Epigallocatechin-3-gallate suppresses proinflammatory cytokines and chemokines induced by Toll-like receptor 9 agonists in prostate cancer cells

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Abstract: Chronic inflammation of the prostate contributes to the increased risk of prostate cancer. Microbial pathogens in the prostate cause inflammation that leads to prostatitis and proliferative inflammatory atrophy frequently associated with the development of prostate cancer. Bacterial lipopolysaccharides and DNA mediate immune responses by engaging Toll-like receptor (TLR) 4 and 9, respectively. Synthetic oligodeoxynucleotides containing CpG motifs (CpG-ODN) mimic bacterial DNA and signal through TLR9 to initiate innate immune responses. Here, we show that stimulation of DU145, PC3, or LnCap prostate cancer cells by the TLR9 agonists, CpG-ODN, induces mRNA expression of IL-6, IL-8, CXCL1, IP-10, CCL5, and TGFβ. In addition, activity of matrix metalloproteinase (MMP)-9 and -2 and cell migration increased on CpG-ODN treatment. Induction of cytokines and chemokines was mediated by NF-kB activation and translocation to the nucleus. Treatment with epigallocatechin-3-gallate (EGCG), the major constituent of green tea, prior to CpG-ODN stimulation, inhibits cytokine and chemokine gene induction, activity of MMP-9 and -2, and cell migration. EGCG treatment sequesters the p65 subunit of transcription factor NF-kB in the cytoplasm and inhibits transcriptional activity of the NF-kB-driven promoter in response to CpG-ODN. Our results suggest that the ability of the TLR9 agonists, CpG-ODN, to induce cytokines, chemokines, and MMP activity, as well as suppression by EGCG, are independent of the androgen receptor and p53 status of the cells. EGCG may provide protective effects against inflammation in the prostate and benefit prostate cancer treatment.

Keywords: CpG-ODN, EGCG, inflammation, NF-kB

Introduction
Prostate cancer is the second leading cause of death among men in the US. Age, hormonal, environmental, and genetic factors contribute to the pathogenesis of the disease. Several epidemiological studies, including meta-analyses, suggest an important role for intraprostatic inflammation in the pathogenesis of prostate cancer.1-6 Several case-controlled studies have reported a statistically significant association of prostate cancer with sexually transmitted infections.7,8 In studies of men with advanced and hormone refractory prostate cancer, high plasma concentrations of inflammatory cytokines like interleukin-6 (IL-6), IL-8, IL-1, tumor necrosis factor alpha (TNFα) and interferon gamma (IFNγ) have been reported.9,10 Microorganisms that cause prostatitis are believed to promote chronic inflammation by inflicting cellular damage, cellular hyperproliferation, and increased production of cytokines.11 While cytokines are required to limit tissue damage by replac-
ing damaged cells, angiogenesis, and tissue repair, uncontrolled response promotes the progression from benign prostatic is to intraepithelial neoplasia (PIN) and cancer. Therefore, preventing or reducing inflammation would serve as an attractive mechanism in the chemoprevention of prostate cancer.

Epigallocatechin-3-gallate (EGCG), a major polyphenolic constituent of green tea, has chemopreventive and chemotherapeutic effects in many tumor models. EGCG selectively induces apoptosis and alters the expression of cell cycle proteins and inhibits cell growth in cancer cells compared to normal cells. Increases in the expression of p21, p27, p18, and p16, and a corresponding decrease in cyclin D1, D2, and cyclin E as well as CDK 2, 4, and 6 were observed in cells treated with EGCG. In prostate cancer cells, EGCG treatment reduced the expression of antiapoptotic proteins like phospho-Pi3K and its substrate Akt, thereby limiting cell growth. In addition, EGCG reduces nuclear localization of nuclear factor kappa-light-chain-enhancer of activated B cells (NF-κB), which has significant roles in cell survival and induction of cytokines. Proteolysis of the extracellular matrix components and induction of neovascularure by angiogenesis are required for tumor development. EGCG inhibits the production and activation of vascular endothelial growth factor (VEGF) in breast and colon cancer, and leukemia cells. Proteolytic activity of matrix metalloproteinases (MMPs) in prostate cancer cells is inhibited in response to EGCG treatment.

Toll-like receptors (TLRs) are evolutionarily conserved transmembrane proteins that recognize conserved molecular motifs specific for microbial components. The mammalian TLR family consists of eleven members, each able to detect a distinct pathogen-derived ligand to induce signaling pathways involved in innate immune responses. TLR signaling pathways utilize TIR-domain-containing adaptors like MyD88 (myeloid differentiation primary response 88), TIRAP/MAL (toll-interleukin 1 receptor [TIR] domain containing adaptor protein/MyD88 adapter-like), TRIF (TIR-domain-containing adapter-inducing IFNβ), and TRAM (translocating chain-associated membrane protein), and differentially use transcription factors like NF-κB or interferon regulatory transcription factor (IRFs) to fine-tune specific responses. Bacterial components such as Gram-negative lipopolysaccharide (LPS) are recognized by TLR4, whereas bacterial DNA stimulates innate immune responses by engaging TLR9. It has been shown that viral or bacterial DNA contain unmethylated CpG dinucleotide motifs (CpG motifs) that have potent immunostimulatory activity. There are two types of CpG DNA that are structurally distinct: type A has greater ability to induce interferon α (IFNα) from plasmacytoid dendritic cells, and type B, which is more conventional and induces inflammatory cytokines. Both types recognize and bind TLR9 to exert antimicrobial responses. On binding a ligand, TLR9 associates with the adaptor, MyD88, and recruits IRAK and TRAF6 to activate the IkB kinase complex (IKK). Phosphorylation of IkB by the IKK complex causes nuclear translocation of NF-κB, which induces expression of inflammatory cytokines. Synthetic phosphorothioate-stabilized oligonucleotides containing the CpG motifs (CpG-ODN) mimic the immunostimulatory activity of microbial DNA and are recognized by TLR9. Expression of TLR9 is not limited to immune cells. Normal epithelial and tumor cells express TLR9, and stimulation with ligands induces production of pro-inflammatory cytokines. TLR9 agonists have been shown to promote migration of breast cancer cells, astrocytes, and prostate cancer cells.

