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Tea Polyphenol (–)-Epigallocatechin-3-Gallate Inhibits DNA Methyltransferase and Reactivates Methylation-Silenced Genes in Cancer Cell Lines

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Abstract

Hypermethylation of CpG islands in the promoter regions is an important mechanism to silence the expression of many important genes in cancer. The hypermethylation status is passed to the daughter cells through the methylation of the newly synthesized DNA strand by 5-cytosine DNA methyltransferase (DNMT). We report herein that (–)-epigallocatechin-3-gallate (EGCG), the major polyphenol from green tea, can inhibit DNMT activity and reactivate methylation-silenced genes in cancer cells. With nuclear extracts as the enzyme source and polydeoxyinosine-deoxycytosine as the substrate, EGCG dose-dependently inhibited DNMT activity, showing competitive inhibition with a K_i of 6.89 μM . Studies with structural analogues of EGCG suggest the importance of D and B ring structures in the inhibitory activity. Molecular modeling studies also support this conclusion, and suggest that EGCG can form hydrogen bonds with Pro¹²²³, Glu¹²⁶⁵, Cys¹²²⁵, Ser¹²²⁹, and Arg¹³⁰⁹ in the catalytic pocket of DNMT. Treatment of human esophageal cancer KYSE 510 cells with 5–50 μM EGCG for 12–144 h caused a concentration- and time-dependent reversal of hypermethylation of *p16^{INK4a}*, retinoic acid receptor β (*RAR β*), *O*⁶-methylguanine methyltransferase (*MGMT*), and human *mutL* homologue 1 (*hMLH1*) genes as determined by the appearance of the unmethylation-specific bands in PCR. This was accompanied by the expression of mRNA of these genes as determined by reverse transcription-PCR. The re-expression of *RAR β* and *hMLH1* proteins by EGCG was demonstrated by Western blot. Reactivation of some methylation-silenced genes by EGCG was also demonstrated in human colon cancer HT-29 cells, esophageal cancer KYSE 150 cells, and prostate cancer PC3 cells. The results demonstrate for the first time the inhibition of DNA methylation by a commonly consumed dietary constituent and suggest the potential use of EGCG for the prevention or reversal of related gene-silencing in the prevention of carcinogenesis.

Introduction

Green tea is a popular beverage worldwide. It has been shown to prevent carcinogenesis in animal models for different organ sites, and many mechanisms have been proposed for this activity (reviewed in Ref. 1). EGCG³, the major polyphenol in green tea, has many interesting activities and is believed to be a key active ingredient (1). In previous studies, we have shown that EGCG is methylated by catechol-*O*-methyltransferase to form MeEGCG and DiMeEGCG both *in vitro* and *in vivo* (2, 3). EGCG is also a potent inhibitor of catechol-

O-methyltransferase activity (3). Catechol-*O*-methyltransferase and DNMT belong to the same superfamily of *S*-adenosylmethionine-dependent methyltransferases. Both enzymes have a common core structure at the catalytic site (4). It is possible that EGCG may also inhibit DNMT by binding to a similar catalytic site.

Hypermethylation of DNA is a key epigenetic mechanism for the silencing of many genes, including those for cell cycle regulation, receptors, DNA repair, and apoptosis (5–8). In this aberrant methylation, the cytosine of the CpG island in or near the promoter region of the newly synthesized DNA strand is methylated by DNMT. The methylated CpG island has higher binding affinity to methyl-CpG binding domain proteins that recruit transcriptional corepressors, such as histone deacetylases and Sin 3A, resulting in chromosome condensation and transcription repression (9, 10). The inhibition of DNMT, especially DNMT1, would block the hypermethylation of the newly synthesized DNA strand, resulting in the reversal of the hypermethylation and the re-expression of the silenced genes (11–13). Indeed, this point has been demonstrated by studies with DNMT inhibitors, DAC (also known as 5-aza-2'-deoxycytidine) and zebularine. These compounds have been shown to inhibit cancer cell growth, induce cancer cell apoptosis, and reduce tumor volume in mice (14–17). There is high potential for developing this group of inhibitors for cancer therapy. However, side effects and toxicity are serious concerns. There is a great need for the development of effective and nontoxic inhibitors of DNMT. We hypothesize that EGCG can be used for this purpose.

In the present study, we tested this hypothesis by examining the inhibition of DNMT by EGCG and the effects of EGCG treatment on four methylation-silenced genes in human esophageal cancer cell line KYSE 510: the tumor suppressor *p16^{INK4a}*, retinoic acid receptor β (*RAR β*), *O*⁶-methylguanine methyltransferase (*MGMT*), and the DNA mismatch repair gene human *mutL* homologue 1 (*hMLH1*). In our previous work, these genes have been shown to be frequently silenced by DNA hypermethylation in human esophageal cancer, and some of these genes can be partially reactivated by treatment with the DNMT inhibitor, DAC (18). The results on the inhibition of DNMT and reactivation of methylation-silenced genes by EGCG are reported herein.

