

Transcriptional targeting of sphingosine-1-phosphate receptor SIP2 by epigallocatechin-3-gallate prevents sphingosine-1-phosphate-mediated signaling in macrophage-differentiated HL-60 promyelomonocytic leukemia cells

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Background: Macrophage chemotaxis followed by blood–brain barrier transendothelial migration is believed to be associated with inflammation in the central nervous system. Antineuroinflammatory strategies have identified the dietary-derived epigallocatechin-3-gallate (EGCG) as an efficient agent to prevent neuroinflammation-associated neurodegenerative diseases by targeting proinflammatory mediator signaling.

Methods: Given that high levels of sphingosine kinase and its product, sphingosine-1-phosphate (S1P), are present in brain tumors, we used quantitative reverse-transcription polymerase chain reaction (qRT-PCR) and immunoblotting to test whether EGCG may impact on *SIP* receptor gene expression and prevent S1P response in undifferentiated and in terminally differentiated macrophages.

Results: Promyelomonocytic human leukemia (HL)-60 cells were differentiated into macrophages, and S1P triggered phosphorylation in extracellular signal-regulated kinase (ERK), c-Jun N-terminal kinase (JNK), and P38 mitogen-activated protein kinase (MAPK) intracellular signaling, as shown by Western blot analysis. Pretreatment of cells with EGCG prior to differentiation inhibited the response to S1P in all three pathways, while EGCG abrogated P38 MAPK phosphorylation when present only during differentiation. Terminally-differentiated macrophages were, however, insensitive to EGCG treatment. Using qRT-PCR, gene expression of the S1P receptors S1P1, S1P2, and S1P5 was predominantly induced in terminally-differentiated macrophages, while the S1P2 was decreased by EGCG treatment.

Conclusion: Our data suggest that diet-derived EGCG achieves efficient effects as a preventive agent, targeting signaling pathways prior to cell terminal differentiation. Such properties could impact on cell chemotaxis through the blood–brain barrier and prevent cancer-related neuroinflammation.

Keywords: chemoprevention, green tea EGCG, neuroinflammation, blood–brain barrier, S1P

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Introduction

Accumulating evidence suggests that neuroinflammation strongly correlates with and characterizes neurodegenerative processes associated with central nervous system (CNS) disorders. Those inflammatory events occur at the blood–brain barrier (BBB) in diseases such as multiple sclerosis, epilepsy, and Alzheimer's disease.¹ Inflammatory response in the CNS is known to mobilize different types of immune cells, including

macrophages, mast cells, T and B lymphocytes, and dendritic cells, as well as CNS-resident cells, such as microglia, astrocytes, and neurons.² During neuroinflammation, responses to proinflammatory cues and subsequent chemotaxis therefore represent important events in the recruitment of these cells to the CNS.³ As a consequence of proinflammatory growth factor- or cytokine-mediated intracellular signaling, passage through the BBB endothelium allows these cells to reach the site of inflammation. If this process is not controlled, the prolonged inflammation can then be the cause of important cerebral damage.

Inflammation has also been closely linked to various forms of cancer and particularly, to brain tumor development.^{4,5} However, very little is known about the role of inflammation in glioma, especially at the initiation stage. It has been suggested that bidirectional communications between immune cells and glioma cells create an immunosuppressed microenvironment that promotes tumor survival and growth.⁶ Such potential tumor-initiating roles of inflammation in glioma prompted researchers to develop several immunotherapy approaches, in order to reverse glioma development and progression and its ability to evade the immune system. In one example of this tumor-initiating role, glioma secrete high levels of the bioactive lipid sphingosine-1-phosphate (S1P),⁷ and high expression levels of the enzyme that forms S1P, sphingosine kinase 1, correlate with shorter survival time for glioblastoma multiforme patients.⁸ Furthermore, S1P is a biologically active metabolite of plasma-membrane sphingolipids, which is essential for immune-cell trafficking. Much of the immune function of S1P results from the engagement of a family of five G-protein-coupled S1P receptors (S1P1-5). Recent findings on the role of S1P in immunosurveillance support regulatory mechanisms in S1P-mediated immune-cell trafficking and suggest that any perturbation in the interactions between S1P and its receptors may affect immunity.⁹

Recently the anti-inflammatory activity of plant and diet-derived compounds, most belonging to the chemical groups of alkaloids, coumarins, flavonoids, polyphenols and terpenoids, was demonstrated.^{10,11} Although flavonoids have been used in inflammatory pathway-targeting,^{12,13} it is not known how these may modulate the recruitment and infiltration of macrophages into the tumor, where the microenvironment subsequently activates them to support the malignant progression of cancer cells. Being the most potent secreted matrix metalloproteinase (MMP) to permeate the BBB,¹⁴ MMP-9 has been ascribed a crucial role and provided the molecular rationale explaining how the disruption of the BBB^{15,16} and

how subsequent transendothelial migration of terminally differentiated human leukemia (HL-60) macrophages occur.¹⁷ Collectively, these processes are thought to lead to cerebral infections and development of brain pathologies. However, while green tea-derived epigallocatechin-gallate (EGCG) was shown to efficiently inhibit the inflammation biomarker cyclooxygenase (COX)-2, it was unable to inhibit MMP-9 secretion in differentiated-macrophages.¹⁸

Given the selective biomarker-targeting of EGCG, we questioned whether pro-oncogenic-mediated differentiation of HL-60 promyelomonocytic leukemia cells into macrophages affected the cellular response to EGCG. In the current study, we therefore explored the capacity of diet-derived EGCG to prevent S1P-mediated signaling, which could possibly contribute to macrophage mobilization through the BBB.

