (Protech, Taiwan), 0.2 µM primers and 1 U Taq DNA polymerase (Fermentas, USA). PCR was performed with a thermal cycler (2720 thermal cycler; Applied Biosystems). The mixture was stored at 4°C and the PCR products were separated by electrophoresis on 2% agarose gel. The gels were stained with ethidium bromide and visualized under UV illumination.

Statistical analysis

Statistical analysis was performed with the SPSS software (SPSS 12, Chicago, IL, USA). The χ^2 test, Fisher's exact test or Mann-Whitney U test were used as appropriate to

Table 1 The PCR primers used in the study

SNP/gene	Primer sequence (5' ♦ 3')	Size (bp)	Restriction enzyme	Reference
MMP-3 ₋₁₆₁₂ 5A/6A	GATTACAGACATGGGTCACG TTTCAATCAGGACAAGACGAAGTTT	120	<i>Xmn</i> I 6A: 120 bp 5A: 97 bp + 23 bp	Shibata et al, 2005
MMP-7 ₋₁₈₁ A/G	TGGTACCATAATGTCTGAAT TCGTTATTGGCAGGAAGCACACAATGAATT	150	<i>EcoR</i> I A: 150 bp G: 120 bp + 30 bp	Jormsjö et al, 2001
MMP-9 _{exon6} A/G	CCATCCATGGGTCAAAGAAC GGGCTGAACCTGGTAGACAG	295	<i>Sma</i> I A: 295 bp G: 192 bp + 103 bp	Shibata et al, 2005 *
TIMP-1 ₃₇₂ C/T	GCACATCACTACCTGCAGTC GAAACAAGCCCACGATTTAG	175	<i>BssS</i> I T: 175 bp C: 152 bp + 23 bp	Wollmer et al, 2002
TIMP-2 ₋₄₁₈ G/C	CGTCTCTTGTGGCTGGTCA CCTTCAGCTCGACTCTGGAG	304	<i>BsoB</i> I C: 253 bp + 51 bp G: 230 bp + 51 bp + 23 bp	Zhou et al, 2004
jhp0917_1	TGGTTTCTACTGACAGAGCGC	307	-	Lu et al., 2005
jhp0917_2	AACACGCTGACAGGACAATCTCCC			
jhp0917_3	GCCTAAGACCTCAAAGTAGC	296	-	This study *
jhp0917_4	CATTCTGTCAAGAGCTACC			
jhp0918_1	CCTATATCGCTAACGCGCGCTC	276	-	Lu et al., 2005
jhp0918_2	AAGCTGAAGCGTTTGTAAACG			
jhp0918_3	GCTAGAAAGATCAACGGAAC	216	-	This study *
jhp0918_4	CACTTGTCTGGCTCTCAT			

*The primers were self-designed by applying Primer 3 and the restriction enzyme was selected based on the reference of Shibata et al, 2005.

validate the *dupA* prevalence rates, histological severity, or SNP genotypes of MMPs and TIMPs between patients with and those without ulcers. A $p < 0.05$ was taken as significant. All test significances were validated by two-tailed analysis. The odds ratios (OR) and 95% confidence intervals (CI) were used to estimate the risk to get gastro-duodenal ulcer in these *H. pylori*-infected subjects.

Results

Demographic features of the study subjects

Of the 470 *H. pylori*-positive dyspeptic patients who provided blood samples for SNP analysis, 276 were females and 194 males, with median age of 47.2 years (range, 16-87 years). Endoscopic diagnoses included 265 with gastritis, 118 with duodenal ulcers, and 87 with gastric ulcers. Their demographics and histology after *H. pylori* infection were listed in Table 2. Patients with gastritis and duodenal ulcer had higher gastric inflammation in the antrum than those with gastric ulcers ($p < 0.05$). Among *H. pylori*-infected patients, males had higher rates of duodenal and gastric ulcers than females (51.7% vs. 30.9% and 58.6% vs. 30.9%, $p < 0.001$, respectively).

Prevalence of *dupA* *H. pylori* infection in patients

One hundred and eighty-one *H. pylori* strains were successfully obtained (Figure 3). The concordance of two

PCR primer pairs was 95.3% (41/43). Only two isolates were *dupA*-positive by single primer pairs. Forty-three isolates (23.8%) were genopositive for *dupA*, of which six (20.0%) were from patients with gastric ulcer, 13 (22.8%) from patients with duodenal ulcer, and 24 (25.5%) from gastritis patients. The prevalence rates of *dupA*-positive *H. pylori* infection were similar between patients with and those without ulcers ($p > 0.05$).

