Glucagon-like peptide-1 exerts anti-inflammatory effects on mouse colon smooth muscle cells through the cyclic adenosine monophosphate/nuclear factor-κB pathway in vitro

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Background: Intestinal smooth muscle cells (SMCs) undergo substantial morphological, phenotypic, and contractile changes during inflammatory bowel disease (IBD). SMCs act as a source and target for different inflammatory mediators, however their role in IBD pathogenesis is usually overlooked. Glucagon-like peptide-1 (GLP-1) is an incretin hormone reported to exert multiple anti-inflammatory effects in different tissues including the gastrointestinal tract through various mechanisms.

 Aim: The aim of this research is to explore the effect of GLP-1 analog exendin-4 on the expression and secretion of inflammatory markers from mouse colon smooth muscle cells (CSMCs) after stimulation with lipopolysaccharide (LPS).

 Materials and methods: Freshly isolated CSMCs from male BALB/c mice were cultured in DMEM and treated with vehicle, LPS (1 µg/mL), LPS+exendin-4 (50 nM), or LPS+exendin-4 (100 nM) for 24 h. Expression of inflammatory cytokines was then evaluated by antibody array membrane.

 Results: CSMCs showed basal expression of several cytokines which was enhanced with the induction of inflammation by LPS. However, exendin-4 (50 and 100 nM) significantly (p<0.05) reduced the expression of multiple cytokines including tumor necrosis factor-α (TNF-α), interleukin-1α (IL-1α), T cell activation gene-3 (TCA-3), stromal cell-derived factor-1 (SDF-1), and macrophage colony stimulating factor (M-CSF). To confirm these results, expression of these cytokines was further assessed by enzyme-linked immunosorbent assay and real-time polymerase chain reaction and similar results were also observed. Moreover, secretion of TNF-α and IL1-α into the conditioned media was significantly downregulated by exendin-4 when compared to LPS-treated cells. Furthermore, LPS increased NF-κB phosphorylation, while exendin-4 significantly reduced levels of NF-κB phosphorylation.

 Conclusion: These data indicate that GLP-1 analogs can exert significant anti-inflammatory effects on CSMCs and can potentially be used as an adjunct treatment for inflammatory bowel conditions.

 Keywords: GLP-1, inflammation, inflammatory bowel disease, smooth muscle

Introduction

Inflammatory conditions of the gastrointestinal tract (GIT) are a growing problem in the industrialized nations, affecting the quality of the patient’s life and having significant financial burden.1,2 Inflammatory bowel disease (IBD) is a group of chronic incurable gastrointestinal inflammatory disorders with undetermined etiology. However, it is thought to result from dysregulated mucosal immune response to intestinal commensal flora in genetically predisposed individuals.1
Ulcerative colitis (UC) and Crohn’s disease (CD) are the most common forms of IBD affecting the colon and small intestine respectively and can involve the mucosa and submucosal layers. These inflammatory conditions impair the ability of affected gastrointestinal tissues to function properly, leading to various symptoms such as diarrhea, malabsorption, abdominal pain, cramping, rectal bleeding, and fatigue. Patients with IBD often alternate between periods of flares and remission. Untreated IBD can result in many intestinal and systemic complications such as intestinal perforations, toxic megacolon, abscesses, and colon cancer. Currently, no cure is available for IBD. Current treatment modalities are focused on reduction of the inflammatory process to alleviate the symptoms and to prevent future complications to improve the patient’s quality of life. Pharmaceutical treatment of IBD includes five major categories: anti-inflammatory drugs, immunosuppressants, biological agents, antibiotics, and drugs for symptomatic relief. However, these treatments are associated with significant side effects and are of limited success in some patients, highlighting the need for novel therapeutic agents with minimal side effects.

At the histological level, the hallmark of IBD is recruitment of inflammatory cells, infiltration and activation within the intestinal mucosa and lamina propria, and enhanced production of pro-inflammatory mediators. Analysis of the mucosa from patients with IBD showed increased expression of several cytokines such as interleukin-1α (IL-1α), IL-6, IL-8, and tumor necrosis factor-α (TNF-α). Cytokines are mainly produced by monocytes and polymorphonuclear cells that have migrated to the inflammatory focus and to a lesser extent by the tissue macrophages and epithelial cells.

The smooth muscle cells (SMCs) within the intestine are found in the longitudinal and circular muscle layers, as well as the muscularis mucosa layer. They are the main contractile nonproliferative apparatus needed for mixing and propelling the intraluminal contents, however they can exhibit phenotypic switching, hyperplasia, hypertrophy, and altered contractile behavior upon intestinal pathologies. Studies conducted using animal models have shown that mucosal inflammation in the gut leads to changes in the growth and contractile properties of SMCs. Other studies have shown that the altered intestinal motility observed in patients with CD is partially attributed to hypertrophy and alterations in the contractility of the intestinal SMCs.

