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Yodthong, T., Kedjarune-Leggat, U., Smythe, C. orcid.org/0000-0001-6526-0565 et al. (4 more authors) (2020) Enhancing activity of Pleurotus sajor-caju (Fr.) Sing β-1,3-Glucanoligosaccharide (Ps-GOS) on proliferation, differentiation, and mineralization of MC3T3-E1 cells through the involvement of BMP-2/Runx2/MAPK/Wnt/β-catenin signaling pathway. Biomolecules, 10 (2). 190.

https://doi.org/10.3390/biom10020190

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Enhancing Activity of *Pleurotus sajor-caju* (Fr.) Sing β-1,3-Glucanoligosaccharide (Ps-GOS) on Proliferation, Differentiation, and Mineralization of MC3T3-E1 Cells through the Involvement of BMP-2/Runx2/MAPK/Wnt/β-Catenin Signaling Pathway

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1. Introduction

Osteoporosis is a global disease resulting from an imbalance of bone homeostasis. An increase in the ratio of bone resorption to bone formation causes low bone mass and strength, which leads...
to increased fragility and susceptibility to fracture [1]. The disease occurs in older (>50 year) men and women [2], with increased likelihood in post-menopausal women. The latter effect is due to the absence of estrogen [3], which plays a crucial role in maintaining bone metabolic balance in bone homeostasis [4,5].

Bone remodeling is a continuous process regulated by two major types of bone cell. Osteoclasts mediate bone resorption to remove small pieces of bone fractures, and osteoblasts mediate bone formation to form new bone tissue [6]. An imbalance involving an increase in bone resorption and/or a decrease in bone formation can cause osteoporosis [7].

Osteoblastogenesis—the process of osteoblast proliferation and differentiation contributing to bone strength and rigidity [8]—is regulated by several signal transduction pathways, including the bone morphogenetic protein (BMP) signaling pathway. BMP-2, one of the most important growth factors in the BMP subfamily, plays an important role by regulating the downstream osteogenic runt-related transcription factor2 (Runx2) [9]. The function of Runx2 is to promote the early stage of osteoblastogenesis, by triggering the expression of bone matrix genes and calcium deposition for mineralization [10]. As inhibition of BMP-2 activity is known to result in a loss of induction of the osteoblast differentiation marker Alkaline Phosphatase (ALP) [11], it follows that BMP-2 is a potential target to seek alternative novel molecular entities that promote bone formation.

In addition to the BMP signaling pathway, both mitogen-activated protein kinase (MAPK) and wingless-type MMTV integration site (Wnt)/β-catenin signaling pathways are involved in the progression of bone remodeling via osteoblast differentiation [12]. MAPK family members extracellular signal regulated-kinases (ERKs), c-Jun N-terminal kinases (JNKs), and p38 kinase perform essential functions in many mammalian cells, including osteoblasts, by controlling proliferation, differentiation, and development [13]. Additionally, Wnt/β-catenin signaling is required for pre-osteoblast proliferation, osteoblastogenesis, and inhibition of osteoblast apoptosis [14,15]. In vivo deletion of β-catenin causes a reduction in bone formation and the induction of osteoclastogenesis [16].

While there are many therapeutics in use for osteoporosis treatment, there are significant risks associated with them. Long-term use of antosteoporotic drugs such as estrogen replacement therapy are associated with breast cancer and ovarian cancer [17,18], while bisphosphonates can cause osteonecrosis of the jaw [19]. It follows there is a unmet need for new agents, with fewer side effects, that can promote bone formation.

Various types of naturally-derived compounds, including glycosides, flavonoids, terpenoids, coumarins, phenols, and phenolates, have been reported to restore bone homeostasis via a range of cellular mechanisms, including the modulation of transcription factors, bone-specific protein, signal pathways, and the OPG/RANKL system [20]. β-Glucans, glucose polymers containing a β-glycosidic link, are derived from natural dietary fiber and are potential candidates for the treatment or prevention of osteoporosis. Numerous β-glucans, with varying degrees of polymerization and branching, including β-1→3-, β-1→4-, β-1→6-, β-1→3/1→4-, and β-1→3/1→6-linkages, are found in a variety of natural sources including cereals, fungi, yeast, bacteria, and algae [21]. β-glucans have been extensively studied, and a wide range of biological activities have been reported, including immune enhancement [22] as well as antitumor, anti-aging, antihypcholesterolemic, antihyperglycemic [23], and antiinflammation activity [24]. Moreover, antosteoporosis activity has also been reported both in vitro and in vivo [25]. For example, β-glucan extracted from P. citrinopileatus was shown to inhibit RANKL-induced osteoclast differentiation [25]. In addition, β-(1,3)-(1, 6)-glucan extracted from Aureobasidium pullulans had a stimulatory effect on BMP-7-associated osteoblast differentiation [26] and the suppression of bone loss in an ovariectomy-induced osteoporosis rat, an animal model for post-menopausal osteoporosis [27]. Nevertheless, the molecular mechanisms by which β-glucans promote bone formation in osteoblasts have not yet been investigated.