MMP and TIMP genotypes and the *H. pylori*-related gastro-duodenal ulcer

The 470 *H. pylori*-infected patients with SNP analysis had > 99% average genotyping success and the distributions of all SNPs were in Hardy-Weinberg equilibrium ($p > 0.05$). Since the ulcer rate had gender differences (Table 2), five genotype distributions were analyzed and separated by gender. There were no significant differences in genotype distributions of MMP-7₋₁₈₁ A/G, MMP-9_{exon 6} A/G, TIMP-1₃₇₂ T/C and TIMP-2₋₄₁₈ G/C between patients with different clinical diagnoses ($p > 0.05$) (Table 3).

There was a higher rate of MMP-3 6A6A genotype in patients with duodenal ulcers than in patients with gastritis (87.7% vs. 74.9%, $p < 0.05$). *H. pylori*-infected subjects with the MMP-3 6A6A genotype had a 2.4-fold

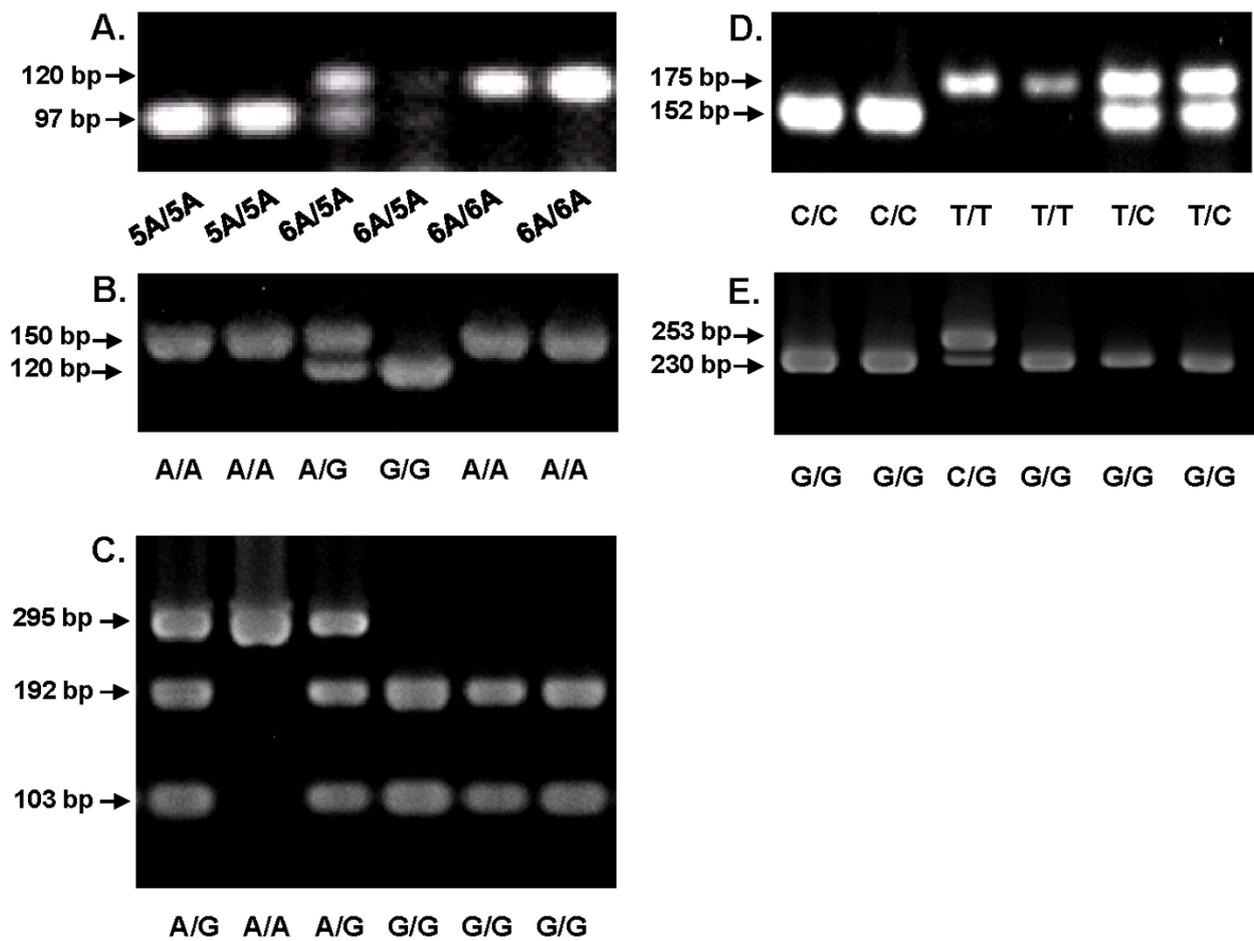


Figure 1 MMP and TIMP genotyping were done by PCR-RFLP and visualized by electrophoresis on 4% agarose gel. The listed genotype patterns were (A) for MMP-3 -1612 as 5A5A or 6A6A; (B) for MMP-7 -181 as AA, AG, or GG; (C) for MMP-9 exon 6 as AA, AG, or GG; (D) for TIMP-1 372; as CC, TC, or TT; and (E) for TIMP-2 -418 as CG or GG.

(95% CI: 1.02-5.66) increased risk of duodenal ulcer in females compared to those with the 5A carrier. Because TIMP-1 genotypes modulated MMP-3 activity, it was further tested whether the MMP-3₋₁₆₁₂/TIMP-1₃₇₂ Combined genotypes contributed to increased risk of duodenal ulcers in females. The combined MMP-3/TIMP-1 genotype as 6A6A/CC had a 3.6-fold ($p < 0.05$) increased risk of duodenal ulcer in *H. pylori*-infected female (Table 4).

