Histone modification alteration coordinated with acquisition of promoter DNA methylation during Epstein-Barr virus infection

Sayaka Funata¹², Keisuke Matsusaka¹, Ryota Yamanaka³, Shogo Yamamoto³, Atsushi Okabe¹, Masaki Fukuyo¹, Hiroyuki Aburatani³, Masashi Fukayama² and Atsushi Kaneda¹

¹Department of Molecular Oncology, Graduate School of Medicine, Chiba University, Chiba, Japan
²Department of Pathology, Graduate School of Medicine, The University of Tokyo, Tokyo, Japan
³Genome Science Division, Research Center for Advanced Science and Technology, The University of Tokyo, Tokyo, Japan

Correspondence to: Atsushi Kaneda, email: kaneda@chiba-u.jp

Keywords: DNA methylation, Epstein-Barr virus, gastric cancer

Received: April 25, 2017  Accepted: July 11, 2017  Published: July 21, 2017

Copyright: Funata et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License 3.0 (CC BY 3.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

ABSTRACT

Aberrant DNA hypermethylation is a major epigenetic mechanism to inactivate tumor suppressor genes in cancer. Epstein-Barr virus positive gastric cancer is the most frequently hypermethylated tumor among human malignancies. Herein, we performed comprehensive analysis of epigenomic alteration during EBV infection, by Infinium HumanMethylation 450K BeadChip for DNA methylation and ChIP-sequencing for histone modification alteration during EBV infection into gastric cancer cell line MKN7. Among 7,775 genes with increased DNA methylation in promoter regions, roughly half were “DNA methylation-sensitive” genes, which acquired DNA methylation in the whole promoter regions and thus were repressed. These included anti-oncogenic genes, e.g. CDKN2A. The other half were “DNA methylation-resistant” genes, where DNA methylation is acquired in the surrounding of promoter regions, but unmethylated status is protected in the vicinity of transcription start site. These genes thereby retained gene expression, and included DNA repair genes. Histone modification was altered dynamically and coordinately with DNA methylation alteration. DNA methylation-sensitive genes significantly correlated with loss of H3K27me3 pre-marks or decrease of active histone marks, H3K4me3 and H3K27ac. Apoptosis-related genes were significantly enriched in these epigenetically repressed genes. Gain of active histone marks significantly correlated with DNA methylation-resistant genes. Genes related to mitotic cell cycle and DNA repair were significantly enriched in these epigenetically activated genes. Our data show that orchestrated epigenetic alterations are important in gene regulation during EBV infection, and histone modification status in promoter regions significantly associated with acquisition of de novo DNA methylation or protection of unmethylated status at transcription start site.

INTRODUCTION

Cancer development and progression occur through accumulation of genetic and epigenetic disruptions [1, 2]. Almost 70% of annotated gene promoters have CpG-rich sequences called “CpG islands”, which are concentrated with non-methylated CpGs in somatic cells. DNA methylation of promoter CpG islands correlates with silencing of the downstream genes. Many tumor suppressor genes were de novo methylated and inactivated in cancer cells, which contributes to carcinogenesis [2, 3]. These aberrant DNA methylations seemed to occur in a non-random manner, and many types of cancer are classified into several epigenotypes based on the degree of DNA methylation. Gastric cancer is classified into at least three DNA methylation phenotypes: low-methylation, high-methylation, and extensively high-methylation epigenotypes [4]. These methylation epigenotypes reflect molecular subtypes such as microsatellite-instable (MSI) gastric cancer or Epstein-Barr virus (EBV)-associated gastric cancer [5, 6].
As the standpoint of the cause for aberrant DNA methylation in gastric cancer, EBV is demonstrated to induce de novo DNA methylation in host gastric epithelial cells during infection, which may be crucial for EBV-positive gastric carcinogenesis [4, 7]. EBV is a γ-herpes virus establishing lifelong asymptomatic infection called “latency” in B-lymphocytes in almost all human populations. The latency is restricted by the host cell types, and EBV is detected in many types of tumor tissues, not only in lymphoid malignancies, but also in many epithelial tumors [8]. Almost 10% of gastric cancer cases are EBV-positive [9]. In gastric cancer cells, EBV forms type I latency, where viral genome undergoes dense DNA methylation compared to other latencies, and the host genome shows extensively high-methylation epigenotype [4], named EBV-CIMP [6]. EBV infection itself was shown to establish this extensive DNA methylation in our previous study [4, 7].

Another epigenetic process, histone modification, which is a post-translational modification of the histone tails, mostly occurs at lysine and arginine residues. The high levels of H3K4 trimethylation (H3K4me3) or H3K27 acetylation (H3K27ac) at the promoters are implicated in the transcriptional activation, whereas H3K27 trimethylation (H3K27me3) is correlated with gene repression and silencing [10]. The establishment of appropriate histone modification patterns are essential for the normal development and tissue differentiation, while mutations of these modulators and aberrations of histone modifications are associated with cancer development [11, 12]. Several gene specific or genome-wide studies of histone modification alterations have been reported in gastric cancer [13–15]. However, alterations of EBV-positive gastric cancer and their association with aberrant DNA methylation are mostly unknown.