Here, we wished to investigate the enhancing activity and underpinning molecular mechanisms of a β-glucan, extracted from the popular edible grey oyster mushroom, Pleurotus sajor-caju [28], on osteoblastogenesis. This mushroom has been widely used in traditional medicine and reported
to possess a variety of biological activities, including antimicrobial, antitumor, antioxidant, anti-hypertension, anti-diabetic, and anti-inflammatory. According to its various biological activities, many biologically active compounds have also been studied regarding its role in therapeutic applications such as β-glucans, proteoglycan, phenolic acids, terpenes, proteins, and sterols. Extracted Pleurotus sajor-caju β-glucan has a high molecular weight and low water solubility. The biological activities of β-glucans usually depend on physicochemical properties, source, purity, primary structure, water solubility, and molecular weight. Insoluble particulate β-glucans have limited potential for medicinal applications, but may have more applicability following partial hydrolysis. For example, curdlan, which is a water-insoluble microbial linear exo-polysaccharide (1→3) β-D-glucan, has recently been effectively digested with a novel recombinant endo-β-1→3-glucanase to obtain a water soluble glucanoligosaccharide. Here, we have used Hevea β-1,3-glucanase isozymes GI and GII, previously reported to specifically hydrolyze glucans such as laminarin, containing a low frequency of β-1,3-D-glucosidic linkages. We have digested particulate β-glucans to obtain a hydrolysate, Pleurotus sajor-caju glucanoligosaccharide (Ps-GOS), with shorter chain length, lower molecular weight, and higher water solubility (Ratthajak et al., manuscript in preparation).

The aim of this study is to investigate the effect of Ps-GOS on the proliferation and differentiation of the murine pre-osteoblastic cell line MC3T3-E1. We show that Ps-GOS promotes MC3T3-E1 cell proliferation, and induction of BMP-2 and Runx-2, which act coordinately, together with components of MAPK and Wnt/β-catenin signaling pathways, to enhance both gene expression of osteogenic bone markers and mineralization.

2. Materials and Methods

2.1. Materials

Alpha modification of Eagle’s minimum essential medium (α-MEM), fetal bovine serum (FBS), RPMI-1640 phenol red-free medium, and tissue culture reagents were purchased from Gibco (Grand Island, NY, USA). β-glycerophosphate, L-ascorbic acid, and MTT (3-(4,5-Dimethylthiazol-2-yl)-2,5-Diphenyltetrazolium Bromide) were purchased from Sigma Aldrich (St. Louis, MO, USA). All specific primers were synthesized from Sigma Aldrich (St. Louis, MO, USA). All other reagents and chemicals were the grade quality available.

2.2. Preparation of Water Soluble P. sajor-caju Glucanoligosaccharide (Ps-GOS)

Ps-GOS was obtained from an investigation under the Center of Excellence in Natural Rubber Latex Biotechnology Research and Development (CERB) (additional details are under patent and will be published elsewhere). Briefly, the insoluble β-glucan fraction was isolated from the fruiting bodies of grey oyster mushroom (Pleurotus sajor-caju) according to modified Freimund’s method. The β-glucan assay was conducted according to the method of McClear and Glennie-Holmes, using the Mushroom and Yeast β-glucan kit (K-YBGL) (Megazyme International Ireland Ltd., Wicklow, Ireland). The soluble β-glucan from P. sajor-caju (Ps-GOS) was prepared by mixing the insoluble β-glucan (mean particle size as 646.6 ± 299.3 μm determined using a Beckman Coulter LS 230 Laser Diffraction Particle Size Analyser) with the concentrate enzyme fraction contained 0.66 nkat ml⁻¹ of glucanase activity (ratio of 1.5%, w/v). The optimized Hevea glucanase conditions of the concentrate enzyme fraction were chosen according to report of Churngchow et al., [33]. After complete digestion, the Ps-GOS derived after centrifugation was freeze-dried (mean particle size as 122.7 ± 147.0 μm). Both insoluble β-glucan and Ps-GOS were subjected to FTIR spectroscopy for structural characterization (data not shown). Data from electrospray ionization mass spectrometry (ESI-MS) demonstrated the mass of insoluble β-glucan, and Ps-GOS was distributed from 20 and 5 kDa, respectively.
2.3. Culture and Differentiation

The murine pre-osteoblast cell line, MC3T3-E1 (subclone 14 CRL-2593), was obtained from American Tissue Culture Collection (ATCC, Manassas, VA, USA). MC3T3-E1 cells were cultured in α-MEM containing 10% FBS with 1% penicillin/streptomycin solution (penicillin 100 U/mL, streptomycin 100 U/mL) at 37 °C in a 5% CO₂ incubator (Thermo Fisher Scientific, Waltham, MA, USA). The culture medium was changed every 2–3 days, and cells of a passage number 23 were used for all experiments. To induce osteoblast differentiation, cells were cultured in osteogenic induction medium containing 50 µg/mL l-ascorbic acid and 1 M β-glycerophosphate.

2.4. Cell Proliferation

MTT assay was performed as an indicator for cell proliferation based on the ability of cells to convert MTT into a formazan reaction product. MC3T3-E1 cells were seeded and then treated with Ps-GOS at various concentrations for 24 and 48 h. At the end of treatment, cultured cells were washed with PBS. Next, 5 mg/mL of MTT and RPMI-1640 phenol red-free medium were added and incubated at 37 °C. After 3 h, the mixture was removed and the formazan product was solubilized by acidic isopropanol. The absorbance was measured by microplate reader at 570 nm.