Glucagon-like peptide-1 (GLP-1) is an incretin hormone synthesized by the enteroendocrine L cells which are located mainly in the distal small intestine and colon. GLP-1 is secreted in response to nutrient ingestion leading to various effects on the body such as increased insulin secretion, decreased glucagon secretion, regulating proximal gut transit, and suppression of appetite signaling in the brain; these ultimately result in improved glucose homeostasis and weight loss. GLP-1 has an extremely short half-life (~1–2 min) due to rapid degradation by the enzyme dipeptidyl peptidase IV (DPP-4). DPP-4 is widely expressed by the gut and vascular endothelium. It cleaves GLP-1 from its N-terminal at position 2 occupied by alanine, resulting in the generation of inactive GLP-1(9–37) and GLP-1(9–36) amide. During the past decade, several GLP-1-based therapies became available for the treatment of type 2 diabetes mellitus and obesity. Recently, GLP-1 was found to exert several beneficial effects beyond glycemic control. It exerts anti-inflammatory and antioxidant effects on various tissues and cells including cardiomyocytes, endothelial cells, and vascular SMCs through various mechanisms via receptor-dependent and receptor-independent pathways. Although the primary function of GLP-1 is to serve as an anorexigenic (appetite-reducing) and incretin hormone, as well as a mediator of the “ileal brake” through inhibition of upper gastrointestinal function to prevent malabsorption and postprandial metabolic disturbances, the influence of GLP-1 on colon function was evaluated by a limited number of studies. Amato et al reported that GLP-1 receptor (GLP-1R) is expressed in the nitrergic neurons of the myenteric plexus in human colon circular muscular strips and, once activated by exogenous GLP-1, mediates an inhibitory effect on large intestinal motility through nitric oxide neural release resulting in reduction of the spontaneous contractile activity of the smooth muscles, which may contribute to delaying of intestinal propulsion to increase water and electrolyte absorption. Moreover, recently that GLP-1R is expressed in pure mouse SMC cultures of the colon and intestine. They showed that organ bath studies identified a role of GLP-1R in the regulation of smooth muscle function by influencing the basal tone in colonic muscle strips and mediating relaxation of acetylcholine-induced phasic and tonic contractions in the colon. Furthermore, recent studies showed that GLP-1R agonists modulate enteric immune responses through the intestinal intraepithelial lymphocyte (IEL) which may influence microbiome composition and intestinal inflammation; therefore, GLP-1R agonists may exert beneficial nonglycemic actions partially through modulation of intraepithelial lymphocyte (IEL) which may influence microbiome composition and intestinal inflammation; therefore, GLP-1R agonists may exert beneficial nonglycemic actions partially through modulation of inflammatory mediators such as cytokines and chemokines.
production; however, the anti-inflammatory effects of GLP-1 on colon smooth muscle cells (CSCMs) have not yet been explored.17 Interestingly, GLP-1 levels are elevated in critically ill patients in different chronic inflammatory diseases including CD, UC, metabolic syndrome, coronary artery disease, and heart failure, and have been found to be increased in animal models of inflammatory conditions in response to bacterial stimuli, suggesting a correlation between inflammation and GLP-1 secretion.18–20 Interestingly, DPP4 inhibitors accelerated mucosal healing in an animal model of experimental colitis, and GLP-1 analog ROSE-010 reduced pain exacerbation in patients with irritable bowel syndrome.21–23

The aim of this research is to explore the anti-inflammatory effects of the GLP-1 analog exendin-4 on the expression and secretion of inflammatory markers from mouse colon smooth muscle under lipopolysaccharide (LPS)-induced inflammation in vitro. LPS is a bacterial endotoxin reported to increase cytokine secretion in different cell types.

Materials and methods

Solutions, drugs, and chemicals

Smooth muscle buffer (SMB) was prepared in house and contained 120 mM NaCl, 4 mM KCl, 2.6 mM KH₂PO₄, 2 mM CaCl₂, 25 mM HEPES (N-2-hydroxyethylpiperazine-N’-2-ethanesulfonic acid), 14 mM glucose, and 2.1% essential amino acid mixture (pH was adjusted to 7.4). The digestion solution contained 0.5 mg collagenase and 0.33 mg trypsin inhibitor per milliliter of SMB. All of these chemicals were purchased from Sigma-Aldrich Co. (St Louis, MO, USA). Cell culture media used during the incubation process were prepared by adding 10% fetal bovine serum (FBS) (GE Healthcare UK Ltd, Little Chalfont, UK), 100 U/mL of penicillin, 100 µg/mL of streptomycin, and 2.5 µg/mL of amphotericin B to DMEM with l-glutamine (Capricorn Scientific GmbH, Ebdsorfergrund, Germany). The remaining reagents were purchased from Euroclone S.P.A. (Pero, Italy). Rabbit GLP-1R polyclonal antibody (H-55; sc-66911) was purchased from Santa Cruz Biotechnology Inc. (Dallas, TX, USA) and anti-calponin antibodies were purchased from Abcam Inc. (Cambridge, UK). LPS was purchased from Sigma-Aldrich Co. GLP-1 analog exendin-4 was purchased from Abcam Inc. The polymerase chain reaction (PCR) primers were purchased from Integrated DNA Technologies, Inc. (Coralville, IA, USA) (Table 1). The protein assay kit, SYBR Green Master Mix, and cDNA synthesis kit were purchased from Bio-Rad Laboratories Inc. (Hercules, CA, USA). Mouse inflammation antibody array membrane (Cat. No. ab133999) was purchased from Abcam Inc. Specific enzyme-linked immunosorbent assay (ELISA) kits for mouse TNF-α, M-CSF, IL-1α, TCA-3, and SDF-1 were purchased from Sigma Aldrich Co. The cyclic adenosine monophosphate (cAMP) ELISA kit was purchased from R&D Systems Inc. (Minneapolis, MN, USA), and the nuclear factor (NF)-κB p65 pS536 ELISA kit was purchased from Abcam Inc. The nuclear protein extraction kit was purchased from Abcam Inc.. The RNA extraction kit was purchased from Zymo Research Corp. (Irvine, CA, USA). The 500-µm Nitex mesh was purchased from Sigma-Aldrich Co.

