Epigallocatechin gallate has pleiotropic effects on transmembrane signaling by altering the embedding of transmembrane domains

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Epigallocatechin gallate (EGCG) is the principal bioactive ingredient in green tea and has been reported to have many health benefits. EGCG influences multiple signal transduction pathways related to human diseases, including redox, inflammation, cell cycle, and cell adhesion pathways. However, the molecular mechanisms of these varying effects are unclear, limiting further development and utilization of EGCG as a pharmacutical compound. Here, we examined the effect of EGCG on two representative transmembrane signaling receptors, integrin αIIbβ3 and epidermal growth factor receptor (EGFR). We report that EGCG inhibits talin-induced integrin αIIbβ3 activation, but it activates αIIbβ3 in the absence of talin both in a purified system and in cells. This apparent paradox was explained by the fact that the activation state of αIIbβ3 is tightly regulated by the topology of β3 transmembrane domain (TMD); increases or decreases in TMD embedding can activate integrins. Talin increases the embedding of integrin β3 TMD, resulting in integrin activation, whereas we observed here that EGCG decreases the embedding, thus opposing talin-induced integrin activation. In the absence of talin, EGCG decreases the TMD embedding, which can also disrupt the integrin–matrix interaction, leading to integrin activation. EGCG exhibited similar paradoxical behavior in EGFR signaling. EGCG alters the topology of EGFR TMD and activates the receptor in the absence of EGF, but inhibits EGF-induced EGFR activation. Thus, this widely ingested polyphenol exhibits pleiotropic effects on transmembrane signaling by modifying the topology of TMDs.

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Green tea has been one of the most popular drinks for thousands of years, both as a beverage and as an herbal medicine. Indeed, green tea has many clinically reported health benefits, including the prevention of cardiovascular diseases (1, 2) and cancer (3). Studies on the beneficial effects of green tea using cellular or animal models have recently converged on EGCG, the most abundant polyphenol considered as a health-promoting phytonutrient in green tea, and have found EGCG to influence multiple signal transduction pathways related to antioxidation, inflammation, cell cycle, and cell adhesion (4). However, the molecular mechanism underlying those effects has remained elusive. Although EGCG has been suggested to have a number of molecular targets (5), only DNA methyltransferase (6) and the 67-kDa laminin receptor (7) have been demonstrated to directly respond to EGCG in an in vitro system. On the other hand, recent nuclear magnetic resonance spectroscopy studies clearly showed that EGCG can interact with model lipid membranes (8, 9), which implies that biological membrane can be a molecular target of EGCG. Furthermore, the EGCG-lipid interaction can cause a deformation of the lipid bilayer, e.g. by inducing an expansion of the lipid bilayer (10) and/or altering the thickness of the membrane (11). Because membrane-receptor interactions are important in maintaining the proper TMD topology, structures, and hence function of the transmembrane receptors (12, 13), chemical and physical alteration in biological membrane may cause changes in activities of those proteins (14). However, whether such lipid-EGCG interaction contributes to cell signaling pathways and how it could account for the broad effects of EGCG on many cell signaling pathways are yet to be elucidated.

We hypothesized that EGCG exerts its effect on transmembrane receptor signaling by interacting with a lipid bilayer and thereby changing the TMD topology and signaling of a broad spectrum of transmembrane proteins. We tested this hypothesis using integrin αIIbβ3 and EGFR, two prototypical signaling receptors. Integrin αIIbβ3 is a heterodimeric glycoprotein that is a key component of the platelet receptor complex (von Willebrand factor receptor), and its interaction with the extracellular matrix (ECM) is an important step in the platelet aggregation cascade. Integrin αIIbβ3 is also expressed on the surface of endothelial cells, where it plays a crucial role in the formation of platelet–endothelial cell complexes (PECs). PECs are believed to mediate the adhesion of platelets to the endothelial surface, and are thought to play a key role in the initiation and propagation of thrombosis.

EGCG binds to several lipid molecules, including phosphatidylcholine (PC) and phosphatidylethanolamine (PE), and alters their physical properties, such as phase transitions and lipid order parameters. It has been shown that EGCG interacts with model lipid membranes and causes a deformation of the lipid bilayer, leading to changes in membrane thickness and fluidity (8). These changes in membrane properties can affect the topology of transmembrane domains (TMDs) and, consequently, the function of transmembrane receptors.

Epidermal growth factor receptor (EGFR) is a prototypical signaling receptor that is involved in many cellular processes, including cell proliferation, survival, and invasion. EGFR is frequently overexpressed in various types of cancer, and a number of EGFR inhibitors have been developed as potential cancer therapeutics. EGFR signaling is regulated by the topology of the transmembrane domain (TMD), which can be altered by lipid-EGCG interactions.