2.5. Alkaline Phosphatase (ALP) Activity Assay

Cells were seeded in 24 well plates and then treated with different concentrations of Ps-GOS for 14 days. After treatment, the cells were washed with PBS and lysed with lysis buffer containing 50 mM Tris-HCl (pH 7.4) and 1% Triton X-100 to collect supernatant for determining ALP activity and protein concentration. ALP activity was determined with 4 mg/mL of 4-nitrophenyl phosphate (4NPP) in 0.2 M of 2-amino-2-methyl-1-propanol with 4 mM of MgCl₂ as a substrate for 30 min at 37 °C. The reaction was stopped by 0.1 M NaOH, and the yellow solution was measured at 405 nm. Protein concentration was measured using the Pierce BCA Protein Assay Kit (ThermoFisher Scientific, Waltham, MA, USA). ALP activity was normalized to the concentration of protein and showed in the term of µmole/min/mg of protein.

2.6. Alizarin Red S Staining

Alizarin red S staining was used to determine calcium accumulation. In brief, cells were grown in 24 well plates and treated with Ps-GOS for 14 and 21 days. After incubation, cells were washed with PBS and then fixed with 10% formaldehyde 15 min. The cells were stained with 40 mM Alizarin red S solution (pH 4.1–4.3) for 30 min with gentle shaking at the room temperature. To quantify the dye, nonspecific staining was removed by washing with distilled water five times (5 min/time) and then solubilizing the stain with cetylpyridinium chloride (CPC) by shaking, and the absorbance at 550 nm was measured.

2.7. Cell Cycle Analysis

Cell cycle assay was performed by flow cytometry to analyze the cell cycle progression while treating with Ps-GOS for 48 h. Cells were trypsinized and fixed in 70% ethanol at 4 °C for 30 min, followed by incubation with propidium iodide containing RNase staining solution (Merck, Branchburg, NJ, USA) for 30 min in the dark to detect DNA content in the cells. The percentage of cells in G0/G1, S, and G2/M phases was analyzed by flow cytometry Guava easyCyteTM HT (Hayward, CA, USA).

2.8. Quantitative Real-Time Polymerase Chain Reaction (qRT-PCR)

Total RNA from growing cells incubated with Ps-GOS for 24, 48, and 72 h was extract using the TriPure Isolation Reagent (Roche, Buonas, Switzerland). To generate cDNA, total RNA (1 µg) was used as a template to transcribe into cDNA using the Transcriptor First Strand cDNA Synthesis Kit (Roche, Buonas, Switzerland) according to the manufacturer’s instruction. This cDNA was used to determine
the gene expression for the genes of interest using the gene specific primers detailed in Yodthong’s
study [37] and GAPDH as the housekeeping control gene. qRT-PCR reactions were amplified using the
EvaGreen HRM Mix (Solis Biodyne, Tartu, Estonia), and then relative expression ratio was calculated
using the $2^{-\Delta\Delta Ct}$ method.

2.9. Western Blot Analysis

Cells were grown and treated with Ps-GOS in six well plates. After indicated time, the cells were
lyzed with lysis buffer and then centrifuged at 14,000×g for 15 min at 4°C to collect supernatant
for using as total protein extract. The concentration of protein was quantified using a protein assay
kit (Bio-Rad, Hercules, CA, USA). Equal amounts of protein from each sample were loaded onto
10% gels for SDS-PAGE, transferred to nitrocellulose membranes (Amersham Pharmacia Biotech,
Amersham Buckinghamshire, UK), blocked with 5% nonfat dry milk solution for 1 h, and then
incubated with antiBMP-2 (Abcam, Milton, UK), antiRunx2 (Cell Signaling Technology, Beverly, MA,
USA), or antiß-Actin (Sigma-Aldrich, St. Louis, MO, USA). After washing three times with TBS-T ween,
the blots were probed with an Alexa infrared dye-conjugated secondary antibody (Invitrogen, Carlsbad,
CA, USA) for 1 h and detected the bands by Odyssey Infrared Imaging System (LI-CORE) in accordance
with the manufacturer’s instruction. The relative of the intensity of the protein interesting bands and
the intensity of ß-Actin band were compared.

2.10. Immunofluorescence Microscopy

The cells treated with Ps-GOS were grown on sterile collagen-coated coverslips in 24 well plates. In brief, coverslips were fixed with 4% paraformaldehyde 20 min, quenched with 50 mM NH$_4$Cl
for 2 × 10 min, washed with PBS, and permeabilized with 0.1% Triton X-100 for 10 min at RT. After
blocking with 0.5% FSG in PBS for 20 min, each coverslip was incubated overnight at 4°C with
polyclonal rabbit antimouse BMP-2 (Abcam, Milton, UK), followed by removing primary antibody
and washing 3 × 5 min in blocking solution before incubation for 1 h with FITC-conjugated secondary
antirabbit IgG (Alexa Fluor 680) (Thermo Fisher Scientific, Waltham, MA, USA) and DAPI counterstain
(Thermo Fisher Scientific, Waltham, MA, USA). Coverslips were washed 3 × 5 min in blocking solution
and mounted on to slides using Prolong® Gold antifade reagent (Invitrogen, Carlsbad, CA, USA).
Fluorescence images were acquired with Nikon Eclipse Ti fluorescence microscope.

