Research Article

Hangeshashinto Inhibits Porphyromonas gingivalis Pathogen-Associated Molecular Patterns-Mediated IL-6 and IL-8 Production through Toll-Like Receptors in CAL27 Cells

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While previous reports have established the anti-inflammatory effects of hangeshashinto, the intracellular signal transduction pathways involved have yet to be elucidated. We aim to employ an experimental system using oral cancer cells to assess the impact of hangeshashinto on intracellular signal transduction pathways in response to stimulation by Porphyromonas gingivalis pathogen-associated molecular patterns (PAMP). Hangeshashinto demonstrated the ability to inhibit the production of interleukin (IL)-6 and IL-8 induced by P. gingivalis PAMP. Furthermore, hangeshashinto suppressed the activation of the IL-6 promoter stimulated by PAMP. Hangeshashinto, like Toll-like receptor (TLR) signaling inhibitors (resatorvid and C29) and an immunosuppressant (dexamethasone), exhibited the ability to suppress TLR-mediated activation of the transcription factor nuclear factor-xB (NF-xB) in response to PAMP stimulation. This study suggests that the anti-inflammatory effects of hangeshashinto may be attributed to the inhibition of TLR signal transduction pathways including NF-xB activation, thereby suppressing NF-xB-dependent gene expression.

1. Introduction

Kampo is a traditional Japanese medicine with its roots in Chinese medicine. It encompasses herbal remedies and is currently prescribed by physicians and dentists in Japan [1]. Among the various kampo medicines, hangeshashinto holds a significant place and can be traced back to ancient Chinese medical texts such as “Shang Han Lun” and “Jin Gui Yao Lue,” compiled in the early 3rd century [2]. Hangeshashinto consists of seven herbal extracts, including pinellia tuber, coptis rhizome, scutellaria root, processed ginger, glycyrhiza, ginseng, and jujube. It has gained approval as a prescribed medicine in both Japan and China. Notably, hangeshashinto has been increasingly prescribed in Japan for the treatment of oral mucositis, and clinical studies employing randomized place-bo-controlled double-blind trials have demonstrated its efficacy in managing oral mucositis caused by chemotherapy for colon and stomach cancers [3]. In clinical practice, hangeshashinto is administered three times a day, with a dosage of 4.5 grams each time for oral mucositis treatment. Our research group has been at the forefront of investigating the anti-inflammatory effects of hangeshashinto. Specifically, we have observed its ability to inhibit lipopolysaccharide (LPS)-induced prostaglandin E2 (PGE2) production, reduce the secretion of interleukin (IL)-6 and IL-8, suppress the activity of cyclooxygenase (COX)-1 and -2, and decrease the expression of cytosolic phospholipase (PL) A2 and LPS-induced COX-2 in human gingival fibroblasts [4]. Subsequent in vitro [5, 6] and in vivo experiments [7–9] have further shed light on the underlying mechanisms by which hangeshashinto improves oral mucositis, highlighting its anti-inflammatory and antibacterial properties. While previous reports have established the anti-inflammatory effects of hangeshashinto, the intracellular signal transduction pathways involved have yet to be elucidated. We aim to employ an experimental system
using oral cancer cells to assess the impact of hangeshashinto on intracellular signal transduction pathways in response to stimulation by Porphyromonas gingivalis (P. gingivalis) pathogen-associated molecular patterns (PAMP). P. gingivalis is a bacterium associated with periodontitis and serves as a reliable stimulus for IL-6 production in oral cancer cells [10]. An experimental system has been established that utilizes P. gingivalis LPS as a model for inflammation, inducing the production of inflammatory cytokines in cultured cells [11]. It is recognized that these inflammatory cytokines exert influence on the development of stomatitis [12]. Our study aims to provide valuable insights into the effects of hangeshashinto on intracellular signaling in response to P. gingivalis PAMP stimulation, using an experimental model involving oral cancer cells. This investigation will contribute to a deeper understanding of the mechanisms underlying the therapeutic potential of hangeshashinto in managing oral mucositis and related inflammatory conditions.

2. Materials and Methods

2.1. Cell Culture. CAL27 cells (oral squamous cell carcinoma, American Type Culture Collection, Virginia, USA) were cultured at 37°C and 5% CO₂ in Dulbecco’s Modified Eagle Medium (DMEM) (manufactured by Nissui Pharmaceutical, Tokyo) containing 10% fetal bovine serum (FBS), 100 units/ml penicillin G, and 100 μg/ml streptomycin.