To clarify epigenomic alteration in promoter regions of host cell during EBV infection, we analyzed DNA methylation using Infinium HumanMethylation 450K BeadChip (Illumina, San Diego, CA, USA) and alteration of histone modifications, e.g., H3K4me3, H3K27ac, and H3K27me3, by ChIP-sequencing using in vitro system of EBV infection into gastric cancer cell line MKN7. A total of 9,275 genes presented newly methylated CpG probes of Infinium BeadChip in promoter regions, which are regarded as DNA methylation-induced genes. These DNA methylation-induced genes were divided into two gene types. Roughly half of the DNA methylation-induced genes were “DNA methylation-sensitive” genes which acquired DNA methylation in whole promoter CpG probes, and the other half represented “DNA methylation-resistant” genes. DNA methylation-resistant genes showed continuous non-methylated CpGs at immediate vicinity of the transcription start site (TSS) and maintained gene expression as if the TSS was protected from the DNA methylation-induction pressure. Histone modifications were also dynamically altered at promoter regions. The activated genes were related to mitotic cell cycle and DNA repair, while inactivated genes were related to apoptosis. It seemed that the protection of unmethylated TSS status at DNA methylation-resistant genes required the existence of active histone marks, whereas the loss of those marks strongly correlated with DNA methylation-sensitive genes.

RESULTS

Distribution of methylated and unmethylated CpG sites around the TSS

If the methylation status of an Infinium probe was altered from the unmethylated state (U; β-score ≤ 0.20) in MKN7_WT cells to a methylated state (M; β-score ≥ 0.40), the probe was called “U to M” probe. If there were ≥ 1 “U to M” probes within ± 1,000 bp from the TSS, we regarded the genes as de novo methylation-induced genes, and 9,275 genes satisfied the criteria. When we calculated the average of β-score of all the probes within ± 1,000 bp from the TSS of the 9,275 genes, based on the distance from the TSS, the elevated level of β-score was lower in the vicinity of the TSS (Figure 1A). Among the 9,275 methylation-induced genes, “U to U” and “U to M” probes were then separately counted, based on the distance from the TSS, and their ratios to all the probes were compared (Figure 1B). The ratio of “U to M” probes was lower in the vicinity of the TSS (within ± 400 bp from TSS), but that of “U to U” probes was inversely higher in the vicinity of the TSS. These findings suggested that unmethylated CpG sites were unevenly maintained or protected under DNA methylation induction during EBV infection.

DNA methylation patterns during EBV infection

We further analyzed DNA methylation induction patterns during EBV infection. Among the methylation-induced genes, 4,043 genes were found to have ≥ 2 consecutive “U to U” probes within ± 500 bp from the TSS and defined as methylation-resistant genes (Figure 2A). In contrast, 3,732 genes did not have consecutive “U to U” probes within ± 500 bp from the TSS and defined as methylation-sensitive genes (Figure 2B). In 3,357 genes without methylation induction, all the probes within ± 1,000 bp from the TSS were “U to U” probes (Figure 2C). In 3,924 originally methylated genes, all the probes within ± 1,000 bp from the TSS were “M to M” probes (Figure 2D). A total of 403 genes had ≥ 1 “M to U” probes within ± 500 bp from the TSS and defined as demethylated genes (Figure 2E).

Features of methylation-sensitive and resistant genes

The methylation-sensitive genes in the two independent EBV-infected MKN7 clones were compared and they overlapped well (Figure 3A). They were
Figure 1: Distribution of DNA methylation status around the TSS after EBV infection. (A) β-score of de novo methylated genes around the TSS. If the methylation status of an Infinium probe was altered from U in MKN7_WT to M in MKN7_EB_A1 cells, the probe was called “U to M” probe. If there were ≥ 1 “U to M” probes within ± 1,000 bp from the TSS, we regarded the genes as de novo methylation-induced genes, and 9,275 genes satisfied the criteria. The average of β-score of all the probes within ± 1,000 bp from the TSS of the 9,275 genes was calculated based on the distance from the TSS, and the distribution of DNA methylation levels around the TSS in MKN7_EB_A1 cells was shown. (B) Distribution of “U to U” and “U to M” probes around the TSS. Among the 9,275 methylation-induced genes, “U to U” and “U to M” probes were separately counted based on the distance from the TSS, and their ratios to all the probes are shown.
markedly repressed in both MKN7_EB_A1 and MKN7_EB_B6 clones (Figure 3B), and gene ontology (GO) terms significantly enriched in the overlapped methylation-sensitive genes were related to differentiation, cell adhesion, development, and regulation of cell proliferation, e.g., CDH1, CDKN2A, and RHOB (Figure 3C). Methylation acquisition in the whole promoter regions of genes related to differentiation and development as well as their repression are consistent with our previous finding that PRC target genes in ES cells were targets of aberrant methylation in EBV-infected gastric cancer cells [4].

The methylation-resistant genes in MKN7_EB_A1 and MKN7_EB_B6 clones were also well overlapped (Figure 3D). Their expression levels in MKN7_WT cells were retained in both EBV-infected clones (Figure 3E), and GO terms significantly enriched in the overlapped methylation-resistant genes were related to DNA repair, e.g., mismatch repair genes, including MLH1, MSH2, and MSH6 (Figure 3F).