Persistent infection in the prostate contributes microbial components, which can induce production of cytokines initiating inflammation. Bacterial or viral unmethylated CpG-DNA released by dying or proliferating pathogens is recognized by TLR9 to initiate immune responses, which in turn can alter the tumor microenvironment. In this study, we compared the role of the TLR9 agonists, CpG-ODN, in cytokine induction in DU145, PC3, and LnCap prostate cancer cells, and examined the effect of EGCG in suppressing cytokine induction. Our results show that stimulation of prostate cancer cells with CpG-ODN induces messenger ribonucleic acid (mRNA) expression of cytokines and chemokines. In addition, the activity of MMP-9 and -2 are increased over basal levels of expression on CpG-ODN treatment, which correlates with increased migration. The induction of cytokine gene transcription is dependent on NF-κB activation and translocation to the nucleus. EGCG treatment inhibits both cytokine gene induction and activity of MMP-9 and -2 by inhibiting translocation of NF-κB to the nucleus and transcriptional activity of the NF-κB-driven promoter in response to TLR9 agonists. Our results have compared cytokine gene induction and MMP activity in three prostate cancer cell lines. DU145 and PC3 are androgen-independent cell lines and LnCap expresses androgen receptor (AR) and is androgen-dependent. In addition, DU145 cells express mutant p53, PC3 cells lack p53 expression, and LnCap cells express functional p53 protein. Our results suggest that the ability of CpG-ODN to induce cytokines, chemokines, and MMPs, and the suppression by EGCG, is independent of the AR and p53 status of the cells, which is an important property of chemoprevention for prostate cancer treatment.

Materials and methods

Chemicals, reagents, and antibodies

Chemicals, unless indicated otherwise, were obtained from Sigma-Aldrich Co, (St Louis, MO, USA). Antibodies to NF-κB...
p65 (C-20) used for immunoblotting and immunofluorescence and 1xIBα were from Santa Cruz Biotechnology Inc., (Dallas, TX, USA). Antibody to β-actin was from Sigma-Aldrich Co, and Histone H2A was from EMD Millipore (Billerica, MA, USA). Anti-mouse immunoglobulin G (IgG) and anti-rabbit IgG horseradish peroxidase (HRP) linked secondary antibodies were from Cell Signaling, Inc., (Danvers, MA, USA) and enhanced chemiluminescence reagents were from GE Healthcare Bio-Sciences Corp (Piscataway, NJ, USA). EGCG was purchased from Cayman Chemical Co, (Ann Arbor, MI, USA). Phosphorothioate-modified, human-specific, class B CpG-ODN (5′-TCGTTTTTTGCTGTTTTGTCGTT-3′) was obtained from InvivoGen Corporation (San Diego, CA, USA) and dissolved in endotoxin-free sterile water as per the manufacturer’s instructions and used at the indicated concentrations. TRIZol reagent, Alexa Fluor® 488 Donkey Anti-Rabbit IgG (H + L) antibody, and Lipofectamine 2000 were from Invitrogen (Thermo Fisher Scientific, Waltham, MA, USA). The luciferase construct containing five copies of NF-κB response elements linked to firefly luciferase complementary DNA (cDNA) (pNF-κB-Luc) was obtained from Stratagene (Agilent Technologies, Santa Clara, CA, USA). Dual luciferase reporter assay reagents and CellTiter 96® AQueous One Solution Cell Proliferation were from Promega Corporation (Fitchburg, WI, USA).

Cell culture and treatment of cells
The human prostate cancer cell lines DU145, PC3 (kindly provided by Aimin Zhou, Cleveland State University, OH, USA), and LnCap (kindly provided by William Taylor, University of Toledo, OH, USA) were grown in Roswell Park Memorial Institute (RPMI) 1640 medium supplemented with streptomycin (100 µg/mL), penicillin (100 units/mL), 2 mmol/L glutamine, and 10% fetal bovine serum (FBS; Invitrogen). Cells were maintained in 95% air, 5% CO₂ at 37°C. EGCG was dissolved in dimethyl sulfoxide (DMSO) at 20 mg/mL, and aliquoted and stored at −20°C. Cells were treated at the indicated dose and time in complete RPMI medium. In some experiments, cells were treated with Bay11-7082 (10 µM; Santa Cruz Biotechnology) for 1 hour at 37°C followed by transfection with CpG-ODN and analyzed. Cells that were used as controls (mock) were incubated with vehicle only.

Cell viability assays
The viability of cells was determined using the CellTiter 96 AQueous Cell Proliferation Assay (3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide [MTT] assay; Promega). Briefly, 2×10⁴ cells were seeded in a 96-well culture plate and treated with varying concentrations of EGCG or CpG-ODN (1 µM). At indicated times after treatment, MTT reagent was added to each well and absorbance was measured at 490 nm with a plate reader (Spectra Max 340; Molecular Devices, LLC, Sunnyvale, CA, USA). Cell viability was normalized to untreated cells. Experiments were performed in triplicate and standard deviation (SD) was calculated.