Animals

Twenty young mature male BALB/c mice (~12 weeks of age, 26.5–30 g) were provided by the animal house of Jordan University of Science and Technology. Mice were euthanized by inhalation of CO₂ and the colon was rapidly excised. The colon was cut into small pieces (2–3 cm in length) and then placed in cold SMB with the composition described previously. All procedures were approved and performed according to the guidelines of the Animal Care and Use Committee at Jordan University of Science and Technology.

Preparation of dispersed smooth muscle cells

CSCMs were isolated by sequential enzymatic digestion of muscle strips, filtration, and centrifugation. The mucosa was scraped off with fine scissors under a dissecting scope, and then the tissues were cut into thin slices (2 mm long; 2 mm thin) and incubated for 20 min in SMB containing 0.5 mg/mL collagenase and 0.33 mg/mL soybean trypsin inhibitor

Table 1 Primers used in real-time/reverse transcription polymerase chain reaction and their sequences

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<th>Gene</th>
<th>Accession number</th>
<th>Forward primer</th>
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<td>NM_007393.5</td>
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in a shaking water bath at 31 °C. The tissues were continuously gassed with 100% oxygen during the entire isolation procedure. The partly digested tissue was washed twice with 50 mL of collagenase-free SMB and the muscle cells were allowed to disperse spontaneously for 10 min in collagenase-free medium. Cells were harvested by filtration through 500-μm Nitex mesh and centrifuged twice at 1000 rpm for 10 min to eliminate broken cells and organelles. The process was repeated four or five times. The cells were counted in a hemocytometer, and it is estimated that 95% are viable using a trypan blue exclusion assay.

Identification of smooth muscle cells

Identification of mouse CSMCs was verified by immunohistochemical staining of paraffin-embedded dispersed colon muscle cells using anti-calponin antibody at 1:100 dilution.

Expression of inflammatory cytokines using antibody array membrane

Ten thousand dispersed mouse CSMCs (n=3/group) were seeded in a 6-well plate in DMEM containing 10% FBS, 100 U/mL of penicillin, 100 μg/mL streptomycin, and 2.5 μg/mL amphotericin B. The cells were incubated in a humidified incubator with 95% air and 5% CO₂ at 37 °C for 24 h with either vehicle, LPS (1 μg/mL), LPS (1 μg/mL)+exendin-4 (50 nM), or LPS (1 μg/mL)+exendin-4 (100 nM). After completion of the incubation period (24 h) the treated samples were centrifuged at 1000 × g for 5 min at 4 °C. The conditioned media were stored at –20 °C for further analysis, and the cell pellets were immediately lysed.

Cell lysates were prepared using BashingBeads Lysis tubes (Zymo Research Corp.) and the cell lysis buffer containing protease inhibitor cocktail provided by the kit according to the manufacturer’s instructions. Then lysates were centrifuged for 10 min at 14000 × g at 4 °C and the supernatant collected for further analysis. Total protein concentration of the supernatants was measured using the DC protein assay kit. The protein concentration was then adjusted to 100 μg/mL in all samples.

To evaluate differential cytokine expression in the treated samples, antibody array membranes were used according to the manufacturer’s instructions. Briefly, the membranes were placed in the 8-well tray provided in the kit, and then 1 mL of each sample was incubated overnight into the designated well at 4 °C. Twenty-four hours later, the samples were aspirated and the membranes were washed with wash buffer and then incubated with 1 mL of biotin-conjugated anti-cytokines overnight at 4 °C. Then the biotin-conjugated anti-cytokines were aspirated and washing performed again as previously described. Finally, 2 mL of horseradish peroxidase-conjugated streptavidin was pipetted into each well and incubated overnight at 4 °C; then it was aspirated and the membranes were washed again by the previously described method. Detection buffer was used to develop a chemiluminescent signal and the C-DIGIT blot scanner (LI-COR Biotechnology, Lincoln, NE, USA) was used to detect the signal intensity. Measurements of the intensity of signals on the array membrane were performed using ImageJ software (National Institutes of Health, Bethesda, MD, USA). Data were normalized to the positive control signals and the relative cytokine expression was determined.

