Periodic Methylation Patterns in the Background Mucosa of Gastric Cancer

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Background/Aims: Gastrointestinal glandular stem cells renew every 8 years. New stem cells with impeded housekeeping gene methylation have unstable phenotypes and are prone to transform into malignant cells. Age-related changes in methylation in the gastric mucosa were evaluated to define the period of cancer-prone stem cell replacement.

Materials and Methods: Endoscopic biopsy specimens of normal-appearing gastric mucosa were obtained from 148 Helicobacter pylori-negative controls, 124 H. pylori-positive controls, and 69 gastric cancer patients with closed-type mucosal atrophy. Methylation-variable sites of two stomach-specific genes (TFF2 and TFF3) and four housekeeping genes (CDH1, ARRDC4, MMP2, and CDKN2A) were analyzed using radioisotope-labeled methylation-specific polymerase chain reaction. Age-related methylation was evaluated depending on the gastric mucosal atrophy at 2-year intervals.

Results: TFF2 methylation peaked periodically at 40 to 41, 48 to 49, 56 to 57, and 64 to 65 years of age in H. pylori-negative controls. Periodic peaks of TFF2 methylation were also found in H. pylori-positive controls. Housekeeping-gene methylation troughed at 48 to 49, 56 to 57, and 68 to 69 years of age in cancer patients. Trough methylation of CDH1 and ARRDC4 was lower in cancer patients than in H. pylori-positive controls.

Conclusions: Methylation peaks of stomach-specific TFF2 in controls and methylation troughs of housekeeping genes in cancer patients were found every 8 years. Periodic methylation patterns may be used to identify individuals at high risk for gastric cancer.

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Key Words: Adult stem cells; DNA methylation; Gastric cancer

INTRODUCTION

Old gastric stem cells that lose their renewal capacity need to be replaced with new stem cells as a part of the aging process.1,2 An age-related methylation study demonstrated that in humans with a relatively long lifespan, old colonic stem cells are replaced with new stem cells every 8 years.3,4 Newly replaced stem cells with unstable phenotypes are stabilized by housekeeping gene methylation.2-5 When phenotype-stabilizing methylation is impeded, new stem cells may experience reactive intrinsic migration and transform into malignant cells. In our previous study of the gastric mucosa, housekeeping gene methylation was impeded in patients with gastric cancer for an average of 8 years.5 The self-renewal capacity of gastric mucosal glandular stem cells appears to be restored periodically. Age-specific methylation patterns could provide a clue to distinguish individuals at high-risk for gastric cancer during the period of stem cell replacement.

Methylation-variable regions are different between housekeeping genes and tissue-specific genes.9 In housekeeping genes containing CpG islands, CpG-island margins display more wide ranges of methylation variations compared to CpG-island centers, and their methylation is influenced by the types and the distances of gene-adjacent retroelements.9,10 Tissue-specific genes lacking CpG islands showed variable methylation near the transcription start sites under the influence of their expression.9 Variable methylation regions of tissue-specific and housekeeping genes are termed transitional–CpG sites.9,11 Previous methylation studies of the stomach have focused on the association between CpG-island center methylation and tumor-suppressor gene inactivation. However,
the inactivation role of CpG-island methylation during gastric carcinogenesis is unclear. For instance, allele-specific hypermethylation of the CDH1 gene is uncommon in familial or sporadic gastric cancer with loss of CDH1 expression. Additionally, housekeeping genes are more frequently methylated in subjects infected with Helicobacter pylori than in gastric cancer patients. Transitional-CpG methylation may be associated with the stabilization of stem cell phenotypes. New stem cells fixed in the gastric mucosa acquire the phenotypes of gastric stem cells through high expression of stomach-specific genes and fine down-regulation of numerous housekeeping genes. Down-regulated housekeeping genes are stabilized by cell division-dependent methylation. Therefore, clonal expansion of new stem cells is likely to be estimated by analyzing the transitional CpG methylation of stomach-specific and housekeeping genes.