This study demonstrates that EGCG exerts its pleiotropic effects on transmembrane signaling by altering the embedding of TMDs. These findings suggest that EGCG may be a promising target for the development of new therapeutic strategies for the treatment of various diseases.
Pleiotropic effect of EGCG on activation of integrin αIIbβ3

Although EGCG is reported to have anti-thrombotic effects (23), the addition of EGCG to platelets, the main cellular mediators of arterial thrombosis, causes complex responses. For example, EGCG inhibited aggregation of thrombin-stimulated platelets, but, paradoxically, caused aggregation of unstimulated platelets at the same dose (24). To better understand the physiological role of this widely consumed polyphenol, we first tested the effect of EGCG on activation of recombinant integrin αIIbβ3 in CHO cells where the integrin is normally in a low affinity state; with the addition of increasing concentrations of EGCG, there was a progressive increase in activation as measured by binding of PAC1, a ligand-mimetic, activation-specific integrin antibody. Specific PAC1 binding was calculated in the absence (green line) or presence of purified THD (10 μM) (blue line). Throughout the panels, error bars represent standard errors (n = 3), and analysis of variance multiple comparison using the Bonferroni’s test was performed to test for significant differences between EGCG-treated and non-treated samples. **, p < 0.01; ***, p < 0.001; ****, p < 0.0001.

Results and discussion

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To directly test the effects of EGCG on talin-induced integrin activation, we introduced talin head domain (THD), the integrin-activating talin fragment (18), into CHO cells expressing integrin αIIbβ3 (CHO/αIIbβ3), and then examined the effects of EGCG on the talin-induced activation. EGCG blocked THD-induced αIIbβ3 activation in a dose-dependent manner, whereas EGCG alone induced integrin activation (Fig. 1B).

Because of the complexity of cellular components that might mediate these paradoxical effects of EGCG, we utilized an in vitro reconstitution system in which purified integrin αIIbβ3 was embedded in nanodiscs, islands of 10-nm lipid bilayer encircled by membrane scaffold protein (29). In the reconstituted system, the addition of purified THD can activate the integrin (30). The integrin nanodiscs were first captured to the surface of assay plate coated with anti-integrin β3 extracellular antibody (AP3), and the degree of integrin activation was measured by PAC1 binding to the immobilized integrin nanodiscs (Fig. 1C). EGCG activated the integrin nanodiscs in a dose-dependent manner (Fig. 1C, green line), as it did in cells. The addition of purified THD increases PAC1 binding in the system as shown previously (30), and the THD-induced increase was inhibited by the addition of an increasing amount of EGCG (Fig. 1C, blue line), showing a similar paradoxical effect of EGCG in the purified system as in cells (Fig. 1B).

To examine the effect of EGCG on activated integrins in cells, we utilized αIIbβ3(D723R) mutant, which is activated in a talin-dependent manner in CHO cells (28). The D723R mutation disrupts the electrostatic interaction between αIIb(R995) and β3(D723), weakening the integrin αIIb-β3 TMD interaction and thus favoring the activated state (31). The activating effect of the D723R mutant is dependent upon integrin-talin interactions, as its activation is abolished by disrupting integrin binding to endogenous talin, e.g. by the β3(Y747A) mutation (28). When we added EGCG to the cells expressing αIIbβ3(D723R), in sharp contrast to the activating effect observed with the wild-type integrin, we observed that EGCG induced an initial suppression of activation that peaked at 200 μM EGCG (Fig. 2A). At higher concentrations, however, EGCG induced activation, exhibiting a distinct biphasic effect (Fig. 2A). Next, we tested another activating mutant, αIIbβ3(L712R), in which TMD is predicted to shorten from 29 to 19 amino acids due to the polar residue in the middle of TMD (32). The activating effect of the αIIbβ3(L712R) mutant is talin-independent, as its activation is not affected by the loss of the talin-talin interactions (28, 33). In contrast to αIIbβ3(D723R), EGCG had no significant effect on the L712R mutant (Fig. 2, B and C).