CSMC viability assay

Freshly isolated CSMCs were cultured in 96-well plates at 1×10⁵ cells/well. Cells were serum-deprived for 24 h before stimulation with vehicle, LPS (1 μg/mL), LPS (1 μg/mL)+exendin-4 (50 nM), or LPS (1 μg/mL)+exendin-4 (100 nM). Cell proliferation was assessed after 24 h using the MTT assay (Intron Biotechnology, Korea) according to the manufacturer’s instructions. Briefly, CSMCs were incubated with MTT reagent for 4 h at 37 °C. MTT reagent was converted to an insoluble formazan. The formazan is then solubilized, and the concentration determined by optical density at 570 nm.

Quantification of protein expression by ELISA

Protein levels of specific cytokines identified previously by the membrane array were further evaluated by ELISA. Specific ELISA kits for TNF-α, IL-1α, M-CSF, TCA-3, SDF-1, and NF-κB p65 (pS536) were used to measure the levels of these cytokines in the lysates and the conditioned media in the control and treated samples according to the manufacturer’s protocol.

Cellular cAMP determination assay

Freshly isolated CSMCs were cultured in DMEM supplemented with 10% FBS, 100 U/mL of penicillin, and 100 μg/mL of streptomycin at 37 °C in 5% CO₂. Cells were seeded at a density of 10000 cells per well into a 96-well plate in the presence of vehicle, LPS (1 μg/mL), LPS (1 μg/mL)+exendin-4 (50 nM), or LPS (1 μg/mL)+exendin-4 (100 nM) for 10 min (n=3/group). Cells were then immediately lysed and assayed for cAMP accumulation using the cAMP determination kit according to the manufacturer’s protocol.
RNA isolation and quantitative real-time/reverse transcription polymerase chain reaction

To extract RNA from CSMCs, a Direct-Zol RNA MiniPrep kit from Zymo Research was used according to the manufacturer’s instructions. The Direct-Zol method assures efficient recovery of small and large RNAs. Dispersed mouse CSMCs were treated with either a vehicle, LPS (1 µg/mL), LPS (1 µg/mL) + exendin-4 (50 nM), or LPS (1 µg/mL) + exendin-4 (100 nM). The samples were incubated for 12 h in a humidified incubator with 95% air and 5% CO₂ at 37 °C. The concentration and purity of RNA were determined using a NanoDrop 2000 spectrophotometer from Thermo Fisher Scientific (Waltham, MA, USA). One microgram of RNA was reverse-transcribed to obtain cDNA using the iScript cDNA synthesis kit (Bio-Rad Laboratories Inc.). Expression of target genes was assayed by quantitative real-time/reverse transcription (qRT)-PCR using Bio-Rad iTaq SYBR Green Master Mix from Bio-Rad Laboratories Inc. contains antibody-mediated hot-start iTaq DNA polymerase, dNTPs, MgCl₂, SYBR green I dye, and fluorescein. The 7500 Fast Real-Time PCR System from Applied Biosystems (Thermo Fisher Scientific) was used to run the reaction. The Master Mix was first activated at 95 °C for 3 min and then 35 cycles of amplification were performed each consisting of 15 sec at 95 °C and 60 sec at 60 °C. A melting point dissociation curve generated by the instrument was used to confirm that only a single product was present.

Immunohistochemistry

Dispersed mouse CSMCs were fixed in 10% neutral-buffered formalin (pH 7.4) overnight and were then embedded in paraffin matrix. Mid-colon tissue from male BALB/c mice was fixed in 10% neutral-buffered formalin (pH 7.4) overnight and were then embedded in paraffin. Five-micrometer sections were prepared, and the sections were dewaxed and rehydrated through a graded alcohol series. Antigen unmasking was performed by boiling the sections in Citra Plus (BioGenex, San Ramon, CA, USA) in a microwave oven for 2 min at power 10 and then for 10 min at power 1, followed by cooling for 20 min. Sections were treated with 3% hydrogen peroxide to quench endogenous peroxidase activity and incubated in blocking solution containing goat IgG (Vectastain Elite ABC kit; Vector Laboratories, Inc., Burlingame, CA, USA) for 30 min. Sections were incubated overnight with rabbit GLP-1R polyclonal antibody (Santa Cruz Biotechnology) and anti-Calponin antibody (Abcam Inc.) followed by incubation with goat anti-rabbit secondary antibody (Vectastain Elite ABC kit; Vector Laboratories) for 30 min. Sections were stained with 3,3’-diaminobenzidine tetra-hydrochloride (Dako Denmark/A/S, Glostrup, Denmark). Slides were dehydrated in an alcohol series and cleared in xylene. Images were obtained with 20× objective by an inverted Nikon TMS-F microscope (Nikon Instruments, Irving, TX, USA).