Discussion

This study surveyed whether the bacterial factor *dupA* in *H. pylori* or single nucleotide polymorphisms of MMPs and TIMPs correlated with the susceptibility of gastroduodenal ulcers after *H. pylori* infection. It shows a rather low prevalence (23.8%) of *dupA*-positive *H. pylori* infection in Taiwan. Moreover, such a low prevalence limits its association to susceptibility to gastroduodenal ulcers after *H. pylori* infection.

The negative finding is consistent with several studies worldwide [8,10,11] but differs from Lu et al., who support the promoting role of *H. pylori dupA* [6]. In their positive report [6], *dupA*-positive prevalence is nearly 50% and it can be attributed as the promoting role of ulceration. Their data suggest that there should be some other bacterial virulence factor of *H. pylori* as *CagA*, *baba2*, *vacA*, or host factors, which determine the susceptibility of ulceration. In Taiwan, there is nearly a 100% prevalence of *CagA*, *baba2*, *vacA* triple-positive infection [4,15]. The current study area should be a good place to validate the host factor predisposing to ulcer risk.

In the *in vitro* promoter functional assay of fibroblasts and vascular smooth muscle cells, the MMP-3 -1612 as 5A allele has greater promoter activities than the 6A allele [19]. This implies that patients carrying the lower promoter activity genotype 6A6A in the MMP-3 promoter are accompanied by lower MMP-3 expressions of the

