

A study on DNA methylation modifying natural compounds identified EGCG for induction of IFI16 gene expression related to the innate immune response in cancer cells

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Abstract. Innate immune sensor IFN-induced protein 16 (IFI16) exhibits anti-inflammatory effects via IFN β and IFN-stimulated gene (ISG)15 induction in cancer cells. Epigallocatechin gallate (EGCG) is a potent natural DNA methyltransferase inhibitor (DNMTi). Previous studies revealed that conventional DNMTis, such as 5-azacytidine (5-aza-dc), induce IFI16 expression and EGCG decreases DNMT mRNA expression and global methylation (5mC) level via promoter demethylation of tumor suppressor genes in cancer cell lines. To the best of our knowledge, however, EGCG-mediated IFI16 promoter methylation status has been overlooked. Here, initial screening was performed to determine IFI16 expression and its correlation with DNMTs in cancer cell lines from various databases. Following treatment of breast cancer cell lines with 5-aza-dc, vitamin C and EGCG, expression levels of IFI16 and its downstream transcription targets IFN β 1 and ISG15 were assessed using RT-qPCR, and the 5mC level was assessed using ELISA. *In silico* molecular docking simulation was performed for all DNMTs to predict the mode of ligands binding with proteins. Finally, promoter methylation level in IFI16 gene was assessed following EGCG treatment. EGCG treatment induced IFI16 expression, interacted with certain amino acids residues in DNMT proteins and decreased 5mC level and promoter methylation of IFI16. The present results may provide a basis for targeting IFI16 expression as a therapeutic option in breast cancer cell lines.

Introduction

IFN-induced protein 16 (IFI16) is a potent innate immune response evoker that amplifies DNA damage-mediated innate

immune response in inflammatory disease (1-3). It also contributes to cell senescence (3). However, previous studies revealed the IFI16, which belongs to the pyrin and HIN domain family, serves a key role as an innate immune sensor and sensitizes foreign or intracellular double-stranded (ds)DNA (4,5). Although IFI16 predominantly sensitizes cytosolic DNA, another study showed that it can also translocate into nucleus to detect dsDNA (5,6).

The alternate splicing of mRNA yields three structural isoforms when IFI16 is transcribed: A, B and C (7,8). IFI16 gene comprises 200-amino acid repeats known as HIN-200 domain, which is classified as HIN-200-A or HIN-200-B. Both A and B domains are separated by the serine-threonine-proline-rich spacer region (9-11). Moreover, the basic region of 1-159 residues binds dsDNA at the N-terminus of the HIN-200-A subunit as well as the N-terminus of the HIN-200-A subunit to form a peptide sequence of amino acids with PYRIN domain that mediates the signal for nuclear localization of IFI16 from cytosol to nucleus (11).

IFI16 exerts its key innate immune responses in diverse manners. For example, upon recognition of cytosolic dsDNA by cyclic GMP-AMP synthase, it activates the downstream STING/TANK-binding kinase 1/IFN regulatory factor 3 pathway to induce transcription of IFN β (12). Moreover, IFI16 binds with dsDNA and induces IFN β transcription, which, in turn, induces IFN-stimulated gene 15 (ISG15) transcription via the common STING axis (13). Another study showed that IFI16 induces IFN β expression while knockdown of IFI16 abrogates the IFN β response (14). Transcriptional cross-talk between IFI16 and IFN β has been proposed in a previous study (15). In addition to acting as a dsDNA sensor, IFI16 also serves as an ISG (16-18).

Previous studies have reported downregulated expression of IFI16 in cancer cells. For example, a study on the cytoplasm of prostate cancer cell lines demonstrated that IFI16 gene was not expressed or expressed in variant forms; the study also showed that functional overexpression of IFI16 halted colony formation (19). Another study showed that in certain cancer cells, loss of IFI16 expression facilitates cell survival even in the presence of low glucose (20). Moreover, IFI16 acts as a tumor suppressor gene, inhibits proliferation of hepatocellular carcinoma and triggers apoptosis (21). However, controversial

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findings have shown an oncogenic role of *IFI16* in certain types of cancer cell, such as oral squamous and renal cell carcinoma and pancreatic adenocarcinoma (22-24).

Natural compounds, such as like vitamin C and green tea extract polyphenol Epigallocatechin gallate (EGCG), exert multiple effects on lipid antioxidation (25) and transcriptional machinery to modulate gene expression by targeting signaling pathways or transcription factors or epigenetically modulating gene expression (26). Moreover, EGCG has been shown to block DNA methyltransferase (DNMT) activity and reactivates methylation-mediated silenced genes, such as p16, in cancer cell lines (27). Additionally, EGCG and vitamin C increase DNA demethylation by inhibiting DNMT1/DNMT3b and modulating ten-eleven translocation enzymes, respectively (28). EGCG exhibits modulatory effects on innate and adaptive immune response stimuli in murine and human models (29,30). In addition, vitamin C also showed innate immune response booster through viral mimicry (pseudo-infection) in breast, colon, leukemia and hepatocellular cancer cell lines (31).

To the best of our knowledge, few studies have been conducted on epigenetic drug-mediated *IFI16* expression. For example, *IFI16* is acetylated in herpes simplex virus-1 (HSV-1) (32). Acetylation and deacetylation regulate nuclear and cytoplasmic localization of *IFI16* (32). Another study showed that histone deacetylase inhibitor trichostatin A and CGK1026 induce *IFI16* gene expression in prostate cancer cells (33). In addition, at a single-cell level, low dose treatment with DNA demethylating agents induces *IFI16* gene expression (34). On the other hand, EGCG is a promoter demethylation-mediated epigenetic drug that induces expression of tumor suppressor genes in colon cancer cells (35). To the best of our knowledge, no previous study has investigated whether EGCG induces *IFI16* expression by decreasing methylation of the *IFI16* promoter. Therefore, the present study aimed to investigate this hypothesis.

Materials and methods

Preliminary screening of mRNA correlation, expression and immune responsiveness of targeted protein. *IFI16* and DNMTs gene (Database of Genotypes and Phenotypes accession no. phs000424.v8.p2) expression was screened out using The Cancer Genome Atlas (TCGA) datasets through the UALCAN web server (ualcan.path.uab.edu/). The correlation between *IFI16* and DNMTs (accession no. ENSG00000163565.17) were screened from the web server and validated by data retrieved from xenabrowser.net/. The immune responsiveness study of *IFI16* was assessed by TIMER2.0 web server (timer.cistrome.org/). Finally, cell lines were screened from the Human Protein Atlas database where the targeted protein *IFI16* gene expression was absent or downregulated. Based on the mRNA expression, SHSY5Y (neuroblastoma), T47D (ductal breast carcinoma), HepG2 (hepatocellular carcinoma), MCF-7 (breast cancer cell) and HeLa (cervical cancer cell) cell lines were identified.