Statistical analysis

Statistical analyses were performed using GraphPad Prism 5.0 (GraphPad Software, Inc., La Jolla, CA, USA). One-way analysis of variance followed by Fisher’s post hoc analysis was used to examine significant differences between groups. A p-value less than 0.05 was considered significant. All data are shown as mean±standard error of the mean.

Results

Colonic smooth muscle cell identification

Freshly isolated mouse cells were viewed at 20× objective on the inverted Nikon TMS-F microscope and they appeared spindle-shaped with varying lengths (Figure 1A). The identity of mouse CSMCs was verified by immunohistostaining with anti-calponin antibody (Figure 1B). The results showed that more than 95% of the cells stained positive for calponin. Calponin is specific for differentiated SMCs.

Inflammatory cytokine expression by antibody array membrane

Antibody array membrane analysis showed a basal expression for several cytokines by mouse CSMCs. Treatment with LPS caused a marked increase (~50–100%) in the expression of several cytokines. Co-treatment with exendin-4 significantly inhibited the overexpression of these cytokines by ~40–50% (Figure 2A and B). This initial screening step showed that five cytokines were especially affected by the LPS and exendin-4 treatment: TNF-α, IL-1α, macrophage colony stimulating factor (M-CSF), T cell activation gene-3 (TCA-3), and stromal cell-derived factor-1 (SDF-1). These data suggest that CSMCs are capable of producing several inflammatory mediators that can play an important role in the pathogenesis of inflammatory conditions of the gastrointestinal system; on the other hand, GLP-1 analog exendin-4 could modulate these inflammatory conditions by affecting smooth muscle’s cytokine and chemokine expression.
Colon smooth muscle cell viability assay

To ensure that CSMC viability is not affected by LPS or exendin-4 treatment, we performed an MTT assay on the treated cells. Our results indicate that cell viability is comparable between all treatment groups, indicating that neither LPS nor exendin-4 is affecting the growth of CSMCs within a 24-h period (Figure 1C).

Assessment of cytokine expression and secretion by ELISA

To further evaluate the levels of differentially expressed cytokines in the treated samples and control, specific ELISAs were used. Data showed that LPS treatment induced ~1–2-fold increase in the expression of TNF-α, IL1-α, M-CSF, TCA-3, and SDF-1 (p<0.05). However, co-treatment with exendin-4 (50 or 100 nM) significantly reduced their expression by ~20–60% compared to LPS alone (p<0.05) (Figure 3).

Inflammatory cytokine expression by real-time PCR

To confirm the results obtained from the antibody array membrane and ELISA at the level of mRNA expression, real-time PCR was performed on mouse CSMCs treated with vehicle, LPS (1 µg/mL), LPS (1 µg/mL)+exendin-4 (50 nM), or LPS (1 µg/mL)+exendin-4 (100 nM). This experiment proved that LPS treatment induced a significant increase (~40–100%, p<0.05) in the level of expression of TNF-α, IL-1α, TCA-3, SDF-1, and M-CSF. On the other hand, exendin-4 co-treatment significantly (p<0.05) inhibited the overexpression of these cytokines by the CSMCs (Figure 5).

GLP-1R expression in CSMCs

GLP-1R is a G protein-coupled receptor that is widely expressed in many tissues including the pancreatic islets, GIT, lung, kidneys, heart, and some regions of the central and peripheral nervous system. GLP-1R is also widely expressed by multiple immune cell populations, particularly by the intestinal intraepithelial lymphocytes. In the pancreatic islets, activation of GLP-1R results in activation of adenyl cyclase which leads to generation of cAMP, which in turn activates various downstream effectors through activation of protein kinase A. Expression of GLP-1R in the CSMCs has not been explored previously, therefore we sought to determine its
Anti-inflammatory effects of GLP-1 on colon smooth muscle cells

Our results indicate that GLP-1R is expressed in isolated single CSMCs (Figure 6A). Furthermore, immunohistochemistry of whole colon sections showed that GLP-1R is expressed in the muscular wall of the colon as well as the epithelial cells (Figure 6B). Moreover, we evaluated the effect of GLP-1R activation by exendin-4 through measuring intracellular cAMP levels in CSMCs. Our results indicate that exendin-4 increased CSMC cAMP levels in a dose-dependent fashion, indicating that GLP-1R mediates its effects in CSMCs through activation of adenylyl cyclase, leading in turn to cAMP-dependent activation of various second messenger pathways (Figure 6C).