Figure 2: Patterns of DNA methylation acquisition during EBV infection. (A) Methylation-resistant genes. Among the methylation-induced genes that had at least one “U to M” probe within ±1000 bp from the TSS, 4,043 genes were found to have ≥2 consecutive “U to U” probes within ±500 bp from the TSS, and were defined as methylation-resistant genes. (B) Among the methylation-induced genes, 3,732 genes did not have consecutive “U to U” probes within ±500 bp from the TSS, and were defined as methylation-sensitive genes. (C) In 3,357 genes without methylation, all the probes within ±1,000 bp from TSS were “U to U” probes. (D) In 3,924 originally methylated genes, all the probes within ±1,000 bp from TSS were “M to M” probes. (E) In 403 demethylated genes, at least one probe was “M to U” probe within 500 bp from the TSS.
Alteration of histone active marks

We next analyzed alteration of histone H3K4me3 and H3K27ac marks during EBV infection by ChIP-seq, and their correlations to gene expression and DNA methylation induction.

H3K4me3 signal around the TSS in MKN7_EB_A1 cells was decreased in 2,363 genes, remained positive in 10,284 genes, remained negative in 7,039 genes, and increased in 913 genes compared to that in MKN7_WT (Figure 4A). Compared to that in MKN7_WT cells, gene expression was markedly downregulated in genes with decreased H3K4me3, but markedly upregulated in genes with increased H3K4me3 (Figure 4B). Consistently, the DNA methylation level was markedly elevated from 0.11 ± 0.00 to 0.35 ± 0.01 (average β-score ± SE) in genes with decreased H3K4me3, which significantly included differentiation- and morphogenesis-related genes (Figure 4C, 4D). However, DNA methylation level was retained at unmethylated state from 0.16 ± 0.01 to 0.16 ± 0.01 (β-score) in genes with increased H3K4me3, which significantly included cell cycle related genes.

H3K27ac signals at promoters were similarly analyzed in MKN7_WT and MKN7_EB_A1 cells. H3K27ac signal around the TSS in MKN7_EB_A1 cells was decreased in 2,350 genes, remained positive in 5,582 genes, remained negative in 10,850 genes, and increased in 1,817 genes compared to that in MKN7_WT cells (Figure 5A). Compared to that in MKN7_WT cells, the gene expression was markedly downregulated in genes with decreased H3K27ac signal, but markedly upregulated in genes with increased H3K27ac signal (Figure 5B). Similar to H3K4me3, DNA methylation level was markedly elevated from 0.09 ± 0.00 to 0.24 ± 0.01 (β-score) in genes with decreased H3K27ac, but retained at unmethylated state from 0.10 ± 0.00 to 0.12 ± 0.00 (β-score) in genes with increased H3K27ac signal (Figure 5C). Significantly enriched GO terms were related to apoptosis and cell death in genes with decreased H3K27ac signal (Figure 5D); repression and DNA methylation of these genes were suggested to contribute to tumorigenesis. Significantly enriched GO terms in increased H3K27ac signal were related to cell cycle and DNA repair; retained expression of these genes was also suggested to contribute to tumorigenesis.

To confirm that these histone modification and gene expression alteration were not due to long-term passage of cells but due to EBV infection, we also analyzed MKN7 cells cultured for the same period (Supplementary Figure 1). Decrease or increase of H3K4me3 and H3K27ac, and the subsequent gene expression alterations, were not generally observed during the long-term passage.

Alteration of histone repressive mark

We next analyzed alteration of histone H3K27me3 mark during EBV infection by ChIP-seq and its correlation to gene expression and DNA methylation induction.

H3K27me3 signal around the TSS in MKN7_EB_A1 cells was decreased in 2,262 genes, and remained negative in 18,222 genes compared to that in MKN7_WT cells (Figure 6A). Genes maintaining positive H3K27me3 signal or showing increased H3K27me3 signal were rarely observed. Compared to that in MKN7_WT cells, gene expression was not upregulated in genes with decreased H3K27me3 mark, but low expression levels were maintained in MKN7_EB_A1. Gene expression was also retained in genes retaining negative H3K27me3 signal (Figure 6B). DNA methylation level was markedly elevated from 0.30 to 0.46 (β-score) in genes with decreased H3K27me3, suggesting a switch of repressive epigenetic modifications from H3K27me3 to DNA methylation.

Integrated analysis of all the histone marks and DNA methylation

As for each histone modification alteration pattern (Figures 4–6), DNA methylation levels in MKN7_WT and MKN7_EB_A1 cells were assessed by calculating the average of β-scores, based on the distance from the TSS (Figure 7). When signals of histone active marks were decreased or retained at lower levels, i.e., were low after EBV infection, these genes were significantly associated with methylation-sensitive genes showing elevated DNA methylation level in whole promoter regions, including the vicinity of the TSS. When signals of histone active marks were increased or retained at high levels, i.e., were high after EBV infection, these genes were significantly associated with methylation-resistant genes, showing protection of unmethylated state in the vicinity of the TSS. Genes losing H3K27me3 were significantly associated with DNA methylation-sensitive genes, as mentioned above.