Materials and Methods

Cell Lines and Cell Culture. The human esophageal squamous cell carcinoma cell lines KYSE 510 and KYSE 150 were gifts from Dr. Yutaka Shimada (Kyoto University, Kyoto, Japan). The cells were maintained in 5% CO₂ atmosphere at 37°C in RPMI 1640 and Ham F12 mixed (1:1) medium containing 5% fetal bovine serum. To determine the dose-dependent changes, KYSE 510 cells were treated with 5, 10, 20, or 50 μM of EGCG (Unilever-Bestfoods, Englewood Cliffs, NJ) or 8.7 μM of DAC for 6 days. EGCG or DAC was added, in new culture medium, to the cells on days 1, 3, and 5. For the time course study, cells were treated with 20 μM of EGCG for 12, 24, 48, 72, or 144 h. Human colon cancer cell line HT-29 and prostate cancer cell line PC3 were obtained from American Type Culture Collection (Manassas, VA), and were grown in McCoy's 5A and RPMI 1640 containing 10% fetal bovine

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³ The abbreviations used are: EGCG, (–)-epigallocatechin-3-gallate; MeEGCG, 4'-*O*-methyl-EGCG; DiMeEGCG, 4', 4''-*O*-dimethyl-EGCG; DNMT, 5-cytosine DNA methyltransferase; DAC, 2'-deoxy-5-azacytidine; *RAR β* , retinoic acid receptor β ; *MGMT*, *O*⁶-methylguanine methyltransferase; *hMLH1*, human *mutL* homologue 1; poly(dI-dC)-poly(dI-dC), polydeoxyinosine-deoxycytosine; ECG, (–)-epicatechin gallate; EGC, (–)-epigallocatechin; EC, (–)-epicatechin; RT-PCR, reverse transcription-PCR.

Table 1 List of primers, annealing temperatures, and product sizes for methylation-specific PCR and RT-PCR

Gene	Chromosome location	Forward primer (5'–3')	Reverse primer (5'–3')	Annealing temperature (°C)	Product size (bp)
Methylation-specific PCR ^a					
<i>RARβ</i>	3p24	N: TTAAG(C/T)TTTGTGAGAATTTTG	CCTATAATTAATCCAATAATCATTTA CC	53	425
		U: TTGAGAATGTGAGTGATTGA	AACCAATCCAACCAAAACAA	62	146
		M: TCGAGAACGCGAGCGATTTCG	GACCAATCCAACCGAAACGA	62	146
<i>MGMT</i>	10q26	N: GGATATGTTG GGATAGTT	CCAAAAACCCCAAACCC	52	289
		U: TTTGTGTTTTGATGTTTGTAGGTTTTTGT	AACTCCACACTCTTCCAAAAACAAAACA	59	93
		M: TTTCGACGTTCTGATGTTTTTCGC	GCACTCTTCCGAAAAACGAAACG	59	81
<i>P16</i>	9p21	N: GGAGAGGGGGAGAGTAGGT	CTACAAACCCTCTACCCACCT	60	208
		U: TGGGGAGTAGTATGGAGTTGGTGGT	CAACCCCAAACCAACCATAA	62	81
		M: CGGGGAGTAGTATGGAGTCGGCGGC	GACCCCGAACCGCAGCCGTA	62	81
<i>hMLH1</i>	3p22	N: GGTATTTTTGTTTTTATGGTTGGAT	AATACCAATCAAATTTCTCAACTCTC	55	185
		U: TAAAAATGAATTAATAGGAAGAGT	CTCTATAAATTAATAAATCTCTTCA	60	117
		M: TAAAAACGAATTAATAGGAAGAGC	CTCTATAAATTAATAAATCTCTTCG	60	117
RT-PCR					
<i>RARβ</i>		GACTGTATGGATGTTCTGTCAG	ATTTGTCTGGCAGACGAAGCA	60	256
<i>MGMT</i>		AAAATGGACAAGGATTGTGAAA	CATCCGATGCAGTGTTACACG	57	568
<i>P16</i>		CGGAAGGTCCTCAGACATC	TCATGAAGTCGACAGTCTCCG	60	385
<i>hMLH1</i>		CAGCGGCCAGCTAATGCTAT	AATCTCAAAGGACTGCAATT	60	195
<i>DNMT1</i>		ACCAAGCAAGAAGTGAAGCC	GCTTCTGCAGAAGAACCTG	60	336
<i>DNMT3a</i>		CACACAGAAGCATATCCAGGAGTG	AGTGGAAACCAAAATACCC	60	550
<i>DNMT3b</i>		AATGTGAATCCAGCCAGGAAAGGC	ACTGGATTACACTCCAGGAACCGT	60	191
<i>MBD2</i>		AACCCTGCTGTTGGCTTAAC	CGTACTTGCTGTAAGCTCTTC	60	101
<i>G3DPH</i>		TTGCAACTGTTTTAGGACTTT	AGCATTGGGAAATGTTCAAGG	60	983

^a N, nest primers; U, unmethylation-specific primers; and M, methylation-specific primers.

serum, respectively. These cells were treated with EGCG for 6 days as described above.

DNA Methyltransferase Assay. Cultured KYSE 510 cells were harvested, and nuclear extracts were prepared with the nuclear extraction reagent (Pierce, Rockford, IL). The DNMT assay was performed according to published methods (19, 20). In brief, the nuclear extracts (4 μg of protein) were incubated for 1.5 or 2 h at 37°C with 0.66 μM of poly(dI-dC)·poly(dI-dC) and 10 μM of *S*-adenosyl-L-[methyl-³H]methionine (2 μCi; Amersham, Piscataway, NJ) in a total volume of 40 μl of a pH 7.4 buffer, containing 20 mM Tris-HCl, 25% glycerol (v/v), 10 mM EDTA, 0.2 mM phenylmethylsulfonyl fluoride, and 20 mM 2-mercaptoethanol. All of the incubations contained 0.02% DMSO, which was used to dissolve EGCG and other inhibitors. Reactions were initiated by the addition of nuclear extracts and stopped by mixing with 300 μl of a solution containing 1% SDS, 3% 4-aminosalicylate, 5% butanol, 2.0 mM EDTA, 125 mM NaCl, 0.25 mg/ml carrier salmon testes DNA (Life Technologies, Inc., Gaithersburg, MD), and 1 mg/ml protease K (Sigma, St. Louis, MO). To remove proteins, the resulting mixture was vortexed with 300 μl of a solution containing 8% phenol, 12% *m*-cresol, and 0.1% 8-hydroxyquinoline (Sigma), and centrifuged. The methylated DNA template was recovered from the aqueous phase by ethanol precipitation and then washed three times with 70% ethanol. The radioactivity in the pellets was counted in a scintillation counter. All of the assays were performed in duplicate. Background levels were determined in incubations without the template DNA.