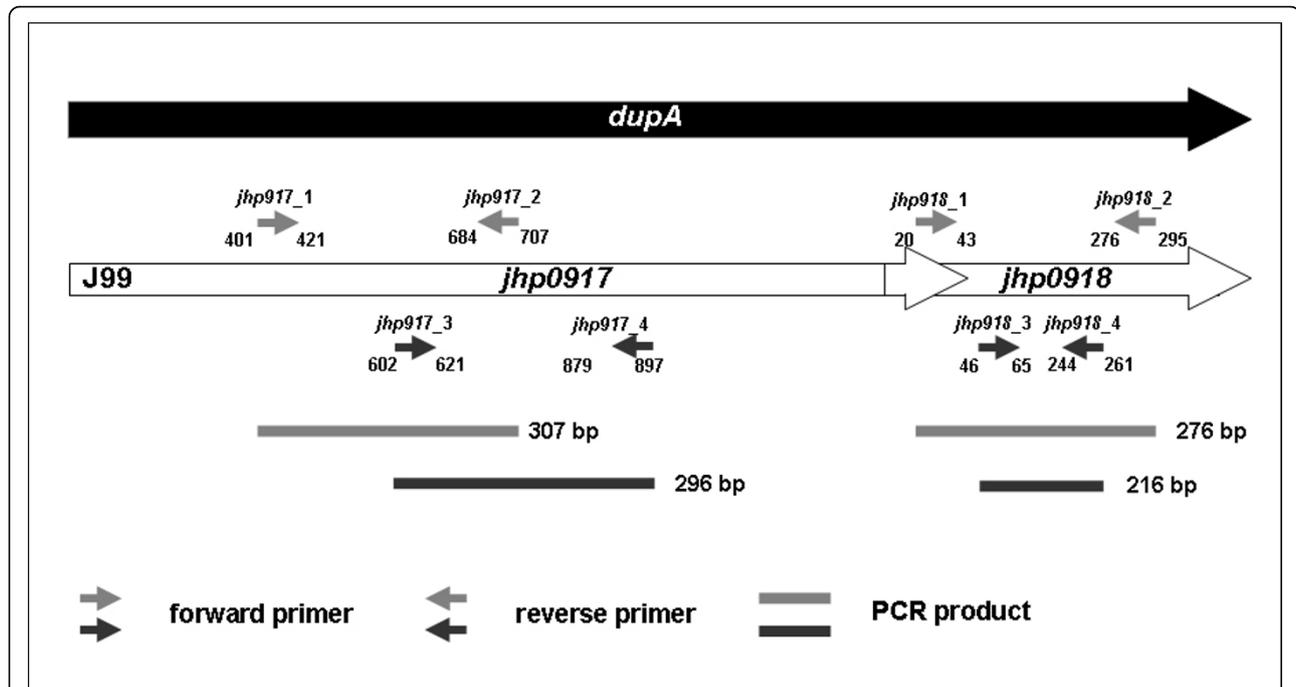


Figure 2 The sites of primer pairs for PCR on the *dupA* regions, including for the *jhp0917* and the *jhp0918* regions, applying the standard *H. pylori* isolates (J99) as reference.

gastric mucosa. This study discloses the host genotype MMP-3 -1612 as 6A6A, which expresses lower MMP-3 carries a 2.4-fold risk of having duodenal ulcers among females after *H. pylori* infection ($p < 0.05$) (Table 3). Moreover, TIMP-1 372, as CC, contributes a higher risk of duodenal ulcers to MMP-3 -1612 6A6A (Table 4). This data suggests that patients with higher MMP-3 expression may have lower ulcer risk, but the reasons remain uncertain.

In general, MMP-3 can degrade a wide range of substrates, including fibronectin, type IV, V, IX, and X

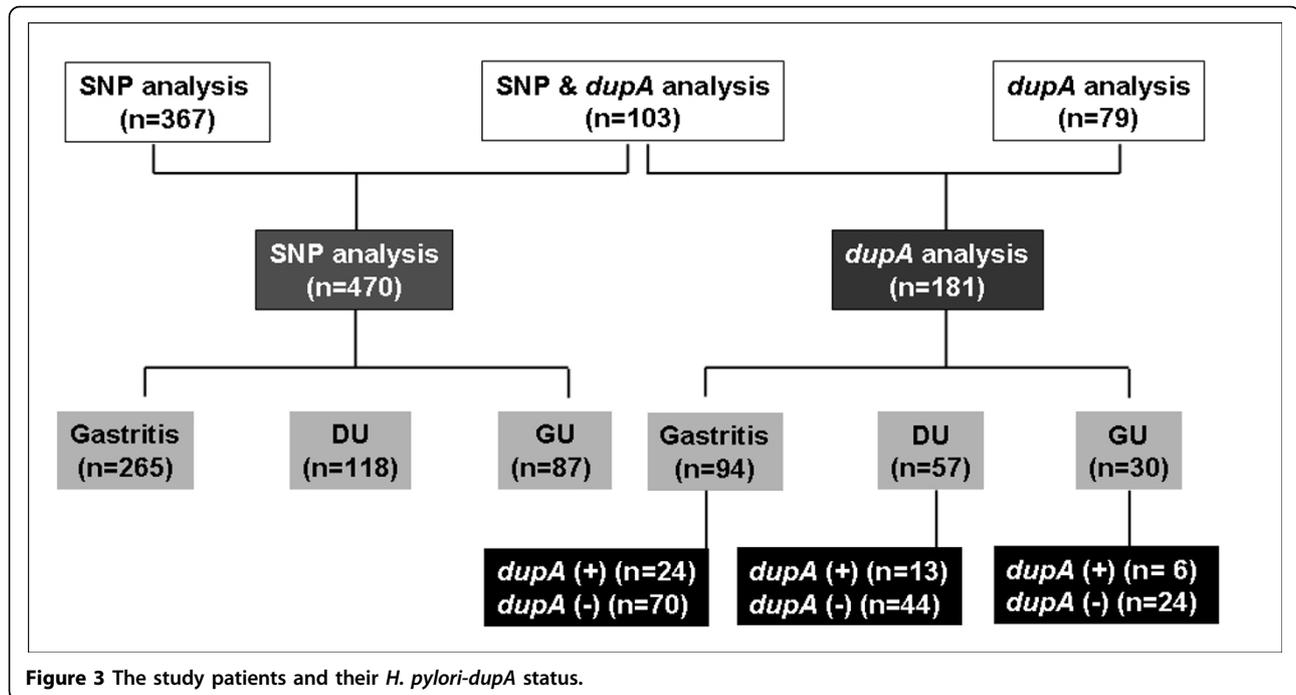
collagens, elastin, laminins, gelatin, and proteoglycan core proteins, and is thus helpful during wound healing of the skin [27-29]. Moreover, the gastric mucosa at the ulcer site also has significantly higher expression of MMP-3 than those in the antrum [30], which suggests MMP-3 is abundant in the ulcer part, and this may help the process of re-epithelialization and contribute to wound healing.