**Opposing changes of integrin β3 TMD topology by EGCG and talin**

To find an explanation for these paradoxical effects, we noted the insensitivity of αIIbβ3(L712R) mutant to EGCG. This mutant activates integrin by shortening the β3 TMD (32), whereas talin does it by increasing the lipid embedding of the β3 TMD (19); both of these changes can alter the β3 TMD tilt angle, thereby disrupting the αIIb-β3 TMD interaction and leading to integrin activation (17). In addition, several studies demonstrated that EGCG can interact with phospholipids and can even decrease the thickness of a lipid bilayer (9–11), which may alter the lipid embedding of TMDs. To test this idea, we investigated whether EGCG can change the embedding of integrin TMD by adapting a β3 TMD embedding assay (19). As EGCG had considerable spectral overlap with bimane, the fluorophore used in the previous study, we used another environment-sensitive fluorophore, mero60, whose fluorescence increases in a more hydrophobic environment and does not overlap with that of EGCG (34). We conjugated the dye to either the N-terminal end (β3(L694C)) or the C-terminal end (β3(L721C)) of β3 TMD and reconstituted the β3 TMD-cytoplasmic tail peptides into phospholipid nanodiscs (Fig. 3A). EGCG decreased the fluorescence of mero60 at either the N-terminal end or the C-terminal end of β3 TMD (Fig. 3, B and C), indicating that EGCG causes both the N-terminal and the C-terminal ends of β3 TMD to become less
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**Figure 3. EGCG induces β3 TMD topological changes in the opposite direction to talin.** A, the environment-sensitive dye, mero60, was conjugated to β3 TMD-tail peptide through the cysteine mutation at Leu⁶⁹⁴ residue to probe the embedding change of β3 TMD at the outer membrane leaflet or at Ile⁷²¹ residue at the inner membrane leaflet. The mero60-labeled β3 TMD-tail peptides were then incorporated into phospholipid nanodiscs. B, EGCG decreases the fluorescence intensity of L694C-mero60 nanodiscs. EGCG is not fluorescent at the wavelength range measured in the analysis. Fluorescence intensities were normalized to the maximum fluorescence intensity in the L694C-mero60 nanodiscs without EGCG. C, the fluorescence of I721C-mero60 nanodiscs was analyzed as in B. D, L694C-mero60 nanodiscs were incubated with 10 μM THD with or without varying concentration of EGCG. EGCG reverses the increase of mero60 fluorescence intensity induced by THD. E, the effect of EGCG on I721C-mero60 nanodiscs was analyzed as in D. F, proposed model of EGCG’s action. Because the association of integrin αIβ and β3 TMD depends on the precise tilt angle of β3 TMD, either increased or decreased embedding of the β3 TMD can disrupt the α-β TMD association and activate the integrin. In some cases, EGCG acts in the opposite direction to talin. When both effects are present, EGCG first neutralizes the effect of talin, but continued reduction in the tilt angle by EGCG activates the integrin.

The figure shows fluorescence intensity changes in response to EGCG and THD treatments. The fluorescence intensity is normalized and plotted against wavelength (nm) for L694C and I721C, with different concentrations of EGCG and THD. The graphs illustrate how EGCG and THD affect the fluorescence intensity of mero60-labeled nanodiscs, indicating changes in the topology of the β3 TMD.

Consistent with our previous report (19) that THD increases the embedding of the β3 TMD domain, THD increased the fluorescence intensities of mero60 conjugated to β3(L694C) or β3(I721C) (Fig. 3, D and E, black dotted lines). The increased fluorescence reflects an altered topology of the β3 TMD (19), which can also disrupt the association of integrin α and β TMDs. Intriguingly, EGCG reversed the THD-induced increase in the fluorescence of mero60 in both β3(L694C) and β3(I721C) (Fig. 3, D and E, green dotted lines), indicating that EGCG alters the topology of the β3 TMD in a manner that opposes the effect of THD (Fig. 3F). We propose that the EGCG-induced change of the β3 TMD topology opposes that induced by talin. Thus, it can offset the talin-induced changes in TMD topology and integrin activation.

The embedding assay using integrin β3 TMD appears more sensitive to EGCG than the integrin activation assay. In the embedding assay, concentrations of EGCG from 25 to 100 μM caused large fluorescence changes both in the absence of talin (Fig. 3, B and C) and in the presence of talin (Fig. 3, D and E), whereas higher concentrations of EGCG were required to affect the integrin activation assay (Fig. 1C). This may be due to the additional stabilizing effects.
**ACCELERATED COMMUNICATION:** *EGCG and transmembrane signaling*