In Silico Molecular Modeling Studies of DNMT1. The protein sequence of DNMT1 (accession no. NP_001370) was retrieved from the National Center for Biotechnology Information Reference Sequence (RefSeq) Collection. A structural model of the catalytic domain of DNMT1 was constructed using the Insight II Homology Module (Accelrys, Inc., San Diego, CA) from the published crystal structure of *HhaI* Mtase (RCSB Protein Data Bank;⁴ PDB ID = 5MHT) as the modeling template. The quality of the model was confirmed by the WHATIF-Check program.⁵ Initial structures of EGCG and the five analogues were built using the Insight II Builder Module and subsequently energy minimized to yield a stable conformation. Each ligand was docked into the putative DNMT1 binding pocket using GOLD (Genetic Optimization for Ligand Docking; Ref. 21), which accommodates ligand flexibility and partial enzyme flexibility to satisfy ligand-enzyme binding requirements. After these docking studies to elucidate stable ligand-binding modes, values of the ligand-enzyme binding energy ($\Delta E_{\text{binding}}$) were calculated

for each of the six ligands using molecular mechanics procedures. Residues within 5.0 Å of the cytosine nucleotide were defined as the “active site” pocket and were permitted full relaxation, while holding all of the other residues fixed. We define $\Delta E_{\text{binding}}$ as the difference between the potential energy of the ligand-enzyme (E_{complex}) and the sum of potential energies of the ligand (E_{ligand}) and enzyme (E_{enzyme}).

$$\Delta E_{\text{binding}} = E_{\text{complex}} - (E_{\text{ligand}} + E_{\text{enzyme}})$$

A favorable (more negative) binding energy is taken as evidence that the ligand possesses high affinity for DNMT1.

Bisulfite Modification and Methylation-Specific PCR. DNA was extracted from the cells using the DNeasy tissue kit (Qiagen, Valencia, CA), and was denatured with 0.2 M NaOH at 37°C for 30 min. The denatured DNA was treated with 3.3 M sodium bisulfite and 0.66 mM hydroquinone according to Herman *et al.* (22), with slight modifications (18). The modified DNA was purified using the Qiaquick gel extraction kit (Qiagen) and eluted with water. NaOH was added to the modified DNA for a final concentration of 0.3 M and maintained for 5 min at room temperature to desulfonate the modified DNA. The resulting DNA was repurified with Qiaquick gel and used immediately or stored at –20°C.

Methylation-specific PCR was carried out using a nested two-stage PCR approach (23). Stage I PCR was performed on bisulfite-modified DNA to amplify a 208-, 289-, 425-, or 185-bp fragment of the CpG-rich promoter regions of the *p16^{INK4a}*, *MGMT*, *RARβ*, or *hMLH1* genes, respectively. The primers recognize the bisulfite-modified template but do not discriminate between methylated and unmethylated alleles. The stage I PCR products were diluted and subjected to a stage II PCR with primer sets specific to methylated or unmethylated template. The primers were designed according to the literature (23–27). The chromosomal locations of all of the genes tested, primer sequences for both the methylated and unmethylated templates, annealing temperatures used, and expected PCR product sizes are summarized in Table 1. Placental DNA and placental DNA treated with *SssI* methyltransferase (New England Biolabs, Beverly, MA) were used as positive controls for unmethylated alleles and methylated alleles, respectively. Amplification was carried out using a 9700 Perkin-Elmer thermal cycler (Applied Biosystems, Foster City, CA).

Reverse Transcription-PCR. Total RNA was isolated from cells by the Tri reagent (Sigma). Reverse transcription of RNA was performed using the Advantage RT-for-PCR kit (Clontech, Palo Alto, CA). PCR was carried out using a 9700 Perkin-Elmer thermal cycler. The primers used in the PCR

⁴ Internet address: <http://www.rcsb.org/pdb>.

⁵ Internet address: <http://www.cmbi.kun.nl/gv/servers/WIWWWI>.

reactions for the genes analyzed (13, 28, 29) are summarized in Table 1. The PCR reactions were run in an approximate linear range (for example, 25–28 cycles for *p16^{INK4a}*, 30–35 cycles for *RAR β* , 20–25 cycles for *hMLH1*, and 30–35 cycles for *MGMT*). The amplified products were subjected to electrophoresis on a 2% agarose gel. The gel was then stained with ethidium bromide and photographed. Negative controls for PCR were run under the same conditions without RNA or reverse transcriptase. Expression of the glyceraldehyde-3-phosphate dehydrogenase gene (*G3PDH*) was used to examine the integrity of RNA in each sample and to standardize the amount of cDNA added to each of the PCR reactions.

Western Blot. The protein samples in SDS-containing lysis buffer were loaded on 4–15% acrylamide gel (40 $\mu\text{g}/\text{lane}$). After electrophoresis, the proteins were blotted onto a nitrocellulose membrane. After incubation in blocking buffer (5% nonfat milk), the membranes were incubated with the primary antibodies (rabbit antihuman *RAR β* and *hMLH1* polyclonal antibodies; Santa Cruz Biotechnology, Santa Cruz, CA) overnight at 4°C. After washing with Tris-buffered saline containing 0.1% Tween 20, the membrane was then incubated with sheep antirabbit horseradish peroxidase-labeled secondary antibody and visualized using the electrochemiluminescence detection kit (Amersham).