Cell culture and treatment. MCF-7 cell line was procured from American Type Culture Collection and sub-cultured in DMEM supplemented with 10% FBS (both UFC Biotech) and

1% penicillin and incubated at 37°C in a 5% CO₂ incubator. Upon 60-80% confluence, cells were trypsinized, seeded (3,000 cells/well) in 6-well plates and incubated at 37°C overnight to ensure that cells were healthy without any contamination. Based on previous studies, half maximal inhibitory concentration doses for EGCG were added to MCF-7 and T47D cell lines at 40 and 20 μM for 48 h, respectively and incubated at 37°C (36,37). At the same time, 5-azacytadine (5-aza-dc) and vitamin C were added at 60 and 240 μM, respectively, and incubated at 37°C for 48 h.

cDNA synthesis and quantitative (q)PCR. Total RNA was extracted using PureLink™ RNA Mini kit (cat. no. 1944999; Thermo Fisher Scientific, Inc.). A total of 100 ng/μl RNA from untreated and treated MCF-7 and T47D cell lines was transcribed into cDNA using High Capacity cDNA Synthesis kit (cat. no. 00656567; Applied Biosystems; Thermo Fisher Scientific, Inc.) according to the manufacturer's protocol. *IFI16*, *IFNβ*, *ISG15* primer sequences were used as previously described (31,38,39) (Table I). DNMTs primers were designed using the UCSC genome browser (genome.ucsc.edu/; Table I). Reverse transcription (RT)-qPCR was performed using PowerUp SYBR Green Master Mix (cat. no. 1805029; Applied Biosystems; Thermo Fisher Scientific, Inc.). Thermocycling conditions were as follows: 50°C for 2 min, 95°C for 2 min, 95°C for 15 sec and 60°C for 1 min. RT-qPCR was performed using ABI 7300 Prism. The numbers of transcripts were normalized to RPLP0 (forward Primer; 5'ATGTGGGCTTTG TGTTCACC3' and Reverse Primer; 5'TCCAGTCTTGAT CAGCTGCA3') and calculated via the 2^{-ΔΔC_q} method (40).

Extraction of genomic DNA. Genomic DNA was extracted from untreated and treated MCF-7 and T47D cell lines using a DNAbler kit (havensci.com/; cat. no. DE95050). A total of 200 μl digestion buffer was added per sample followed by 20 μl Proteinase K and RNase A. The sample was vortexed and centrifuged at 1,000 x g for 10 min in 25°C followed by 5 min incubation in a heat block (55-60°C). A total of 200 μl lysis buffer was added, vortexed and centrifuged at 10,000 x g for 5 min in 25°C. Then, 99% ethanol was added, followed by short vortex and centrifugation at 10,000 x g for 15 sec in 25°C. The ethanol-lysis buffer content was transferred into the nuclease-free spin column and centrifuged at 10,000 x g for 1 min in 25°C. Then, 500 μl wash buffer was added according to the manufacturer's instructions and, 50 μl elution buffer was added to the spin column and centrifuged at 8,000 x g for 2 min at 25°C to elute genomic DNA.

Determination of global methylation (5mC) levels. MethylFlash™ 5mC ELISA Easy kit (cat. no. P-1030, EpiGentek) was used to assess 5mC level in MCF-7 and T47D cell lines in untreated and treated conditions. According to manufacturer's protocol, binding solution was added to 200 ng genomic DNA/sample in 8 wells followed by addition of 5 mC antibody and developer and stop solution. Finally, optical density was measured at 450 nm using BioTek ELISA microplate reader.

Protein, ligand retrieval and molecular docking simulation. The crystallographic protein structure of DNMT1, DNMT3a

Table I. Reverse transcription-quantitative PCR primers.

Gene	Forward primer, 5'-3'	Reverse primer, 5'-3'
<i>IFI16</i>	CTCGGAGAGCTCGGACAG	TACCTATGACGACGCTGCTG
<i>ISG15</i>	GCCTCAGCTCTGACACC	CGAACTCATCTTTGCCAGTACA
<i>IFNB1</i>	TCTGGCACAACAGGTAGTAGGC	GAGAAGCACAACAGGAGAGCAA
<i>DNMT1</i>	CAGCAACGGGCAGATGTTTC	CGGAGGGTGCTTTGTAGATG
<i>DNMT3a</i>	CTACGCACCACCTCCACCAG	CAATGTTCCGGCACTTCTGC
<i>DNMT3b</i>	GAGTCCATTGCTGTTGGAACCG	ATGTCCCTTTGTTCGCCAACCT
Methylated <i>IFI16</i>	TTCGAGTAGTTGGGATTATAGGC	TAATACAAAATTAATAACGCGAT
Unmethylated <i>IFI16</i>	TTTTTTGAGTAGTTGGGATTATAGGT	AAATAATACAAAATTAATAACACAAT

IFI, IFN-induced protein; ISG, IFN-stimulated gene; DNMT, DNA methyltransferase.

and DNMT3b (ID nos. 4wxx, 6brr, 6kdl) was retrieved from the Protein Database Bank web database (rcsb.org/). The 2D structure of 5-aza-dc (PubChem ID 9444), vitamin C (PubChem ID 54670067), EGCG (PubChem ID 65064), S-adenosyl methionine (SAM) (PubChem ID 34755) and S-adenosyl homocysteine (SAH) (PubChem ID 439155) were retrieved from the PubChem (<https://pubchem.ncbi.nlm.nih.gov/>) database as sdf format. The 5-aza-dc was selected as a positive control for DNMT proteins. The proteins were preprocessed using Biovia Discovery Studio visualizer (3ds.com/products-services/biovia/products/molecular-modeling-simulation/biovia-discovery-studio/visualization/) version (16.1.0) (41). For molecular docking, PyRX software version 0.8 (<https://pyrx.sourceforge.io/>) (42) was used via integrated Open Babel to optimize ligands. Autodock wizards were used to create maximum grid box dimensions. After running docking via Run Autodock in Vina wizard, the ligand in a complex with the protein was visualized using Biovia Discovery Studio visualizer version (16.1.0) (41).

Bisulfite modification and methylation-specific qPCR. Using EpiJET DNA Bisulfite Conversion kit (cat. no. 00596381; Thermo Fisher Scientific, Inc.), bisulfite conversion of genomic DNA was performed according to the manufacturer's protocol. The bisulfite-modified DNA template was used for qPCR using kit (Hi-Tech Green 2X qPCR Universal Mix, cat. no. GQM-M-001-10, Molecule-On). Whole gene sequences were selected for *IFI16* and submitted to Metprimer online software (urogene.org/methprimer/) v1.1 beta for methylated and unmethylated primer design (Table I). During primer design, CpG islands were selected. Bisulfite-converted DNA was used for qPCR using Methylation specific PCR primers and thermocycling conditions as follows: 50°C for 2 min, 95°C for 2 min, 95°C for 15 sec and 60°C for 1 min. For gel electrophoresis, 1.7% (w/v) agarose gel was prepared in 1X Tris acetate EDTA with ethidium bromide staining. qPCR product was mixed with 6X DNA gel loading dye for separation via electrophoresis. Amplified products were visualized using GelDoc Biorad imaging system.