If the decrease in H3K4me3 occurred in H3K27ac(-) genes, the genes likely showed decrease of H3K27ac (Figure 8, subgroup 2). If decrease in H3K4me3 occurred in H3K27ac(-) genes, the genes maintained H3K27ac(-) status and hardly gained H3K27ac (Figure 8, subgroups 3–4). Decrease in H3K4me3 thus hardly coexists with high level of H3K27ac after EBV infection. Whereas the decreases of these active marks were correlated with de novo DNA methylation, the genes losing H3K4me3, but keeping H3K27ac(+) status were significantly correlated with methylation-resistant genes, implying that not only H3K4me3, but also modulators for H3K27ac might also work as resistant factors for DNA methylation induction (Figure 8).

Genes maintaining H3K4me3(+) status with decreased H3K27ac (Figure 8, subgroup 6) or maintained H3K27ac(-) status (Figure 8, subgroups 8–9) were mostly DNA methylation-resistant genes. However, genes maintaining both H3K4me3(+) and H3K27ac(+) status, abundantly included DNA methylation-resistant genes.
as well as unmethylated genes (Figure 8, subgroup 5). The unmethylated genes are mostly H3K4me3-positive genes, whose H3K27ac status is mostly retained positive (Figure 8, subgroup 5) or increased (Figure 8, subgroup 7).

Genes with increased H3K4me3 signal generally related to DNA methylation-resistant genes (Figure 8, subgroups 10–12).

Among genes maintaining H3K4me3(−) status, those with H3K27me3 pre-mark likely lost H3K27me3, but acquire de novo DNA methylation instead (Figure 8, subgroup 14). Those without H3K27me3 pre-mark markedly correlated with already methylated genes; these genes were generally negative for all three histone modifications (Figure 8, subgroup 13).

Figure 3: Characteristics of methylation-sensitive and methylation-resistant genes. (A) The methylation-sensitive genes in MKN7_EB_A1 cells and those in MKN7_EB_B6 cells overlapped well. (B) The expression of methylation-sensitive genes in MKN7_WT, MKN7_EB_A1, and MKN7_EB_B6 cells was compared and is shown by fold expression change against WT. Methylation-sensitive genes were repressed in both MKN7_EB_A1 and MKN7_EB_B6 cells. (C) GO terms enriched in the overlapped methylation-sensitive genes were analyzed, and terms related to differentiation, cell adhesion, development, and regulation of cell proliferation were significant. (D) The methylation-resistant genes in MKN7_EB_A1 and MKN7_EB_B6 cells overlapped well. (E) The expression of methylation-resistant genes in WT, MKN7_EB_A1, and MKN7_EB_B6 cells was compared and is shown by fold expression change against WT. Expression levels of methylation-resistant genes in MKN7_WT cells were maintained in both MKN7_EB_A1 and MKN7_EB_B6 cells. (F) GO terms enriched in the overlapped methylation-resistant genes were analyzed, and terms related to DNA repair were significant.
Figure 4: H3K4me3 alteration during EBV infection and its correlation to gene expression and DNA methylation induction. (A) H3K4me3 signals at promoters in MKN7_WT and MKN7_EB_A1 cells. Compared with H3K4me3 signal around the TSS in MKN7_WT cells, that in MKN7_EB_A1 was decreased in 2,363 genes (decreased), remained positive in 10,284 genes (positive), increased in 913 genes (increased), and remained negative in 7,039 genes (negative). The heatmap shows the distribution of H3K4me3 signals of all the genes included in each subset within ± 2 kb from the TSS (left). The average profile plot shows the distribution of averaged H3K4me3 signal within ± 4 kb from the TSS (middle). Distribution of ChIP-seq reads of a representative gene is shown (right). (B) The gene expression was compared between MKN7_WT and MKN7_EB_A1 cells, and presented as fold expression change against WT. Gene expression was markedly downregulated in “decreased” genes (top), retained in “positive” genes (second), markedly upregulated in “increased” genes (third), and retained in “negative” genes (bottom). (C) As for DNA methylation alteration, β-score of each gene at the probe nearest to the TSS was compared between MKN7_WT and MKN7_EB_A1 cells, and presented as the average and SE. The methylation level was markedly elevated in “decreased” genes (top), retained at unmethylated state in “positive” (second) and “increased” genes (third), and retained at methylated state in “negative” genes (bottom). (D) GO term enrichment in genes with increased and decreased H3K4me3 signal. Genes related to differentiation and morphogenesis were significantly enriched in genes with decreased H3K4me3 signal, whereas genes related to cell cycle were significantly enriched in genes with increased H3K4me3 signal.
Figure 5: H3K27ac alteration during EBV infection, and its correlation to gene expression and DNA methylation induction. (A) H3K27ac signals at promoters in MKN7_WT and MKN7_EB_A1 cells. Compared with H3K27ac signal around the TSS in MKN7_WT cells, that in MKN7_EB_A1 cells was decreased in 2,350 genes (decreased), remained positive in 5,582 genes (positive), increased in 1,817 genes (increased), and remained negative in 10,850 genes (negative). The heatmap shows the distribution of H3K27ac signals of all the genes included in each subset within ±2 kb from the TSS (left). The average profile plot shows the distribution of averaged H3K27ac signal within ±4 kb from the TSS (middle). Distribution of ChIP-seq reads of a representative gene is shown (right). (B) The gene expression was compared between MKN7_WT and MKN7_EB_A1 cells and presented as fold expression change against WT. Gene expression was markedly downregulated in “decreased” genes (top), retained in “positive” genes (second), markedly upregulated in “increased” genes (third), and retained in “negative” genes (bottom). (C) As for DNA methylation alteration, β-score of each gene at the probe nearest to the TSS was compared between MKN7_WT and MKN7_EB_A1 cells, and presented as the average and SE. The methylation level was markedly elevated in “decreased” genes (top), retained at unmethylated state in “positive” (second) and “increased” genes (third), and retained at high levels in “negative” genes (bottom). (D) GO term enrichment in genes with increased and decreased H3K27ac signal. Genes related to cell cycle and DNA repair were significantly enriched in genes with increased H3K27ac signal, whereas genes related to apoptosis and cell death were significantly enriched in genes with decreased H3K27ac signal.
Figure 6: H3K27me3 alteration during EBV infection and its correlation to gene expression and DNA methylation induction. (A) H3K27me3 signals at promoters in MKKN7_WT and MKKN7_EB_A1 cells. Compared with H3K27me3 signal around the TSS in MKKN7_WT cells, that in MKKN7_EB_A1 cells was decreased in 2,262 genes (decreased), remained positive in as few as 84 genes (positive), increased in as few as 31 genes (increased), and remained negative in 18,222 genes (negative). The heatmap shows the distribution of H3K27me3 signals of all the genes included in each subset within ± 2 kb from the TSS (left). The average profile plot shows the distribution of averaged H3K27me3 signal within ± 4 kb from the TSS (middle). Distribution of ChIP-seq reads of a representative gene is shown (right). (B) The gene expression was compared between MKKN7_WT and MKKN7_EB_A1 cells and presented as fold expression change against WT. Gene expression was not upregulated in “decreased” genes despite of loss of repressive H3K27me3 histone mark (top) and was retained in “negative” genes (bottom). (C) As for DNA methylation alteration, β-score of each gene at the probe nearest to the TSS was compared between MKKN7_WT and MKKN7_EB_A1 cells and presented as the average and SE. The methylation level was markedly elevated to methylated state in “decreased” genes during the loss of repressive H3K27me3 histone mark (top). (D) GO term enrichment in genes with decreased H3K27me3 signal. Genes related to differentiation and morphogenesis were significantly enriched in genes with increased H3K27me3 signal.
DISCUSSION

