Research Article

Anti-inflammatory Effects of Empagliflozin and Gemigliptin on LPS-Stimulated Macrophage via the IKK/NF-κB, MKK7/JNK, and JAK2/STAT1 Signalling Pathways

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Background. Sodium-glucose cotransporter 2 (SGLT2) and dipeptidyl peptidase-4 (DPP-4) inhibitors are glucose-lowering drugs whose anti-inflammatory properties have recently become useful in tackling metabolic syndromes in chronic inflammatory diseases, including diabetes and obesity. We investigated whether empagliflozin (SGLT2 inhibitor) and gemigliptin (DPP-4 inhibitor) improve inflammatory responses in macrophages, identified signalling pathways responsible for these effects, and studied whether the effects can be augmented with dual empagliflozin and gemigliptin therapy. Methods. RAW 264.7 macrophages were first stimulated with lipopolysaccharide (LPS), then cotreated with empagliflozin, gemigliptin, or empagliflozin plus gemigliptin. We conducted quantitative RT-PCR (qRT-PCR) to determine the most effective anti-inflammatory doses without cytotoxicity. We performed ELISA and qRT-PCR for inflammatory cytokines and chemokines and flow cytometry for CD80, the M1 macrophage surface marker, to evaluate the anti-inflammatory effects of empagliflozin and gemigliptin. NF-κB, MAPK, and JAK2/STAT signalling pathways were examined via Western blotting to elucidate the molecular mechanisms of anti-inflammation. Results. LPS-stimulated CD80⁺ M1 macrophages were suppressed by coincubation with empagliflozin, gemigliptin, and empagliflozin plus gemigliptin, respectively. Empagliflozin and gemigliptin (individually and combined) inhibited prostaglandin E₂ (PGE₂) release and COX-2, INOS gene expression in LPS-stimulated RAW 264.7 macrophages. These three treatments also attenuated the secretion and mRNA expression of proinflammatory cytokines, such as TNF-α, IL-1β, IL-6, and IFN-γ, and proinflammatory chemokines, such as CCL3, CCL4, CCL5, and CXCL10. All of them blocked NF-κB, JNK, and STAT1/3 phosphorylation through IKKα/β, MKK4/7, and JAK2 signalling. Conclusions. Our study demonstrated the anti-inflammatory effects of empagliflozin and gemigliptin via IKK/NF-κB, MKK7/JNK, and JAK2/STAT1 pathway downregulation in macrophages. In all cases, combined empagliflozin and gemigliptin treatment showed greater anti-inflammatory properties.

1. Introduction

Certain aspects of chronic low-grade inflammation in obesity and metabolic syndrome-related diseases, such as hypertension, atherosclerosis, and diabetes mellitus, have been studied [1]. Some antidiabetic drugs exert anti-inflammatory properties that can be mediated by directly or indirectly regulating the inflammatory response, and recent studies show that sodium-glucose cotransporter 2 (SGLT2) and dipeptidyl peptidase-4 (DPP-4) inhibitors may exert potential anti-inflammatory functions [2–4].

Sodium-glucose cotransporters are a family of active glucose transporter proteins expressed in bacteria and animals, including 12 human genes [5, 6]. SGLT1 is expressed in numerous organs, including the small intestine, kidney, brain, heart, and immune cells [7–11], while SGLT2 is predominantly expressed in the kidney [12]. SGLT2 inhibitors were initially known to improve hyperglycemia through...
glycosuria by competitively inhibiting both SGLT1 and SGLT2 [13]. These antidiabetic agents have been recently reported to promote M2 macrophage polarization with an anti-inflammatory phenotype and to attenuate the production of proinflammatory cytokines on diabetic cardiomyopathy and nephropathy in mice [14,15]. Two recent studies have shown evidence of SGLT2 protein expression in RAW 264.7 (mouse macrophage cell line) [16] and Kupffer (hepatic stellate macrophage) cells [17].