2.11. Statistical Analysis

Representative data are presented as the mean ± standard error of the mean (SEM). Statistical
analysis was performed using the SPSS 23 statistical software (SPSS Inc., Chicago, IL, USA). The
significant results were analyzed by one-way analysis of variance (ANOVA), followed by Duncan’s
multiple range test. Different significances are indicated (* = p < 0.05, ** = p < 0.01).

3. Results

3.1. Effect of Ps-GOS on Proliferation of MC3T3-E1 Cells

To determine the dose dependence of Ps-GOS on proliferation, MTT assays were used to measure
the metabolic activity of osteoblastic cells during proliferation. MC3T3-E1 cells were treated with
0.001–1000 µg/mL of Ps-GOS for 24–72 h (Figure 1). Compared to control (cells without treatment),
exposure to Ps-GOS for 48 h significantly increased cell proliferation at concentrations up to 10 µg/mL
(p < 0.05) by 48 h, although proliferation was reduced at all concentrations after 72 h. Ps-GOS appeared
to have a cytotoxic effect at higher concentrations (100 and 1000 µg/mL (p < 0.01)) after treatment
for 72 h. Subsequently, only concentrations in the range 0.001 to 10 µg/mL were used in further
experiments. This results indicated that low concentrations of Ps-GOS stimulate osteoblast proliferation
without toxicity.
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**Figure 1.** The effect of glucanoligosaccharide (\( \text{Ps-GOS} \)) on osteoblastic cell proliferation. MC3E3-E1 cells were treated with the indicated concentrations of \( \text{Ps-GOS} \) for 24, 48, and 72 h. Cell proliferation was measured by MTT assay. The result shows percentage of metabolically active cells relative to untreated cells (control = 100% viability). Each point of data represents the means of four replicate samples ± SEM. * = \( p < 0.05 \), ** = \( p < 0.01 \) when compared to the control.

3.2. Effect of \( \text{Ps-GOS} \) on Cell Cycle Distribution on MC3T3-E1 Cells

Flow cytometry was performed to investigate the effect of \( \text{Ps-GOS} \) on cell cycle progression. Cell cycle analysis was undertaken by measuring cellular DNA content corresponding to each of the three principle cell cycle stages; G0/G1, S, and G2/M. Actively proliferating cells were treated with indicated concentrations of \( \text{Ps-GOS} \) for 48 h. At higher concentrations (1 and 10 μg/mL), \( \text{Ps-GOS} \) significantly decreased the proportion of cells in G0/G1 phase while concomitantly increasing that in S phase when compared to control cells (Figure 2). This study demonstrates that \( \text{Ps-GOS} \) can promote cell proliferation by accelerating progression into S phase from G1.

**Figure 2.** The cell cycle distribution of MC3T3-E1 cells treated with various concentrations of \( \text{Ps-GOS} \).
3.3. Effects of Ps-GOS on Osteoblastic Differentiation of MC3T3-E1 Cells

Mineralization is the last stage of bone formation that results in the formation of calcified nodules that contribute to bone strength [38]. To examine the effect of Ps-GOS, MC3T3-E1 cells were treated with Ps-GOS for 14 and 21 days in an osteogenic induction medium to induce differentiation. Calcium production in MC3T3-1 cells was determined by staining with Alizarin red S dye (Figure 3A). Extraction and quantification of Alizarin red from treated cells showed that Ps-GOS significantly enhanced the calcium deposit formation at both day 14 and 21, with the former time-point showing the maximal increase (175%) when compared to control (Figure 3B) at 1 and 10 μg/mL Ps-GOS. In addition, ALP a
marker of mature osteoblastic differentiation [39] was determined at day 14. All Ps-GOS treatments markedly promoted ALP activity when compared to control (Figure 3C). This result also correlated with the study of Yazid et al.: that the highest ALP level was produced from MC3T3-E1 cells at day 14 during osteoblast differentiation [40]. These results imply that Ps-GOS can promote osteoblast differentiation and mineralization in the process of bone formation.

**Figure 3.** The effect of different concentrations of Ps-GOS on osteoblastic differentiation and mineralization of MC3T3-E1 cells. (A) Cells were treated with Ps-GOS for 14 and 21 days for observing the calcium accumulation measured by Alizarin red S staining. (B) Quantitative data of mineralization process relate to the intensity of Alizarin red S stains extracted by cetylpyridinium chloride. (C) Alkaline phosphatase (ALP) activity of MC3T3-E1 cells after treatment with Ps-GOS for 14 days. Each point of data represents the means of four replicate samples ± SEM. The data in columns with different letters in each group were significantly different at p < 0.05.