2.2. Reagents. Hangeshashinto was provided by Tsumura (Tokyo, Japan). P. gingivalis LPS (InvivoGen, California, USA, referred to as "PAMP" in this study) [10], 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) and dexamethasone (Sigma-Aldrich, Missouri, USA); resatorvid and C29 (Selleck Bio, Wisconsin, USA); anti-IL-6 antibody and biotinylated anti-IL-6 antibody (Thermo Fisher Scientific (BioSource)) [10].

2.3. MTT Assay. CAL27 cells (1 × 10⁴) were cultured for 24 hours in concentrations of 10, 100, and 1000 μg/mL of hangeshashinto. Subsequently, the cells were cultured with 0.5 mg/ml MTT for 4 hours. Then, 0.02 N HCl-isopropanol was added to the medium to dissolve and solubilize the formazan product completely. Samples were measured using a Microplate Reader model 550 (Bio-Rad Labs, Inc., Hercules, California, USA) with dual wavelengths of 595/655 nm (test/reference), as previously described [10].

2.4. Enzyme-Linked Immunosorbent Assays (ELISAs). CAL27 cells (1 × 10⁴) were cultured for 24 hours in a mixture of P. gingivalis PAMP (100 ng/mL) and hangeshashinto (1000 μg/mL). The levels of IL-6 in the culture medium were measured using anti-IL-6 (1 μg/mL) and biotinylated anti-IL-6 (0.6 μg/mL) antibodies. The levels of IL-8 in the culture medium were also measured using anti-IL-8 (2.5 μg/mL) and biotinylated anti-IL-8 (0.2 μg/mL) antibodies. Samples were measured using a Microplate Reader model 550 with dual wavelengths of 450/655 nm (test/reference). The ELISA procedure was carried out as described in the CytoSet Kit User Manual (Thermo Fisher Scientific (BioSource)) [10].

2.5. Transfection and Luciferase Assay. One microgram of phIL-6pro-Luc (a fusion of the human IL-6 promoter region spanning from position 2636 to 5035 of GenBank Accession Number NG_011640 linked to the luciferase gene (Figure 2(a)) [10], along with 0.1 μg of pRSV-β-gal (standard plasmid), was mixed with TransIT-LT1 reagent (Mirus Bio, Wisconsin, USA). Additionally, 1 μg of plgx-Luc (Figure 3(a)) [10] and 0.1 μg of pRSV-β-gal were also mixed with TransIT-LT1 reagent. The mixtures were then added to the cells (3 × 10⁵), which were subsequently cultured for 24 hours. Afterward, the cells were stimulated with hangeshashinto (1000 μg/mL) for 24 hours, followed by stimulation with P. gingivalis PAMP (100 ng/mL) for 6 hours. The cells were harvested and lysed. The luciferase and β-galactosidase activities in the lysates were measured as previously described [10].

2.6. Statistical Analysis. Quantitative data were analyzed using one-way analysis of variance (ANOVA). Data were then statistically analyzed using the Tukey’s test (StatMate software (ATMS, Chiba, Japan)). Significant differences were considered statistically significant at p < 0.05.

3. Results

3.1. Effect of Hangeshashinto on Viability of CAL27 Cells. We examined the impact of hangeshashinto on the viability of CAL27 cells. CAL27 cells were subjected to a 24-hour culture with hangeshashinto, followed by an MTT assay. As depicted in Figure 4, the presence of hangeshashinto at concentrations up to 1000 μg/mL did not exert any discernible influence on the viability of CAL27 cells. Therefore, the experiments described below were carried out with a concentration of 1000 μg/mL hangeshashinto.

3.2. Effects of Hangeshashinto on Proinflammatory Cytokine Production by PAMP Stimulation. CAL27 cells were cultured with PAMP stimulation in the presence of hangeshashinto. The amounts of IL-6 and IL-8 in the culture medium were measured using the ELISA method. The results indicate that PAMP stimulation significantly enhanced the production of IL-6 (Figure 5(a), p < 0.001) and IL-8 (Figure 5(b), p < 0.001) in CAL27 cells. The levels of production approximately 2.5 times for IL-6 and approximately 3 times for IL-8 higher compared to that of unstimulated cells. Interestingly, the presence of hangeshashinto resulted in an approximate 60% reduction in IL-6 production induced by PAMP stimulation. These results suggest that
Figure 1: Three-dimensional high-performance liquid chromatography profile of hangeshashinto (provided by Tsumura, Tokyo, Japan).