RNA isolation and quantitative reverse transcription polymerase chain reaction
RNA was isolated using TRIzol reagent as per the manufacturer’s instructions and used for cDNA synthesis using random decamers and a RETROscript cDNA synthesis kit (Life Technologies; Thermo Fisher Scientific, Waltham, MA, USA). Expression of cytokines and chemokines was determined by quantitative reverse transcription polymerase chain reaction (qRT-PCR) using SYBR Green PCR Master Mix (Bio-Rad Laboratories Inc., Hercules, CA, USA) using the gene-specific primers listed below and normalized to GAPDH expression. The primers used are: IL-8F 5′ ACC ACA CTG CGA CGA CAC AGA AAT 3′, IL-8R 5′ AAA CTT CTC CAC AAC CCT CTG CAC 3′, CXCL1 F 5′ TCC AAA GTG TGA ACG TGA AGT CCC 3′, CXCL1 R 5′ CAA GCT TTC CGC CCA TTC TTG AGT 3′, CCL5 R 5′ TCT TTC CGC CCA TTC TTG AGT 3′, TGFβ F 5′ CGA GAG GAG CGA GGA AGA GT 3′, TGFβ R 5′ AGG GCG GCA TGT CTA TTT TG 3′, CCL5F 5′ TTT CTA CAC CAG TGG CAA GTG CTC 3′, CCL5 R 5′ TCT TCT CTG GGT TGG CAC ACA CTT 3′, IL-6F 5′ TGT GAA AGC AGC AAA GAG GCA CTG 3′, IL-6R 5′ CAC CAG GCA AGT CTC CTC ATT GAA 3′, IP-10F 5′ ACC GTA CGC TGT ACC TGC AT 3′, IP-10R 5′ TCT TGA TGG CCT CTG ATT CT 3′, GAPDH 5′ TCG ACA GTC AGC CGC ATC TTC TTT 3′, and GAPDH 5′ ACC AAA TCC GTT GAC TCC GAC CTT 3′.

Cell lysates and immunoblotting
Cells were lysed in buffer and separated into nuclear and cytoplasmic extracts using a nuclear/cytosol fractionation kit (MBL International Corporation, Woburn, MA, USA) using the protocol provided by the manufacturer. Protein concentrations in the supernatants were determined using the protein assay (Bio-Rad Laboratories Inc., Hercules, CA, USA) and β-actin was from Sigma-Aldrich Co, and Histone H2A was from EMD Millipore (Billerica, MA, USA). Antibody to β-actin was from Sigma-Aldrich Co, and Histone H2A was from EMD Millipore (Billerica, MA, USA) and enhanced chemiluminescence reagents were from GE Healthcare Bio-Sciences Corp (Piscataway, NJ, USA). EGCG was purchased from Cayman Chemical Co, (Ann Arbor, MI, USA). Phosphorothioate-modified, human-specific, class B CpG-ODN (5′-TCGTTTTTTGCTGTTTTGTCGTT-3′) was obtained from InvivoGen Corporation (San Diego, CA, USA) and dissolved in endotoxin-free sterile water as per the manufacturer’s instructions and used at the indicated concentrations. TRIZol reagent, Alexa Fluor® 488 Donkey Anti-Rabbit IgG (H + L) antibody, and Lipofectamine 2000 were from Invitrogen (Thermo Fisher Scientific, Waltham, MA, USA). The luciferase construct containing five copies of NF-κB response elements linked to firefly luciferase complementary DNA (cDNA) (pNF-κB-Luc) was obtained from Stratagene (Agilent Technologies, Santa Clara, CA, USA). Dual luciferase reporter assay reagents and CellTiter 96® AQueous One Solution Cell Proliferation were from Promega Corporation (Fitchburg, WI, USA).

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Cell lysates and immunoblotting
Cells were lysed in buffer and separated into nuclear and cytoplasmic extracts using a nuclear/cytosol fractionation kit (MBL International Corporation, Woburn, MA, USA) using the protocol provided by the manufacturer. Protein concentrations in the supernatants were determined using bovine serum albumin as a standard (Bio-Rad protein assay kit). The protein in the cell lysates were separated in 10% or 15% polyacrylamide/sodium dodecyl sulfate (SDS) gels and transferred to a nitrocellulose membrane (Bio-Rad).
Membranes were probed with different primary antibodies according to the manufacturer’s protocols. The membranes were washed with Tris-buffered saline with 1% Tween 20 and incubated with goat anti-mouse or goat anti-rabbit antibody tagged with HRP for 1 hour. Immunoreactive bands were detected by enhanced chemiluminescence (GE Healthcare Bio-Sciences Corp).

**Luciferase reporter assays for NF-κB**

DU145 or PC3 cells (2×10⁴ cells per well) were cotransfected with 1.0 μg of pNF-κB-Luc plasmid along with 0.1 μg of pRL-TK (Renilla luciferase vector) using lipofectamine 2000 reagent according to the protocol provided by the manufacturers. The IL-8 promoter fused to luciferase (IL-8 WT-Luc) and constructs with mutations in NF-κB (IL-8mNF-κB) or AP-1 binding sites (IL-8mAP-1; kindly provided by Dr Gary Wu, University of Pennsylvania, PA, USA) were cotransfected with the pRL-TK plasmid. After 24 hours, some samples were transfected with CpG-ODN (1 μM) using lipofectamine 2000. EGCG (40 μg/mL) was added to some experiments prior to CpG-ODN transfection. Cells were lysed in lysis buffer after 16–18 hours and luciferase activity was determined and normalized to Renilla luciferase activity using a dual luciferase assay kit (Promega).