Hellmig et al. disclose the positive associations of MMP-7 promoter -181 and MMP-9 exon 6 SNPs to the presence of gastric ulcer among Germans [17]. However,

Table 2 Demographic and histologic characteristics of *H. pylori*-infected patients with single nucleotide polymorphism analysis (n = 470)

Parameters	Gastritis (n = 265)	Duodenal ulcer (n = 118)	Gastric ulcer (n = 87)	p value
Age, mean (SD) (yr)	46.9 (13.7)	47.6 (14.0)	47.8 (11.7)	NS
Gender, n (%)				
Female: Male	183 (69.1): 82 (30.9)	57 (48.3) : 61 (51.7)	36 (41.4) : 51 (58.6)	$p^a < 0.05; p^b < 0.05$
Histology score, mean (SD)				
(Antrum)				
AIS (range 1-3)	1.18 (0.99)	1.39 (0.95)	0.99 (1.03)	$p^c < 0.05$
CIS (range 0-3)	2.34 (1.01)	2.56 (0.89)	2.05 (1.17)	$p^a < 0.05; p^b < 0.05; p^c < 0.05$
(Corpus)				
AIS (range 1-3)	0.85 (0.99)	0.72 (0.95)	0.86 (1.06)	NS
CIS (range 0-3)	2.24 (0.86)	2.15 (0.83)	2.13 (0.89)	NS

Abbreviations: AIS, acute inflammation; CIS, chronic inflammation. The p value was determined by t test (age), χ^2 test (gender) or Mann-Whitney U test (histology score). ^a indicated significance with $p < 0.05$ of such parameter between gastritis and duodenal ulcer; ^b between gastritis and gastric ulcer; ^c between duodenal ulcer and gastric ulcer. NS: no significant difference.



this may be due to distinct ethnic or racial variations and such positive linkage is not disclosed in the current study from Taiwan.

This is the first report to show that there is no direct association between the genotypes of TIMP-1 372 at exon 5 and TIMP-2 at promoter -418, and the presence of gastroduodenal ulcers (Table 3). However, because TIMP-1 genotypes may modulate MMP-3 activity,

further testing if the MMP-3₋₁₆₁₂ /TIMP-1₃₇₂ Combined genotypes contribute to increased risk of duodenal ulcers shows that the combined MMP-3/TIMP-1 genotype as 6A6A/CC has a 3.6-fold increased risk of duodenal ulcer ($p < 0.05$) in *H. pylori*-infected females. This data suggests that TIMP-1 may also have a supportive role in interacting with MMP-3 during ulcerogenesis by *H. pylori* infection, especially in females.

Table 3 The SNP genotypes of MMPs and TIMPs in the both genders with different clinical diagnoses