**Figure 4. EGCG has a dual effect on EGFR signaling.** A. HEK293 cells expressing FLAG-tagged human EGFR (HEK/EGFR) were serum-starved overnight and pretreated with different concentrations of EGCG. 30 min after EGCG treatment, cells were stimulated with 50 ng/ml EGF for an additional 30 min. EGFR was immunoprecipitated (IP) and analyzed by Western blotting (WB) with anti-phosphotyrosine antibody (4G10) and anti-EGFR antibody. The degree of phosphotyrosine signal per precipitated EGFR band intensities was normalized to unstimulated control (0%) and EGF-treated control (100%), and shown as a bar graph for each sample. **Left**, effects of EGCG on lipid embedding of EGFR TMD. **Right,** purified EGFR TMD peptide (EGFR(F667C)) labeled with mero60 was reconstituted into nanodiscs, and the effect of EGCG on the embedding of EGFR TMD was analyzed as in Fig. 3B. *Effects of EGCG on transmembrane signaling through a receptor tyrosine kinase*

If the EGCG-induced topological change of integrin β3 TMD is due to an EGCG-lipid interaction, we reasoned that EGCG should also have effects on other transmembrane proteins with signaling functions. To test this hypothesis, we focused on the receptor tyrosine kinases because the dimerization of their TMDs in the lipid bilayer may play a role in activation of those receptors (21, 22). Indeed, EGCG was reported to inhibit activation of receptor tyrosine kinases such as EGFR (35, 36), possibly due to effects on lipid order (36). Pretreatment of HEK293 cells stably expressing EGFR with EGCG inhibited EGF-induced EGFR phosphorylation (Fig. 4A). In contrast, EGCG treatment in the absence of EGF induced EGFR phosphorylation in a concentration-dependent manner (Fig. 4B). Furthermore, EGCG decreased the fluorescence intensity of mero60 attached to the C-terminal region of the nanodisc-embedded EGFR TMD (Fig. 4C). These data suggest that a topological change of EGFR TMD induced by EGF (37) can be reversed by the action of EGCG, and that EGCG-induced topological change in the absence of EGF may favor dimerization of EGFR TMDs, leading to activation.

Our results show that EGCG can change the TMD topologies of receptors and activate those receptors. Conversely, when physiological activation involves shifts in TMD topology, then EGCG can oppose those shifts and inhibit transmembrane signaling. We propose that such a dual effect can account for the conflicting reported effects of EGCG. Recent studies showed that EGCG can bind to lipid bilayers and reside near the phosphate head groups of phospholipids, and that the interaction is further stabilized by cation-π interaction between one of the ring structures in EGCG and the quaternary amine of the phospholipid head group (8). Because the interactions between

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TMDs and phospholipids can influence the topology of TMDs in a lipid bilayer by mechanisms such as snorkeling of the basic lysine side chain into the phosphate head groups of the phospholipids (16), EGCG may alter the interaction of the TMD with the phosphate head group, thus leading to changes in TMD topology. Alternatively, the rigidity of lipids induced by insertion of EGCG into hydrophobic lipid bilayer, as suggested by molecular simulation study (38), or the rigidity of lipid-inserted EGCG itself, due to its less flexible aromatic rings, might alter the tilt angle of TMDs, causing less embedding. Future studies will be required to address these hypotheses; however, our observation that EGCG has a dual effect on transmembrane signaling by modulating lipid embedding of TMDs provides an attractive mechanism to explain some of EGCG’s pleiotropic effects on transmembrane signaling.

Experimental procedures

Reagents, cell lines, and plasmids

1,2-Dimyristoyl-sn-glycery-3-phosphocholine (DMPC) and 1,2-dimyristoyl-sn-glycero-3-phospho-1'-rac-glycerol (DMPG) were purchased from Avanti Polar Lipids, Inc. Membrane scaffold protein (MSPI1D1) was kindly provided by Dr. Stephen Sliker (University of Illinois at Urbana-Champaign). PAC1 and D57 were described previously (16). Anti-FLAG antibody (M2) and anti-phosphotyrosine antibody (4G10) were purchased from Sigma-Aldrich and Merck Millipore, respectively. CHO/α1β cells and CHO/α1β3 cells were generated by infecting CHO cells with lentivirus encoding α1β and/or β3 as described previously (16). HEK/EGFR cells were kindly provided by Dr. Seung-Taek Lee (Yonsei University). The β3 TMD-tail fused with N-terminal His6 and ketosteroid isomerase (KSI) in the pET-31 expression vector was described previously (19). Similarly, the EGFR TMD (Pro637–Gln701) construct containing N-terminal His6 and KSI with cysteine mutation at Phe667 was generated by ligation of the PCR-amplified EGFR TMD region into the pET-31 expression vector.