DPP-4, which exists both as a type II cell surface protein (CD26) and a soluble molecule lacking intracellular and membrane-anchoring domains, is ubiquitously expressed by numerous cells in rats, mice, and humans [18–24]. It acts as an enzymatic and nonenzymatic multifunctional protein, depending on the expressing cell type and cellular conditions, by regulating peptides or influencing cell signals [18,25,26]. DPP-4 aids macrophage and dendritic cell maturation, consequently inducing T-cell activation, and has recently emerged as an important inflammatory response regulator [24,27,28]. DPP-4 inhibitors, which have been initially approved as an antidiabetic drug, are now proposed as a non-diabetic drug for treatment of several inflammatory diseases [29–31]. DPP-4 inhibitors also attenuated NOD-like receptor pyrin domain-containing protein 3 (NLRP3) inflammation, reducing proinflammatory cytokine productions in diabetic mouse kidney and heart [32,33].

Combination therapy of SGLT2 and DPP-4 inhibitors has a stronger glucose-lowering effect than either monotherapy in type 2 diabetes patients due to different mechanisms and complementary effects [34]. This treatment strategy is expected to provide additional anti-inflammatory effects to reduce disease progression and risk of complications. However, the molecular mechanisms of SGLT2 and DPP-4 inhibitors for inflammatory disease have not been fully verified. Furthermore, it is rare to investigate the anti-inflammatory effect after direct treatment of macrophages with a combination of these two agents. Therefore, we evaluated the impact of SGLT2 inhibition, DPP-4 inhibition, and simultaneous SGLT2 and DPP-4 inhibition on proinflammatory response in LPS-stimulated macrophages and their potential mechanism.

2. Materials and Methods

2.1. Cell Culture and Reagents. RAW 264.7 murine macrophage cells were obtained from the American Type Culture Collection (Manassas, VA, USA). The macrophages were cultured in high-glucose Dulbecco’s Modified Eagle’s medium (DMEM; Welgene Inc., Daegu, South Korea) supplemented with 10% (v/v) fetal bovine serum (FBS; Grand Island, NY, USA) and antibiotics (10 μg/mL streptomycin, and 100 IU/mL penicillin) at 37°C and 5% CO2 in a humidified atmosphere of 95% air. These cells were cultured to ≥85% confluence before treatment with empagliflozin (AdooQ Bioscience, Irvine, CA), gemigliptin (LG Life Sciences Ltd., Seoul, South Korea), lipopolysaccharide (LPS) from Escherichia coli 055:B5 (Sigma-Aldrich, St. Louis, MO, USA), or dexamethasone (Sigma-Aldrich). Empagliflozin and gemigliptin were dissolved in dimethyl sulfoxide (DMSO; Sigma-Aldrich) and added to the cell culture at the desired concentrations. The final DMSO concentration did not exceed 0.1%, and all samples were incubated with the same amounts of DMSO. LPS and dexamethasone were dissolved in phosphate-buffered saline (PBS; Welgene Inc.), and all additives were used as cotreatments following a 2-h period of cell starvation.

2.2. Cell Viability Assay. A Cell Counting Kit-8 (CCK-8) assay (Dojindo Laboratories, Kumamoto, Japan) was used to measure the cytotoxicity of the regents on RAW 264.7 macrophages. After incubation at 37°C for 24 h in 96-well plates at a density of 5 × 104 cells/well, the cells were treated with reagents at 37°C for 48 h. Then, the cells were washed, and 10 μL CCK-8 solution was added to each well, followed by incubation at 37°C for 2 h. The absorbance at 450 nm was determined by a microplate absorbance reader (Bio-Rad, Hercules, CA, USA). Treated cell viability was assessed as a percentage of the absorbance values compared to the control (untreated) cells.