**NF-κB nuclear translocation by immunofluorescence assay**

DU145 cells were grown on glass coverslips and mock treated or treated with CpG-ODN (1 μM) or TNFα (10 ng/mL; R&D Systems, Inc., Minneapolis, MN, USA) for 1 hour with or without pretreatment with EGCG for 24 hours. Cells were rinsed with phosphate-buffered saline (PBS) and permeabilized with 0.1% Triton-X100 for 15 minutes. After 1 hour of incubation with NF-κB p65 subunit antibody (1:250), cells were washed once with 0.01% Tween 20 followed by two washes with PBS. Cells were incubated for 1 hour with Alexa Fluor 488 donkey anti-rabbit secondary antibody (1:400). Cells were washed with 0.01% Tween 20 and PBS, mounted in VECTASHIELD® with 4',6-diamidino-2-phenylindole (DAPI; Vector Labs, Burlingame, CA, USA). Cells were imaged with a Leica CS SP5 multi-photon laser scanning confocal microscope (Leica Microsystems, Weitzler, Germany).

**MMP-9 and -2 gelatin zymography**

Activity of MMP-9 and -2 was determined in culture supernatants by gelatin zymography. DU145, PC3, or LnCaP cells (2×10⁴) were treated as indicated with CpG-ODN (1 μM) with or without EGCG (40 μg/mL) pretreatment. Culture supernatant was electrophoresed in 8% SDS-polyacrylamide gel electrophoresis (PAGE) containing 0.1% gelatin (Sigma-Aldrich Co). Gels were soaked in 2.5% Triton-X100 for 2 hours, followed by incubation in digestion buffer (10 mM CaCl₂, 50 mM Tris, pH 7.4, 1% Triton-X100) for 20 hours at 37°C. The gels were then stained with Coomassie Brilliant Blue R-250 (0.2% in 40% methanol, 10% acetic acid) and destained in 20% methanol, 10% acetic acid solution. Clear bands representing MMP-9 and -2 activity were imaged.

**Cell migration assay**

Transwell cell migration assays were performed using a modified Boyden chamber (Corning Inc., Corning, NY, USA) containing a fibronectin-coated polycarbonate membrane filter (6.5 mm diameter, 8 μm pore size). DU145 or PC3 cells (2×10⁴) were pretreated or not with EGCG (40 μg/mL) for 24 hours followed by transfection with CpG-ODN (1 μM). Cells were plated in the upper chamber and the lower chamber contained culture medium with 10% FBS. Cells were incubated for 24 hours at 37°C in 5% CO₂. Nonmigrated cells were scraped from the upper surface of the membrane with a cotton swab, and migrated cells remaining on the bottom surface were trypsinized and counted with a hemocytometer.

**Statistical analysis**

All values are presented as mean± SD. Student’s t-tests were used for determining statistical significance between groups. *P<0.05 was considered significant.

**Results**

**Effect of CpG-ODN and EGCG on viability of prostate cancer cells**

CpG-ODN, like bacterial DNA, has substantial immunostimulatory effects and serves as TLR9 agonists. The prostate cancer cells DU145, PC3, and LnCaP cells have been shown to express TLR9 and its adaptor, MyD88. To determine if CpG-ODN affects the viability of prostate cancer cells, DU145, PC3, and LnCaP cells were transfected with varying concentrations of CpG-ODN (1 μM or 10 μM) or 1 μM of CpG-ODN for 8 or 24 hours using lipofectamine 2000. Cell viability was determined by MTT colorimetric assays (Figure 1A and B). A high concentration (10 μM) of CpG-ODN transfection induced significant cell death in all three cell lines. LnCaP cells showed more resistance compared to DU145 and PC3 cells, whereas treatment with 1 μM of CpG-ODN had very little effect on viability; 85%-91% of the cells
were viable after 24 hours of treatment. All experiments were therefore performed with 1 µM CpG-ODN, which induced less cytopathicity.

EGCG inhibits the growth of several types of cancer cells, including prostate cancer cells. To determine the effect of EGCG on DU145, PC3, or LnCap cells, we treated cells with 20–100 µg/mL EGCG for 24 hours. Cell viability was determined by MTT colorimetric assays (Figure 2). At 40 µg/mL, 79% of DU145 cells and 85% of PC3 or LnCap cells were viable. A high concentration of EGCG (100 µg/mL) reduced cell viability of DU145 cells to 51%, compared to 61% of PC3 and 76% of LnCap cells. These results indicate that EGCG inhibits cell growth of the prostate cancer cell lines tested in a dose-dependent manner.

**CpG-ODN induces expression of proinflammatory cytokines and chemokines**

To determine if CpG-ODN can induce proinflammatory cytokines and chemokines in DU145, PC3, and LnCap cells, CpG-ODN (1 µM) was transfected for 6 or 24 hours. Levels of mRNAs for IL-6, IL-8, CXCL1, IP-10, CCL5, and TGFβ were determined by qRT-PCR and normalized to GAPDH levels. Induction of IL-6 and IL-8 followed similar kinetics in all the cell lines (Figure 3). CXCL1 levels dropped significantly 24 hours postinduction in LnCap cells compared to DU145 and PC3 cells. In PC3 cells, the kinetics of induction of IP-10, CCL5, and TGFβ were significantly different compared those in DU145 and LnCap cells. These results suggest that CpG-ODN induced proinflammatory cytokines and chemokines in prostate cancer cells, albeit with different kinetics involving a mechanism that is independent of expression of AR and p53 status of the cells.