This study aimed to evaluate the period of cancer-prone stem cell replacement. Transitional CpG methylation will change in accordance with the periodic replacement of new stem cells. Because H. pylori-associated gastric mucosal atrophy spreads from the gastric antrum to the body, the body sites are less affected by atrophy-related methylation inhibition. Age-specific changes in transitional CpG methylation in both the antrum and the body were analyzed in H. pylori-negative controls, H. pylori-positive controls, and patients with gastric cancer at 2-year intervals.

MATERIALS AND METHODS

1. Endoscopic biopsy specimens

Endoscopic biopsy specimens of normal gastric mucosa were obtained from St. Paul Hospital (Seoul, Korea), Bucheon St. Mary’s Hospital (Bucheon, Korea), and Incheon St. Mary’s Hospital (Incheon, Korea) between March 2008 and June 2016 and included in a methylation study. Paired mucosa specimens were biopsied from the lesser curvature of the antrum and the greater curvature of the middle body and stored immediately at −70°C. H. pylori infection was determined using the Warthin-Starry silver impregnation method. The extent of gastric mucosal atrophy was classified into closed- and open-type gastric mucosal atrophy based on the endoscopic atrophic border score proposed by Kimura and Takemoto. The subjects with open-type mucosal atrophy were excluded in this study in order to discriminate past or long-term H. pylori infection.

2. DNA extraction and bisulfite treatment

DNA extraction and bisulfite treatment were performed as described elsewhere. In brief, biopsy specimens were dissolved in DNA extraction buffer (0.5% Tween 20, 1 mM EDTA, 50 mM Tris, and 0.5 mg/mL proteinase K) for 24 hours at 37°C. The extracted DNA samples were modified as follows: 21.5 μL of genomic DNA were denatured in 3 M NaOH at 37°C for 15 minutes and subsequently treated with 2.5 M sodium bisulfite and 10 mM hydroquinone at 50°C for 12 hours. DNA was purified using Wizard DNA purification resin (Promega, Madison, WI, USA) and desulfonated with 3 M NaOH at 37°C for 15 minutes, precipitated with ethanol. DNA was dissolved in 40 μL of 5 mM Tris buffer.

3. Methylation analysis of biopsy specimens

Two stomach-specific genes (TFF2 and TFF3), two housekeeping genes adjacent to Alu retroelements (CDH1 and ARRDCA), and two housekeeping genes adjacent to LTR retroelements (MMP2 and CDKN2A) were selected for the methylation analysis of small biopsy specimens. Methylation primers were designed to target the methylation-variable transitional-CpG sites between the transcriptional start sites and gene-adjacent retroelements.

Methylation-specific polymerase chain reaction (MSP) was performed under stringent conditions with a hot start condition and radioisotope labeling to compensate for the low guanine-cytosine contents and repetitive sequences of transitional-CpG sites. One microliter of bisulfite-treated DNA was mixed with 1 mCi of α-32P dTTP (PerkinElmer, Boston, MA, USA), 62.5 mM each dATP, dCTP, and dGTP, 25 mM dTTP, 1 pM of primers, 0.1% Tween 20, and 0.5 unit of Taq DNA polymerase (Takara, Kyoto, Japan). The reactions were subjected to thermal cycling as follows: denaturation at 99°C for 5 minutes followed by the addition of DNA polymerase and dNTP mixture; amplification for 32 cycles of 93°C for 50
seconds, 53°C to 62°C for one minute, and 72°C for one minute; and extension at 72°C for 3 minutes. MSP products were separated by polyacrylamide gel electrophoresis and visualized by autoradiography with a radioluminograph scanner (PMI 170-9400; BioRad, Hercules, CA, USA). The band intensities were measured using TINA image software (Raytest Isotopenmeβgerate, Straubenhardt, Germany). Methylation values were scored as the proportion of band intensities of the methylation primer to the total band intensities of the methylation and unmethylation primers, and were stratified into 10 levels at 10% increments.