We previously reported that EBV-positive gastric cancer is distinguished by extensive DNA hypermethylation of gene promoters, and EBV itself is the cause of de novo DNA methylation [4]. Some genes, named common marker genes, are commonly methylated in low-methylation, high-methylation/MSI+, and extensively high-methylation/EBV+ gastric cancer cases; MKN7, a low-methylation gastric cancer cell line, shows methylation in these genes, while GES1, a normal gastric epithelial cell immortalized with SV40, is not methylated in these genes. EBV infection was shown to cause genome-wide methylation and thus induce EBV-CIMP in both MKN7 and GES1 [4, 7]. Here, we show that DNA methylation-induced genes in EBV-infected MKN7 were generally classified into “DNA methylation-sensitive” and “DNA methylation-resistant” depending on methylation status around the TSS, that histone modification was also dynamically altered with EBV infection, and that histone modification alteration occurred in a coordinated manner with DNA methylation alteration.

Among the de novo methylated promoter regions, roughly half acquired complete methylation in the whole promoter region (Figure 2A), which led to repression of the downstream genes (Figure 3B). These DNA methylation-sensitive genes included known anti-oncogenic genes acting in cell adhesion or apoptosis (Figure 3C). The inactivation of \textit{CDH1} was observed with TSS methylation, and this may contribute to the characteristic feature of EBV-positive gastric cancer that they often appear as diffuse-type gastric cancer [16, 17]. \textit{CDKN2A} was also...

<table>
<thead>
<tr>
<th>K4me3</th>
<th>K27ac</th>
<th>K27me3</th>
</tr>
</thead>
<tbody>
<tr>
<td>Decreased</td>
<td>WT E8</td>
<td>n=2,363</td>
</tr>
<tr>
<td>Positive</td>
<td>n=10,284</td>
<td>924,2,805</td>
</tr>
<tr>
<td>Increased</td>
<td>n=913</td>
<td>37,208</td>
</tr>
<tr>
<td>Negative</td>
<td>n=7,039</td>
<td>184,904</td>
</tr>
<tr>
<td></td>
<td>n=2,350</td>
<td>570,638</td>
</tr>
<tr>
<td></td>
<td>n=5,582</td>
<td>269,1,402</td>
</tr>
<tr>
<td></td>
<td>n=84</td>
<td>11,6</td>
</tr>
<tr>
<td></td>
<td>n=1,817</td>
<td>68,404</td>
</tr>
<tr>
<td></td>
<td>n=31</td>
<td>3,3</td>
</tr>
<tr>
<td></td>
<td>n=10,850</td>
<td>1,193,2,009</td>
</tr>
<tr>
<td></td>
<td>n=18,222</td>
<td>2,157,3,459</td>
</tr>
</tbody>
</table>

### Figure 7: Ratio of methylation-sensitive and resistant genes and its association with histone modification alteration patterns.