Results

Inhibition of DNA Methyltransferase Activity by EGCG. With nuclear extracts from the KYSE 510 cells as the enzyme source and poly(dI-dC)·poly(dI-dC) as the substrates for DNMT, under our assay conditions, the production of methylated poly(dI-dC)·poly(dI-dC) was linear for 2 h. A dose-dependent inhibition by EGCG was observed, with an IC_{50} of $\sim 20 \mu\text{M}$ (Fig. 1A). The structural analogues of EGCG: ECG, EGC, EC, and methylated EGCG all had inhibitory activities, with potency in the rank order of $\text{EGCG} > \text{ECG}$, $\text{MeEGCG} > \text{EGC}$, and $\text{DiMeEGCG} > \text{EC}$. Kinetic studies indicated that EGCG increases the K_m of DNMT without affecting the V_{max} , suggesting a competitive inhibition with a K_i of $6.89 \mu\text{M}$ (Fig. 1B). The same conclusion was reached when this experiment was repeated.

Molecular Modeling of the Interaction between EGCG and DNMT. To investigate the possible inhibition mechanism of EGCG to DNMT1, we used information available from the X-ray crystal structure of *HhaI* Mtase (30) to model the three-dimensional structure of the DNMT1 catalytic domain. This structural model revealed a substantial interaction region with hemimethylated DNA and a cytosine active pocket for subsequent methylation, which was folded by evolutionarily conserved sequence motifs identified from multiple sequence alignments (Fig. 2A). Docking of EGCG into the putative cytosine pocket indicated that the gallate moiety of this compound formed potential hydrogen bonds with two catalytically important residues, Glu¹²⁶⁵ and Pro¹²²³, which were the same residues that appear to stabilize the flipped cytosine through hydrogen bonding. Interestingly, the D ring was oriented at approximately the same position as the pyrimidyl ring of cytosine in the structural model of DNMT1. Furthermore, Ser¹²²⁹ and Cys¹²²⁵ are positioned to form hydrogen bonds with the hydroxy groups in the A and B rings of EGCG, respectively, thereby contributing to high-affinity DNMT1 binding. The interactions are depicted in Fig. 2B. These findings suggest that EGCG exerts its inhibitory effect on DNMT1 function by blocking entry of the key nucleotide cytosine into its active site and, thus, prevents methylation. A consensus-binding mode was obtained from 25 independent docking procedures for EGCG and each of the five analogues bound to DNMT1. Accordingly, calculated values of $\Delta E_{\text{binding}}$ (shown in parentheses, in units of kcal/mol) for each ligand in its consensus-binding mode yielded the following order of binding affinity to DNMT1: $\text{EGCG} (-49.75) > \text{ECG} (-46.53) > \text{Me-EGCG} (-44.91) > \text{diMe-EGCG} (-41.64) > \text{EGC} (-40.79) > \text{EC}$

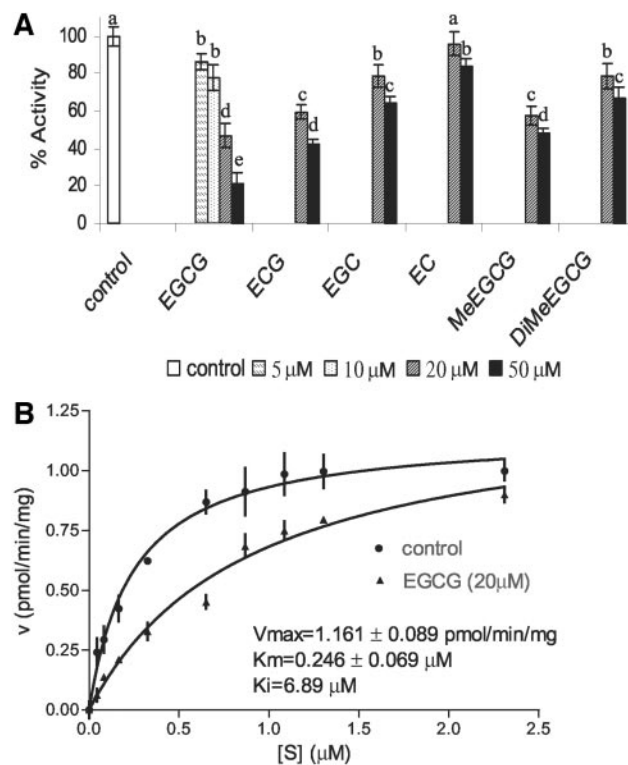


Fig. 1. Inhibition of 5-cytosine DNA methyltransferase activity by EGCG. A, inhibition of DNMT by different concentrations of EGCG and 20 or 50 μM of other compounds. The reaction mixture contained nuclear extracts (4 μg protein), poly(dI-dC)·poly(dI-dC); 0.66 μM , and *S*-adenosyl-L-[methyl-³H]methionine (10 μM , 2.0 μCi) in 40 μl . The incubation time was 2 h. The results are mean of 6–10 determinations, and are analyzed by one-way ANOVA (SigmaStat 1.01) using Student-Newman-Keuls pairwise test; bars, \pm SD. Bars identified by different letters signified that they are significantly different ($P < 0.05$). B, kinetic studies on the inhibition of DNMT by EGCG. The reaction mixture contained 10 μM of *S*-adenosyl-L-[methyl-³H]methionine, and different concentrations of poly(dI-dC)·poly(dI-dC), and was incubated for 1.5 h. Each data point represents the mean \pm SD of 4 determinations, and the data were analyzed by GraphPad Prism 4. In the absence of EGCG, the mean \pm 95% confidence interval of the V_{max} was 1.161 ± 0.089 pmol/min/mg and of the K_m was $0.246 \pm 0.070 \mu\text{M}$. In the presence of 20 μM EGCG, the V_{max} was 1.235 ± 0.205 pmol/min/mg, and K_m was $0.897 \pm 0.210 \mu\text{M}$, showing a competitive inhibition with a K_i of 6.89 μM .

(–38.32). These predicted results are generally consistent with the experimental data.