4. Ethics statement

The study of biopsy specimens was reviewed and approved by St. Paul’s Hospital Institutional Review Board, the Catholic University of Korea (IRB no. PCMC08BR005 and XC15TIMI0044P). All participating subjects provided written informed consent prior to the study.

RESULTS

A total of 148 H. pylori-negative controls, 124 H. pylori-positive controls, and 69 cancer patients from 38 to 81 years of age were enrolled to perform methylation study on the normal gastric mucosa (Fig. 1). Cancer patients were not grouped by H. pylori infection because there were few differences in transitional-CpG methylation between H. pylori-positive and -negative cancer patients in previous studies.6

Age-related changes in methylation were plotted at 2-year age intervals, and the periodic curves were evaluated during 8-year cycles (Fig. 2A). The TFF2 methylation curve tended to peak every 8 years, regardless of sex or H. pylori status. In samples from H. pylori-negative men, the TFF2 methylation curve exhibited five regular peaks at 40 to 41, 48 to 49, 56 to 57, 64 to 65, and 72 to 73 years of age. TFF2 methylation peaks were detected at 40 to 41, 46 to 47, 54 to 55, and 64 to 65 years of age in H. pylori-positive men. The methylation of TFF3, which is specifically expressed in the antrum, was known

Fig. 1. Age distribution of subjects included in the transitional-CpG methylation analysis. Samples are collected at 2-year intervals.
Fig. 2. Age-related changes in methylation in *Helicobacter pylori*-negative controls, *H. pylori*-positive controls, and gastric cancer patients with closed-type atrophy. (A) Individual methylation of TFF2 and TFF3 in the antrum and concurrent methylation of CDH1, ARRDC4, MMP2, and CDKN2A in the body are depicted. (B) The mean levels of CDH1 and ARRDC4 methylation in the body are shown in the presence or absence of TFF3 overmethylation in the antrum. TFF3 overmethylation is defined using level 6 as the cutoff point. Samples were collected at 2-year intervals. Vertical dashed lines indicate 8-year intervals after 40 years of age. Horizontal dashed lines indicate level 3 methylation.
to be impeded under the influence of gastric mucosal atrophy. In *H. pylori*-positive controls, TFF3 methylation was increased at 50 to 51 and 56 to 57 age groups in women but not in men (Fig. 2A).

*CDH1* and *ARRDC4* that are adjacent to *Alu* retroelements were frequently methylated in the body of *H. pylori*-negative male controls. Mean methylation levels of these two genes were less than 3, with weak increases at ages 46 to 47 and 56 to 57 in the body of *H. pylori*-negative males. Concurrent methylation of *CDH1* and *ARRDC4* was increased in the 66 to 77 age group in the body of *H. pylori*-negative male controls (Fig. 2A). *CDH1* and *ARRDC4* were concurrently methylated in the 50 to 51 and 58 to 59 age groups in the body of both *H. pylori*-positive male and female controls. The mean methylation levels of *CDH1* and *ARRDC4* troughed at 48 to 49, 56 to 57, and 68 to 69 years of age in male cancer patients; troughs were lower in cancer patients than in *H. pylori*-negative and *H. pylori*-positive controls (2.0 vs. 2.5 and 2.5 at 48 to 49 years; 1.0 vs. 2.4 and 2.6 at 56 to 57 years; and 1.7 vs. 2.9 and 4.0 at 68 to 69 years).

*MMP2* and *CDKN2A* that are adjacent to *LTR* retroelements were gradually methylated in the body of both *H. pylori*-negative and *H. pylori*-positive controls. Mean methylation levels of *MMP2* and *CDKN2A* troughed at 48 to 49, 56 to 57, and 66 to 67 years of age in the body of men with gastric cancer; the last two troughs were lower in cancer patients than in *H. pylori*-negative and *H. pylori*-positive controls (3.5 vs. 3.8 and 2.8 at 48 to 49 years; 3.0 vs. 4.5 and 3.6 at 56 to 57 years; and 3.5 vs. 4.9 and 6.0 at 66 to 67 years) (Fig. 2A). Concurrent methylation of *MMP2* and *CDKN2A* methylation was increased in the 50 to 53 and 58 to 61 age groups in men with gastric cancer, and that was higher than in *H. pylori*-negative and *H. pylori*-positive male controls (5.4 vs. 3.5 and 4.5 at 50 to 51 years; 5.8 vs. 3.3 and 4.9 at 52 to 53 years; 5.2 vs.
3.2 and 4.7 at 58 to 59 years; and 6.0 vs. 3.4 and 4.4 at 60 to 61 years).