Figure 2: Continued.
Figure 2: (a) Schematic representation of a reporter plasmid pIL-6pro-Luc containing the IL-6 promoter linked to the luciferase gene (luc). Putative binding sites of transcriptional factors are indicated. Plasmids of pIL-6pro-Luc and pRSV-β-gal (standard) were cotransfected into CAL27 cells. Twenty-four hours after transfection, the cells were cultured with hangeshashinto (b), resatorvid and C29 (c), and dexamethasone (d) for 24 hours. Then, the cells were stimulated with PAMP for 6 hours. The cells were lysed, and luciferase and β-galactosidase assays were performed. The transcriptional activities indicated fold values of those with no stimulators. Bars represent the means and range of triplicate samples. ***, p < 0.001 versus stimulation with PAMP alone (one-way ANOVA with Tukey’s test).

Figure 3: Continued.
hangeshashinto suppresses the production of proinflammatory cytokines by *P. gingivalis* PAMP in CAL27 cells.

### 3.3. Transcriptional Regulation of the IL-6 Promoter by PAMP in the Presence of Hangeshashinto.

The effect of hangeshashinto on the transcriptional activity of the IL-6 promoter was investigated. A reporter plasmid containing the IL-6 promoter linked to the luciferase gene (Figure 2(a)) was transfected into CAL27 cells. The cells were stimulated with PAMP in the presence or absence of hangeshashinto, and a luciferase assay was performed. As shown in Figure 2(b), PAMP enhanced the transcriptional activity of the IL-6 promoter, but hangeshashinto alone did not enhance it. When CAL27 cells were stimulated with PAMP in the presence of hangeshashinto, the level of transcriptional activity significantly decreased by approximately 50% compared to that of PAMP stimulation alone (*p* < 0.001). Resatorvid and C29, Toll-like receptor (TLR) 4 and TLR2 signaling inhibitors, respectively, also significantly repressed the transcriptional activity of the IL-6 promoter upon PAMP stimulation (Figure 2(c), *p* < 0.001). In addition, dexamethasone, which is an immunosuppressant and an anti-inflammatory agent, repressed that activity (Figure 2(d), *p* < 0.001). These results suggest that hangeshashinto inhibits the transcriptional activation of the IL-6 promoter via the TLR signaling pathways.

### 3.4. Nuclear Factor-κB (NF-κB) Activation by PAMP with or without Hangeshashinto.

Next, the effect of hangeshashinto on the activation of NF-κB by PAMP stimulation was investigated. An NF-κB-dependent reporter plasmid (Figure 3(a)) was transfected into CAL27 cells. The cells were stimulated with PAMP in the presence or absence of hangeshashinto, and luciferase assays were performed. As shown in Figure 3(b), the level of NF-κB-dependent transcriptional activation by PAMP stimulation was approximately 4.5 times higher than that without PAMP stimulation. When CAL27 cells were stimulated with PAMP in the presence of hangeshashinto, the level of transcriptional activation significantly decreased compared to that in the absence of hangeshashinto (*p* < 0.001). The transcriptional activation with PAMP in the presence of resatorvid or C29 was also significantly suppressed.
compared to that with PAMP alone (Figure 3(c), \( p < 0.001 \)). Moreover, dexamethasone significantly suppressed the transcriptional activation by stimulating with PAMP in CAL27 cells (Figure 3(d), \( p < 0.001 \)). These results suggest that hangeshashinto inhibits NF-\( \kappa \)B activation through the TLR signaling pathways.