Genotype	Female					Male					
	Gastritis	DU	GU	P^a	P^b	Gastritis	DU	GU	P^a	P^b	
MMP-3											
5A carrier	46 (25.1)	7 (12.3)	8 (22.2)	0.04	0.71	28 (34.1)	15 (24.6)	11 (21.6)	0.22	0.12	
6A/6 ^a	137 (74.9)	50 (87.7)	28 (77.8)			54 (65.9)	46 (75.4)	40 (78.4)			
MMP-7											
A/A	150 (82.9)	45 (80.4)	32 (91.4)	0.67	0.31	72 (88.9)	52 (86.7)	39 (76.5)	0.69	0.06	
G carrier	31 (17.1)	11 (19.6)	3 (8.6)			9 (11.1)	8 (13.3)	12 (23.5)			
MMP-9											
A carrier	78 (42.6)	19 (33.3)	15 (41.7)	0.21	0.92	25 (30.5)	22 (36.1)	22 (43.1)	0.48	0.14	
G/G	105 (57.4)	38 (66.7)	21 (58.3)			57 (69.5)	39 (63.9)	29 (56.9)			
TIMP-1											
♀ T carrier	♂ T	148 (80.9)	41 (73.2)	29 (80.6)	0.22	0.97	46 (56.8)	31 (50.8)	26 (51.0)	0.48	0.51
C/C		C	35 (19.1)	15 (26.8)			7 (19.4)	35 (43.2)	30 (49.2)		
TIMP-2											
C carrier		54 (30.2)	17 (32.1)	10 (28.6)	0.79	0.85	27 (33.3)	20 (32.3)	13 (25.5)	0.89	0.34
G/G		125 (69.8)	36 (67.9)	25 (71.4)			54 (66.7)	42 (67.7)	38 (74.5)		

Abbreviations: DU, duodenal ulcer; GU, gastric ulcer. The p value was determined by Fisher's exact test or χ^2 test. ^a indicated significance with $p < 0.05$ of such parameter between gastritis and duodenal ulcer; ^b between gastritis and gastric ulcer. Genotype distribution of SNP in cases and control was in Hardy-Weinberg equilibrium ($p < 0.05$).

Table 4 Risks of combined MMP-3/TIMP-1 genotype for developing duodenal ulcer in females

MMP-3 ₋₁₆₁₂ -TIMP-1 ₃₇₂	Gastritis n (%)	Duodenal ulcer n (%)	OR (95% CI)	P
5A carrier - T carrier	39 (88.6)	5 (11.4)	1	-
5A carrier - C/C	7 (77.8)	2 (22.2)	2.23 (0.36 - 13.85)	0.59
6A/6A - T carrier	109 (75.2)	36 (24.8)	2.58 (0.94 - 7.03)	0.06
6A/6A - C/C	28 (68.3)	13 (31.7)	3.62 (1.16 - 11.32)	0.03

The p value was determined by Fisher's exact test. OR, odds ratio; 95% CI, 95% confidence interval.

However, the exact reason why such a combined MMP-3/TIMP-1 genotype as 6A6A/CC has just an increased risk of duodenal ulcer in *H. pylori*-infected females, but not in male, remains uncertain. If such a regulation between MMP-3 and TIMP-1 play a role in female's ulceration, our data at least imply some other host or environment factors in male should be more dominant than such combined genotype.

The study has a limitation of just providing 181 isolates for the analysis of the *dupA* status of *H. pylori*, which disclose a rather low 20% *dupA*-positive prevalence rate. Accordingly, the study became limited to only 103 patients to provide both analyses on the infected isolate's *dupA* status and the host's SNPs (Figure 2). It thus cannot provide an adequate statistical power to determine the exact impact of MMP-3 SNPs under *dupA*-negative specific conditions.

Conclusions

In conclusion, this study provides evidence that host promoter polymorphisms of MMP-3 contribute to increased individual susceptibility to duodenal ulcers in females after *H. pylori* infection in Taiwan. The MMP-3 promoter genotypes may serve to screen out patients at risk and target for *H. pylori* eradication in order to stop the ulceration process among *H. pylori*-infected patients without ulcers yet.

Acknowledgements

This study was supported by grants from the National Science Council, Taiwan (95-2314-B-006-029-MY3 and 98-2628-B-006-013-MY3), NHRI-EX99-9908BI from the National Health Research Institute, and DOH99-TD-C-111-003 from Department of Health, Taiwan. The authors also thank Miss Hunt-Wen Wu for her assistance.

Author details

¹Institute of Basic Medical Sciences, National Cheng Kung University Hospital, Medical College, Tainan. ²Institute of Clinical Medicine, National Cheng Kung University Hospital, Medical College, Tainan. ³Department of Internal Medicine, National Cheng Kung University Hospital, Medical College, Tainan. ⁴Department of Pathology, National Cheng Kung University Hospital, Medical College, Tainan. ⁵Department of Pathology, Ton-Yen General Hospital, Hsin-Chu, Taiwan.

Authors' contributions

YCY prepared the manuscript, and carried out the molecular genetic studies to the host SNPs and *dupA* genotyping for the collected isolates of *H. pylori*. HCC and WLC carried out the SNP analysis and clinical specimen collection during endoscopy. HBY participated in the design of the study, performed

the analysis of pathology, and statistical analysis. BSS conceived of the whole study, and participated in its design and coordination. All authors read and approved the final manuscript.