**EGCG suppresses proinflammatory cytokines induced by CpG-ODN**

EGCG modulates multiple signal transduction pathways, providing growth inhibitory effects. To investigate if EGCG regulates CpG-ODN-induced gene expression, prostate cancer cells were pretreated with EGCG (40 µg/mL) for 24 hours followed by transfection with CpG-ODN (1 µM) for 6 or 24 hours. Expression of mRNA for cytokines and chemokines
Figure 3 CpG-ODN stimulation induces the expression of proinflammatory cytokines and chemokines in prostate cancer cells.

Notes: DU145, PC3, and LnCap cells were transfected with CpG-ODN (1 μM) for 6 or 24 hours. Total RNA was prepared from each sample, converted to cDNA, and used for qRT-PCR in triplicate using the following gene-specific primers: (A) IL-6, (B) IL-8, (C) CXCL1, (D) IFN-γ, (E) CCL5, and (F) TGFβ1. RNA expression was normalized to GAPDH mRNA levels. The data shown are means ± SD of experiments performed in triplicate. Student’s t-test: *P<0.05; **P<0.01; #P>0.05; not significant.

Abbreviations: CpG-ODN, CpG oligodeoxynucleotides; qRT-PCR, quantitative reverse transcription polymerase chain reaction; SD, standard deviation; cDNA, complementary DNA; mRNA, messenger RNA; GAPDH, glyceraldehyde 3-phosphate dehydrogenase; IL, interleukin; CXCL1, chemokine (C-X-C motif) ligand 1; IFN-γ, interferon gamma-induced protein 10; TGFβ1, tumor growth factor beta 1; CCL5, chemokine (C-C motif) ligand 5.

was determined by RNA isolation and qRT-PCR, and compared to cytokine gene induction without EGCG treatment (Figure 3). Our results show that EGCG potently inhibits gene induction of most cytokines in response to CpG-ODN in all three prostate cancer cell lines. However, suppression of IL-8 and TGFβ1 induction in DU145 cells and CCL5 in PC3 was not significant at 24 hours. Similarly, no significant suppression of TGFβ1 in PC3 cells was observed at 6 hours.
Effect of EGCG on CpG-ODN-induced NF-κB activation

TLR9 agonists signal through the adaptor protein MyD88 and activate transcription factor NF-κB to induce target gene expression. Other studies have shown that CpG-ODN activates NF-κB signaling pathways in several cancer cell lines. To determine the role of NF-κB in gene induction of chemokines and cytokines by CpG-ODN, DU145 or PC3 cells were pretreated with a pharmacological inhibitor of NF-κB and Bay 11-7083 (10 μM), and a level of mRNAs as determined by RNA isolation and qRT-PCR (Figure 4). Inhibiting the activity of NF-κB potently reduced mRNA levels induced by CpG-ODN suggesting requirement of NF-κB transcriptional activity. We further evaluated the role of NF-κB more specifically in the induction of IL-8 by CpG-ODN in promoter luciferase assays. The activity of the −135bp promoter containing NF-κB and AP-1 transcription factor binding sites fused to luciferase cDNA was determined in response to CpG-ODN transfection. To determine the contribution of NF-κB or AP-1 to gene induction, promoter constructs with point mutations abolishing the binding of mutant NF-κB (mNF-κB) or mutant AP-1 (mAP-1) were tested in luciferase assays (Figure 5). Loss of the NF-κB binding site reduced induction of IL-8 in both DU145 and PC3 cells, whereas the AP-1 site mutation did not significantly affect IL-8 gene induction by CpG-ODN.

To further determine if CpG-ODN stimulated NF-κB transcriptional activity, we cotransfected DU145 or PC3 cells with promoter constructs containing five copies of NF-κB binding sites fused to luciferase cDNA along with Renilla luciferase vector for normalization. We observed an increase in NF-κB-dependent luciferase activity following transfection with CpG-ODN (Figure 6). Further, pretreatment with EGCG for 24 hours prior to CpG-ODN transfection resulted in a decrease in NF-κB-dependent transcriptional activity; the suppression was greater in DU145 cells compared to PC3 cells.

CpG-ODN-induced nuclear translocation of NF-κB

NF-κB is made of p65 and p50 subunits that are sequestered in the cytoplasm in an inactive form through interaction with an inhibitory subunit, IκBα. Phosphorylation of IκBα by the upstream kinases, IKKs, leads to rapid ubiquitination and degradation of IκBα by the 26S proteasome system, which allows NF-κB to translocate to the nucleus to activate expression of target genes. Our results along with others demonstrate that EGCG inhibits activation of NF-κB and prevents degradation of IκBα. As shown in Figure 7, in mock-treated DU145 cells, NF-κB is predominantly cytoplasmic. Transfection with CpG-ODN for 1 hour resulted in translocation of NF-κB to the nucleus, similar to treatment with TNFα. Pretreatment with EGCG inhibited CpG-ODN-induced translocation of NF-κB, which accumulates in the cytoplasm. We further evaluated the translocation of NF-κB in nuclear and cytosolic extracts of cells treated with CpG-ODN with or without EGCG pretreatment. On immunoblots, we observed less cytosolic NF-κBp65 protein in CpG-ODN-treated cells and more in the nuclear fraction (Figure 8). Cytosolic levels of IκBα are low as would be expected in cells where phosphorylation byIKK