### 3.4. Ps-GOS Up-Regulated Osteogenic-Related Gene Marker Expression in MC3T3-E1 Cells

During osteoblast differentiation, many osteogenic gene markers are expressed; these genes play important roles in matrix formation and calcium accumulation during bone formation. We determined the expression levels of OPN, ALP, COL1, OCN, RUNX2, and BMP-2, in MC3T3-E1 cells, after treating with Ps-GOS for 24 and 72 h using qRT-PCR. The results shown in Figure 4 indicate that
the expression of ALP, COL1, OCN, and RUNX2 were up to two times higher than control levels after 24 h exposure to Ps-GOS. Moreover, Ps-GOS at concentrations of 0.1 and 1.0 µg/mL sharply enhanced BMP-2 expression at 24 and 72 h, respectively. These results are consistent with the notion that Ps-GOS promotes osteoblastic differentiation by upregulating osteogenic-related gene expression via the BMP signaling pathway.

Figure 4. Analysis of mRNA level expressions of OPN, OCN, COL1, ALP, RUNX2, and BMP-2 by quantitative real-time polymerase chain reaction (PCR). MC3T3-E1 cells were treated with Ps-GOS in various concentrations for 24 and 72 h. The data were obtained from three independent experiments in four replicates as mean ± SEM. The data in columns with different letters in each group were significantly different at p < 0.05.

3.5. Ps-GOS Stimulates Osteoblast Differentiation via BMP Signaling Pathway

The BMP signaling pathway is responsible for osteoblast differentiation through up-regulation of the key proteins BMP-2 and Runx2. In the present study, we observed that Ps-GOS exposure resulted in an increase in BMP-2 and Runx2 mRNA expression (Figure 4). We thus examined the protein expression of BMP-2 and Runx2 to further validate the mechanism of action of Ps-GOS. Ps-GOS treatment at concentrations up to 10 µg/mL for 48 h resulted in elevated levels of BMP-2, as judged by both indirect immunofluorescence (IF) microscopy (Figure 5A), as well as by Western blotting (Figure 5D). Quantitative analysis of BMP-2 staining (Figure 5A), following treatment with 0.01, 0.1, and 10 µg/mL Ps-GOS for two days, indicated a significant increase in BMP-2 positive staining cells
compared to the control (Figure 5B). Similarly, western blotting analysis showed that all Ps-GOS treatments significantly enhanced protein expression level of BMP-2 at day 2, with maximal increase at concentrations >0.1 µg/mL (Figure 5C). Consistent with the notion that Runx operates downstream of BMP-2, Western blotting of Runx1 showed that levels of this protein were significantly elevated after exposure for 10 days to Ps-GOS concentrations > 0.1 µg/mL (Figure 5D).

Figure 5. Effect of Ps-GOS on the bone morphogenetic protein (BMP) signaling pathway in osteoblast cells. Osteoblastic MC3T-E1 cells were treated with various concentrations of Ps-GOS for 2 days. (A) The image of BMP-2 expressing cells observed by immunofluorescence staining (original magnifications 20×). (B) Quantitation of immunofluorescence staining. (C) Protein level expression of BMP-2 was determined by Western blot analysis at day 2. (D) Protein level expression of Runx2 at day 10. The quantitation was calculated according to the referent bands of β-actin (C) and nucleolin (D). The data were obtained from four replicates as mean ± SEM. The data in columns with different letters in each group were significantly different at p < 0.05.

3.6. Stimulating Effect on Osteoblastic Differentiation by Ps-GOS Could Affect through MAPK and Wnt/β-Catenin Signaling Pathways

While BMP signaling plays a critical role in osteoblast differentiation and bone formation, additional pathways, participating in a range of cross-talk signaling, are also involved. The expression of a range of MAPK family members was investigated by qRT-PCR. As shown in Figure 6, Ps-GOS significantly promoted the expression of ERK1, ERK2, JNK1, and JNK2 following 12 and 24 h exposure, compared to control, and in all cases, expression levels reverted to control levels or below after 48 h. A similar pattern was observed with p38α, although Ps-GOS-enhanced expression was reduced after 48 h exposure and did not revert to control levels. We also investigated expression of genes involved in Wnt/β-catenin signaling (Figure 7). Wnt5a, Fzd4, β-catenin, and LRP5 were significantly up-regulated with Ps-GOS after 12 and 24 h exposure, and expression levels then slightly decreased at 48 h (the same time observed with MAPK). These outcomes suggest that the stimulatory effect of Ps-GOS on osteoblastic differentiation additionally involves stimulation via MAPK and Wnt/β-catenin signaling pathways.
Figure 6. Analysis of mRNA level expressions of MAPK pathway genes ERK1, ERK2, JNK1, JNK2, and p38α by quantitative real-time PCR. MC3T3-E1 cells were treated with Ps-GOS in various concentrations for 24, 48, and 72 h. The data were obtained from three independent experiments in four replicates as mean ± SEM. The data in columns with different letters in each group were significantly different at \( p < 0.05 \).