Materials and methods

Materials

Sodium dodecyl sulfate (SDS) and bovine serum albumin (BSA) were purchased from Sigma-Aldrich Corp (St Louis, MO, USA). Cell culture media was obtained from Life Technologies Corp (Carlsbad, CA, USA). Electrophoresis reagents were purchased from Bio-Rad Laboratories (Hercules, CA, USA). The HyGLO™ Chemiluminescent HRP (horseradish peroxidase) Antibody Detection Reagents were from Denville Scientific Inc. (Metuchen, NJ, USA). Micro bicinchoninic acid (BCA) protein assay reagents were from Pierce (Micro BCA™ Protein Assay Kit; Thermo Fisher Scientific, Waltham, MA, USA). The polyclonal antibodies against extracellular signal-regulated kinase (ERK), phosphorylated (P)-ERK, c-Jun N-terminal kinase (JNK), P-JNK, P38, and P-P38 were all purchased from Cell Signaling Technology, Inc (Danvers, MA, USA). HRP-conjugated donkey anti-rabbit and anti-mouse immunoglobulin (Ig) G secondary antibodies were from Jackson ImmunoResearch Laboratories (West Grove, PA, USA). EGCG was from MP Biomedicals (Santa Ana, CA, USA). All other reagents were from Sigma-Aldrich Corp.

Cell culture and treatments

The HL-60 promyelomonocytic cell line was purchased from American Type Culture Collection (Manassas, VA, USA) and maintained in Iscove's Modified Dulbecco's Medium (Life Technologies Corp) containing 20% (v/v) fetal bovine serum (FBS) (HyClone™; Thermo Fisher Scientific), 2 mM glutamine, 100 units/mL penicillin, and 100 µg/mL streptomycin (WISENT Inc., Québec, QC, Canada), and were cultured at 37°C, under a humidified atmosphere containing 5% CO₂. The tumor-promoting agent phorbol-12-myristate-13-acetate

(PMA) was used to trigger macrophage differentiation. Slides of PMA-treated HL-60 cells were then mounted for light microscopy and air-dried, stained with Diff-Quick Solution (Baxter Healthcare Corp, Deerfield, IL, USA), and examined for morulas, as previously validated.¹⁹ Given that numerous protocols can be found in the literature to differentiate resting HL-60 cells into “macrophage-like cells” using PMA (between 2 to 8 days with various PMA concentrations, alone or in combination with other molecules), we wish to emphasize that, throughout the text the “HL-60 macrophage differentiation” condition represents the adherent subpopulation of HL-60 cells immediately harvested upon PMA treatment. We use “terminally differentiated macrophages” to refer to those same adherent cells, which were subsequently maintained in culture for another 24–48 hours. Cell starvation is defined as: treatments of the cells, either with PMA, EGCG, or SIP, in medium from which only FBS was removed.

Gelatin zymography

Gelatin zymography was used to assess the extent of pro-MMP-9 gelatinolytic activity, as previously described.¹⁸ Briefly, an aliquot (20 μ L) of the culture medium was subjected to SDS–polyacrylamide gel electrophoresis (PAGE) in a gel containing 0.1 mg/mL gelatin, a substrate that is efficiently hydrolyzed by pro-MMP-9. The gels were then incubated in 2.5% Triton™ X-100 and rinsed in nanopure distilled H₂O. The gels were further incubated at 37°C for 20 hours in 20 mM NaCl, 5 mM CaCl₂, 0.02% Brij®-35, and 50 mM tris(hydroxymethyl)aminomethane (Tris)-HCl buffer, pH 7.6, then stained with 0.1% Coomassie Brilliant blue R-250 and destained in 10% acetic acid and 30% methanol in H₂O. Gelatinolytic activity was detected as unstained bands on a blue background.

Immunoblotting procedures

Proteins from control and treated cells were separated by SDS-PAGE. After electrophoresis, proteins were electrotransferred to polyvinylidene difluoride membranes (Thermo Fisher Scientific), which were then blocked for 1 hour, at room temperature, with 5% nonfat dry milk in Tris-buffered saline (150 mM NaCl, 20 mM Tris–HCl, pH 7.5) containing 0.3% Tween™ 20 (TBST). The membranes were further washed in TBST and incubated with the primary antibodies (1/1,000 dilution) in TBST containing 3% BSA and 0.1% sodium azide, followed by a 1-hour incubation with HRP-conjugated anti-rabbit or anti-mouse IgG (1/2,500 dilution) in TBST containing 5% nonfat dry milk. Immunoreactive material was visualized by chemiluminescence.

Total RNA isolation, cDNA synthesis, and qRT-PCR

Total ribonucleic acid (RNA) was extracted from cell suspensions or monolayers using TRIzol® reagent (Life Technologies Corp). For complementary deoxyribonucleic acid (cDNA) synthesis, 2 μ g of total RNA were reverse-transcribed using a high capacity cDNA reverse-transcription kit (Life Technologies Corp). cDNA was stored at –80°C prior to quantitative reverse-transcription polymerase chain reaction (qRT-PCR). Gene expression was quantified by qRT-PCR using iQ™ SYBR® Green Supermix (Bio-Rad Laboratories). DNA amplification was carried out using an iCycler iQ™5 (Bio-Rad Laboratories), and product detection was performed by measuring binding of the fluorescent dye SYBR Green I to double-stranded DNA. The following QuantiTect primer sets were obtained from Qiagen (Venlo, the Netherlands): *SIP1* (QT00208733), *SIP2* (QT00230846), *SIP3* (QT00244251), *SIP4* (QT01192744), and *SIP5* (QT00234178). *Glyceraldehyde 3-phosphate dehydrogenase (GAPDH)* primer sets were synthesized by Biocorp (Dollard-des-Ormeaux, QC, Canada), with the following sequences: forward CCATCACCATCTTCCAGGAG and reverse CCTGCTTACCACCTTCTTG. The relative quantities of target gene messenger (m)RNA, compared against two internal controls, *GAPDH* and β -actin mRNA, were measured by following a Δ cycle threshold (C_t) method employing an amplification plot (fluorescence signal versus [vs] cycle number). The difference (ΔC_t) between the mean values in the triplicate samples of target gene and those of *GAPDH* and β -actin mRNAs were calculated using iQ5 Optical System Software (Version 2.0; Bio-Rad Laboratories), and the relative quantified value was expressed as $2^{-\Delta C_t}$.