Figure 4 Effect of inhibition of NF-κB on CpG-ODN-induced expression of chemokines and cytokines.
Notes: DU145 and PC3 cells were pretreated or not (No Inh) with the NF-κB inhibitor, Bay11-7083 (10 μM) for 1 hour prior to transfection with CpG-ODN (1 μM) for 6 hours. Total RNA was prepared from each sample, converted to cDNA, and used for qRT-PCR in triplicate using the following gene-specific primers: IL-6, IL-8, CXCL1, IP-10, CCL5, and TGFβ1. RNA expression was normalized to GADPH mRNA levels. The data shown are means ± SD of experiments performed in triplicate. Student’s t-test: *P < 0.001; **P < 0.05.
Abbreviations: NF-κB, nuclear factor kappa-light-chain-enhancer of activated B cells; CpG-ODN, CpG oligodeoxynucleotides; qRT-PCR, quantitative reverse transcription polymerase chain reaction; RLU, relative luciferase units; SD, standard deviation; IL, interleukin; CXCL1, chemokine (C-X-C motif) ligand 1; IP-10, interferon-gamma-induced protein 10; TGFβ1, tumor growth factor beta; cDNA, complementary DNA; GADPH, glyceraldehyde 3-phosphate dehydrogenase; mRNA, messenger RNA; Inh, inhibitor.
Figures 5 and 6: Induction of the IL-8 promoter by CpG-ODN requires NF-κB transcriptional activity.

Notes: (A) DU145 and (B) PC3 cells were cotransfected with 1.0 µg of IL-8 promoter fused to luciferase (IL-8 WT-Luc) or constructs with mutations in NF-κB (IL-8mκB) or AP-1 binding sites (IL-8mAP-1), along with 0.1 µg of Renilla luciferase plasmid. After 24 hours, cells were transfected with CpG-ODN (1 µM) with or without EGCG pretreatment (40 µg/mL) for 24 hours. IL-8-luciferase activity was normalized to Renilla luciferase activity and represented as fold induction. Data shown are means ± SD of experiments performed in triplicate. Student’s t-test: values as indicated.

Abbreviations: CpG-ODN, CpG oligodeoxynucleotides; EGCG, epigallocatechin-3-gallate; RLU, relative light units; SD, standard deviation; IL, interleukin; NF-κB, nuclear factor kappa-light-chain-enhancer of activated B cells; WT, wild-type.

Figure 7: EGCG inhibits CpG-ODN-induced nuclear translocation of NF-κB.

Notes: DU145 cells were grown on glass cover slips and mock treated or transfected with CpG-ODN (1 µM) for 1 hour with or without EGCG pretreatment (40 µg/mL) for 24 hours. Cells were incubated with goat anti-rabbit polyclonal NF-κB p65 subunit antibody (green) for 1 hour followed by alexa Fluor 488 donkey anti-rabbit secondary antibody and mounted in Vectashield® (Vector Labs, Burlingame, CA, USA) with DAPI to stain nuclei (blue). Images were merged to determine nuclear localization. Representative images of cells visualized under confocal microscope at 60× are shown.

Abbreviations: CpG-ODN, CpG oligodeoxynucleotides; DAPI, 4′,6-diamidino-2-phenylindole; EGCG, epigallocatechin-3-gallate; NF-κB, nuclear factor kappa-light-chain-enhancer of activated B cells; TNF, tumor necrosis factor.
would lead to degradation by the proteosomal system (Figure 8). However, increase in cytosolic accumulation of the p65 subunit was observed in cytosolic lysates of cells pretreated with EGCG prior to CpG-ODN transfection. EGCG has been shown to inhibit degradation of IκBα in many cells, including both normal and tumor cells, by inhibiting phosphorylation of IκBα.16 We observed inhibition of IκBα degradation in cells pretreated with EGCG prior to CpG-ODN as well as EGCG alone. The accumulation of the p65 subunit was more significant in EGCG treatment alone. Our results show that EGCG suppresses the induction of cytokines and chemokines induced by CpG-ODN by inhibiting the transcriptional activity of NF-κB.

CpG-ODN induces MMP activity and cell migration

TLR9 agonists have been shown to promote invasion of prostate cancer cells and MMP-13 activity in breast cancer cells. MMPs degrade the extracellular matrix and promote cell invasion. To test if CpG-ODN regulates MMP activity, in addition to inducing proinflammatory cytokines, culture supernatants of prostate cancer cells transfected with CpG-ODN with or without EGCG pretreatment was assessed by gelatin zymography. Both PC3 and DU145 cells produce significant amounts of MMP-9 and MMP-2, compared to LnCap cells. Treatment with CpG-ODN induced MMP-9 and -2 activity 2.5-fold each in DU145 and LnCap cells and 3.5-fold in PC3 cells (Figure 9). In bladder cancer cells and pancreatic cancer cells, EGCG inhibits the activity of MMP-9 by inhibiting NF-κB activity.56–58 Our results show that pretreatment with EGCG inhibited the induction of MMP-9 and -2 by CpG-ODN, but did not alter the basal level of MMP-9 and -2 activities in all three prostate cancer cell lines.

MMP activity causes remodeling of cell–matrix and cell–cell interactions and promotes cell migration. To determine if the upregulation of MMP-9 and -2 activity by CpG-ODN translates into increased migration, DU145 or PC3 cells were pretreated or not with EGCG followed by CpG-ODN transfection. Substantially higher numbers of cells treated with CpG-ODN migrated across the membrane, which correlates with an increase in MMP-9 and -2 activity (Figure 10). Cells pretreated with EGCG showed significant inhibition of migration and MMP-9 and -2 activity. EGCG suppressed

Figure 8 EGCG sequesters NF-κB p65 subunit in the cytosol.