As for each of 12 histone modification alteration patterns (See Figures 4, 5, and 6), DNA methylation levels were assessed by calculating the average of \( \beta \)-scores based on the distance from the TSS. The Pie chart shows the number of the methylation-sensitive genes (black) and methylation-resistant genes (white).
Figure 8: Integrated analysis of all the histone marks and DNA methylation. Genes decreasing H3K4me3 signal generally related to DNA methylation-sensitive genes and gene repression, but were further classified into four subgroups (1-4). Whereas genes also losing H3K27ac (2) and retaining H3K27ac-negative status (3) strongly associated with DNA methylation-sensitive genes and gene repression, genes retaining H3K27ac-positive status (1) rather associated with DNA methylation-resistant genes and were thus less downregulated. Genes retaining H3K4me3-negative status generally related to DNA methylation-resistant genes, but were further classified into five subgroups (5-9). Whereas genes without H3K27me3 mark (5-8) strongly associated with DNA methylation-resistant genes regardless H3K27ac status, genes losing H3K27me3 mark (9) strongly associated with DNA methylation-sensitive genes. Gene expression alteration in subgroups 5-8 depended on H3K27ac alteration patterns. Genes with increased H3K4me3 signal generally related to DNA methylation-resistant genes (10-12). Genes retaining H3K27ac-negative and H3K27me3-negative status (13) strongly associated with originally methylated genes, genes losing H3K27me3 mark associated with DNA methylation-sensitive genes. Regardless of H3K4me3 status, genes losing H3K27me3 mark at promoters (4, 9, 14) associated with DNA methylation-sensitive genes.
a DNA methylation-sensitive gene, being consistent with previous reports that CDKN2A is hypermethylated and silenced in EBV-positive gastric cancer [6].

In contrast, the other half acquired de novo methylation in the surrounding region of the promoters, but protected unmethylated status at the immediate vicinity of the TSS (Figure 2B). These genes maintained gene expression (Figure 3E), and were regarded as DNA methylation-resistant genes. These were significantly enriched with DNA repair genes (Figure 3F), e.g., mismatch repair genes, including MLH1, MSH2, and MSH6. This supports the previous report that EBV-positive gastric cancer does not show MLH1 methylation and is thus microsatellite stable [5, 18]. The existence of DNA methylation-resistant genes and its reproducibility among EBV-infected clones suggest that there might be some mechanisms protecting TSS from de novo DNA methylation pressure of EBV infection [19].

Dynamic alteration was detected in both active and repressive histone marks. Approximately 2,000 to 4,000 genes showed alteration of H3K4me3, H3K27ac, and H3K27me3 levels at promoter regions. The activated genes with increase of H3K27ac were significantly enriched with DNA repair genes (Figure 5D). As similar GO terms were enriched in DNA methylation-resistant genes, genes with gain of H3K27ac significantly correlated with DNA methylation-resistant genes (Figure 7). BIRC5, which negatively regulates apoptosis and was reportedly upregulated by EBV latent protein LMP2 [20], showed increased H3K27ac signal with upregulated expression. In contrast, decrease in H3K27ac was observed in 2,350 genes (Figure 5A), which significantly included genes related to apoptosis and cell death (Figure 5D). These genes were significantly repressed, and strongly correlated with DNA methylation-sensitive genes (Figure 7). However, 161 among the 2,350 genes did not acquire de novo DNA methylation, and were downregulated during EBV infection. For example, genes related to significantly enriched GO terms of cell adhesion and apoptosis, included COL4A3 [21, 22], PHLP1 [23], and TIA1 [24], which were reportedly downregulated in cancer. These genes did not acquire de novo DNA methylation, but showed decrease of H3K27ac and downregulation of gene expression. It was suggested that histone modification alteration induced by EBV infection might regulate critical tumor-related genes without necessarily accompanying DNA methylation alteration. While EBV reportedly established its latency by changing histone modifications of viral genome as well as DNA methylation of viral genome [25, 26], EBV might also alter histone modification of host genome and contribute to tumorigenesis.

Genes with decrease of H3K4me3 or H3K27me3 signals significantly included differentiation-related genes (Figure 4D and 6D) and were accompanied with de novo DNA methylation with gene repression (Figures 4B and 6B). In contrast, the methylation-resistant genes and the unmethylated genes were marked with both or one of active histone marks (Figure 8). It is reported that active promoters with H3K4me3 are protected from de novo methylation by occupying transcription factors and histone methyltransferases at promoter regions. De novo methyltransferases, DNMT3A, 3B and 3L, are known to have ADD domain that recognize H3K4me0 [27, 28]. The polycomb repressive complex, for example, establishes and maintains H3K27me3 marks, which were suggested to be pre-marks for de novo DNA hypermethylation [29–32]. The switch of epigenetic repression marks from H3K27me3 to DNA methylation was observed in H3K4me3(+) genes as well as H3K4me3(−) genes in this study, suggesting that H3K27me3 pre-marks might be strong predisposed promoters rather than H3K4me0 status that recruits DNA methyltransferases [27–32].