In a previous study, body-site housekeeping genes and antrum-site TFF3 tended to be coincidentally methylated at an early stage. CDH1 and ARRDC4 body-site methylation were evaluated in the presence and absence of antrum-site TFF3 overmethylation of level ≥6 (Fig. 2B). In H. pylori-negative samples from men, four regular peaks of CDH1 and ARRDC4 methylation (level >3) were observed at 48 to 49, 56 to 57, 66 to 67, and 72 to 73 years of age in the absence of TFF3 overmethylation.

DISCUSSION

Stomach-specific TFF2 is strongly expressed in gastric glands in both the antrum and body. Notably, our analysis of normal gastric mucosa samples revealed peaks in the methylation of TFF2 at regular intervals of approximately 8 years, and these peaks corresponded to similar peaks in housekeeping gene methylation in H. pylori-negative subjects. These regular peaks in age-related methylation likely represent the periodic replacement of gastric glandular stem cells, some of which may be transformed into cancer stem cells. Housekeeping genes tended to be less methylated in cancer patients than in H. pylori-positive controls during the periods of TFF2 methylation peaks. Previous studies of the background mucosa of gastric cancer have demonstrated that insufficient methylation of some housekeeping genes leads to an early unstable stage of stem cell replacement. The appearance of cancer may vary during the extended cancer risk stage during each period of stem cell replacement. The methylation type related to cancer risk is expected to be informative when evaluating the individuals at risk for cancer.

H. pylori-positive subjects had high-level methylation but no periodic methylation changes. H. pylori-induced active inflammation can recruit many new mesenchymal cells around the gastric glands. In such cases, glandular epithelial cells obtained by endoscopic biopsy were contaminated with nearby mesenchymal cells. Both the mesenchymal and epithelial cells adapting to a new tissue environment can be stabilized by methylation of housekeeping genes, and the mixture of them in the biopsy specimens may make no cyclic changes and no sex-based differences in the biopsy specimens. Periodic fluctuation in housekeeping gene methylation was more evident in the body than in the antrum of H. pylori-negative subjects. In a body with mild inflammation, endoscopic biopsy specimens contain a relatively low number of new mesenchymal cells, which can disclose cyclic methylation changes.

Cyclic TFF2 methylation seems to result from rapid demethylation of transitional-CpG sites influenced by strong gene expression. Cyclic TFF3 methylation was somewhat ambiguous in the antrum, possibly because antrum-specific TFF3 expression is weak. TFF3 and housekeeping genes tended to be concurrently methylated at an early stage and afterward TFF3 methylation decreased at a late stage. Housekeeping-gene methylation regularly fluctuated in the absence of TFF3 overmethylation in the body of H. pylori-negative subjects. Because there were a low number of new stem cells in H. pylori-negative body, housekeeping gene methylation might be increasingly detected as methylation-inducible cells expanded sufficiently and TFF3 methylation decreased.

Retroelements dispersed throughout the genome are thought to be involved in the epigenetic suppression of numerous housekeeping genes during phenotype stabilization of new stem cells. Humans with a long lifespan have higher numbers of SINE retroelements around housekeeping genes than mice with a short lifespan. This helps to prompt phenotype-stabilizing methylation of new stem cells in humans. Housekeeping genes adjacent to Alu retroelements, human SINES, are frequently methylated in the gastric mucosa of subjects in their early 40s. Previous studies of the background mucosa of gastric cancer have demonstrated that impeded methylation of Alu-adjacent genes leads to an early unstable stage of new stem cells. LTR retroelements also participate in the phenotypic stabilization of stem cells. LTR-adjacent housekeeping gene methylation increased slowly compared to Alu-adjacent gene methylation. LTR-adjacent genes were more frequently methylated in gastric cancer patients than in H. pylori-positive subjects. Sequential methylation changes in Alu-adjacent and LTR-adjacent
genes may provide information on the formation of cancer-prone stem cells in the gastric mucosa.