Statistical data analysis

Unless otherwise stated, data are representative of three or more independent experiments. Statistical significance was assessed using Student’s unpaired *t*-test. Probability values of less than 0.05 were considered significant, and an asterisk identifies such significance in the figures. Error bars in all figures represent standard error of the mean (SEM) values.

Results

PMA triggers promyelocytic HL-60 cell differentiation into adherent macrophages

Among the agents shown to induce differentiation of HL-60 cells, the tumor-promoting agent PMA triggers a terminal differentiated monocytic/macrophage phenotype.²⁰

Here, we treated HL-60 cells with increasing concentrations of PMA for 24 hours, then discarded any cells that remained in suspension and took phase contrast pictures (Figure 1A). We validated that the optimal dose of PMA that triggered optimal cell adhesion was 3 μM (Figure 1B). Finally, we demonstrated, by zymography, increased secretion of MMP-9, characteristic of the adhesive phenotype of terminally differentiated macrophages, in agreement with previous reports.^{18,19}

Macrophage differentiation correlates with significant increases in *SIP1*, *SIP2*, and *SIP5* gene expression

We next evaluated the gene expression profiles of the *SIP* receptor family, in macrophage/monocyte terminally differentiated HL-60 cells. Promyelocytic HL-60 cells were treated with increased PMA concentrations, total RNA extracted from adherent cells, and cDNA synthesis performed, as described in the “Methods” section. We first

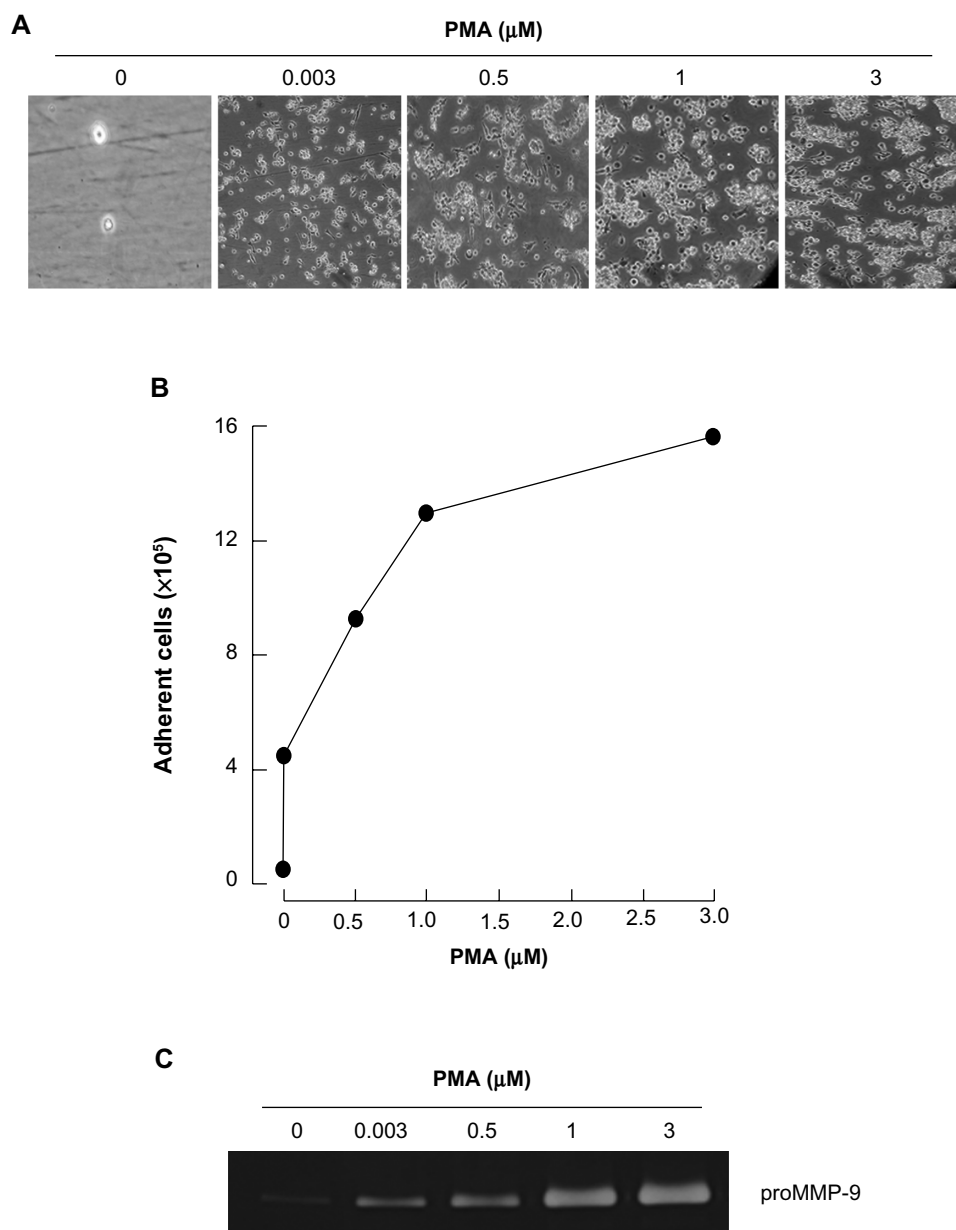


Figure 1 PMA triggers promyelocytic HL-60 cell differentiation into adherent macrophages.

Notes: Serum-starved HL-60 cells in suspension (10^6 cells/flask) were treated with various PMA concentrations for 18 hours. **(A)** Cells that remained in suspension were discarded, and phase contrast pictures were taken of the macrophage-differentiated adherent cells, and **(B)** the numbers of adherent cells per well counted. The data is representative of one out of four independent PMA treatments performed at different HL-60 cell passages. **(C)** Conditioned media were also collected from each condition, and gelatin zymography was performed.