While DNA methylation induction was found to be preferentially accompanied with loss of histone active marks or repressive mark, it is yet to be investigated which epigenetic alteration occurs first and could causally induce other alterations. The efficiency of in vitro EBV infection into gastric cells is as low as < 0.001% [7], so that time series analysis of histone modification alteration is difficult at early time points. Improvement of infection efficiency and ChIP technique using smaller number of cells is now underway [Okabe et al. unpublished data], which will enable us elucidate the order of epigenetic alterations in host genome, as well as viral genome [7]. The analysis using non-neoplastic gastric epithelial cell GES1 should also help us understand the whole mark of epigenomic alteration in gastric epithelial cell induced by EBV infection.

In conclusion, histone modification altered coordinately with DNA methylation alteration during EBV infection. H3K27me3 pre-marks and decrease of active histone marks were strongly correlated with DNA methylation-sensitive genes.

**MATERIALS AND METHODS**

**Cells**

We obtained MKN7, a low DNA methylation phenotype gastric cancer cell line, from Riken BioResource Center (Ibaraki, Japan). We infected recombinant EBV into MKN7 cells (MKN7_WT) using the Akata system [33] to establish two independent EBV-infected MKN7 clones (MKN7_EB_A1 and MKN7_EB_B6) in our previous study [4]. MKN7_WT and EBV-infected MKN7 clones were cultured at 37°C in a 5% CO2 incubator in RPMI 1640 medium (Wako, Tokyo, Japan) with 10% fetal bovine serum (HyClone SH30910.03; GE Healthcare, Chicago, IL, USA), streptomycin sulfate (100 mg/L) (Sigma-Aldrich, St. Louis, MO, USA), and penicillin G sodium (100 mg/L) (Sigma-Aldrich). EBV-infected MKN7 clones were selected by 300 g/mL G418 (Life technologies, Carlsbad, CA, USA). These clones were cultured for 18 weeks after EBV infection.
for selection and cloning. EBER in situ hybridization was performed, as previously described [4], to confirm existence of EBV in each cancer cell (Supplementary Figure 2). These cells underwent the subsequent analyses of DNA methylation and histone modification. MKN7 cells cultured for the same period of 18 weeks, named MKN7_EB(−) cells, also underwent histone modification analyses, while we previously reported that MKN7_EB(−) cells did not acquire aberrant DNA methylation or gene silencing [4]. MKN7 cells were transfected with a vacant vector pcDNA3.1 (Invitrogen, Carlsbad, CA, USA) with neomycin resistant gene, and it was also already confirmed that long exposure to neomycin for selection did not cause aberrant DNA methylation or gene silencing [7].

**Infinium assays**

High-resolution methylation analysis was conducted on the Illumina Infinium HumanMethylation450 BeadChip array. Illumina Infinium HumanMethylation450 BeadChip array contains over 485,000 CpG sites and covers 99% of RefSeq genes. There were 177,656 CpG sites within 1,000 bp (from − 500 bp to + 500 bp) around the TSS of 25,593 RefSeq genes on autosomal chromosomes, with an average of 6.8 CpG sites and a median of 7.0 CpG sites covered in Infinium 450K.

Five hundred nanograms of DNA from cultured cells were extracted by using QIAamp DNA Mini Kit (Qiagen, Hilden, Germany) and underwent bisulfite conversion using EZ DNA Methylation Kit (Zymo Research, Irvine, CA, USA) according to the manufacturer’s protocol. Bisulfite-converted DNA was subsequently amplified, fragmented, and hybridized to the BeadChip. After single base extension and staining, the BeadChip was scanned by using the Illumina HiScan SQ scanner following the manufacturer’s instructions. Infinium data were submitted to the GEO DataSets, and the accession number is GSE89269.

For each CpG site, the methylation rate was calculated as the value ranging from 0.00 (completely unmethylated) to 1.00 (completely methylated), based on the fluorescence signal intensity of the probe. When β-score was ≤ 0.20, the CpG site was considered unmethylated (U) and the probe was defined as a “U” probe. When β-score was ≥ 0.40, the CpG site was considered methylated (M), and the probe was defined as an “M” probe. When the methylation status was altered from U to M, or from M to U, the probe was called a “M to U” probe or a “M to U” probe, respectively. When the U probe still showed unmethylated state after EBV infection, the probe was called a “U to U” probe. Similarly, when the M probe still showed methylated state after EBV infection, the probe was called “M to M” probe.