Statistical analysis. Unpaired t-test was used to compare two groups; >2 groups were compared using one-way ANOVA followed by post hoc Tukey's post hoc test. Data were analyzed using GraphPad Prism software (version 9; GraphPad Software,

Inc.). $P \leq 0.05$ was considered to indicate a statistically significant difference. The data are presented as the mean \pm SD of one independent experiment performed in triplicate.

Results

IFI16 is differentially expressed in different tumor cell lines, inversely correlated with differential DNMT mRNA expression and correlated with immune responsiveness. *IFI16* expression was screened in various tumor and normal cell lines. *IFI16* expression was not found in MCF-7, T4D, SHSY5, HepG2 cell lines but was observed in HeLa (Fig. 1A). DNMT mRNA expression was inversely associated with *IFI16* mRNA expression in most cell lines, except HeLa (Fig. 1A-D). Moreover, TGCA omics data showed that the breast cancer (BRCA) has a significantly lower expression of *IFI16* (Fig. 1E) but higher expression of DNMT1 (Fig. 1F), DNMT3 (Fig. 1G) and DNMT3B (Fig. 1H). An inverse correlation between expression of *IFI16* and DNMTs was observed in the breast cancer cell line however this was not statistically significant (Fig. S1A). The immune responsiveness study of *IFI16* showed a moderate correlation with different immune infiltrating cells such as CD8+/4+ T cells, macrophages, dendritic cells and neutrophils (Fig. S1B). Hence, the screening results indicated *IFI16* had impact on immune infiltrating cells.

DNMTi analog EGCG induces IFI16 and IFN-associated gene expression in breast cancer cell lines. The extracted RNAs from EGCG-treated cell lines were used to investigate *IFI16* expression. The results showed that 40 μ M EGCG induced expression of *IFI16* in the MCF-7 cell line and 20 μ M EGCG treatment induced expression in the T47D cell line (Fig. 2A and B). As *IFI16* acts as an upstream target for transcribing IFN β 1 and ISG15 (43), expression of IFN β 1 and ISG15 was assessed in both cell lines. In MCF-7 cells, 40 μ M treatment induced IFN β 1 expression but did not affect ISG15 gene expression (Fig. 2A). By contrast, 20 μ M EGCG induced IFN β 1 and ISG15 gene expression in the T47D cell line (Fig. 2B). These data suggested that EGCG treatment induced expression of *IFI16* and its downstream targeted genes expression.

Gene expression levels in both cell lines were also assessed following treatment with conventional DNMTis, such as 5-aza-dc and vitamin C. *IFI16*, IFN β 1 and ISG15 gene expression in