Notes: DU145 cells were transfected with CpG-ODN (1 μM) for 1 hour with or without EGCG pretreatment (40 μg/mL) for 24 hours. Levels of NF-κB p65 subunit in the (A) nuclear extract normalized to histone H2A or (B) cytosol normalized to β-actin levels was determined by immunoblot analysis and compared to levels of inhibitory subunit IκBα.

Abbreviations: CpG-ODN, Cp oligodeoxynucleotides; EGCG, epigallocatechin-3-gallate; NF-κB, nuclear factor kappa-light-chain-enhancer of activated B cells.

Figure 9 EGCG inhibits CpG-ODN-induced activity of MMP-9 and -2.

Notes: DU145, PC3, and LnCap cells were transfected with CpG-ODN (1 μM) for 8 hours with or without EGCG pretreatment (40 μg/mL) for 24 hours. Activity of MMP-9 and -2 in culture supernatants was determined by electrophoresis in 8% SDS-PAGE gels containing 0.1% gelatin. Gels were stained and clear areas representing MMP activity were imaged. Results are representative of three independent experiments.

Abbreviations: CpG-ODN, Cp oligodeoxynucleotides; EGCG, epigallocatechin-3-gallate; MMP, matrix metalloproteinase; SDS-PAGE, sodium dodecyl sulfate polyacrylamide gel electrophoresis.
both MMP-9 and -2 activity and migration of cells induced by CpG-ODN.

**Discussion**

Inflammation caused by microbial pathogens, host genetics, and the environment contribute to the development of prostate cancer. Persistent inflammation causes increased proliferation to replace damaged tissue accompanied by elaborating production of cytokines, which alters the prostatic microenvironment and promotes tumorigenesis. Host innate immune response allows recognition of invading pathogens by TLRs to activate inflammatory and antimicrobial responses. The inflammatory response is transmitted by the TLR–MyD88 pathway, leading to activation of NF-κB and AP-1 responsive genes. Bacterial DNA, contributed by infection of the prostate, is a potent immunostimulator and produces inflammatory cytokines by engaging TLR9 and its adaptor, MyD88. Synthetic CpG-ODN mimics bacterial DNA and has been shown to induce TLR9-dependent cytokines in many cell lines. In this study, we have evaluated the role of the TLR9 agonist, CpG-ODN, in inducing proinflammatory cytokines and MMPs in DU145, PC3, and LnCap prostate cancer cells. Further, we determined the role of EGCG, a major constituent of green tea, in the suppression of cytokines and MMPs and its antiproliferative effects in prostate cancer cells. Our results presented here, demonstrate that CpG-ODN induces mRNA expression of IL-6, IL-8, CXCL1, IP-10, CCL5, and TGFβ with variable kinetics in all three prostate cancer cell lines. Induction of the mRNAs involved the activity of NF-κB, since the pharmacological inhibitor Bay11-7082 significantly decreased mRNA levels. Analysis of the induction of the IL-8 promoter in response to CpG-ODN using constructs with mutations in the NF-κB or AP-1 binding site shows the requirement of the NF-κB binding site. Activity of MMP-9 and -2 was induced by CpG-ODN 2.5- to 3.5-fold over the basal levels expressed in these cells, which correlates with increased migration. EGCG significantly suppressed cytokine gene induction, activity of MMP-9 and -2, and cell migration by inhibiting the transcriptional activity and nuclear translocation of NF-κB, as well as inhibiting the degradation of IκBα. Our results suggest that TLR9 agonists induce inflammatory cytokines and MMPs independent of the expression of AR or p53, but involving transcriptional activity of NF-κB.

TLR9 is expressed in all types of immune cells and a variety of cancer cells. Several studies have shown that in addition to protective roles during pathogen infection, TLR9 engagement promotes cell survival and tumor progression by facilitating NF-κB activation. Stimulation of neutrophil granulocytes, B cells, and macrophages induced production of proinflammatory cytokines, such as IL-6, IL-8, TNFα, IL-12, MIP1α, MIP1β, IgM, IgG, and type II IFN, respectively. Expression in RAW264.7 macrophages of CpG-DNA-induced cytokines was significantly decreased in cells with knockdown of TLR9. In various cancer cells, CpG-ODN promotes the release of cytokines and chemokines through the TLR9 pathway. For instance, lung cancer cells stimulated with CpG-ODN produced elevated levels of IL-1, IL-8, and MMP-2, which correlated with increased metastasis. In murine astrocytes and microglia, CpG-ODN induces expression of adhesion molecules, MMP-9, and inflammatory cytokines involving activation of NF-κB and Jun N-terminal kinase (JNK) pathways. In PC3 cells, CpG-ODN can induce IL-8 and TGFβ1 release through the TLR9 pathway and promotes

![Figure 10](image-url)  
*Figure 10* EGCG inhibits CpG-ODN-induced migration of DU145 and PC3 cells.  
**Notes:** Cell migration assays were performed using a modified Boyden chamber containing fibronectin-coated polycarbonate membrane filter (8 μm pore size). DU145 or PC3 cells (2×10⁴) were pretreated or not with EGCG (40 µg/mL) for 24 hours followed by transfection with CpG-ODN (1 µM). Cells were plated in the upper chamber and the lower chamber contained culture medium with 10% FBS. Cells were incubated for 24 hours at 37°C in 5% CO₂. Nonmigrated cells were scraped from the upper surface of the membrane with a cotton swab, and migrated cells remaining on the bottom surface were trypsinized and counted with a hemocytometer. Data shown are means ± SD of experiments performed in triplicate. Student’s t-test: *p*<0.001.  
**Abbreviations:** CpG-ODN, CpG oligodeoxynucleotides; EGCG, epigallocatechin-3-gallate; FBS, fetal bovine serum; SD, standard deviation.
EGCG suppresses TLR9 ligand-induced cytokines