2.3. Flow Cytometry Analysis. RAW 264.7 macrophages were collected after treatment for 48 h, and the cells were washed in PBS/2% FCS (flow cytometry staining buffer). The Fc region was blocked by incubating with anti-FcRII/III monoclonal antibodies (mAbs, clone 2.4G2; Invitrogen, Carlsbad, CA) and 10% normal mouse serum for 15 min on ice. Then, the cells were stained for 45 min on ice with FITC anti-CD80 (B7-1; Invitrogen) and fixed in PBS/1% paraformaldehyde. Twenty thousand cells were analysed on a FACSaria III cell sorter (BD Biosciences, San Jose, CA, USA), and data were analysed using FlowJo Vx (TreeStar Inc., Ashland, OR, USA).

2.4. Western Blot Assay. The cells were lysed in radioimmunoprecipitation assay (RIPA) buffer (150 mM NaCl, 1% (v/v) NP-40, 0.5% (w/v) deoxycholate, 0.1% (w/v) sodium dodecyl sulphate (SDS), 50 mM Tris-HCl (pH 7.5), and a protease inhibitor (Pancreas extract, Pronase, Thermolysin, Chymotrypsin, Papain) cocktail; Roche Applied Science, Mannheim, Germany) and incubated for 20 min on ice. Total proteins were then extracted by differential centrifugation at 13,000×g for 10 min, and the protein concentrations in lysates were determined using a protein assay kit (Bio-Rad, Hercules, CA, USA). Subsequently, the cell lysates were mixed with equal volumes of 2× SDS sample buffer (125 mM Tris-HCl (pH 6.8), 4% (w/v) SDS, 4% (v/v) 2-mercaptoethanol, and 20% (v/v) glycerol). The cell lysates, containing equivalent amounts of 20 μg protein, were subjected to 8–12% (w/v) sodium dodecyl sulphate-polyacrylamide gel electrophoresis (SDS-PAGE) and transferred to polyvinylidene fluoride (PVDF) membranes (Millipore, Billerica, MA, USA). The membranes were blocked in 5% (w/v) skim milk or bovine serum albumin (BSA) for 30 min and then incubated overnight with primary antibodies, against cyclooxygenase-2 (COX-2), inducible nitric oxide synthase (iNOS), phosphor-nuclear factor kappa-light-chain-enhancer of activated B cells (NF-κB), total NF-κB, phosphor- inhibitor of NF-κB kinase (IKKa/β, total IKKα, total IKKaβ, phosphor-c-Jun N-terminal kinase (JNK), total JNK,
phosphor-p38 mitogen-activated protein kinase (p-38), total p-38, phosphor-mitogen-activated protein kinase 4 (MKK4), total MKK4, phosphor-MKK7, total MKK7, phosphor-signal transducer and activator of transcription 1 (STAT1), total STAT1, phosphor-STAT3, total STAT3, phosphor-njanus family tyrosine kinase 2 (JAK2), total JAK2, glyceraldehyde 3-phosphate dehydrogenase (GAPDH) (Cell Signaling Technology, Danvers, MA, USA), phosphor-extracellular signal-regulated kinase (ERK), and total ERK (Santa Cruz Biotechnology, Santa Cruz, CA, USA) at 4°C. The membranes were then incubated with the secondary antibodies (horseradish peroxidase-conjugated anti-goat IgG and anti-rabbit IgG (Bethyl Laboratories, Montgomery, TX, USA)). Finally, the blots were measured and visualized using an enhanced chemiluminescence kit (Amersham Pharmacia Biotech, Piscataway, NJ, USA).