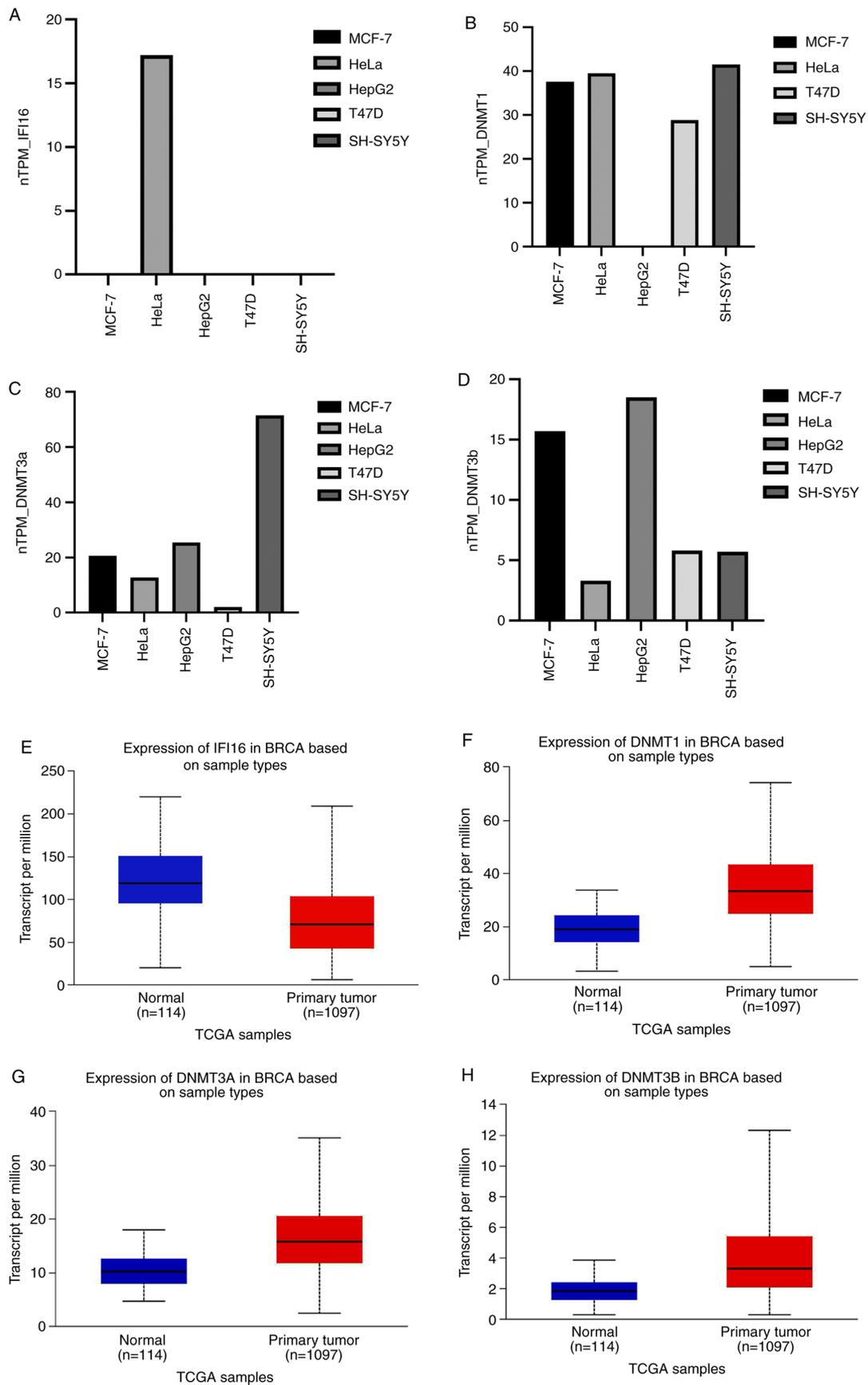


Figure 1. Differential expression of *IFI16* and DNMTs in cancer cell lines and immune response of *IFI16*. (A) Different cancer cell lines showing *IFI16* gene expression. (B) Different cancer cell lines showing (B) DNMT1, (C) DNMT3a and (D) DNMT3b gene expression. (E) TCGA samples data showing *IFI16* gene expression ($P=1.8 \times 10^{-12}$) in breast cancer cell line. (F) TCGA samples data showing DNMT1 gene expression ($P < 10^{-12}$) in breast cancer cell line. (G) TCGA samples data showing DNMT3A gene expression ($P=1.6 \times 10^{-12}$) in breast cancer cell line. and (H) TCGA samples data showing DNMT3B gene expression ($P=1.6 \times 10^{-12}$) in breast cancer cell line. IFI, IFN-induced protein; DNMT, DNA methyltransferase; TPM, transcripts per million; TCGA, The Cancer Genome Atlas.

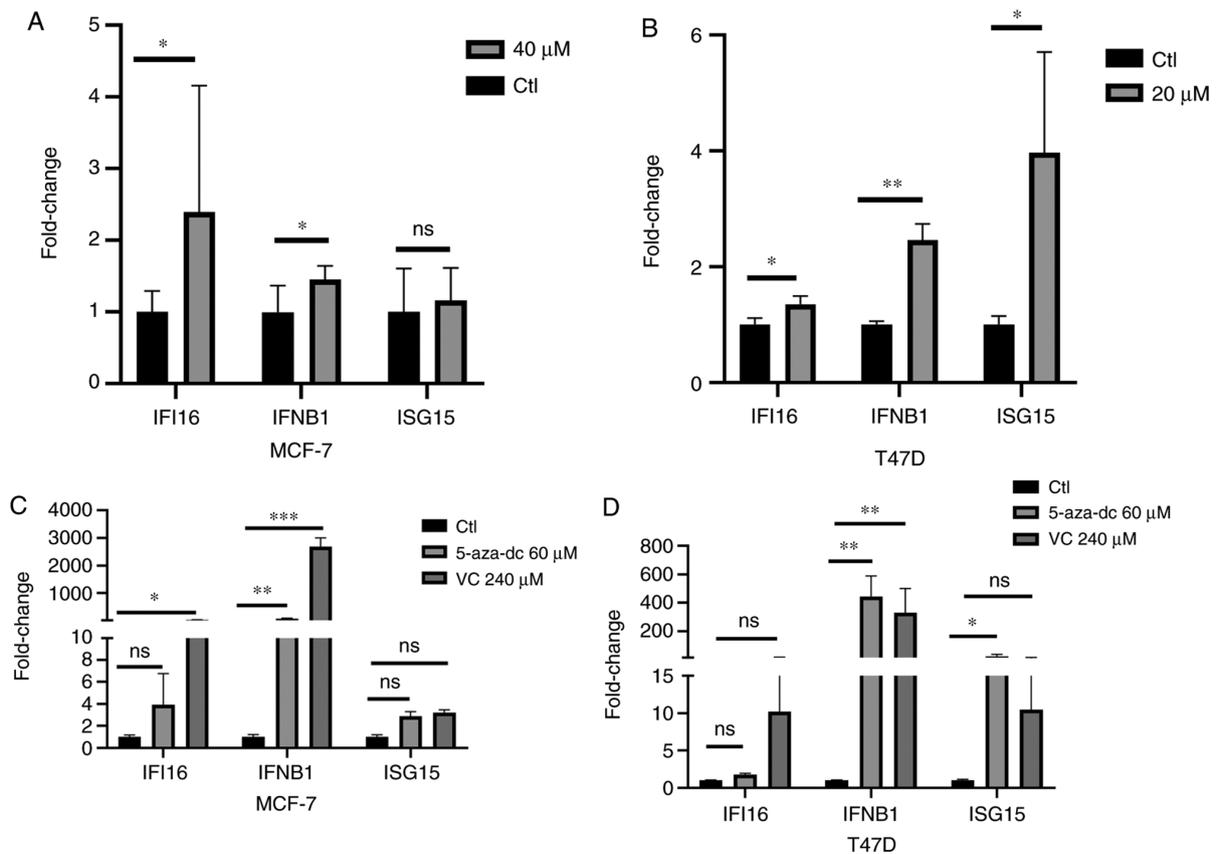


Figure 2. Differential expression of IFI16 and IFN-associated genes following EGCG and DNMTi treatment. IFI16 and IFN-associated gene expression in EGCG-treated (A) MCF-7 and (B) T47D cell lines. IFI16 and IFN-associated gene expression in DNMTi-treated (C) MCF-7 and (D) T47D cell lines. The data are presented as the mean \pm SD. * P <0.05, ** P <0.01 and *** P <0.001. IFI, IFN-induced protein; DNMTi, DNA methyltransferase inhibitor; EGCG, Epigallocatechin gallate; Ctl, control; ISG, IFN-stimulated gene; 5-aza-dc, 5-Azacytadine; VC, vitamin C; ns, not significant.

MCF-7 cell line was induced following treatment with 60 μ M 5-aza-dc or 240 μ M vitamin C (Fig. 2C) although the slight induction of ISG15 gene expression results were not statistically significant. Similar results were also observed in the T47D cell line in which IFI16 gene expression was induced following treatment with 5-aza-dc and vitamin C but this was not statistically significant (Fig. 2D). Moreover, both treatments significantly induced the IFN β 1 gene expression in T47D cell line, while ISG15 gene expression was significantly induced with 5-aza-dc treatment (Fig. 2D). Hence, DNMTi or vitamin C may induce the IFI16 gene and its downstream targeted genes IFN β 1 and ISG15.

DNMTi analog EGCG decreases DNMT expression in breast cancer cell lines. 5-aza-dc and vitamin C are potent DNMTis that modify the epigenome (31,44). Hence, DNMT gene expression was assessed following treatment with EGCG, 5-aza-dc or vitamin C in MCF-7 and T47D cell lines. DNMT1, DNMT3a and DNMT3b gene expression levels were notably decreased following treatment with 60 μ M 5-aza-dc or 240 μ M vitamin C in both cell lines (Fig. 3A and B). Additionally, 40 and 20 μ M EGCG treatment decreased expression of all DNMTs in both MCF-7 and T47D cell lines (Fig. 3A and B). These data indicated that DNMTi, vitamin C and EGCG may affect DNMT gene expression.