Abbreviations: HL, human leukemia; MMP, matrix metalloproteinase; PMA, phorbol-12-myristate-13-acetate.

validated that each of our specific primer sets amplified a single amplicon corresponding to the *S1P1*, *S1P2*, *S1P3*, *S1P4*, and *S1P5* genes, as seen on an ethidium bromide-stained agarose gel (Figure 2A). qRT-PCR then allowed us to assess the extent of changes in gene expression; we

concluded that *S1P1*, *S1P2*, and *S1P5* were significantly increased in adherent cells, while only moderate increases were observed for *S1P3* and *S1P4* (Figure 2B). Collectively, these results suggest that macrophage-like cells could be responsive to S1P.

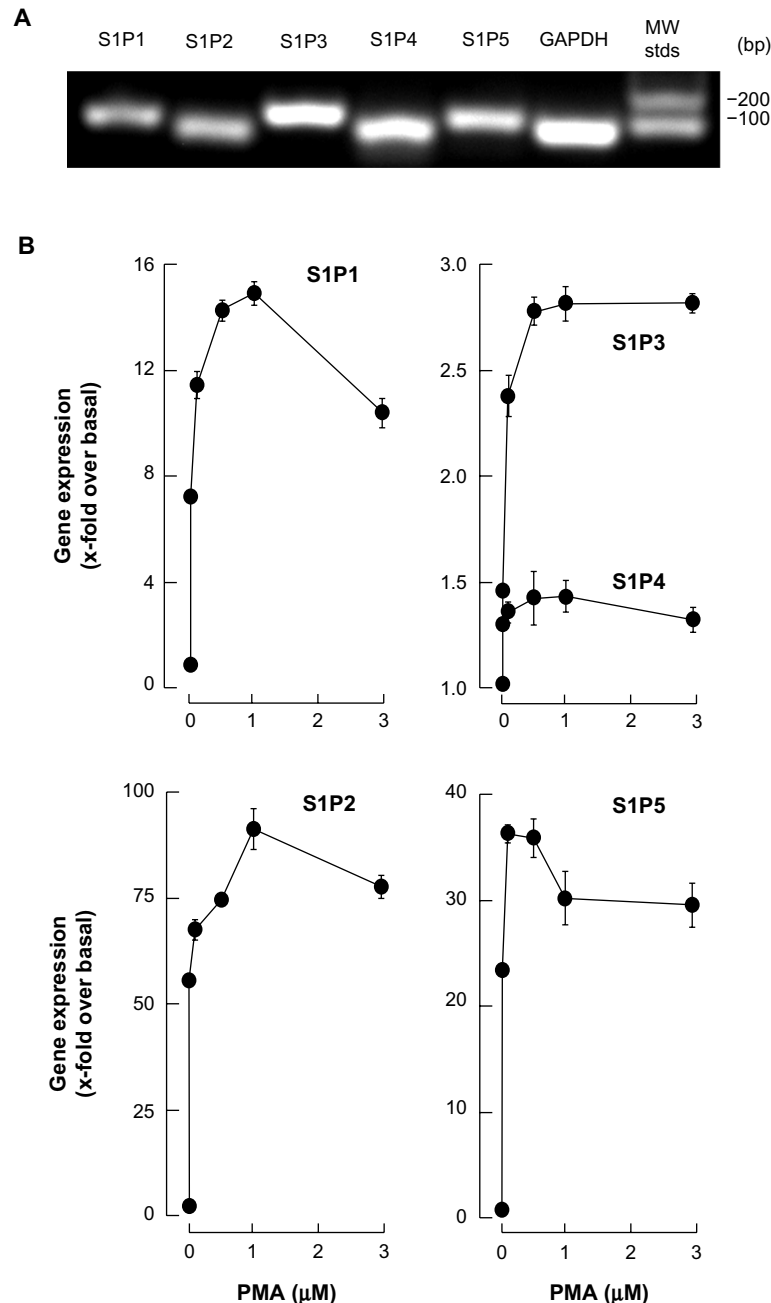


Figure 2 Macrophage differentiation correlates with significant increases in *S1P1*, *S1P2*, and *S1P5* gene expression.

Notes: Serum-starved HL-60 cells were treated for 18 hours with 3 μM PMA to induce HL-60 cell differentiation and adhesion. Total RNA was then isolated from adherent cells, transcribed into cDNA, and qRT-PCR was performed to assess the expression of the S1P receptors *S1P1*, *S1P2*, *S1P3*, *S1P4*, and *S1P5*. (A) A representative agarose gel electrophoresis is shown to confirm the generation of single amplicons. (B) The qRT-PCR gene expression assay for each of the five genes assessed was performed and expressed as the mean of triplicates. This experiment is a representative quantification of one out of four PMA treatments performed at different HL-60 cell passages.

Abbreviations: cDNA, complementary deoxyribonucleic acid; HL, human leukemia; GAPDH, glyceraldehyde 3-phosphate dehydrogenase; MW stds, molecular weight standard deviations; PMA, phorbol-12-myristate-13-acetate; qRT-PCR, quantitative reverse-transcription polymerase chain reaction; RNA, ribonucleic acid; S1P, sphingosine-1-phosphate.

SIP triggers rapid intracellular signaling through multiple pathways in macrophage-differentiated HL-60 cells

In order to assess whether differentiated HL-60 cells were capable of responding to SIP, we measured the extent of phosphorylation of ERK, P38 MAPK, and JNK. We observed that SIP rapidly triggered the phosphorylation of all the aforementioned signaling intermediates (Figure 3A, C, and E), typically reaching an optimal effect between 30 seconds and 1 minute of stimulation (Figure 3B, D, and F). A prior assessment of dose–response ERK phosphorylation, by increasing concentrations of SIP, found that 1 μ M SIP triggered an optimal signal (not shown).