Reversal of Hypermethylation and Reactivation of *RAR β* , *MGMT*, *p16^{INK4a}*, and *hMLH1* by EGCG. In KYSE 510 cells, *p16^{INK4a}*, *RAR β* , *MGMT*, and *hMLH1* had hypermethylation status with the loss of the respective mRNA expression. After treating the cells with 5, 10, 20, or 50 μM of EGCG for 6 days, the methylation-specific bands of these four genes still existed. The unmethylation-specific bands of these four genes, however, appeared after treatment with 20 or 50 μM of EGCG for 6 days (Fig. 3A). Corresponding to the appearance of the unmethylation-specific band was the re-expression of mRNA of all four of the genes. The unmethylation-specific bands and mRNA expression were both observed with 20 and 50 μM of EGCG. For *hMLH1*, the unmethylation-specific band was observed even with 10 μM of EGCG in comparison to the faint bands observed with 0 and 5 μM EGCG (Fig. 3A). The reversal of hypermethylation and re-expression of these four genes by EGCG was similar to that produced by the classical DNMT inhibitor, DAC. After treating the cells with 20 μM of EGCG for 0, 12, 24, 48, 72, and 144 h, strong unmethylation-specific bands for all four of the genes began to appear at 48 h (Fig. 3A). The mRNA expression of these four genes was also observed at 48 h and, depending on the genes, higher levels were observed at either 72 or 144 h. Again, a faint *hMLH1* mRNA band

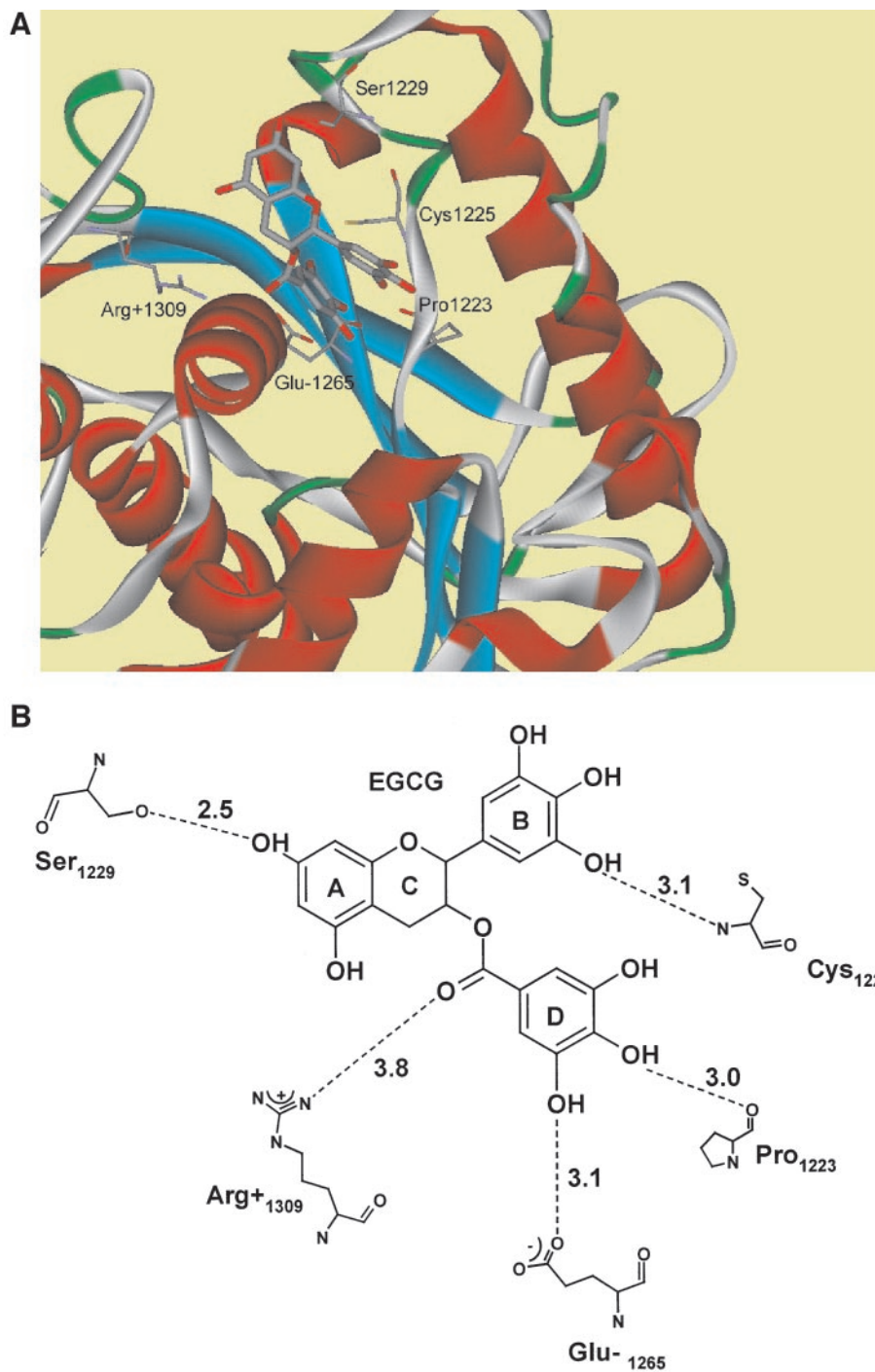


Fig. 2. Molecular modeling of the interaction between EGCG and DNMT. **A**, binding orientation of EGCG in DNMT1. The close-up view of the consensus orientation for EGCG. The protein is depicted in ribbon representation and colored by secondary structures (*i.e.*, helix, strand, and loop). Both EGCG and ligand contact residues are represented in stick form and colored by atom type with carbon in gray, oxygen in red, and sulfur in yellow. **B**, hydrogen-binding network of EGCG in DNMT1. ---- represents hydrogen bond. Figure depicts all potential H-bonding X-H...Y interactions for which the X...Y distance is <4.0 Å.

was observed at 0, 12, and 24 h. Similar results were obtained in two repeated experiments.