Receptor grid box generation and molecular docking simulation: In silico interaction of EGCG with DNMT. Molecular

docking-based computational techniques have been considered as a mechanistic tool (45) for *in silico* drug design. For DNMT1_A, grid box comprised X=205.34, Y=88.76 and Z=145.45 points and spaced dimension was centered on DNMT1_A protein at X=-23.06, Y=47.81 and Z=-23.78. The binding energy of SAM, SAH, vitamin C, EGCG and 5-aza-dc with DNMT1_A was -8.5, -7.6, -5.8, -10.4 and -6.9 kJ/mol, respectively (Fig. 4). For DNMT1_B, grid box comprised X=120.97, Y=89.06 and Z=95.42 points and spaced dimension was centered on the DNMT1_B protein at X=17.63, Y=21.36 and Z=-51.18. The binding energy of SAM, SAH, vitamin C, EGCG and 5-aza-dc with DNMT1_B was -7.7, -7.4, -6.4, -10.0 and -7.1 kJ/mol, respectively (Fig. 4). For DNMT3a, grid box comprised of X=83.85, Y=87.88 and Z=60.33 points and spaced dimension was centered on X=-20.31, Y=-56.03 and Z=20.19. The binding energy of SAM, SAH, vitamin C, EGCG and 5-aza-dc with DNMT3a was -7.2, -6.6, -5.8, -9.5 and -7.2 kJ/mol, respectively (Fig. 4). For DNMT3b, the grid box comprised X=83.85, Y=87.88 and Z=60.33 points and spaced dimension was centered on the DNMT3b protein at X=-20.31, Y=-9.01 and Z=-3.88. The binding energy of SAM, SAH, vitamin C, EGCG and 5-aza-dc with DNMT3b was -8.7, -8.4, -5.8, -9.6 and -7.3 kJ/mol, respectively (Fig. 4).

In silico molecular docking simulation predicted that SAM and EGCG compounds bind in similar positions. SAM was bound at Glu:1168, Phe:1145, Glu:1266, Arg:1310, Thr:1526, Thr:1528 and Asn:1578 residue (Fig. S2), whereas EGCG bound at Met:1169, Pro:1225, Glu:1168, Glu:1266 and Arg:1310 residue (Fig. S2).

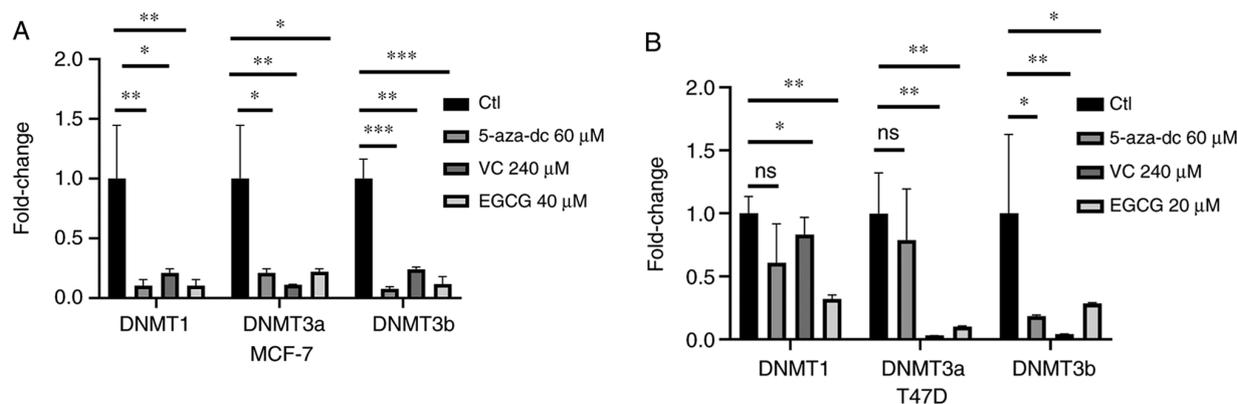


Figure 3. Differential expression of DNMTs following EGCG and DNMTi treatment. DNMT expression in EGCG- or DNMTi-treated (A) MCF-7 and (B) T47D cell lines. The data are presented as the mean \pm SD. * P <0.05, ** P <0.01 and *** P <0.001. DNMTi, DNA methyltransferase inhibitor; EGCG, Epigallocatechin gallate; Ctl, control; 5-aza-dc, 5-Azacytidine; VC, vitamin C; ns, not significant.

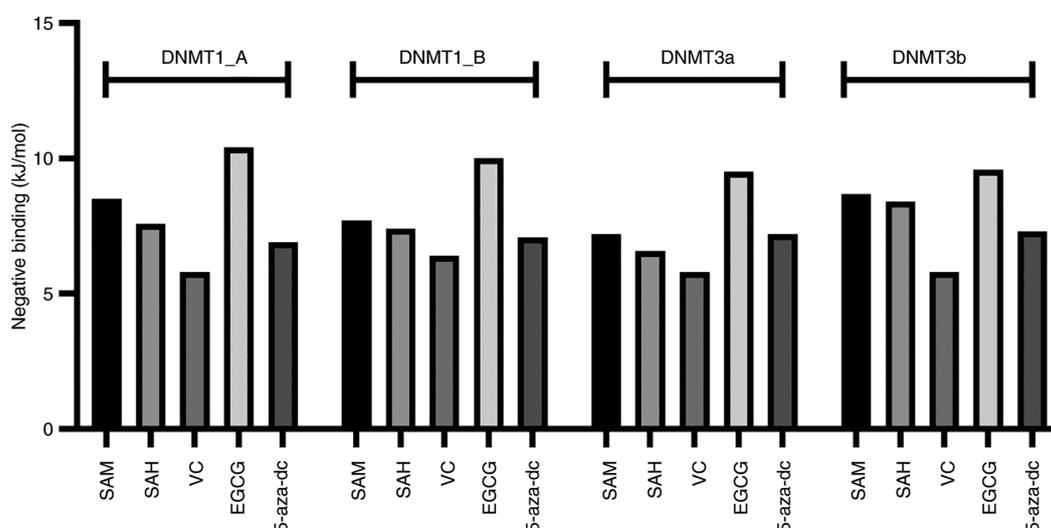


Figure 4. *In silico* docking simulation of different DNMTs with SAM, SAH, 5-aza-dc, VC and EGCG. Docking score as binding energy of all compounds with DNMT1_A chain, DNMT1_B chain, DNMT3a and DNMT3b. DNMT, DNA methyltransferase; 5-aza-dc, 5-Azacytidine; VC, vitamin C; EGCG, Epigallocatechin gallate; SAM, S-adenosyl methionine; SAH, S-adenosyl homocysteine.