Figure 7. Analysis of mRNA level expressions of Wnt/β-catenin signaling pathway genes LRP5, β-catenin, Wnt5a, and Fzd4 by quantitative real-time PCR. MC3T3-E1 cells were treated with Ps-GOS in various concentrations for 24, 48, and 72 h. The data were obtained from three independent experiments in four replicates as mean ± SEM. The data in columns with different letters in each group were significantly different at \( p < 0.05 \).
4. Discussion

It is well known that osteoblasts are critical players in the regulation of bone formation and bone remodeling. The reduction of osteoblast activity results in low bone mass and osteoporosis. Recently, it has been shown that β-glucan has a therapeutic effect on osteoporosis by promoting the function of osteoblasts. Lee et al. have suggested that β-(1-3),(1-6)-glucan has a stimulatory effect on BMP-7 promoting bone formation by inducing the production of ALP and calcium accumulation on mineralization [26]. However, the mechanism by which small molecular mass β-glucans, such as Ps-GOS, promote bone formation has not yet been determined.

This study has demonstrated that Ps-GOS has the potential to prevent osteoporosis. At low concentrations (<10 µg/mL), Ps-GOS induced osteoblastic proliferation in MC3T3-E1 cells without toxicity for up to 48 h of treatment. The elevated rate of proliferation correlated with an increase in the proportion of cells in S-phase, suggesting that Ps-GOS promotes proliferation by stimulating the G1-S transition within the cell cycle.

The production of ALP and calcium accumulation is essential steps in the process of mineralization leading to bone formation. ALP, a marker of early-stage osteoblastic differentiation, can be detected on the cell surface and in matrix vesicles of mature osteoblasts [41]. High level expression of ALP initiates and plays a fundamental role during the process of mineralization process [42], where it has been suggested that it hydrolyses pyrophosphate, a natural inhibitor of mineralization, concomitantly increasing local phosphate concentration [43,44]. We found that Ps-GOS stimulated the expression of ALP and promoted the accumulation of calcium, consistent with a potential role for Ps-GOS in promoting bone formation.

In normal development, mesenchymal stem cells function as progenitors of osteoblasts and osteocytes, as well as adipocytes and myocytes [45]. Differentiation of stem cells into an osteoblast lineage requires specific osteogenic signaling proteins and transcription factors. BMP-2, a member of the transforming growth factor-beta (TGF-β) superfamily, is synthesized by osteoblasts and is a critical inducer regulating osteoblastic differentiation and bone formation [46]. BMP-2-induced osteogenesis is absolutely dependent on the formation and transcriptional activity of the downstream osteogenic complex Runx2-SMAD. Runx2 has been reported to mediate the expression of many osteogenic gene markers, including ALP, COL1, OCN, and OPN [47]. Mutations on one allele of human Runx2 gene cause cleidocranial dysplasia [48], while deletion of Runx2 in mice leads to a complete failure of bone formation and ossification [49]. We found that Ps-GOS promoted bone formation by upregulating both gene and protein expression levels of BMP-2 and Runx2. As a consequence of Runx2 up-regulation, Ps-GOS also promoted expression of ALP, COL1, and OCN. Surprisingly, Ps-GOS had no effect on OPN expression. The significance of this observation is not yet clear.

The Wnt/β-catenin signaling pathway works cooperatively with BMP signaling to regulate osteoblast differentiation and bone formation. Crosstalk between Wnt/β-catenin and BMP signaling pathways results in an elevation in BMP-2 and Runx2 expression; the target of this signaling in osteoblasts [50]. Wnt/β-catenin signaling is activated when Wnt ligand binds to the receptors Frizzled (Fzd) and low-density lipoprotein receptor-related protein 5 or 6 (LRP5/6), which in turn inhibits β-catenin phosphorylation. The translocation of β-catenin into nucleus leads to the activation of members of the TCF/LEF transcription factor family in order to promote osteogenic gene expression [51]. Mutation of LRP5 can cause osteoporosis pseudoglioma syndrome (OPPG) in humans [52] and decrease osteoblast numbers in mice [53]. In addition, Wnt5a is an important component in the regulation of Wnt/β-catenin signaling in osteoblast linear cells, as lack of Wnt5a leads to a reduction in LRP5 and LRP6 expression [54]. To identify additional molecular targets of Ps-GOS that might be associated with the up-regulation of BMP-2 and Runx2, we examined expression levels of Wnt/β-catenin-related genes. Our data revealed that Ps-GOS significantly up-regulates the expression of Wnt5a, LRP5, β-catenin, and Fzd4 mRNA, in MC3T3-E1 cells, suggesting that Ps-GOS may prevent bone loss in osteoporosis through Wnt/β-catenin together with BMP-2 signaling pathways.
The JNK-ERK-p38 MAPK signaling pathways also play an essential role in the control of bone formation and bone homeostasis during osteoblast differentiation [55]. Specifically, JNK is important for late-stage osteoblast differentiation, as inactivation of JNK by a specific small molecule inhibitor causes a lack of mineralization in MC3T3-E1 cells [56], while deletion of *JNK1* and *JNK2* in mice (*Jnk1-2osx*) results in severe osteopenia arising from bone loss [57]. Phosphorylation and consequent activation of Runx2 in osteoblasts is dependent on the expression of both ERK1 and ERK2 [58], while inactivation of ERK1 and ERK2 reduces β-catenin expression [59]. In addition, p38 is required for early-stage of osteoblast differentiation by promoting osteogenic differentiation markers, including ALP, osteocalcin, and mineralization [60].