Received: 24 March 2010 Accepted: 13 August 2010

Published: 13 August 2010

References

- Labigne A, de Reuse H: Determinants of *Helicobacter pylori* pathogenicity. *Infect Agents Dis* 1996, **5**(4):191-202.
- Maeda S, Mentis AF: Pathogenesis of *Helicobacter pylori* infection. *Helicobacter* 2007, **12**(Suppl 1):10-14.
- Prinz C, Schwendy S, Voland P: *H. pylori* and gastric cancer: shifting the global burden. *World J Gastroenterol* 2006, **12**(34):5458-5464.
- Sheu BS, Odenbreit S, Hung KH, Liu CP, Sheu SM, Yang HB, Wu JJ: Interaction between host gastric Sialyl-Lewis X and *H. pylori* SabA enhances *H. pylori* density in patients lacking gastric Lewis B antigen. *Am J Gastroenterol* 2006, **101**(1):36-44.
- Lai CH, Kuo CH, Chen YC, Chao FY, Poon SK, Chang CS, Wang WC: High prevalence of *cagA*- and *babA2*-positive *Helicobacter pylori* clinical isolates in Taiwan. *J Clin Microbiol* 2002, **40**(10):3860-3862.
- Lu H, Hsu PI, Graham DY, Yamaoka Y: Duodenal ulcer promoting gene of *Helicobacter pylori*. *Gastroenterology* 2005, **128**(4):833-848.
- Schmidt HM, Andres S, Nilsson C, Kovach Z, Kaakoush NO, Engstrand L, Goh KL, Fock KM, Forman D, Mitchell H: The *cag* PAI is intact and functional but HP0521 varies significantly in *Helicobacter pylori* isolates from Malaysia and Singapore. *Eur J Clin Microbiol Infect Dis* 2010, **29**(4):439-451.
- Nguyen LT, Uchida T, Tsukamoto Y, Kuroda A, Okimoto T, Kodama M, Murakami K, Fujioka T, Moriyama M: *Helicobacter pylori dupA* gene is not associated with clinical outcomes in the Japanese population. *Clin Microbiol Infect* 2010, **16**(8):1264-1269.
- Hussein NR: The association of *dupA* and *Helicobacter pylori*-related gastroduodenal diseases. *Eur J Clin Microbiol Infect Dis* 2010, **29**(7):817-821.
- Gomes LI, Rocha GA, Rocha AM, Soares TF, Oliveira CA, Bittencourt PF, Queiroz DM: Lack of association between *Helicobacter pylori* infection with *dupA*-positive strains and gastroduodenal diseases in Brazilian patients. *Int J Med Microbiol* 2008, **298**(3-4):223-230.
- Argent RH, Burette A, Miendje Deyi VY, Atherton JC: The presence of *dupA* in *Helicobacter pylori* is not significantly associated with duodenal ulceration in Belgium, South Africa, China, or North America. *Clin Infect Dis* 2007, **45**(9):1204-1206.
- Chang YT, Wu MS, Shun CT, Lin MT, Chang MC, Lin JT: Association of polymorphisms of *interleukin-1 beta* gene and *Helicobacter pylori* infection with the risk of gastric ulcer. *Hepatogastroenterology* 2002, **49**(47):1474-1476.
- Visse R, Nagase H: Matrix metalloproteinases and tissue inhibitors of metalloproteinases: structure, function, and biochemistry. *Circ Res* 2003, **92**(8):827-839.
- Yeh YC, Sheu BS, Cheng HC, Wang YL, Yang HB, Wu JJ: Elevated serum matrix metalloproteinase-3 and -7 in *H. pylori*-related gastric cancer can be biomarkers correlating with a poor survival. *Dig Dis Sci* 2010, **55**(6):1649-1657.
- Mori N, Sato H, Hayashibara T, Senba M, Gelezianus R, Wada A, Hirayama T, Yamamoto N: *Helicobacter pylori* induces matrix metalloproteinase-9 through activation of nuclear factor kappaB. *Gastroenterology* 2003, **124**(4):983-992.
- Crawford HC, Krishna US, Israel DA, Matrisian LM, Washington MK, Peek RM Jr: *Helicobacter pylori* strain-selective induction of matrix