Though more conventional H bond interactions were found in the case of SAM, other molecular bonding interactions, such as π -anion and π -alkyl, in addition to four conventional H bonds, were also observed in the EGCG-DNMT1_A complex (Fig. S2). Furthermore, 5-aza-dc was found to interact with DNMT1_A by interacting with Glu:562, Glu:566, Asp:565, Ser:570 and Gln:687 residue with 3 conventional H bonds (Fig. S2). Vitamin C bound at Gln:369, Cys:409, Ser:436 and Glu:494 residue with 4 conventional H bonds (Fig. S2). SAH bound at Glu:562, ASP:565, Glu:566, Pro:574 and Arg:690 with conventional H bond and alkyl interactions (Fig. S2). The predicted site of SAM and EGCG interaction is presented in Fig. 5A.

For DNMT1_B, 5-aza-dc interacted with Phe:1145, Ser:1146, Leu:1151 and Asn:1578 with four conventional H bonds and three C-H bonds. SAM interacted with Phe:1145, Glu:1168, Glu:1266, Arg:1310, Gln:1536, Arg:1574 and Asn:1578 with eight conventional H bonds. EGCG interacted with DNMT1_B at Ser:1146, Glu:1168, Pro:1224/1225, Glu:1266 and Ala:1579 with four conventional H bonds, and one each of π -anion, π -alkyl and C-H bonds (Fig.S3). SAH and vitamin C interacted with DNMT1_B

at Glu:562, Asp:565, Ser:570, Asp:571, Glu:572, Gln:594, Arg:595 Val:658 and Met:1232, Arg:1276, Ser:1277, Val:1344 residues, respectively. Nine conventional H and one alkyl bond were found in the case of SAH, while vitamin C exhibited four conventional H bonds (Fig. S3). The predicted site of 5-aza-dc, SAM and EGCG interaction is presented in Fig. 5B.

Docking study of DNMT3a showed that SAM, SAH, EGCG and vitamin C interacted with DNMT3a as follows: SAM, Gly:707, Cys:710, Arg:792, Arg:891 with four conventional H bonds; SAH, Phe:640, Pro:709, Glu:765, Arg:891 with four conventional H bonds and one each π -sulfur and alkyl bond; EGCG, Phe:640, Gly:642, Val:665, Ser:663, Gly:685, Asp:686, Gly:707, Pro:709, Asn:711, Arg:891 with five H, one C-H, one π - σ , one π -alkyl, one π -cation, two π - π T-shaped bond and vitamin C, Phe:640, Pro:709, Glu:765 and Arg:891 (Fig. S4). 5-aza-dc interacted with Phe:640, Gly:707, Glu:765, Arg:891, Trp:893 residues with five H and one π - σ bond (Fig. S4). The predicted site of SAM SAH, EGCG and vitamin C interaction is presented in Fig. 5C.

Docking study of DNMT3b showed that 5-aza-dc, SAM, SAH and EGCG interacted with DNMT3b as follows:

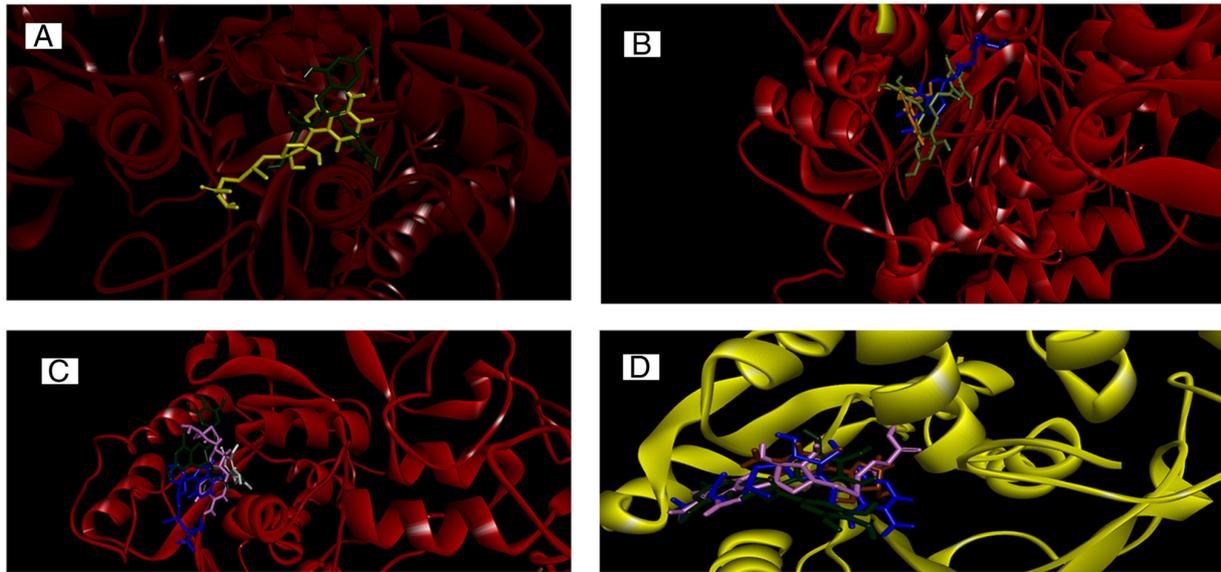


Figure 5. *In silico* interaction of EGCG, SAM, SAH, 5-aza-dc and vitamin C with DNMTs. (A) Predicted binding sites for SAM (yellow) and EGCG (green) in DNMT1_A chain. (B) Predicted binding sites for 5-aza-dc (orange), SAM (blue) and EGCG (green) in DNMT1_B chain. (C) Predicted binding sites for SAM (blue), SAH (pink), EGCG (green) and vitamin C (white) in DNMT3a. (D) Predicted binding sites for SAM (blue), SAH (pink), EGCG (green) and 5-aza-dc (orange) in DNMT3b. DNMT, DNA methyltransferase; 5-aza-dc, 5-Azacytidine; VC, vitamin C; EGCG, Epigallocatechin gallate; SAM, S-adenosyl methionine; SAH, S-adenosyl homocysteine.

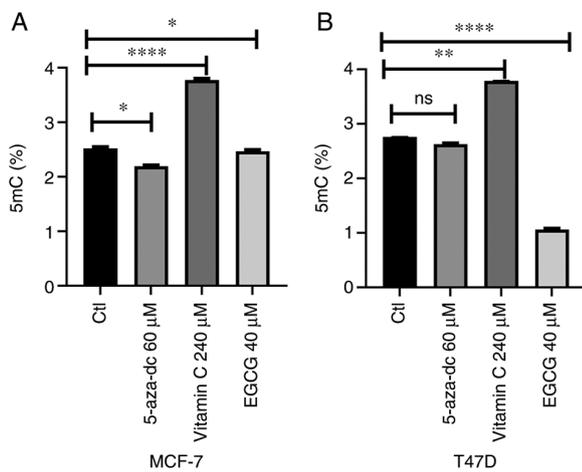


Figure 6. 5mC level following treatment with DNA methyltransferase inhibitors. 5mC following 5-aza-dc, vitamin C and EGCG treatment in (A) MCF-7 and (B) T47D cell lines. The data are presented as the mean \pm SD. * P <0.05, ** P <0.01 and **** P <0.0001. 5mC, global methylation; EGCG, Epigallocatechin gallate; ns, not significant; Ctl, control.