2.5. RNA Isolation and Quantitative Real-Time Reverse Transcripase-Polymerase Chain Reaction (qRT-PCR) Analysis. The total RNAs were extracted using the RNAiso Plus reagent (Takara Bio, Otsu, Japan) according to the manufacturer’s protocol. Complementary DNA (cDNA) was synthesized from RNA using avian myeloblastosis virus (AMV) reverse transcriptase (BEAMS Biotechnology, Seongnam, South Korea), and random 9-mer primers, then amplified qPCR using primer sets (Takara Bio Inc., Shiga, Japan) specific for mouse COX-2: CAG CAA ATC CTT GCT GTT CC (forward, F) and TGG GCA AAG AAT GCA AAC ATC (reverse, R); mouse iNOS: AGA CCT CAA CAG AGC CCT CA (F) and GGC TGG ACT TTT CAC TCT GC (R); mouse tumor necrosis factor-α (TNF-α): TCG TAG CAA ACC ACC AAG TG (F) and AGA TAG CAA ATC GGC TGA CG (R); mouse interleukin (IL)-1β: TCT CGC AGC AGC ACA TCA ACA (F) and CCT GGA AGG TTC ACG GGA AA (R); mouse IL-6: GAC CTG TCT ATA CCA CTT CAC (F) and GTG CAT CAT CGT TGT TCA TAC (R); mouse C-C motif chemokine ligands (CCL) 3: ACT GCC CTT GCT GTT CT (F) and GTG TCT TTG GAG TCA GCG CA (R); mouse CCL4: CTC TCT CTT CTT CTT CTG CG (F) and CTC ACT GGG GTT AGC ACA GA (R); mouse CCL5: CCA TCA TCC TCA CTG CAG CC (F) and CTC TGG GTT GGC ACA CAC TT (R); and mouse C-X-C motif chemokine ligand (CXCL) 10: CCA AGT GCT GCC GTC ATT TT (F) and TCA TCA TTC TTT TTC ATC GTG GCA (R). Quantitative real-time PCR was performed using SYBR Green Master Mix according to the manufacturer’s instructions.

2.7. Statistical Analysis. All experiments were conducted in triplicates, and the data were presented as the means ± standard deviation. The results were analysed using Student’s t-test, and results with p values of ≤0.05 were considered statistically significant.

3. Results

3.1. Effect of Empagliflozin and Gemigliptin on LPS-Stimulated Inflammation in RAW 264.7 Macrophages. To determine the effective concentration of empagliflozin and gemigliptin without cell toxicity, RAW 264.7 macrophages were treated with the various doses of empagliflozin (40, 60, and 80 μM) or gemigliptin (100, 250, and 500 μM) for 4 h (Figures 1(a) and 1(b)) [16, 35]. Treatment of LPS-stimulated RAW 264.7 macrophages with 80 μM empagliflozin or 500 μM gemigliptin significantly inhibited mRNA expression of proinflammatory cytokines, without affecting cell viability for 48 h (Figure 1(c)). Thus, these concentrations were used in subsequent experiments. We confirmed that empagliflozin and gemigliptin inhibited the proinflammatory response by reducing the number of M1 macrophage surface marker, CD80, expressing cells in LPS-induced RAW 264.7 macrophages after 48 h treatment (Figure 1(d)) [36].

3.2. Empagliflozin and Gemigliptin Inhibited PGE2 Production and Attenuated COX-2 and iNOS mRNA Expression in LPS-Activated RAW 264.7 Macrophages. LPS induces the macrophage activation and the release of proinflammatory mediators, such as PGE2, and nitric oxide (NO), by upregulation of COX-2 and iNOS mRNA expression [37–39]. We confirmed that PGE2 production increased significantly in RAW 264.7 macrophages stimulated with LPS compared to cells without LPS induction. In contrast, empagliflozin and gemigliptin (individually and combined) inhibited PGE2 production in LPS-activated RAW 264.7 macrophages (Figure 2(a)). COX-2 mRNA and protein expression also increased markedly after LPS stimulation. Consistent with the decrease in PGE2 protein, the LPS-induced upregulation of COX-2 mRNA expression was attenuated after treatment with empagliflozin or gemigliptin (Figure 2(b)). mRNAs of another important inflammatory enzyme, iNOS, was also downregulated by empagliflozin or gemigliptin (Figure 2(c)). Consequently, COX-2 and iNOS proteins were reduced in LPS-stimulated RAW 264.7 macrophages treated with empagliflozin or gemigliptin, indicating that the reduction of PGE2 protein was due to the suppression of the gene responsible for expression (Figures 2(d) and 2(e)).