- metalloproteinase-7 in vitro and within gastric mucosa. *Gastroenterology* 2003, **125**(4):1125-1136.
17. Hellmig S, Ott S, Rosenstiel P, Robert Folsch U, Hampe J, Schreiber S: **Genetic variants in matrix metalloproteinase genes are associated with development of gastric ulcer in *H. pylori* infection.** *Am J Gastroenterol* 2006, **101**(1):29-35.
 18. Jormsjo S, Whatling C, Walter DH, Zeiher AM, Hamsten A, Eriksson P: **Allele-specific regulation of matrix metalloproteinase-7 promoter activity is associated with coronary artery luminal dimensions among hypercholesterolemic patients.** *Arterioscler Thromb Vasc Biol* 2001, **21**(11):1834-1839.
 19. Ye S, Eriksson P, Hamsten A, Kurkinen M, Humphries SE, Henney AM: **Progression of coronary atherosclerosis is associated with a common genetic variant of the human stromelysin-1 promoter which results in reduced gene expression.** *J Biol Chem* 1996, **271**(22):13055-13060.
 20. Shipley JM, Doyle GA, Fliszar CJ, Ye QZ, Johnson LL, Shapiro SD, Welgus HG, Senior RM: **The structural basis for the elastolytic activity of the 92-kDa and 72-kDa gelatinases. Role of the fibronectin type II-like repeats.** *J Biol Chem* 1996, **271**(8):4335-4341.
 21. Clark IM, Swingle TE, Sampieri CL, Edwards DR: **The regulation of matrix metalloproteinases and their inhibitors.** *Int J Biochem Cell Biol* 2008, **40**(6-7):1362-1378.
 22. Sheu BS, Sheu SM, Yang HB, Huang AH, Wu JJ: **Host gastric Lewis expression determines the bacterial density of *Helicobacter pylori* in *babA2* genopositive infection.** *Gut* 2003, **52**(7):927-932.
 23. Dixon MF, Genta RM, Yardley JH, Correa P: **Classification and grading of gastritis. The updated Sydney System. International Workshop on the Histopathology of Gastritis, Houston 1994.** *Am J Surg Pathol* 1996, **20**(10):1161-1181.
 24. Shibata N, Ohnuma T, Higashi S, Usui C, Ohkubo T, Kitajima A, Ueki A, Nagao M, Arai H: **Genetic association between matrix metalloproteinase MMP-9 and MMP-3 polymorphisms and Japanese sporadic Alzheimer's disease.** *Neurobiol Aging* 2005, **26**(7):1011-1014.
 25. Zhou Y, Yu C, Miao X, Tan W, Liang G, Xiong P, Sun T, Lin D: **Substantial reduction in risk of breast cancer associated with genetic polymorphisms in the promoters of the matrix metalloproteinase-2 and tissue inhibitor of metalloproteinase-2 genes.** *Carcinogenesis* 2004, **25**(3):399-404.
 26. Wollmer MA, Papassotiropoulos A, Streffer JR, Grimaldi LM, Kapaki E, Salani G, Paraskevas GP, Maddalena A, de Quervain D, Bieber C, *et al*: **Genetic polymorphisms and cerebrospinal fluid levels of tissue inhibitor of metalloproteinases 1 in sporadic Alzheimer's disease.** *Psychiatr Genet* 2002, **12**(3):155-160.
 27. Bullard KM, Mudgett J, Scheuenstuhl H, Hunt TK, Banda MJ: **Stromelysin-1-deficient fibroblasts display impaired contraction in vitro.** *J Surg Res* 1999, **84**(1):31-34.
 28. Saarialho-Kere UK: **Patterns of matrix metalloproteinase and TIMP expression in chronic ulcers.** *Arch Dermatol Res* 1998, **290**(Suppl):S47-54.
 29. Madlener M: **Differential expression of matrix metalloproteinases and their physiological inhibitors in acute murine skin wounds.** *Arch Dermatol Res* 1998, **290**(Suppl):S24-29.
 30. Tomita M, Ando T, Minami M, Watanabe O, Ishiguro K, Hasegawa M, Miyake N, Kondo S, Kato T, Miyahara R, *et al*: **Potential role for matrix metalloproteinase-3 in gastric ulcer healing.** *Digestion* 2009, **79**(1):23-29.

doi:10.1186/1471-2180-10-218

Cite this article as: Yeh *et al.*: Matrix metalloproteinase-3 promoter polymorphisms but not *dupA-H. pylori* correlate to duodenal ulcers in *H. pylori*-infected females. *BMC Microbiology* 2010 **10**:218.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit