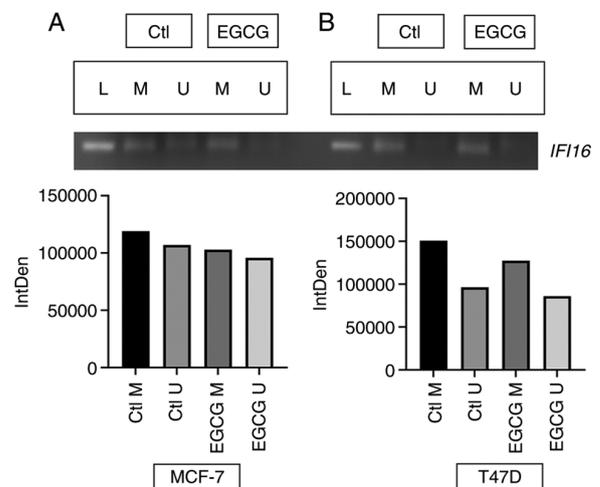


Figure 7. EGCG treatment demethylates *IFI16* promoter. *IFI16* gene promoter demethylation in (A) MCF-7 and (B) T47D cell lines following EGCG treatment. L, ladder; IntDen, Integrated density; Ctl, control; M, methylated; U, unmethylated; IFI, IFN-induced protein; EGCG, Epigallocatechin gallate.

5-aza-dc, Asp:582, Ser:610, Arg:832, Trp:834 with two H and one π - σ bond; SAM, Glu:605, Val:606/628, Ser:649, Glu:697 residues with eight H bonds and one attractive charge interaction; SAH, Phe:581, Gly:583, Thr:586, Glu:605, Cys:607, Asp:627, Val:628, Arg:832, Trp:834 with seven H and one each C-H and alkyl bond and EGCG, Phe:581, Ser:610, Val:606/628, Pro:650, Cys:651, Arg:832 with four H and one each π - σ , π -sulfur, π -cation, π -alkyl and π - π T-shaped interactions (Fig. S5). DNMT3b and vitamin C interacted with Asp:582, Ile:584, Thr:586, Gly:648, Arg:832 with five H and one π -donor H bond (Fig. S5). The predicted site of SAM SAH, 5-aza-dc and EGCG interaction is presented in Fig. 5D.

EGCG, but not vitamin C, decreases 5mC level. As EGCG is a potent DNMTi (46), its effect on 5mC level in both cell lines. In both cell lines, 60 μ M 5-aza-dc decreased 5mC level but this was only significant in MCF-7 cells (Fig. 6A and B). Additionally, 40 μ M EGCG significantly decreased 5mC level in MCF-7 cell line (Fig. 6A). Furthermore, 20 μ M EGCG induced a greater 5mC decrease in the T47D cell line (Fig. 6B). Vitamin C treatment significantly increased 5mC level in both cell lines (Fig. 6A and B). These data suggested that EGCG decreased 5mC level in breast cancer cell lines.

EGCG induces IFI16 gene expression by decreasing DNA methylation. Since EGCG decreased 5mC level in both cell

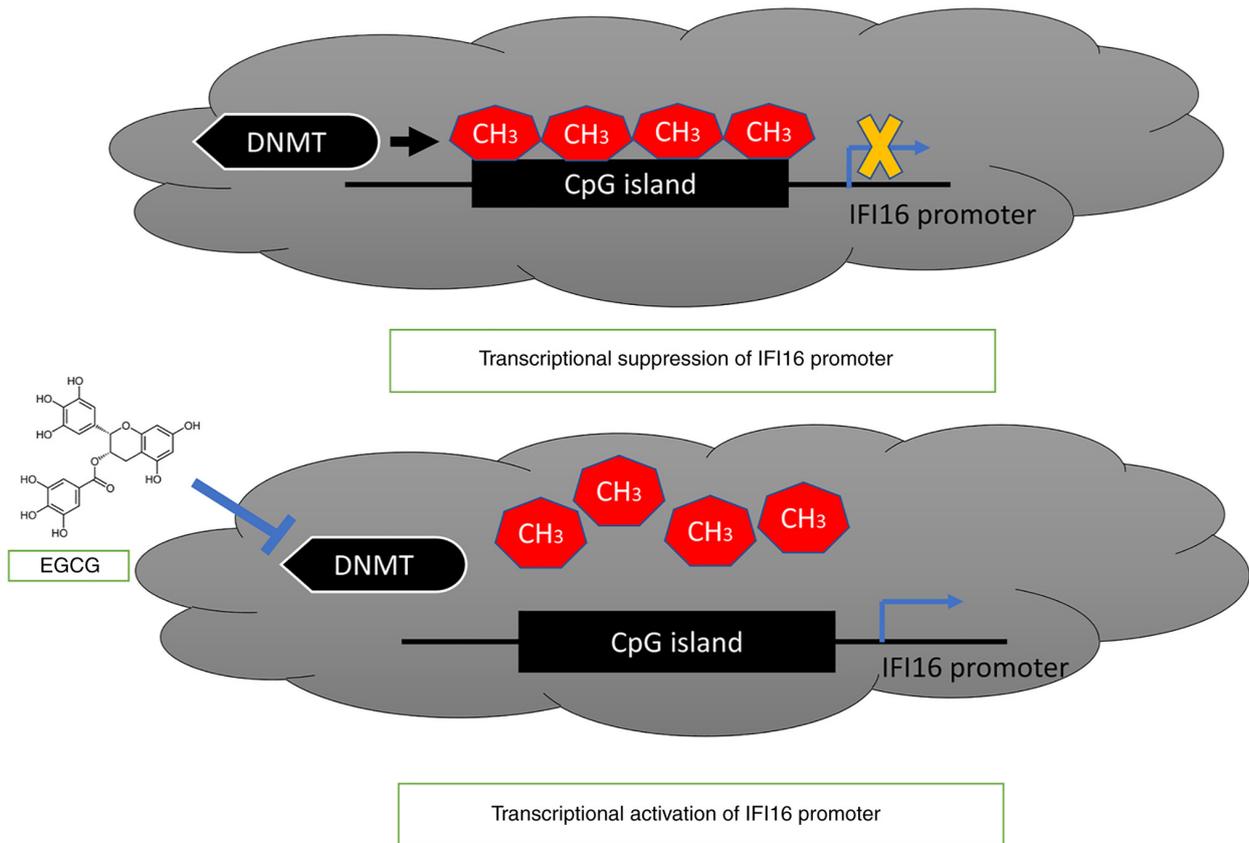


Figure 8. EGCG effect on IFI16 gene promoter. The proposed model showed IFI16 gene promoter is methylated by DNMT-mediated catalysis of methyl group incorporation. EGCG treatment can block the DNMT activity and in turn decrease methylation level in IFI16 gene promoter, leading to transcriptional activation of IFI16 gene. IFI, IFN-induced protein; EGCG, Epigallocatechin gallate; DNMT, DNA methyltransferase.