Using the reactivation of *RARβ* mRNA as an assay, ECG was found to be slightly less active than EGCG; EGC, MeEGCG, and EC had lower activities than EGCG; and DiMeEGCG had no activity when the cells were treated with $20 \mu\text{M}$ of the test agents for 6 days (Fig. 3B). In another experiment, demethylation and reactivation of *RARβ* was observed with DiMeEGCG at $50 \mu\text{M}$, but not at $20 \mu\text{M}$ (data not shown).

Western blot analysis indicated that the *RARβ* protein expression was increased in the cells treated with 20 and $50 \mu\text{M}$ of EGCG for 6 days (Fig. 4). A time-dependent increase of hMLH1 protein expression level was observed in the cells treated with $20 \mu\text{M}$ of EGCG for

3 and 6 days. Because of the lack of suitable antibodies, we were unable to accurately analyze the re-expression of MGMT and *p16^{INK4a}* proteins.

To determine whether the reactivation of methylation-silenced genes by EGCG is a general phenomenon that also occurs in other cell lines, we studied the effects of EGCG treatment on the methylation status and mRNA levels of *p16^{INK4a}* or *RARβ* in three other human cancer cell lines (Fig. 3C). The appearance of unmethylation-specific bands and reactivation of the mRNA of *p16^{INK4a}* in HT-29 cells, and *RARβ* in KYSE 150 and PC3 cells, were observed after treating the cells with $20 \mu\text{M}$ EGCG for 6 days. In the KYSE 150 cells, a faint *RARβ* mRNA band was also observed in cells without EGCG treatment, but the band was much

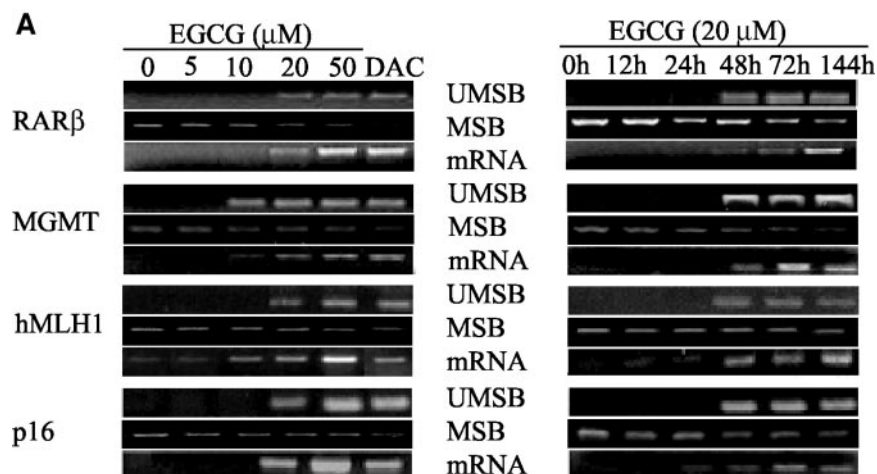
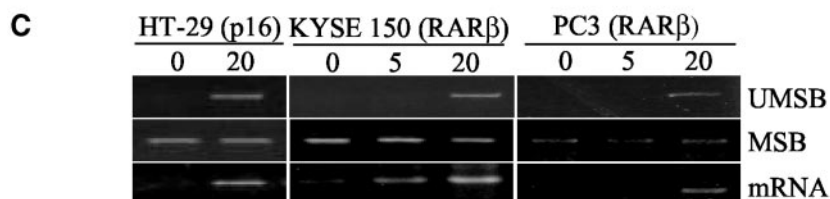
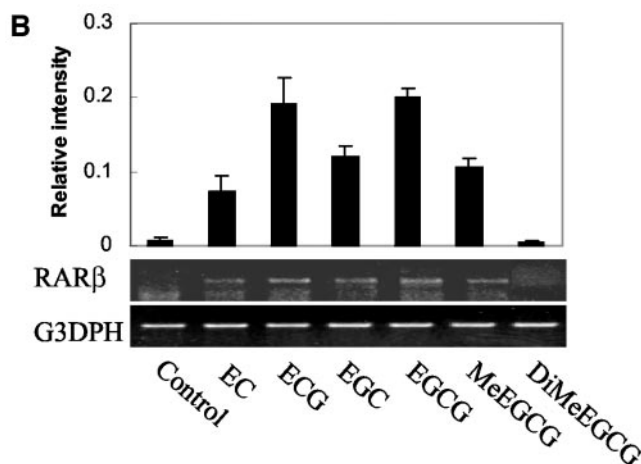


Fig. 3. Alterations of methylation status and mRNA expression levels of *RARβ*, *MGMT*, *p16*, or *hMLH1* genes after treatment with EGCG or other polyphenols. For each gene, the *top panel* presents the unmethylation-specific bands (*UMSB*), the *middle panel* presents the methylation-specific bands (*MSB*), and the *bottom panel* presents the mRNA expression levels (*mRNA*). *A, left*: KYSE 510 cells were treated with 5, 10, 20, or 50 μM of EGCG or 8.7 μM of DAC for 6 days as described in "Materials and Methods;" *right*: the cells were treated for 12, 24, 48, 72, and 144 h with 20 μM of EGCG added in new culture medium at 0, 48, and 96 h. *B*, *RARβ* mRNA levels of KYSE 510 cells treated with 20 μM of different polyphenols for 6 days. The *RARβ* mRNA level was determined with RT-PCR with *G3DPH* as an internal control. The band intensity was determined with densitometry. The values presented are mean ($n = 3$); bars, \pm SD. *C*, HT-29, KYSE150, and PC3 cells were treated with 5 or 20 μM of EGCG for 6 days under conditions described in "Materials and Methods."



more intense after EGCG treatment. These results were reproduced in three experiments with the three cell lines. The results, although still preliminary in a quantitative sense, demonstrate that the reactivation of methylation-silenced genes does occur in different cell lines, suggesting it is a general phenomenon.