EGCG pretreatment of promyelocytic HL-60 inhibits SIP response in macrophage-differentiated HL-60 cells

We next assessed the potential ability of EGCG to affect the SIP cellular response. To this end, we evaluated EGCG effects either prior (predifferentiation) to inducing HL-60 cells differentiation with PMA, simultaneously (codifferentiation) with HL-60 differentiation, or on cells that were already differentiated (postdifferentiation). Cell lysates were isolated following stimulation with SIP, and the extents of ERK, P38 MAPK, and JNK phosphorylation were evaluated by Western blotting (not shown).

Scanning densitometry of the autoradiograms showed that PMA-differentiated cells that had been pretreated with EGCG had an abrogated global SIP response as ERK, P38 MAPK, and JNK phosphorylation were inhibited (Figure 4A–C). Where EGCG had been added to cells simultaneously with PMA, only the SIP-mediated phosphorylation of P38 MAPK was inhibited (Figure 4D–F). Finally, where EGCG was added to terminally differentiated cells, none of the SIP-induced phosphorylated intermediates were diminished (Figure 4G–I). Altogether, these observations suggest that EGCG efficiently prevents subsequent responses to SIP in macrophages, while it selectively abrogates the cell response during differentiation and is completely inefficient in terminally differentiated cells.

EGCG selectively alters *SIP* receptor gene expression in macrophage-differentiated HL-60 cells

We next examined whether expression of the induced *SIP1*, *SIP2*, or *SIP5* genes were affected by EGCG. Total RNA was extracted from cells treated as described above and gene expression assessed by qRT-PCR. EGCG was found to only significantly downregulate *SIP2* gene expression in pre- and codifferentiation conditions, while it did not alter the increased expression of all three receptors in the postdifferentiation condition (Figure 5).

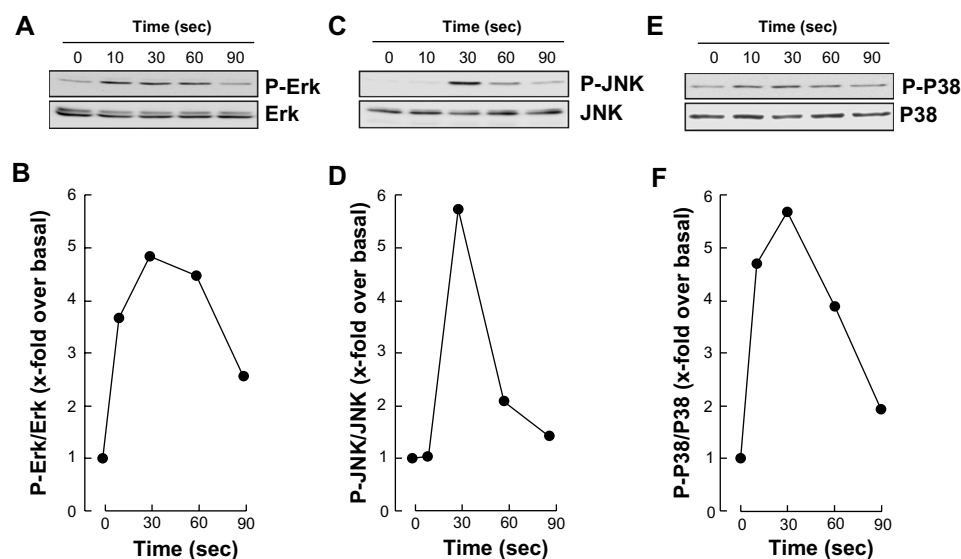


Figure 3 SIP triggers rapid intracellular signaling through multiple pathways in macrophage-differentiated HL-60 cells.

Notes: Serum-starved HL-60 cells were treated for 18 hours with 3 μ M PMA to induce HL-60 cell differentiation and adhesion. Adherent cells were then treated with 1 μ M SIP for up to 1.5 minutes, following which cell lysates were isolated, electrophoresed via SDS-PAGE, and immunodetected for (A) P-ERK, ERK, (C) P-JNK, JNK, (E) P-P38, and P38 proteins. Quantification was performed by scanning densitometry of the autoradiograms (B, D, and F).

Abbreviations: ERK, extracellular signal-regulated kinase; HL, human leukemia; JNK, c-Jun N-terminal kinase; P-ERK, phosphorylated ERK; P-JNK, phosphorylated JNK; P-P38, phosphorylated P38; PMA, phorbol-12-myristate-13-acetate; SDS-PAGE, sodium dodecyl sulfate–polyacrylamide gel electrophoresis; SIP, sphingosine-1-phosphate; sec, seconds.