3.3. Empagliflozin and Gemigliptin Reduced LPS-Induced Proinflammatory Cytokines and Chemokines in RAW 264.7 Macrophages. We confirmed increased production of inflammatory cytokines, such as TNF-α, IL-1β, IL-6, and IFN-γ, in the supernatant of LPS-treated RAW 264.7 macrophages [40]. These cytokines were reduced when the macrophages were treated with empagliflozin or gemigliptin, and an additional reduction was observed in empagliflozin plus gemigliptin-treated macrophages (Figure 3(a)). LPS-stimulated RAW 264.7 macrophages naturally upregulated the mRNA expression of proinflammatory cytokines (TNF-
standard errors CCK-8 assay. (d) M1 macrophage marker, CD80, was assessed via μ with LPS alone, plus 500 and Gemi-, Empa-, or Gemi+Empa-treated groups.

α, IL-1β, and IL-6) and chemokines (CCL3, CCL4, CCL5, and CXCL10). Treatment with empaglifl ozin or gemigliptin reversed the mRNA overexpression of these cytokines and chemokines in LPS-induced RAW 264.7 macrophages (Figures 3(b) and 3(c)). The effect of the empaglifl ozin plus gemigliptin combination was significantly greater than that of empaglifl ozin or gemigliptin alone.

3.4. Impact of Empaglifl ozin and Gemigliptin on NF-κB, MAPK, and STAT Signalling Pathways in RAW 264.7 Macrophages. LPS activated various transcription factors associated with the inflammatory response in RAW 264.7 macrophages. Therefore, we conducted Western blot analysis of NF-κB, MAPKs, and STAT signalling pathways to clarify the potential mechanisms of action of SGLT2 and DPP-4 inhibitors in the LPS-induced release of proinflammatory mediators, such as cytokines and chemokines [41–43]. We also used dexamethasone, a well-known anti-inflammatory reagent, as a positive control to compare the efficacy of empaglifl ozin and gemigliptin [44, 45]. NF-κB is a crucial transcription factor for proinflammatory mediators in macrophages [46]. NF-κB phosphorylation was suppressed in RAW 264.7 macrophages treated with empaglifl ozin or gemigliptin. We also confirmed that empaglifl ozin and gemigliptin (individually and combined) inhibited IKK phosphorylation, which is a central regulator of NF-κB activation, in LPS-activated RAW 264.7 macrophages (Figure 4(a)). STATs are another important group of transcription factors in a variety of cytokines and growth factors for mediating pro- and anti-inflammatory responses [43]. STAT1 phosphorylation was attenuated in RAW 264.7 macrophages treated with empaglifl ozin or gemigliptin. However, gemigliptin was the more potent suppressor of STAT1/3 phosphorylation compared with empaglifl ozin (p < 0.05). Empaglifl ozin and gemigliptin (individually and combined)
Figure 2: Continued.
inhibited LPS-induced JAK2 phosphorylation in macrophages (Figure 4(b)).

MAPKs mediate biological processes and cellular responses to external stimuli, such as LPS [42, 47]. The JNK, p38, and ERK phosphorylation significantly increased in RAW 264.7 macrophages after LPS induction. Neither empagliflozin nor gemigliptin affected p38 and ERK phosphorylation; meanwhile, empagliflozin and gemigliptin (individually and combined) markedly attenuated JNK phosphorylation in RAW 264.7 macrophages activated by LPS. This effect of JNK phosphorylation was more pronounced in empagliflozin treatment (Figure 4(c)). To further determine the MAPK kinases involved in the JNK pathway inhibition, we examined MKK4 and MKK7, which cooperate in JNK activation. Empagliflozin significantly suppressed both MKK4 and MKK7 phosphorylation in RAW 264.7 macrophages stimulated by LPS, while gemigliptin mainly inhibited MKK7 activation (Figure 4(d)).