Japan and Korea, where gastric cancer is common, conduct biennial endoscopic screening to detect early gastric cancer, as a 2-year screening interval has been proposed to reduce gastric cancer mortality. However, the interval of cancer screening has been determined empirically, and there is no clear evidence as to the optimal screening interval. In this study, periodic methylation was observed in consistent with the cyclic stabilization of cancer-prone stem cells every 8 years. A previous gastric cancer screening at 5-year intervals revealed a decrease in the incidence of gastric cancer at the first follow-up and an increase at the second follow-up. The results of the 5-year interval screening, as well as the 8-year-periodic fluctuation in methylation, indicate that the chances of gastric cancer arise at 8-year intervals and that a screening interval <8 years is necessary for early cancer detection.

Gastric mucosal atrophy is evaluated using endoscopic examination of the atrophic border and histopathologic examination based on the Sydney system. These two atrophy-staging systems demonstrate relatively good agreements. Our previous studies revealed that Alu-adjacent methylation was impeded in cancer patients with closed-type atrophy and LTR-adjacent methylation was increased in cancer patients with open-type atrophy compared with controls. This suggests that cancer-related methylation changes are distinct from the progression of gastric mucosal atrophy. Active H. pylori inflammation detected in this study may induce the recruitment of new stem cells intermittently in the gastric mucosa. However, stomach-specific TFF2 methylation peaked at regular intervals in H. pylori-positive subjects. Stem cell replacement appears to occur regularly regardless of H. pylori infection. Gastric mucosal atrophy and H. pylori infection may influence the phenotype-stabilizing methylation during stem cell replacement.

This study had some potential limitations. First, inherent factors may affect methylation patterns. Thus, it is challenging to analyze periodic methylation patterns depending on the family history of gastric cancer patients. Second, because of a relatively small sample size, the authors could not conclude the significant decreases in trough methylation of housekeeping genes in cancer patients. Longitudinal studies with a larger sample size are required to investigate periodic methylation levels and the risk of gastric cancer. Third, there may be interobserver differences in the extent of gastric mucosal atrophy based on endoscopic examination. To ensure the reliability of the atrophic border classification, gastric mucosal atrophy was measured repeatedly by two endoscopy specialists. Fourth, the histopathologic types of gastric cancer were not evaluated in this study. In human and mouse gastric cancers, phenotype shift from diffuse to intestinal types is clearly defined. New stem cells may be unstable at an early stage and acquire gastric stem cell traits at a late stage. Periodically replaced stem cells seem to develop into diffuse-type cancer, followed by intestinal-type cancer, depending on phenotypic stability. Further studies are needed to clarify the relationship between the histologic types of gastric cancer and the phenotype-stabilizing methylation.

In summary, methylation peaks of stomach-specific TFF2 in controls and methylation troughs of housekeeping genes in gastric cancer patients appeared at similar intervals of 8 years. This suggests that new gastric stem cells with high self-renewal capacity replace old stem cells at certain periods rather than irregular periods and that new stem cells displaying impeded housekeeping gene methylation are prone to transform into malignant cells. To our knowledge, stem cell periodicity in terms of gastric cancer evolution has not been previously reported. Even though gastric glandular stem cells were periodically replaced, the initiation of stem cell replacement could be diverse among individuals. Age-related transitional CpG methylation provides a clue to the initiation point of stem cell replacement. Individuals at high risk for gastric cancer may be effectively distinguished by tracing the phenotype-stabilizing methylation of housekeeping genes after the period of stem cell replacement.

**CONFLICT OF INTEREST**

No potential conflict of interest relevant to this article was reported.
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