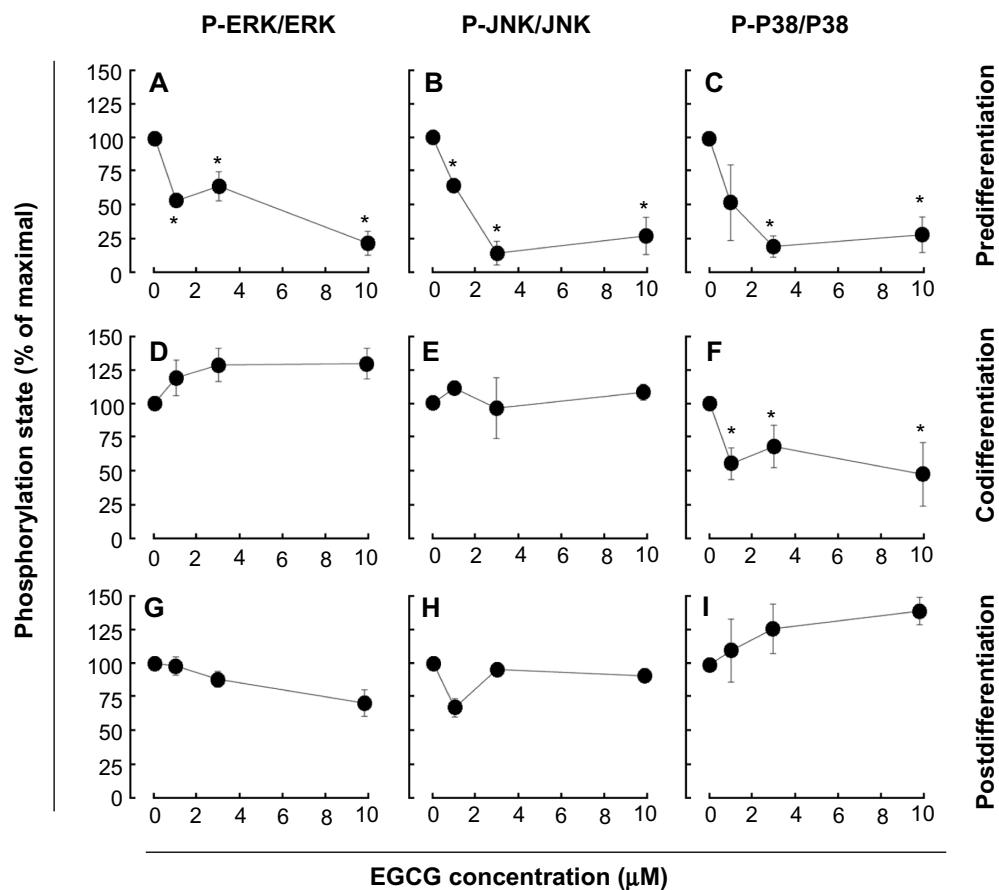


Figure 4 EGCG pretreatment of promyelocytic HL-60 inhibits SIP response in macrophage-differentiated HL-60 cells.

Notes: Three different conditions were used to evaluate the EGCG impact on the SIP response. Predifferentiation conditions: serum-starved HL-60 cells were treated for 24 hours with various concentrations of EGCG (0–10 μM), then treated for 18 hours with 3 μM PMA, to induce HL-60 cell differentiation and adhesion (A–C). Codifferentiation conditions: serum-starved HL-60 cells were treated for 18 hours with either 3 μM PMA alone or with a combination of 3 μM PMA and various EGCG concentrations (D–F). Postdifferentiation conditions: serum-starved HL-60 cells were treated for 18 hours with 3 μM PMA, then adherent cells were treated for 24 hours with various EGCG concentrations (G–I). In all three conditions, adherent cells were then treated with 1 μM SIP for 30 seconds, then cell lysates were isolated, electrophoresed via SDS-PAGE, and immunodetection performed. Scanning densitometry of the autoradiograms was performed, and ratios of P-ERK/ERK, P-JNK/JNK, and P-P38/P38 were obtained. The data shown is representative of three independent experiments.

Abbreviations: EGCG, epigallocatechin-3-gallate; ERK, extracellular signal-regulated kinase; JNK, c-Jun N-terminal kinase; HL, human leukemia; P-P38, phosphorylated P38; P-ERK, phosphorylated ERK; P-JNK, phosphorylated JNK; PMA, phorbol-12-myristate-13-acetate; SDS-PAGE, sodium dodecyl sulfate–polyacrylamide gel electrophoresis; SIP, sphingosine-1-phosphate.

Discussion

The BBB concept, originally suggested to limit access of blood-borne immune cells into the healthy CNS,^{21,22} has now been challenged by demonstration of leukocyte penetration and by increased recruitment of microglia, which resemble immature antigen-presenting cells.²³ Less documented are secreted factors that may contribute to such recruitment and enable modulation of the immunosuppressive environment of glioma. While factors such as transforming growth factor (TGF)- β , COX-2 and interleukin (IL)-10 have been suggested to contribute to the recruitment and expansion of regulatory T cells,²⁴ we now provide new evidence that SIP, which can originate from brain tumor cells where sphingosine kinase 1 is highly expressed, can also be a major actor in the regulation of undifferentiated and terminally differentiated

macrophage functions. Given that recruitment of macrophages plays a critical role in cancer and tumor angiogenesis,^{25,26} we hypothesized that diet-derived molecules, such as the green tea-derived polyphenol EGCG, could sustain a pharmacological intervention targeting cell transformation, affect SIP-mediated signaling through various cell differentiation processes, and ultimately prevent the brain tumor proinflammatory phenotype occurrence.

Chemopreventive assessment of EGCG has, in fact, established this class of diet-derived molecule as a signal transduction inhibitor. In support of this, a screening of 89 nuclear factor (NF)- κB -regulated genes was performed in carcinogen-induced macrophage differentiation of HL-60 cells, where several gene expression profiles were found to be induced during the cell differentiation, immune response,