lines, promoter methylation status was assessed in IFI16 promoter. EGCG treatment decreased methylation signaling in the promoter of the IFI16 gene in MCF-7 cell line (Fig. 7A). The decrease in methylation signaling in IFI16 was greater in the T47D cell line than the MCF-7 cell line (Fig. 7B). The present data suggesting that EGCG decreases methylation level in the IFI16 gene promoter.

Discussion

Nucleic acid sensors, such as IFI16 DNA sensor, generate an innate immune response in tumor cells (47). IFI16 senses ds nucleic acid ligands and relays molecular signaling to induce inflammatory innate immune response via IFN stimulation. Previous epigenetic studies of these sensors in various cancer cell lines have shown that DNMTi and vitamin C induce their expression (31,38,48). EGCG-mediated epi-transcriptomic based regulation of IFI16 sensor is still not well studied. Hence, the aim of the present study was to investigate this in breast cancer lines.

IFI16 gene expression was screened in cancer cell lines, including MCF-7, T47D, SHSY5, HepG2 and HeLa, from the Human Protein Atlas. The present study aimed to investigate epigenetic mechanisms, such as DNA methylation-mediated epigenetic transcriptional regulation of IFI16 gene; therefore, DNMT mRNA expression was assessed in cancer cell lines. The TCGA dataset for IFI16 and DNMT genes was validated in breast cancer cell line from online data sources (UALCAN

and UCSC Xena browser). The present study found a negative association between IFI16 and DNMT gene expression. Hence, MCF-7 and T47D cell lines were treated with DNMTis, like EGCG, 5-aza-dc and vitamin C and expression of IFI16 and DNMTs was assessed. EGCG treatment induced IFI16 gene expression in both cell lines and this was greater in the MCF-7 cell line. As IFI16 stimulates IFN and ISG response, IFN β 1 and ISG15 expression was assessed; EGCG upregulated IFN β 1 gene expression in both cell lines and this was greatest in MCF-7 cell line. To the best of our knowledge, the effect of EGCG on expression of these genes has not been reported previously. Expression of these genes was measured following treatment with 60 μ M 5-aza-dc or 240 μ M vitamin C, which resulted in significant upregulation of mRNA expression for these genes. Previously, it has been shown that IFI16 and ISG15 gene expression is upregulated following treatment with vitamin C and 5-aza-dc alone or combination in HCT116, SNU398 and HL60 cancer cell lines (31). Additionally, 5-aza-dc treatment induces IFI16, ISG15 and IFN β 1 expression in ovarian cancer cell lines (48).

In addition, 5-aza-dc, vitamin C and EGCG treatment decreased DNMT1, DNMT3a and DNMT3b mRNA expression. A previous study showed that 20 μ M EGCG and 5 μ M 5-aza-dc treatment decrease DNMTs mRNA expression and DNMT activity and reactivates tumor suppressor genes by decreasing promoter methylation levels in various breast cancer lines (49).

As 5-aza-dc, vitamin C and EGCG decreased DNMTs mRNA expression, 5mC levels were assessed post-treatment. In both MCF-7 and T47D cell lines, 5-aza-dc slightly decreased 5mC%

level. Moreover, EGCG decreased 5mC levels in MCF-7 cells and a significant decrease in 5mC level in T47D cells was observed. A previous study on skin cancer cells revealed EGCG mediated decline of 5mC level and suggested that EGCG-mediated methylation is slow and might be more effective as a treatment when used for long periods (>72 h) (50). In the present study, no effect of vitamin C on the 5mC level was observed in any cell line. Binding of vitamin C outside of the binding pocket for DNMT1 and DNMT3b might be the reasons or any other factors that might be involved in the regulation of vitamin C-mediated 5mC regulation. A previous study showed the effect of vitamin C on methylation is reversible (51). DNMT3A activity regulation is governed by 5mC-binding protein MeCP2 and histone 3 tail modification (52). Ubiquitin-like With PHD and Ring Finger Domains 1 can regulate DNMT1 (53). Vitamin C may not functionally affect all these factors.

In silico molecular docking simulation showed that EGCG exhibited the greatest binding energy with DNMT proteins. 5-aza-dc showed the second highest binding energy with DNMTs. SAM and SAH were used as a control for docking simulation. The lowest binding energy was observed with vitamin C for all DNMTs. Additionally, docking interaction and predicted binding sites for all compounds showed that SAM and EGCG bound in a similar position across all DNMTs. Binding pocket was not specified during docking simulation to avoid bias docking; docking simulation demonstrated that compounds bind with certain amino acid residues in the binding pocket site. Similar amino acid residues interactions were also found in several previous molecular docking simulations of EGCG and DNMT (27,54,55).

As EGCG mediated a decrease in 5mC level and induction of IFI16 gene expression, the effect of EGCG on status of IFI16 gene promoter was assessed. Following EGCG treatment, methylation signaling was decreased in both cell lines. MCF-7 cell lines showed low decrease in methylation signaling following EGCG treatment, whereas T47D showed a greater decrease in methylation signaling following EGCG treatment. This may be due to low decrease in 5mC level in EGCG-treated MCF-7 cell line and greater 5mC decrease in EGCG-treated T47D cell line. However, a recent study revealed that IFI16 is hypomethylated in glioblastoma (56). To the best of our knowledge, EGCG-mediated reactivation of the IFI16 gene by decreasing promoter methylation has not previously been reported. IFI16 gene promoter methylation is DNMT-mediated, whereas EGCG treatment can block the DNMTs activity and reduce the methylation of IFI16 gene promoter, thus activating transcription of IFI16 gene (Fig. 8).

As a potent natural DNMTi, EGCG induces expression of innate immune sensor IFI16 by decreasing promoter methylation in breast cancer cells. The present findings provide a basis to investigate whether natural DNMTis (such as EGCG) exert beneficial effects by inducing immune response.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

MIK designed and conceptualized the study. SMN and WHA performed experiments, analyzed the data and wrote the manuscript. MIK and WHA confirm the authenticity of all the raw data. All authors have read and approved the final manuscript.

Ethics approval and consent to participate

Not applicable.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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