In summary, this study demonstrated for the first time that the short-chain-length, lower-molecular-weight, higher-water-solubility *Pleurotus sajor-caju* glucanoligosaccharide (*Ps*-GOS) promotes osteoblast differentiation and mineralization, which are accompanied by the increase of ALP, COL1, and OCN expression through the induction of BMP-2 and Runx2. Furthermore, we have found that *Ps*-GOS also promotes bone formation via Wnt/β-catenin and MAPK signaling pathways. Thus, our findings show that *Ps*-GOS could be developed as a candidate supplement promoting bone formation. As such, this represents an example of both value creation as well as value addition in the biotechnological exploitation and use of natural rubber latex serum. Future work will be directed at determining the precise mechanism of action of *Ps*-GOS for comparison with other known modulators of osteoblastogenesis. Given the important role of the RANKL/RANL/OPG system in regulating osteoclast function, it will be interesting to establish whether *Ps*-GOS can impact this system also.

5. Conclusions

In the present study, we report for the first time that small size *Pleurotus sajor-caju* glucanoligosaccharide (*Ps*-GOS) has a stimulatory effect, promoting bone formation by stimulating osteoblast proliferation, differentiation, and mineralization possibility through BMP-2, Wnt/β-catenin, and MAPK signaling pathways. *Ps*-GOS has potential as a useful therapeutic drug to prevent osteoporosis.

**Author Contributions:** T.P. and T.Y. conceived and designed the experiments with additional help from C.S., R.W. and U.K.-L. T.Y. performed the experiments. T.P., P.S., and T.Y. drafted the original manuscript. U.K.-L., A.A., R.W., and C.S. modified the manuscript. All authors have read and agreed to the published version of the manuscript.

**Funding:** This research was funded by grants from Prince of Songkla University, contract no. SCI6202084S and through the government budget, Scholarship Awards for Thai Ph.D. Students under Thailand’s Education Hub for the Southern Region of ASEAN Countries (Grant No. PHD/2557 to Thanintorn Yodthong) and Scholarhip for an Overseas Thesis Research Study (to Thanintorn Yodthong), the Graduate School and the Academic Strengthening Program in Biochemistry, Faculty of Science. It was also funded by the Thailand Centre of Excellence for Life Sciences (Public Organization).

**Conflicts of Interest:** The authors declare no conflict of interest.

**References**

1. Sozen, T.; Ozisik, L.; Calik Basaran, N. An overview and management of osteoporosis. *Eur. J. Rheumatol.* 2016, 4, 46–56. [CrossRef] [PubMed]
2. Wright, N.C.; Looker, A.C.; Saag, K.G.; Curtis, J.R.; Delzell, E.S.; Randall, S.; Dawson-Hughes, B. The recent prevalence of osteoporosis and low bone mass in the United States based on bone mineral density at the femoral neck or lumbar spine. *J. Bone Miner. Res.* 2014, 29, 2520–2526. [CrossRef] [PubMed]
3. Ji, M.; Yu, Q. Primary osteoporosis in postmenopausal women. *Chronic Dis. Transl. Med.* 2015, 1, 9–13. [CrossRef] [PubMed]
4. Weitzmann, M.N.; Pacifi, R. Estrogen deficiency and bone loss: An inflammatory. *J. Clin. Investig.* 2006, 116, 1186–1194. [CrossRef]
5. Riggs, B.L. The mechanisms of estrogen regulation of bone resorption. *J. Clin. Investig.* 2000, 106, 1203–1204. [CrossRef]
6. Rucci, N. Molecular biology of bone remodeling. *Clin. Cases Miner. Bone Metab.* 2008, 5, 49–56.
7. Feng, X.; McDonald, J.M. Disorders of bone remodeling. *Annu. Rev. Pathol.* 2011, 6, 121–145. [CrossRef]

8. Zuo, C.; Huang, Y.; Bajis, R.; Sahih, M.; Li, Y.P.; Dai, K.; Zhang, X. Osteoblastogenesis regulation signals in bone remodeling. *Osteoporos. Int.* 2012, 23, 1653–1663. [CrossRef]

9. Yamaguchi, A.; Sakamoto, K.; Minamizato, T.; Katsube, K.; Nakanishi, S. Regulation of osteoblast differentiation mediated by BMP, Notch, and CCN3/NOV. *Jpn. Dent. Sci. Rev.* 2008, 44, 48–56. [CrossRef]

10. Komori, T. Regulation of bone development and maintenance by Runx2. *Front. Biosci.* 2008, 1, 898–903. [CrossRef]

11. Hashimi, S.M. Exogenous noggin binds the BMP-2 receptor and induces alkaline phosphatase activity in osteoblasts. *J. Cell Biochem.* 2019, 120, 13237–13242. [CrossRef] [PubMed]

12. Chau, J.F.; Leong, W.F.; Li, B. Signaling pathways governing osteoblast proliferation, differentiation and function. *Histol. Histopathol.* 2009, 24, 1593–1606. [PubMed]