**Expression array analysis**

Gene expression in MKN7_WT, MKN7_EB_A1, and MKN7_EB_B6 cells was analyzed by Affymetrix GeneChip Human Genome U133 plus 2.0 oligonucleotide arrays (Affimetrrix, Fremont, CA, USA) as previously described [4]. Data were collected and analyzed by GeneChip Scanner 3000 (Affimetrrix). The GeneChip data were analyzed using the Affymetrix GeneChip Operating Software v1.3 by MAS5 algorithms, to obtain signal value (GeneChip score) for each probe set. For global normalization, the average signal in an array was made equal to 100. Expression array data were submitted to the GEO DataSets, and the accession number is GSE31789.

**GO analysis**

Gene annotation enrichment analysis was performed using the Functional Annotation tool at DAVID 6.7 Bioinformatics Resources (https://david.ncifcrf.gov/home.jsp). The GO terms were filtered to select terms from the biological process (BP) subontology.

**Chromatin immunoprecipitation (ChIP)**

ChIP was performed as previously described [34]. MKN7_WT and MKN7_EB_A1 cells were cross-linked with 1% formaldehyde for 10 min, followed by quenching with 125 mM glycine for 5 min. DNA fragmentation was performed by using UD 201 (Tomy, Tokyo, Japan) with the following settings: output level 2, 50% duty, 15 sec, 5 cycles, on floating ice to obtain 200–500 bp DNA fragments. An aliquot of chromatin was removed as input and the remainder of the chromatin was used for ChIP analysis using anti-H3K4me3 (ab8580, Abcam, Cambridge, MA, USA, 2 µL per ChIP), anti-H3K27ac (CMA309, MBL, Nagoya, Japan, 2 µL per ChIP), and anti-H3K27me3 (07–142, Millipore, Darmstadt, Germany, 2 µL per ChIP) antibodies.

The fragmented samples were incubated with antibodies bound to protein A-sepharose beads (GE Healthcare) or Dynabeads (Life technologies) at 4°C overnight. The beads were then washed eight times and eluted with elution buffer. The eluates and input were treated with pronase at 42°C for 2 h and then underwent incubation at 65°C overnight to reverse the cross-linking. The immunoprecipitated and input DNA were purified by 2 extractions with phenol:chloroform, followed by ethanol precipitation. Purified DNA samples were resuspended in 20 µL of Tris-HCl (pH 8.0). The concentration of the DNA sample was measured with a Qubit fluorometer (Life Technologies).

**ChIP-seq analysis**

The ChIP DNA was used to prepare library samples following the manufacturer’s instructions (Illumina). Deep sequencing was performed on the Solexa Genome Analyzer II to obtain 36-bp single reads. Sequencing tags were mapped to UCSC build hg19 (GRCh37) assembly of the human genome using ELAND. Window sizes...
of 300 bp for H3K4me3 and H3K27ac and 500 bp for H3K27me3 were used to calculate the number of mapped reads per million reads (RPM) via the Galaxy workflow system [35]. The highest RPM within ± 1,000 bp around the TSS was regarded as the represented RPM.

Then, represented RPM of H3K4me3 and H3K27ac were normalized by per-sample scale factors based on the previous report [36]. Briefly, a set of genes were selected which expressed high level (array score > 200) and had low variance (variance in $-0.1 \leq \log_2$ fold change $\leq 0.1$) in both MKN7 and MKN7_EB_A1 cells. A total of 1,023 genes meet the criteria and the average of represented RPM of those genes was considered to reflect the variability in read count due to technical differences in sample preparation. The normalized RPM were visualized using IGV browser and ngs.plot [37].

A histone mark was recognized as increased when the number of represented RPM of MKN7_WT cells was greater than the cut-off RPM and MKN7_EB_A1/MKN7_WT was $\geq 1.5$. A histone mark was recognized as lost when the number of represented RPM of MKN7_EB_A1 cells was greater than the cut-off RPM and MKN7_EB_A1/MKN7_WT < 2/3. The cut-off RPM was 2.0 for H3K4me3 and H3K27ac, and 1.0 for H3K27me3.

**Statistical analysis**

Gene expression levels by GeneChip score and methylation levels by $\beta$-score are presented as mean ± SE. Student’s $t$-test was conducted for the comparison of two groups. Correlation between DNA methylation status and histone modification status was analyzed by $\chi^2$ test. R program (www.r-project.org/) was used in these analyses.

**Abbreviations**


**Authors’ contributions**

SF, KM, OA performed the experiments. SF, RY, SY, MF analyzed the data. SF, KM, OA, HA, MF, AK interpreted the data. SF, OA, AK wrote the manuscript.

**ACKNOWLEDGMENTS**

The authors thank Kyoko Fujinaka, Kaori Shina, Hiroko Meguro, and Hiromi Tanaka for technical assistance, and Editage (www.editage.jp) for English language editing.

**CONFLICTS OF INTEREST**

All authors have no potential conflicts of interest to disclose.

**FUNDING**

This study was supported by AMED-CREST program 16gm0510010h0005 (AK), Practical Research for Innovative Cancer Control 17ck010623h0001 (AK), and P-CREATE 16cm0106510h0001 (AK) from Japan Agency for Medical Research and Development, Grant-in-Aid for Scientific Research 16H05412 (AK) from Japan Society for the Promotion of Science, a grant from Terumo Foundation for Life Science and Arts, and a grant from Suzuken Memorial Foundation.

**REFERENCES**