Inhibition of Cell Growth and Other Effects. After treatment with EGCG for 6 days, significant inhibition of cell growth was not observed at 5 or 10 μM , but 36 and 54% inhibition were observed at 20 and 50 μM of EGCG, respectively (Fig. 5A). Many damaged cells and some polynuclear cells were observed with 50 μM of EGCG, and to a lesser extent with 20 μM of EGCG, in this 6-day experiment (Fig. 5C). In experiments in which cells were treated for 2 days with EGCG, growth inhibition and signs of toxicity were not apparent at 5, 10, or 20 μM ; but at 50 μM , some damaged cells with vesicles were observed (Fig. 5C). After treatment of the cells with 50 μM of EGCG for 2 days, the colony formation (measured after an additional 7 days of culturing) was inhibited by 55%, but significant inhibition was not observed with EGCG at 20 μM or lower concentrations (Fig. 5B). Preliminary experiments with RT-PCR also indicated that EGCG treatment did not affect the mRNA expression level of DNMT1, DNMT3a, DNMT3b, and MBD2 (methyl-CpG binding domain

protein 2), which has been reported to have DNA demethylase activity (31).

Discussion

The present study clearly demonstrates that EGCG inhibits DNMT activity, and causes CpG demethylation and reactivation of methylation-silenced genes. To our knowledge, this is the first demonstration of such an activity of a commonly consumed dietary constituent. Kinetic studies indicate that EGCG inhibits nuclear DNMT activity competitively with a K_i of 6.89 μM . We believe that most of the activity is because of the dominant form of DNMT, DNMT1, although DNMT 3a and 3b may also be present in smaller quantities in the nuclear extracts (14). Molecular modeling studies indicate that EGCG is well accommodated in a hydrophilic pocket of DNMT1. As depicted in Fig. 2B, EGCG is effectively tethered within the DNMT1 binding site by at least four hydrogen bonds involving Ser¹²²⁹ (2.5Å), Glu¹²⁶⁵ (3.1Å), Pro¹²²³ (3.0 Å), and Cys¹²²⁵ (3.1Å). An additional hydrogen bond is possible with the protonated side chain of Arg¹³⁰⁹, although the N...O distance (3.8Å) is at the outer limit of the typical hydrogen bond and, thus,

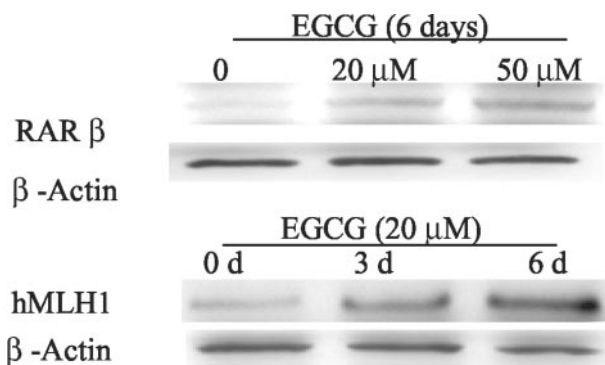


Fig. 4. Re-expression of *RARβ* and *hMLH1* proteins after EGCG treatment. KYSE 510 cells were treated with 20 and 50 μM EGCG for 6 days, or 20 μM for 3 and 6 days; protein levels were determined with Western blot analysis using β -actin as an internal control.

would imply a weak interaction. This hydrogen-bonding scheme is consistent with the observed reduction in DNMT1 inhibitory activity for the five EGCG structural analogues. The absence of the B-ring 3'-hydroxy group in ECG abolishes the hydrogen bond with Cys¹²²⁵. Likewise, the absence of the 4"-hydroxy group in MeEGCG eliminates the hydrogen bond with Pro¹²²³. Addition of another methyl group at the B ring as in DiMeEGCG additionally interferes with the binding. The importance of galloyl moiety in EGCG-like compounds is evident from interactions with the two conserved residues, Glu¹²⁶⁵ and Pro¹²²³, which are associated with the catalytic activity of DNMT1. This observation concurs with the

experimental results for EGC and EC, two poor DNMT1 inhibitors in which the galloyl moiety is conspicuously absent.

The activities of these compounds to reactivate *RARβ* mRNA expression are also correlated with their ability to inhibit DNMT activity. With EGCG, demethylation and reactivation of all four of the genes are observable after 48 h of treatment, and 20 μM is the effective concentration that activates all four of the genes. All of the results are consistent with our hypothesis that reactivation of methylation-silenced genes is due to the inhibition of DNMT by EGCG. More work is needed to determine the quantitative reactivation of the different genes as well as the extents of the demethylation in different cell lines under different treatment conditions. It would be interesting to determine whether the different hypermethylated genes respond similarly or differently to the EGCG treatment. The reactivated *p16^{INK4a}* is expected to regain its function in cell cycle regulation, and re-expressing *RARβ* is expected to regain responsiveness to retinoic acids. It is not clear whether the presently observed growth-inhibitory effect after 2 or 6 days of treatment with EGCG are related to the reactivation of these genes. The functional importance of the reactivation of those genes remains to be determined.