4. Discussion

This study confirmed potent anti-inflammatory functions and concurrently demonstrated an anti-inflammatory pathway of SGLT2 and DPP-4 inhibition in LPS-activated RAW 264.7 macrophages. Empagliflozin and gemigliptin concurrently reduced proinflammatory cytokine and chemokine release and gene expression via the IKK/NF-κB, JAK2-STAT1/3, and MKK4/7-JNK pathways in LPS-stimulated RAW 264.7 macrophages (Figure 5).

During inflammation, a host defense mechanism against various harmful stimuli, activated macrophages play an important role by producing several proinflammatory cytokines, such as TNF-α, IL-1β, IL-6, and IFN-γ, and inflammatory mediators, including NO and PGE₂ [48]. However, chronic low-grade inflammation may result in metabolic syndrome, diabetes, cardiovascular disease, and cancer [49–52]. Currently, antidiabetic drugs are in the spotlight for their anti-inflammatory properties [53]. There is a meta-analysis to show C-reactive protein and proinflammatory cytokine IL-6 reduction in people treated with at least one SGLT2 or DPP-4 inhibitor among patients confirmed with coronavirus infection [54].

One SGLT2 inhibitor, canagliflozin, but neither dapagli-flozin nor empagliflozin, inhibited proinflammatory cytokine IL-6 in an AMP-activated protein kinase (AMPK) dependent manner in human endothelial cells (HUVECs) [55]. Canagliflozin also diminished proinflammatory cytokines, such as TNF-α, IL-1β, and IL-6, released by AMPK activation in LPS-treated RAW 264.7 macrophages [55]. In our study, empagliflozin decreased secretion of the proinflammatory mediators and downregulated the mRNA levels of COX-2, iNOS, TNF-α, IL-1β, and IL-6 in LPS-induced RAW 264.7 macrophages compared to LPS alone. Additionally, we confirmed its inhibitory effect on CCL3, CCL4, CCL5, and CXCL10 mRNA expression.
This divergence of empaglifoxin according to the cell lines could be explained by the different distributions of SGLT1 and SGLT2, and the different concentrations used 1 μM in HUVECs vs. 80 μM in RAW 264.7 macrophages. Xu et al. [17] recently confirmed SGLT2 protein expression in RAW 264.7 macrophages, and in another study, the selectivity for SGLT2 of empaglifoxin (half maximal inhibitory concentration, IC_{50} 3.1 nM) was similar to that of canaglifoxin (IC_{50} 2.7 nM), but the selectivity for SGLT1 (IC_{50} 8,300 nM) was much weaker than that of canaglifoxin (IC_{50} 710 nM) [56].

According to other study groups [57], canaglifoxin reduced proinflammatory cytokines by lowering hexose kinase II (HKII) and blocking ERK phosphorylation, but not affecting NF-κB in LPS-stimulated human coronary artery endothelial cells (HCAECs). Dapaglifoxin also suppressed iNOS, TNF-α, IL-1β, and IL-6 mRNA expression by attenuating the NF-κB transcription factor in diet-induced atherosclerosis in rat aortic arteries [58]. In the mouse hepatic inflammation model [59], empaglifoxin reduced the number of F4/80⁺ M1 macrophages and the mRNA expression of proinflammatory cytokines by attenuating NF-κB and JNK phosphorylation.

We also confirmed that empaglifoxin inhibited NF-κB phosphorylation like other SGLT2 classes. Additionally, we investigated another inflammatory signalling STATs pathway to elucidate an anti-inflammatory mechanism. Empaglifoxin inhibited STAT1 and STAT3 phosphorylation and also suppressed its upstream JAK2 activation. Subsequently, we verified that empaglifoxin blocked IKK, which is upstream kinase of NF-κB. However, it only suppressed JNK signalling without affecting p38 and ERK and naturally inhibited MKK4 and MKK7, which are upstream molecules of JNK. Empaglifoxin inhibited the MKK4/JNK pathway more significantly than gemigliptin.