Consistent with these studies, our results show that CpG-ODN induces IL-6, IL-8, CXCL1, IP-10, CCL5, and TGFβ genes. IL-8 is produced constitutively by prostate cancer cells and its expression correlates with tumorigenicity and metastasis. Following CpG-ODN treatment, a 17- to 26-fold increase is observed in induced levels of IL-8, and IL-8 promoter activity is reduced when the NF-κB binding site is mutated. Increased plasma IL-6 levels in metastatic and castration-resistant prostate cancer correlate inversely with tumor survival and chemoresistance. In all the cell lines, levels of IL-6 increased sharply at 6 hours and decreased by 24 hours. Induction of CXCL1 in prostate cancer cells increases migration and invasion and correlates with high-grade malignancy of tumors. CpG-ODN induced CXCL1 levels 3- to 4-fold in our experiments. Proinflammatory chemokine IP-10 was induced more significantly in DU145 and PC3 cells compared to LnCap cells. CCL5 and its receptor, CCR5, are expressed in all prostate cancer cells and increase in CCL5 stimulates proliferation and invasion in vitro. TGFβ1 regulates cell proliferation, apoptosis, and invasion. In our studies, the levels of TGFβ1 increased at 6 hours in DU145 and LnCap cells, and at 24 hours in PC3 cells. MMPs degrade the extracellular matrix and promote tumor invasion and metastasis. Expression of MMP-9 is regulated by NF-κB, and the MMP-9 promoter has binding sites for NF-κB. We observed increased activity of MMP-9 and -2 in the supernatants of DU145, PC3, and LnCap cells following stimulation with CpG-ODN. Accordingly, migration increased 3.5-fold in DU145 and 10-fold in PC3 cells, which is reduced in cells pretreated with EGCG. LnCap cells have very low levels of MMP, which correlates with reduced migration and invasion in vitro. In all our results show that expression of cytokines, chemokines, and MMPs is induced significantly in prostate cancer cells in response to CpG-ODN. It is conceivable that infection in the prostate may promote the release of cytokines, chemokines, and MMPs with roles in propagating inflammation and promoting invasion that could lead to progression of prostate cancer.

EGCG has anti-inflammatory and antitumorigenic effects in several cell types in cell culture and animal models. These effects are exerted by inducing apoptosis, cell cycle arrest, decreasing the expression of antiapoptotic proteins such as PI3K and AKT, suppressing prosurvival transcription factors such as NF-κB, and inhibiting the activity of proteins that promote invasion. Treatment of prostate cancer cells decreases cell viability with increasing doses of EGCG. Studies in LnCap cells showed that increase in cellular p53 levels accompanied a decrease in viability. We did not observe significant differences in cell viability between DU145, PC3, and LnCap cells, which differ in the expression of p53. Since CpG-ODN stimulates cytokines involving NF-κB, and EGCG inhibits NF-κB translocation, we tested to see if EGCG suppresses NF-κB target genes following CpG-ODN stimulation. We show that pretreatment with EGCG prior to CpG-ODN stimulation inhibits the induction of cytokines and MMP-9 and -2 activities. NF-κB is an antiapoptotic protein that also increases cell proliferation and is constitutively expressed in many cancers, including prostate cancer. We demonstrate that EGCG decreases NF-κB transcriptional activity in DU145 and PC3 cells, which can be explained by inhibition of nuclear translocation of the p65 subunit of NF-κB. In addition, 1xBα degradation by ubiquitination is inhibited by EGCG. We observed an increase in cytosolic 1xBα in our experiments, possibly due to inhibition of degradation. It is also likely that EGCG decreases levels of antiapoptotic proteins such as bcl2 and increases proapoptotic proteins such as BAX to increase apoptosis as shown in LnCap cells. These effects are predicted to involve p53. In our studies, the induction of proinflammatory cytokines and MMPs are independent of the expression of AR and p53, since DU145 and PC3 cells lack AR and are mutant, or lack p53 compared to LnCap cells, which express AR and p53. EGCG has broad antitumorigenic roles specifically in cancer cells and anti-inflammatory roles independent of p53, which is frequently mutated in cancers. These attributes suggest that EGCG can be used for the chemoprevention of tumors, more specifically, prostate tumors, which are promoted by inflammation.

Conclusion

Like bacterial DNA, TLR9 agonists like CpG-ODN induce proinflammatory cytokines in DU145, PC3, and LnCap prostate cancer cells. The mRNA for cytokines and chemokines are induced in all three cell lines but with variable kinetics. Activity of matrix MMP-9 and -2, which degrade the extracellular matrix to promote cell invasion, increased in cells stimulated with CpG-ODN. The cytokine transcriptional response and MMP activity induced by CpG-ODN was mediated by a transcription factor, NF-κB. EGCG pretreatment suppressed cytokine gene induction and MMP-9 and -2 activities. The inhibitory effect of EGCG was mediated by inhibiting the translocation of the p65 subunit of NF-κB to the nucleus and transcriptional activity. These studies demonstrate that TLR9 agonists induce proinflammatory cytokines and MMPs, and that EGCG suppresses inflammatory cytokines independent of the expression of AR and p53 status of the cells. EGCG may provide protection...
against inflammation, which contributes to the development of prostate cancer.

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Disclosure
The authors report no conflicts of interest in this work.

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