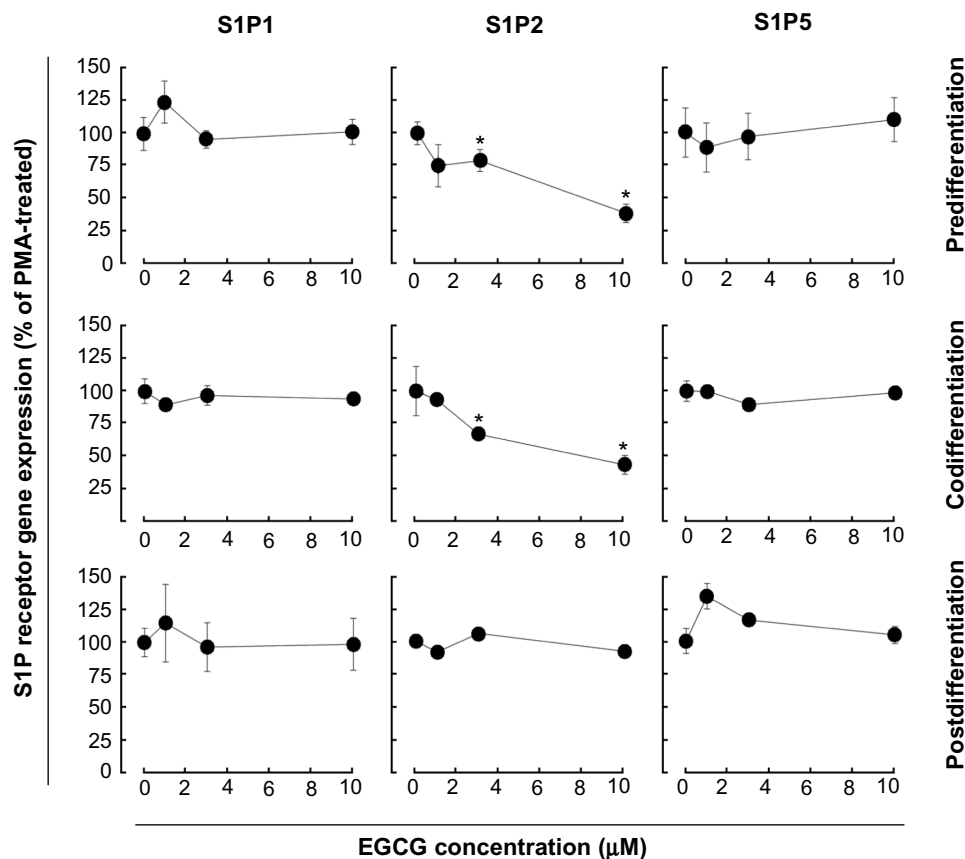


Figure 5 EGCG selectively alters S1P receptor gene expression in macrophage-differentiated HL-60 cells. Three different conditions (predifferentiation, codifferentiation, and postdifferentiation) were used to evaluate EGCG impact on *S1P1*, *S1P2*, and *S1P5* gene expression. In all three conditions, total RNA was extracted from adherent cells, and qRT-PCR performed.

Abbreviations: EGCG, epigallocatechin-3-gallate; HL, human leukemia; qRT-PCR, quantitative reverse-transcription polymerase chain reaction; RNA, ribonucleic acid; S1P, sphingosine-1-phosphate; PMA, phorbol-12-myristate-13-acetate.

and inflammation functions, and to be significantly reversed by EGCG.¹⁸ However, although leukemic cell-derived MMP-9 increased the BBB permeability,¹⁶ EGCG was found inefficient to decrease MMP-9 secretion in terminally differentiated macrophages from HL-60 promyelomonocytic cells,¹⁸ suggesting alternate transendothelial migration and cell response events could be preferentially targeted. In the current study, we confirm that S1P effectively triggers multiple signaling cascades, in agreement with previous reporting,²⁷ in both undifferentiated and terminally differentiated macrophages. We show that the S1P receptor expression profile was significantly inducible upon macrophage differentiation for S1P1, S1P2, and S1P5, with very marginal inductions for S1P3 and S1P4. Finally, this expression profile was found to be altered by EGCG only for S1P2, under both predifferentiation and codifferentiation treatment conditions. More importantly, the inhibition by EGCG of ERK, JNK, and P38 MAPK signaling pathways in the predifferentiation condition could be explained by either direct downstream targeting of each S1P receptor, or by the reduced S1P2

receptor expression that we report (Figure 6A) and which perfectly matches those functions reported for S1P2.²⁷ Interestingly, although S1P2 expression remained reduced in the codifferentiation condition, EGCG was unable to inhibit S1P-mediated ERK and JNK phosphorylation (Figure 6B), suggesting that those two pathways may be relayed through a S1P receptor signaling function²⁷ and their gene expression is not affected by EGCG, as in the case for S1P1 and S1P5 in the current study. S1P2 downstream functions remain controversial as they relay both a repressive signal in macrophage recruitment,²⁸ while they also potentiate human lung fibroblast chemotaxis.²⁹ In fact, the HL-60 promyelomonocytic cell model was chosen in the current study to address the impact of EGCG treatments on the cellular response of a given cell differentiation status. This model is well recognized to reflect the molecular signature of both the macrophage-differentiated and of the undifferentiated circulation leukemia cells. Given that terminally differentiated macrophages (postdifferentiation condition) became unresponsive to EGCG inhibition (Figure 6C), it becomes