Discussion

The intestinal epithelial barrier is usually the first site to be involved in the inflammatory conditions affecting the gut,24 where microorganisms or allergens are first encountered, leading to activation of the local immune cells and recruitment of new immune cells which alter the local environment resulting in the major hallmarks of inflammation;25 however, emerging evidence supports the notion that intestinal SMCs

expression in mouse CSMCs by immunohistochemistry. Our results indicate that GLP-1R is expressed in isolated single CSMCs (Figure 6A). Furthermore, immunohistochemistry of whole colon sections showed that GLP-1R is expressed in the muscular wall of the colon as well as the epithelial cells (Figure 6B). Moreover, we evaluated the effect of GLP-1R activation by exendin-4 through measuring intracellular cAMP levels in CSMCs. Our results indicate that exendin-4 increased CSMC cAMP levels in a dose-dependent fashion, indicating that GLP-1R mediates its effects in CSMCs through activation of adenylyl cyclase, leading in turn to cAMP-dependent activation of various second messenger pathways (Figure 6C).

Discussion

The intestinal epithelial barrier is usually the first site to be involved in the inflammatory conditions affecting the gut,24 where microorganisms or allergens are first encountered, leading to activation of the local immune cells and recruitment of new immune cells which alter the local environment resulting in the major hallmarks of inflammation;25 however, emerging evidence supports the notion that intestinal SMCs
Figure 3 Effect of exendin-4 treatment on the expression of inflammatory cytokines by mouse CSMCs using ELISA. Freshly isolated CSMCs from 20 male Balb/C mice were cultured in DMEM in 6-well plates and divided into four treatment groups (n=3/group): control (Ctrl), LPS, LPS+exendin-4 (50 nM), and LPS+exendin-4 (100 nM). After 24 h of incubation the cells were collected and lysed. Levels of different cytokines in the lysate were assayed using specific ELISA kits. Bar graphs represent levels of: (A) tumor necrosis factor-α (TNF-α), (B) macrophage colony stimulating factor (M-CSF), (C) interleukin-1α (IL-1α), (D) T cell activation gene-3 (TCA-3), and (E) stromal cell-derived factor-1 (SDF-1). Different lowercase letters indicate significant statistical difference between groups where \( p < 0.05 \). Values shown are representative of three independent experiments performed in triplicates. Bars represent mean±standard error of the mean.

Abbreviations: CSMC, colon smooth muscle cell; ELISA, enzyme-linked immunosorbent assay; Ex 50, exendin-4 (50 nM); Ex 100, exendin-4 (100 nM); LPS, lipopolysaccharide.
are affected functionally and structurally by most inflammatory conditions involving the intestine leading to altered morphology and contractile behavior, and secretion of various inflammatory mediators. Some studies have shown that the intestinal SMCs are capable of producing several cytokines such as IL-6 and TNF-α under different pathological conditions. A study showed that TNF-α binds to receptors on the SMCs, leading to activation of NF-κB and expression of a number of chemotactic genes such as MCP-1, IL-8, and intercellular adhesion molecule-1. Moreover, human CSMCs can express IL-1α, IL-6, IL-8, cyclooxygenase-2, and regulated on activation, normal T cell expressed and secreted (RANTES) in response to exposure to TNF-α, IL-1α, and interferon-γ.

The aim of this study was to evaluate the expression of inflammatory mediators by normal and inflamed CSMCs, and to evaluate the effects of GLP-1 analog exendin-4 on the expression of inflammatory mediators by CSMCs after induction of inflammation by LPS.

GLP-1 is an incretin hormone synthesized and secreted by the intestinal L cells in response to the presence of nutrients within the gut. It improves glucose homeostasis through various mechanisms including activation of insulin secretion and inhibition of glucagon secretion, delaying gastric emptying, reducing hepatic glucose production, and decreasing food intake. GLP-1 has been shown to exert various anti-inflammatory actions on different tissues and can potentially be used for the treatment of several chronic inflammatory conditions like atherosclerosis, neurodegenerative disorders, diabetic nephropathy, asthma, psoriasis, and nonalcoholic steatohepatitis. Moreover, GLP-1 has been shown to improve the inflammatory milieu, vascular endothelial function, lipid profile, and cholesterol in diabetic patients. Several synthetic forms of GLP-1 are now available for treatment of diabetes mellitus such as exenatide. Treatment with GLP-1 analogs can suppress multiple inflammation parameters in the vascular and bronchial SMCs including a reduction in the production of certain cytokines.

In our study, we used an antibody array membrane as a screening tool to assess the effect of inflammation and GLP-1 co-treatment on the expression of cytokines and chemokines in mouse CSMCs. After careful analysis and comparison between the four treatment groups, we found that normal CSMCs are capable of producing several cytokines and chemokines at the basal level, while inducing inflammation by LPS stimulation caused a marked increase in the expression of TNF-α, IL-1α, M-CSF, TCA-3, and SDF-1. On the other hand, co-treatment with GLP-1 analog exendin-4 caused a significant reduction in the expression of these cytokines. We further evaluated these results by ELISA and RT-PCR and evaluated their secretion into the media. Our results indicate that exendin-4 is effective in suppressing secretion of TNF-α and IL-1α from CSMCs.