Consequently, we revealed that empaglifoxin markedly decreased the number of CD80⁺ M1 macrophages and...
Figure 4: Empagliflozin and gemigliptin attenuated the NF-κB, JNK, and STAT signalling pathways in RAW 264.7 macrophages. LPS-stimulated RAW 264.7 macrophages were treated with empagliflozin (Empa), gemigliptin (Gemi), empagliflozin plus gemigliptin (Empa+Gemi), or 50 μM dexamethasone (DEXA) for 4 h. Here, DEXA served as the positive control for the anti-inflammatory reagent. (a) IKKa/β, NF-κB, (b) JAK2, STAT1/3, (c) MAPKs, and (d) MKK4/7 signalling pathways in the cell lysates prepared were measured using Western blotting. Relative expression of each protein was compared. Data are presented as the means ± standard errors using Western blotting. Relative expression of each protein was compared. Data are presented as the means ± standard errors.

![Figure 4](image1.png)

Figure 5: Proposal of a proinflammatory pathway inhibited by empagliflozin (SGLT2 inhibitor) and gemigliptin (DPP-4 inhibitor) in LPS-activated RAW 264.7 macrophages.

![Figure 5](image2.png)
and MKK7 are known to be upstream signalling components that are required for JNK, and we verified that inhibition of the JNK signal may result from blocking MKK7 signalling by gemigliptin. Notably, we newly proved that gemigliptin inhibits proinflammatory cytokine and chemokine production by reducing STAT1/3 phosphorylation, achieved by blocking the JAK2 signalling pathway, in LPS-induced RAW 264.7 macrophages.

These results suggest that gemigliptin improves inflammation in macrophages by inhibiting the activation of the IKK and NF-κB, MKK7 and JNK, and JAK2 and STAT1/3 pathways. Gemigliptin was more potent in inhibiting the JAK2-STAT1/3 pathways than empagliflozin.

Only a few in vivo study reports on the anti-inflammatory effects of the combination of SGLT2 and DPP-4 inhibitors. Dual therapy, empagliflozin and linagliptin, synergistically ameliorated hepatic fibrosis and inflammation by reducing mRNA expression of proinflammatory cytokines in mouse liver [59]. Another combination therapy with dapagliflozin and saxagliptin ameliorated diabetic cardiomyopathy and diabetic nephropathy in mice by inhibiting proinflammatory cytokines and NLRP3/ASC inflammasome production [15, 64]. However, no studies have investigated whether the combination of SGLT2 and DPP-4 inhibitors directly affects the inflammatory response and its signalling pathways involved in macrophages. We clarified for the first time that the dual therapy, empagliflozin and gemigliptin, has stronger anti-inflammatory effects in RAW 264.7 macrophages.

5. Conclusions

In summary, we verified that empagliflozin and gemigliptin individually possess anti-inflammatory activity by reducing PGE$_2$ and proinflammatory cytokine production, resulting from the inhibitory effects of COX-2, iNOS, cytokine, and chemokine mRNA expression in RAW 264.7 macrophages. These results were mediated by blocking the IKK and NF-κB, MKK4/7 and JNK, and JAK2 and STAT1/3 signalling pathways. MKK4/JNK phosphorylation was prominently inhibited by empagliflozin, while JAK2-STAT1/3 activation was primarily suppressed by gemigliptin. All these anti-inflammatory effects were enhanced when empagliflozin was combined with gemigliptin compared to empagliflozin or gemigliptin treatment alone. This study is the first to show the direct effect of the anti-inflammatory mechanisms of empagliflozin and gemigliptin on macrophages and presents the stronger anti-inflammatory properties of combination therapy.

Data Availability

The data used to support the findings of this study are available from the corresponding authors upon reasonable request.

Conflicts of Interest

All authors have no conflicts of interest to declare.

Authors’ Contributions

HJK, NL, YJH, and SEC performed the conceptualization. NL, YJH, and HJK performed the data curation. YJH and NL contributed to the experiments. HJK, KWL, and YK did the supervision. NL and YJH wrote the original draft. NL, YJH, HJK, YYJ, SJH, and DJK wrote, reviewed, and edited the manuscript. All authors agreed with the conclusion and approved the final version of manuscript.

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