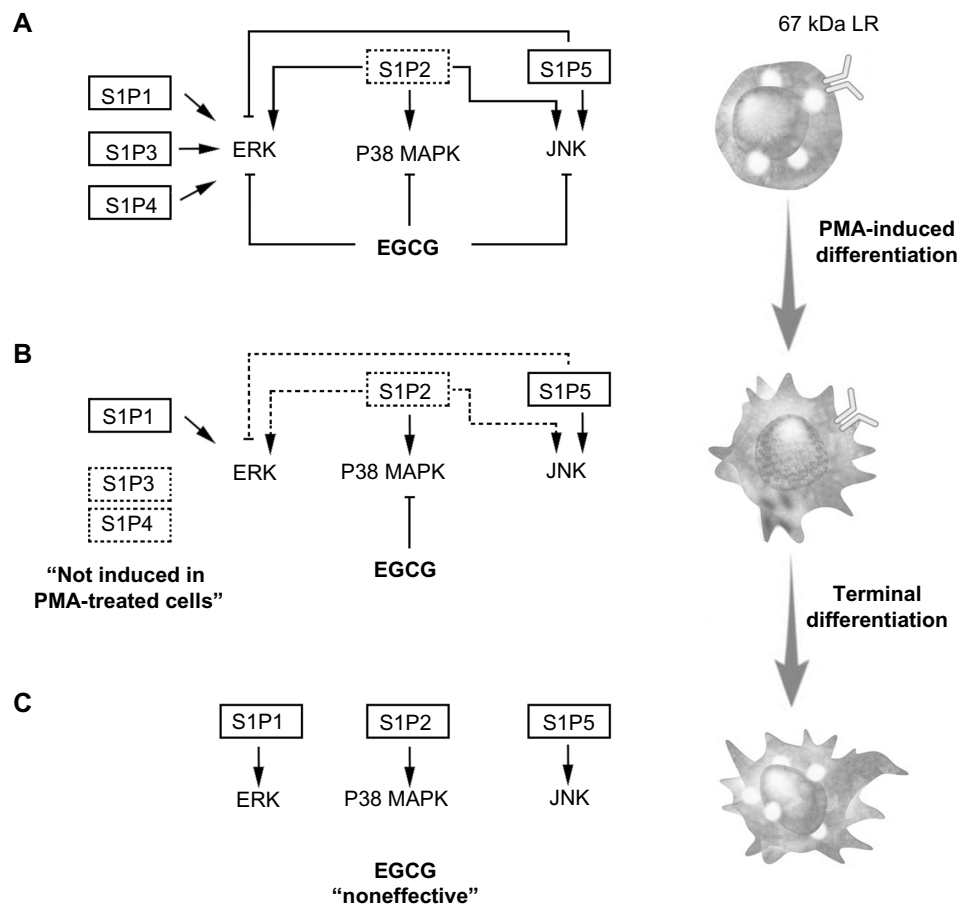


Figure 6 Schematic representation of pretreatment, cotreatment, or postdifferentiation cell treatments with EGCG, on SIP-mediated signaling pathways.

Notes: (A) Undifferentiated HL60 cells express the putative EGCG receptor 67LR as well as all five SIP receptors. SIP triggers phosphorylation of ERK, P38 MAPK, and JNK. When cells were pretreated with EGCG, SIP-mediated signaling was abrogated and *SIP2* gene expression inhibited. (B) Differentiation into adherent macrophages was induced with PMA, and only *SIP1*, *SIP2*, and *SIP5* gene expression was increased. When EGCG was added simultaneously with PMA-induced cell differentiation, *SIP2* gene expression was inhibited, and only SIP-mediated P38 MAPK phosphorylation was abrogated. (C) Once terminal differentiation was achieved, EGCG was ineffective at inhibiting SIP receptor expression or SIP-mediated signaling, partly because the 67LR was no longer expressed.

Abbreviations: 67LR, 67kDa laminin receptor; EGCG, epigallocatechin-3-gallate; ERK, extracellular signal-regulated kinase; JNK, c-Jun N-terminal kinase; P38 MAPK, P38 mitogen-activated protein kinase; PMA, phorbol-12-myristate-13-acetate; SIP, sphingosine-1-phosphate.

tempting to suggest that efficient EGCG impact can only be achieved in a chemopreventive context, where tumor-derived growth factor-mediated chemotaxis, such as can be triggered by SIP, could still be abrogated. Furthermore, given the inability of EGCG to also inhibit MMP-9 expression/function in terminally differentiated HL-60 macrophages, our current data further confirm that such a pharmacological targeting strategy may only affect predifferentiated cells.

SIP fulfills tasks in the immune system through auto- and/or paracrine mechanisms, ultimately regulating macrophage trafficking.^{30,31} Accordingly, our current assessment of SIP responsiveness in all pre-, co-, and postdifferentiation conditions will shed light and impact on events in macrophage biology, such as apoptosis, migration and cell proliferation,³² but will require further investigation. In our cellular model, SIP treatment was reported to not induce macrophage cell death.³² Therefore, the impact

of SIP signaling and of its inhibition by EGCG can be anticipated on mechanisms involved in cell proliferation and/or chemotaxis. Interestingly, modern approaches in cancer therapy are currently aimed at neutralizing cancer stem cell responsiveness to circulating growth factors.³³ To this end, a cluster of differentiation (CD)133(+) cell population isolated from U87 glioblastoma cells was recently shown to be highly responsive to SIP signaling and to require the cooperation of membrane-associated, type-I (MT1)-MMP,³⁴ a cell surface MMP shown to be efficiently targeted by EGCG.^{35,36} Whether such cooperative MT1-MMP to SIP signaling is physiologically relevant was further explored in alternate circulating cell models, such as T cells³⁷ and bone marrow-derived mesenchymal stromal cells,³⁸ where MT1-MMP functions were targeted by EGCG. Evidence possibly supporting such a cooperative MT1-MMP to SIP signaling axis was recently based on the fact that SIP induced the

association of MT1-MMP with p130Cas in endothelial cells³⁹ and that this interaction represents a novel mechanism in the macrophage cell fusion machinery.⁴⁰

Conclusion

The fact that only 5%–10% of all cancer cases are due to genetic defects and that the remaining 90%–95% are due to environment and lifestyle provides opportunities for preventing cancer.¹⁰ Accordingly, among the various phytochemicals identified in fruits, vegetables, spices, and grains that exhibit chemopreventive potential, our current study on the impact of EGCG clearly demonstrates its safety and ability to act in preventive rather than therapeutic settings. Accordingly, its lack of cytotoxicity in undifferentiated HL-60 cells is in good agreement with previous studies.⁴¹ Tea polyphenols have been ascribed the ability to cross the BBB,^{42–44} but given the lack of EGCG action on S1P-mediated signaling in terminally differentiated macrophages (Figure 6C) in our study, one possible explanation may be provided by the downregulation of the putative 67-kDa laminin receptor (67LR) EGCG receptor.¹⁸ Such adaptive processes may, in part, explain the molecular basis for the cancer-preventive activity of EGCG prior to oncogenic cell transformation in response to carcinogens, and will help in the promotion of new diet strategies to prevent brain tumor-associated neuroinflammation consequent, in part, to macrophage infiltration of the CNS.

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Disclosure

The authors report no conflicts of interest in this work.

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