TNF-α is a potent pro-inflammatory cytokine primarily secreted from activated macrophages; however, many
Figure 5 Effect of exendin-4 treatment on mRNA abundance of inflammatory cytokines in mouse CSMCs using RT-PCR. Freshly isolated CSMCs from 20 male Balb/C mice were cultured in DMEM in 6-well plates and divided into four treatment groups (n=3/group): control (Ctrl), LPS, LPS + exendin-4 (50 nM), and LPS + exendin-4 (100 nM). After 12 h of incubation the cells were collected and RNA was extracted as mentioned previously. Relative gene expression of different cytokines in the treated cell was assayed using RT-PCR. Bar graphs represent gene expression levels of: (A) tumor necrosis factor-α (TNF-α), (B) macrophage colony stimulating factor (M-CSF), (C) interleukin-1α (IL-1α), (D) T cell activation gene-3 (TCA-3), and (E) stromal cell-derived factor-1 (SDF-1). Different lowercase letters indicate significant statistical difference between groups where p<0.05. Values shown are representative of two independent experiments performed in triplicates. Bars represent mean ± standard error of the mean.

Abbreviations: CSMC, colon smooth muscle cell; Ex 50, exendin-4 (50 nM); Ex 100, exendin-4 (100 nM); LPS, lipopolysaccharide; RT-PCR, real-time/reverse transcription polymerase chain reaction.
other cell types are capable of producing TNF-\(\alpha\), while LPS enhanced this capability and GLP-1 co-treatment significantly inhibited its expression and secretion, suggesting an important role for CSMC-driven TNF-\(\alpha\) in the pathogenesis of gastrointestinal inflammatory diseases through recruitment and activation of different inflammatory cells.

IL-1\(\alpha\) is a central mediator of innate immunity and inflammation;\(^7\) in experimental colitis, it is proven to be a key mediator and an early marker of inflammation and its tissue levels correlate with the severity of inflammation. High levels of IL-1\(\alpha\) have been reported in freshly homogenized colonic biopsies taken from IBD patients.\(^36\) Moreover,
specific blockade of IL-1α receptors reduces the inflammatory responses associated with experimental colitis.37 Some of the effects of TNF-α are mediated through IL-1α. The IL-1α precursor is constitutively expressed by the epithelial layer of the entire GIT and several other cell types outside the GIT including the VSMCs, however its expression is greatly enhanced during inflammatory conditions. If cell death occurs by necrosis, this active precursor is released, thereby initiating a cascade of inflammatory cytokines and chemokines, hence accounting for the early phases of sterile inflammation. Moreover, increased expression and secretion of both IL-1α and TNF-α together are considered an alarm signal that initiates inflammatory responses by inducing a cascade of other pro-inflammatory genes.38

M-CSF is a cytokine produced by several types of cells in the body. It controls survival, proliferation, and differentiation of macrophages. It is proven that both M-CSF and macrophages are increased in the GIT in the presence of IBD, hence it is postulated that M-CSF may play a significant role in the pathogenesis of IBD.39,40 Expression of many cytokines is dependent on M-CSF. Marshall et al 39 found that blockade of M-CSF with a neutralizing anti-M-CSF antibody inhibits dextran sulfate sodium-induced colitis in a mouse model. Our experiment has clearly shown that exendin-4 significantly inhibited M-CSF expression by the inflamed CSMCs.

Chemokines are chemotactic cytokines that play an essential role in the inflammatory response through recruitment and activation of various types of leukocytes. Several studies support the notion that chemokines play an essential role in the pathogenesis of IBD; rectal biopsies taken from patients during the active phase of IBD show high levels of IL-8 (CXCL8) and RANTES (CCL5).41–43 Moreover, MCP-1 (CCL2) mRNA expression was found to increase during the active disease state in endothelial cells and SMCs.44 In our study, expression of two chemokines, TCA-3 (CCL1) and SDF-1 (CXCL12), increased after induction of inflammation, while exendin-4 co-treatment significantly inhibited the production of these chemokines, suggesting that GLP-1 analogs may interfere in attraction and recruitment of inflammatory cells by inhibition of CSMC-derived chemokines.

TCA-3 is a β-chemokine produced from activated T lymphocytes at the site of inflammation; it functions as a chemotactic factor for neutrophils and macrophages. According to a study conducted by Scheerens et al,45 mRNA levels encoding TCA-3 and several other chemokines were significantly upregulated in the chronically inflamed colons of IL-10−/− mice when compared with wild-type mice. Furthermore, reversal of colitis in these mice was accompanied by inhibition in the expression of TCA-3.