4. Discussion

Hangeshashinto which has an anti-inflammatory effect is indicated for stomatitis [5, 6]. Recent review studies have demonstrated the high effectiveness of hangeshashinto in the treatment of stomatitis. Currently, hangeshashinto is widely used for preventing and treating oral mucositis [13], as it has anti-inflammatory properties. However, the anti-inflammatory mechanism of action of hangeshashinto is not known in detail. TLR signaling induces the activation of some transcription factors and thereby expresses various genes. Upon binding of ligands (PAMP) to TLRs, in the myeloid differentiation factor 88 (MyD88)-dependent pathway, adaptor proteins such as MyD88 and Toll/IL-1R domain–containing adaptor inducing IFN-\( \beta \) (TRIF) are recruited to TLRs [14], resulting in forming myddosome, which contains MyD88 and interleukin 1 receptor associated kinase (IRAK) family [15]. IRAK1 activated by autophosphorylated IRAK4 activates TNF receptor–associated factor 6 (TRAF6) [16]. TRAF6 signaling induces the activation of mitogen-activated protein kinase MAPK (c-Jun N-terminal kinase (JNK) and p38) and NF-\( \kappa \)B [17]. Subsequently, activator protein-1 (AP-1, dimers of FOS and JUN) and NF-\( \kappa \)B stimulate the expression of proinflammatory cytokines and chemokines [18]. The present finding showed that the NF-\( \kappa \)B activation through TLRs by PAMP was observed, which was significantly suppressed by hangeshashinto (Figure 3(b)).

Figure 5: Effects of hangeshashinto on the production of proinflammatory cytokines in CAL27 cells by PAMP. Hangeshashinto (1000 \( \mu \)g/ml) and PAMP (100 ng/ml) were added to CAL27 cells and the cells were cultured for 24 hours. The levels of IL-6 (a) and IL-8 (b) in the culture media were measured by ELISAs. ***, \( p < 0.001 \) versus stimulation with PAMP alone (one-way ANOVA with Tukey’s test).
expression of the COX-2 gene is suppressed by baicalin and berberine, which are constituents of scutellaria root. Additionally, it is suggested that wogonin, another component of scutellaria root, inhibits the MAPK pathway and suppresses the transcription factor activation factor NF-κB [9]. Animal studies have suggested that constituents of hangeshashinto, such as baicalin from scutellaria root and berberine from coptis rhizome, may have anti-inflammatory effects by suppressing the production of pro-inflammatory cytokines [23]. Moreover, hangeshashinto has been reported to ameliorate chemotherapy-induced oral mucositis by reducing oxidative stress and inhibiting the expression of inflammatory mediators [23]. Further research is needed to elucidate the molecular mechanisms underlying the pharmacological effects of hangeshashinto. In recent years, we conducted a randomized double-blind placebo-controlled trial of hangeshashinto in 16 patients with head and neck cancer who developed oral mucositis due to chemical radiation therapy. The trial demonstrated that gargling with hangeshashinto shortened the healing time of oral mucositis and reduced the severity of mucositis grade [24]. In addition, a study involving 22 terminally ill cancer patients reported that hangeshashinto improved oral dryness and may prevent the development of oral mucositis [25]. Notably, in addition to its anti-inflammatory effects, hangeshashinto has been reported to selectively inhibit the growth of Gram-negative bacteria, demonstrating potential antibacterial properties [26]. The importance of prevention has been recognized in traditional Chinese medicine for over 2000 years, as evidenced by the statement “a wise man treats the disease before it occurs” in The Yellow Emperor’s Classic of Medicine. In Oriental medicine, the term “pre-disease” refers to a state of mild symptoms that do not progress to full-blown disease [27]. Therefore, the anti-inflammatory effects of kampo medicine may be viewed as a means of preventing pre-disease from progressing to full-blown disease.

5. Conclusions

In conclusion, this study suggests that the anti-inflammatory effects of hangeshashinto may be attributed to the inhibition of TLR signal transduction pathways including NF-κB activation, thereby suppressing NF-κB-dependent gene expression.

Abbreviations

- P. gingivalis: Porphyromonas gingivalis
- PAMP: Pathogen-associated molecular patterns
- NF-κB: Nuclear factor-κB
- TLR: Toll-like receptor
- LPS: Lipopolysaccharide
- PGE2: Prostaglandin E2
- IL: Interleukin
- COX: Cyclooxygenase
- PL: Phospholipase
- DMEM: Dulbecco’s modified eagle medium
- FBS: Fetal bovine serum
- MTT: 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide
- ANOVA: Analysis of variance
- MyD88: Myeloid differentiation factor 88
- TRIF: Toll/IL-1R domain-containing adaptor inducing IFN-β
- IRAK: Interleukin 1 receptor-associated kinase
- TRAF6: TNF receptor-associated factor 6
- MAPK: Mitogen-activated protein kinase
- JNK: c-Jun N-terminal kinase
- AP-1: Activator protein-1

Data Availability

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

Conflicts of Interest

The authors declare that there are no conflicts of interest regarding the publication of this paper.

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References