Flow cytometry

Transfection and flow cytometry were performed with a similar procedure as described before (23). Briefly, CHO or CHO/α1β cells were transfected with various integrin constructs using Lipofectamine LTX and Plus reagents (Life Technologies) or Lipofectamine 2000 (Life Technologies). CHO/α1β3 cells were transfected with a total of 10 µg of plasmids, including 1 µg of tdTomato cDNA as a transfection marker. At 24 h after transfection, cells were detached by trypsinization and treated with EGCG for 10 min. Those cells were stained with PAC1 followed by allophycocyanin-conjugated anti-mouse IgM antibody. When integrin constructs were transfected, cells were co-stained with D57 to gate cells with similar high α1β3 expression.

EGFR phosphorylation assay

Serum-starved subconfluent HEK/EGFR cells were treated with varying concentrations of EGCG for 30 min before EGF treatment (final 50 ng/ml). Cells were lysed by a lysis buffer (20 mM HEPES, pH 7.4, 150 mM NaCl, 1% Triton X-100, 10 mM EDTA, pH 7.4, supplemented with PhosSTOP (Roche Applied Science) and protease inhibitor cocktail (Roche Applied Science)). After clarification by centrifugation at 17,000 × g for 30 min, the clarified lysates were incubated at 4 °C overnight in the presence of 3 µg of anti-FLAG antibody, and the bound proteins were precipitated with protein G-Sepharose. The bound proteins were analyzed by SDS-PAGE and subsequent Western blotting with anti-phosphotyrosine antibody and anti-EGFR antibody.

Expression and purification of TMD peptides

The preparation of integrin nanodiscs was performed essentially as described previously (19). Briefly, His6-KSI-fused TMD proteins were expressed in Escherichia coli BL21(DE3) and purified using a HiTrap Chelating HP column charged with Ni2+. The Asp-Pro bond between KSI and TMD peptide in the purified TMD proteins was cleaved in 10% formic acid for 120 min at 80 °C (24). The resulting TMD peptide was then dialyzed against a buffer containing 50 mM Tris-HCl, pH 7.4, 500 mM NaCl, and 6 mM urea, and then passed through a Ni2+-nitrilotriacetic acid column again to absorb the KSI, leaving the purified TMD peptide in the solution. The TMD peptide was labeled with excess mero60 (1:5 molar ratio). 0.1% Triton X-100 was added to the labeled TMD peptide, and the labeled TMD peptide was then dialyzed extensively against 0.1% Triton X-100 in Tris-buffered saline (20 mM Tris-HCl, pH 7.4, 150 mM NaCl).

Preparation of integrin nanodiscs

The preparation of integrin nanodiscs was previously described (25). Briefly, DMPC and DMPG lipids were dissolved in chloroform or chloroform/methanol mixture, mixed into a 1:1 ratio, and dried onto a glass tube with a steady flow of argon. The lipid mixture was dissolved in 100 mM cholate in TBS. To assemble nanodiscs, 360 µl of the 1:1 lipid mixture (50 mM), 1 ml of 200 µM MSP1D1, and the purified TMD peptides (10 µM) or the purified integrin α1β3 from human platelets (10 µM) were mixed. The mixture was added with 2 volumes of Bio-Beads SM-2 (Bio-Rad) to initiate nanodisc assembly and incubated overnight at room temperature in the dark. The assembled nanodiscs were further purified with a size-exclusion column (HiLoad 16/60 Superdex 200) with TBS as the column buffer. When necessary, the nanodiscs were concentrated using an Amicon Ultra centrifugal filter unit with Ultracel-30 membrane (Millipore).

Fluorescence spectroscopy

200 µl of the purified nanodiscs were mixed with 50 µl of various concentrations of EGCG (and THD in the case of β3 TMD nanodiscs). After 30 min of incubation at room temperature, the emission spectrum (from 605 to 655 nm) at the excitation wavelength, 593 nm, was scanned with a 1-nm interval using a FluoroMax-2 Spectrofluorometer (HORIBA Scientific). The fluorescence from unlabeled TMD peptide, talin, empty nanodiscs, or insertion of EGCG into hydrophobic lipid bilayer, as suggested by molecular simulation study (38), or the rigidity of lipid-inserted EGCG itself, due to its less flexible aromatic rings, might alter the tilt angle of TMDs, causing less embedding. Future studies will be required to address these hypotheses; however, our observation that EGCG has a dual effect on transmembrane signaling by modulating lipid embedding of TMDs provides an attractive mechanism to explain some of EGCG’s pleiotropic effects on transmembrane signaling.

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Author contributions—F. Y., M. H. G., and C. K. designed the research. F. Y., C. Y., and J. K. performed experiments. C. J. M. and K. M. H. synthesized the fluorescent dye. F. Y., C. Y., D. P., M. H. G., and C. K. analyzed the data. F. Y., C. Y., and C. K. wrote the manuscript, which was edited by M. H. G.
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