SDF-1 is a chemokine secreted by various types of cells; it serves as a chemotactic factor for lymphocytes and monocytes. Several studies linked SDF-1 expression to IBD pathogenesis. SDF-1 is expressed by normal intestinal epithelial cells and its expression is upregulated in intestinal epithelial cells in IBD.41 In a mouse model of colitis, using an antagonist for SDF-1 receptor resulted in a reduction in colonic inflammation manifested as a reduction in the production of pro-inflammatory cytokines and an improvement in colonic pathology.46

Although CSMCs are important in the pathogenesis of IBD, their role is rarely explored compared to the role of inflammatory cells. Their secretory functions are also overlooked since most studies focus on the morphological changes and altered contractility that occur during colonic inflammation.

GLP-1 exerts its various functions by interacting with GLP-1R, which is a G protein-coupled receptor that is widely distributed in the pancreatic islets, GIT, lungs, kidneys, heart, and some regions of the central and peripheral nervous system. GLP-1R is also widely expressed by multiple immune cell populations, particularly by the intestinal intraepithelial lymphocytes.47 In our study, immunohistochemical staining revealed that GLP-1R is expressed in mouse CSMCs, suggesting that GLP-1 can directly influence CSMC function. In the pancreatic islets, activation of GLP-1R results in activation of adenyl cyclase and generation of cAMP, activating various downstream mediators through protein kinase A. GLP-1R activation is also coupled to increased intracellular calcium concentration, inhibition of voltage-dependent potassium currents, and activation of gene expression through effects on Erk1/2, protein kinase C, and phosphatidylinositol 3-kinase.48 In the current study, treatment with GLP-1R agonist exendin-4 significantly increased cAMP levels, suggesting that mouse CSMCs utilize the protein kinase A signaling pathway to mediate GLP-1 effects.

It has been shown that agents that increase intracellular cAMP can exert multiple anti-inflammatory effects through reduction of release of cytokines and lysosomal enzymes from neutrophils, histamine, and leukotrienes from mast cells, and other inflammatory cell functions.49 It has been shown that inhibitors of phosphodiesterase type 4 can elevate cAMP intracellular levels, and can downregulate the release of pro-inflammatory cytokines in the mucosa of IBD patients.50

The exact mechanism through which GLP-1 was able to affect the production of cytokines in the CSMCs was beyond
We speculate that increased intracellular cAMP levels after binding of GLP-1 on GLP-1R on the surface of CSMCs results in downregulation of NF-κB phosphorylation and nuclear translocation which could ultimately result in a reduction in the expression of cytokines and chemokines (Figure 8). NF-κB is a transcription factor that controls the transcription of DNA, cytokine production, and cell survival, and has been shown to be suppressed by cAMP.51 Several studies have proved the inhibitory effect of GLP-1 on NF-κB. Dai et al52 found that incubating human umbilical vein endothelial cells with long-acting GLP-1 analog liraglutide was associated with an inhibition of NF-κB phosphorylation and translocation from the cytoplasm to the nucleus. In another study performed on a rat model of diabetes, it was found that exendin-4 treatment significantly inhibited the phosphorylation of NF-κB in the kidneys of diabetic rats.53 Chaudhuri et al32 found that 12-week treatment with exenatide for diabetic patients suppressed the generation of reactive oxygen species and the intranuclear NF-κB binding by mononuclear cells.

Gut inflammation is associated with alterations in intestinal motility; this might result from changes in the SMC function and/or the enteric nervous system. It was observed through various studies that intestinal inflammation is associated with

The scope of this study.
either an increase or a decrease in SMC contractility; this most probably reflects a difference in the cytokine profiles depending on the inflammatory stimuli.10 Vermillion et al10 found that the contraction of the smooth muscle tissue taken from the inflamed small intestine of patients with CD is abnormal compared with noninflamed tissues from a control group. In another study, IL-1α was found to be significantly elevated in the human sigmoid circular smooth muscle layer taken from patients with UC and that the circular smooth muscle from these samples does not contract normally.10

In an experiment performed on human colon circular muscular strips taken from normal subjects, it was found that GLP-1R is expressed in the human colon and when activated by exogenous GLP-1 it results in an inhibitory effect on the large intestine motility through nitric oxide neural release.55 If it proves to be beneficial in reversing the inflammation-induced effects on SMC contraction, GLP-1 can prove to be of value as a potential therapeutic option in inflammatory conditions involving the colon.

**Conclusion**

GLP-1 analog exendin-4 was successful in inhibiting the overexpression of cytokines associated with the induction of inflammation in CSMCs. Several studies are needed to evaluate the true value of GLP-1 analogs and the possibility of using them as adjunct therapy in the treatment of inflammatory bowel conditions.

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**Author contributions**

AA: conception and design of study, acquisition of data, analysis and interpretation of data, drafting the manuscript, revising the manuscript critically for important intellectual content. TEA: acquisition of data (RT-PCR and ELISA), analysis and interpretation of data, drafting the manuscript, revising the manuscript critically for important intellectual content. OA, AGM, MAA, and MA: analysis and interpretation of data, revising the manuscript critically for important intellectual content. All authors agree to be accountable for all aspects of the work.

**Disclosure**

The authors report no conflicts of interest in this work.

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