Hypermethylation of DNA is a key epigenetic mechanism for the silencing of many genes including those for tumor suppressors, DNA repair enzymes, and receptors (5–8). In theory, this epigenetic process is reversible if the newly synthesized DNA strands are not methylated. Therefore, DNMT inhibitors such as DAC and zebularine have been actively explored as cancer therapeutic agents, especially when they are used in combination with a

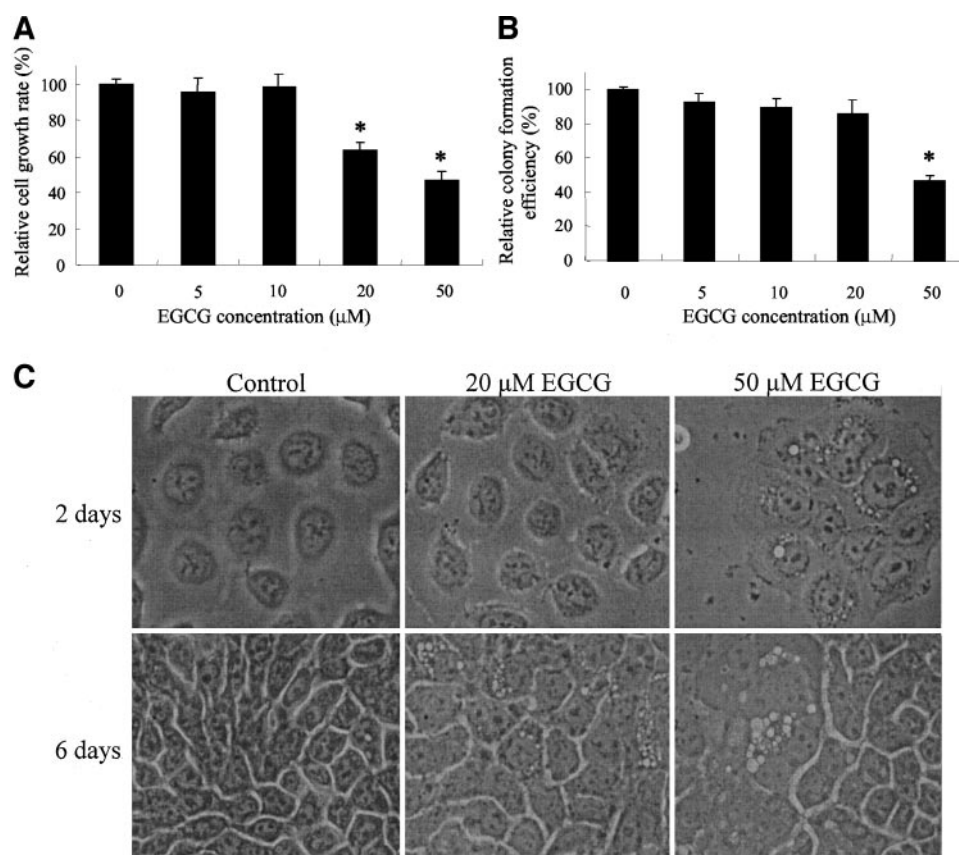


Fig. 5. Effects of EGCG treatment on cell growth, colony formation, and morphological change. **A**, KYSE 510 cells were treated with EGCG, and after 6 days, its effect on cell growth was determined by counting living cell numbers by the trypan blue exclusion assay. **B**, after the cells were treated with EGCG for 2 days, the cells were washed, cultured with new medium (without EGCG) for another 7 days, and colony number was analyzed. **C**, morphological changes (magnification, $\times 400$) after 2 and 6 days of EGCG treatment are shown. The values presented are mean ($n = 3$); bars, \pm SD. * indicates statistically significant ($P < 0.05$).

histone deacetylase inhibitor, such as trichostatin A (15, 26, 32–36). It has also been suggested that methylation is not the initial event in triggering gene silencing in cancer; rather, the methylation of the promoter CpG islands is a consequence of prior gene inactivation, and it is a mechanism for locking the chromatin in a repressed state (5, 13, 37). Many tumor suppressor and receptor genes have been reported to be hypermethylated and transcriptionally silenced during the development of different types of cancers. It is likely that inhibition of DNMT and histone deacetylase can also prevent the hypermethylation and silencing of these key genes and, therefore, this inhibition would contribute to the prevention of carcinogenesis. The prevention of intestinal tumorigenesis by *Dnmt1* deficiency and by DAC has been demonstrated in the *Min* mice, which carry a mutated *Apc* gene (38, 39).

Oral administration of green tea has been shown to inhibit tumorigenesis in different organs, and multiple mechanisms may be involved (1). It remains to be determined whether and to what extent this cancer-preventive activity is due to the inhibition of DNMT by EGCG. The presently observed effective dose of EGCG, K_i of 6.89 μM , or IC_{50} of 20 μM , is achievable in the oral cavity (in the saliva) after drinking green tea, and perhaps in the stomach, esophagus, and intestines where there is direct contact between EGCG and the epithelial cells. This effective concentration, however, is higher than those in the internal organs, which depend on the systemic bioavailability of EGCG (1). Therefore, the extent of DNMT inhibition *in vivo* would depend on the bioavailability of EGCG in a particular organ site. Although inhibition of DNMT is expected to prevent hypermethylation, severe inhibition of DNMT activity, as suggested by recent genetic studies, may cause DNA hypomethylation, genomic instability, and early development of cancers such as T-cell lymphomas and sarcomas (40, 41). There is no evidence for such adverse effects due to regular consumption of tea; however, it could be a concern if large doses of EGCG are given to humans.

On the basis of this analysis, for the practical application of tea in the prevention of cancer, it may be desirable to use doses to produce levels of EGCG in the tissue that would cause a moderate inhibition of DNMT. The possible synergy generated between EGCG with histone deacetylase inhibitors or other agents will hold great promise. For example, butyrate, which is a histone deacetylase inhibitor produced in the colon (42), may work synergistically with EGCG in the prevention of colon cancer. The application of this concept in animal models or humans remains to be demonstrated.

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