13. Higuchi, C.; Myoui, A.; Hashimoto, N.; Kuriyama, K.; Yoshioka, K.; Yoshikawa, H.; Itoh, K. Continuous inhibition of MAPK signaling promotes the early osteoblastic differentiation and mineralization of the extracellular matrix. *J. Bone Miner. Res.* 2002, 17, 1785–1794. [CrossRef] [PubMed]

14. Duan, P.; Bonewald, L.F. The role of the Wnt/β-Catenin signaling pathway in formation and maintenance of bone and teeth. *Int. J. Cell Biol.* 2016, 77, 23–29. [CrossRef] [PubMed]

15. Krishnan, V.; Bryant, H.U.; Macdougald, O.A. Regulation of bone mass by Wnt Signaling. *J. Clin. Investig.* 2006, 116, 1202–1209. [CrossRef]

16. Chen, J.; Long, F. β-catenin promotes bone formation and suppresses bone resorption in postnatal growing mice. *J. Bone Miner. Res.* 2013, 28, 1160–1169. [CrossRef]

17. Clemons, M.; Goss, P. Estrogen and the risk of breast cancer. *N. Engl. J. Med.* 2001, 344, 276–285. [CrossRef]

18. Lacey, J.V.; Brinton, L.A.; Leitzmann, M.F.; Mouw, T.; Hollenbeck, A.; Schatzkin, A.; Hartge, P. Menopausal hormone therapy and ovarian cancer risk in the national institutes of Health-AARP Diet and Health Study Cohort. *J. Natl. Cancer Inst.* 2006, 98, 1397–1405. [CrossRef]

19. George, E.L.; Lin, Y.L.; Saunders, M.M. Bisphosphonate-related osteonecrosis of the jaw: A mechanobiology perspective. *Bone Rep.* 2018, 8, 104–109. [CrossRef]

20. An, J.; Yang, H.; Zhang, Q.; Liu, C.; Zhao, J.; Zhang, L.; Chen, B. Natural products for treatment of osteoporosis: The effects and mechanisms on promoting osteoblast-mediated bone formation. *Life Sci.* 2016, 15, 46–58. [CrossRef]

21. Bashir, K.M.I.; Choi, J.S. Clinical and physiological perspectives of β-glucans: The past, present, and future. *Int. J. Mol. Sci.* 2017, 18, 1906. [CrossRef] [PubMed]

22. Kawata, K.; Iwai, A.; Muramatsu, D.; Aoki, S.; Uchiyama, H.; Okab, M.; Hayakawa, S.; Takaoka, A.; Miyazaki, T. Stimulation of macrophages with the β-glucan produced by *Aureobasidium pullulans* promotes the secretion of tumor necrosis factor-related apoptosis inducing ligand (TRAIL). *PLoS ONE* 2015, 13, e0124809. [CrossRef] [PubMed]

23. Tiwari, U.; Cummins, E. Meta-analysis of the effect of β-glucan intake on blood cholesterol and glucose levels. *Nutrition* 2011, 27, 1008–1016. [CrossRef] [PubMed]

24. Kanagasabapathy, G.; Kuppusamy, U.R.; Nurestri, S.; Malek, A.; Abdulla, M.A.; Chua, K.; Sabaratnam, V. Glucan-rich polysaccharides from *Pleurotus Sajor-Caju* (Fr.) Singer prevents glucose intolerance, insulin resistance and inflammation in C57BL/6j mice fed a high-fat diet. *BMC Complement. Altern. Med.* 2012, 21. [CrossRef]

25. Jang, J.H.; Lee, J.; Kim, J.H.; Lee, Y.H.; Ju, Y.C.; Lee, J.S. Isolation and identification of RANKL-induced osteoclast differentiation inhibitor from *Pleurotus citrinopileatus*. *Mycoscience* 2013, 54, 265–270. [CrossRef]

26. Lee, D.H.; Han, D.W.; Park, B.J.; Baek, H.S.; Takatori, K.; Aihara, M.; Tsubaki, K.; Park, J. The influences of β-glucan associated with BMP7 on MC3T3-E1 proliferation and osteogenic differentiation. *Key Eng. Mater.* 2005, 288–289, 241–244. [CrossRef]

27. Shin, H.D.; Yang, K.J.; Park, B.R.; Son, C.W.; Jang, H.J.; Ku, S.K. Antioestrogenic effect of polycyan, β-glucan from *Aureobasidium*, in ovariectomized osteoporotic mice. *Nutrition* 2007, 23, 853–860. [CrossRef]

28. Breene, W.M. Nutritional value of specialty mushrooms. *J. Food Prot.* 1990, 53, 883–894. [CrossRef]

29. Finimundy, T.C.; Barros, L.; Calhelha, R.C.; Alves, M.J.; Prieto, M.A.; Abreu, R.M.V.; Dillon, A.J.P.; Henriques, J.A.P.; Roesch-Ely, M.; Ferreira, I.C.F.R. Multifunctions of *Pleurotus sajor-caju* (Fr.) Singer: A highly nutritious food and a source for bioactive compounds. *Food Chem.* 2018, 245, 150–158. [CrossRef]
30. Regand, A.; Tosh, S.M.; Wolever, T.M.S.; Wood, P.J. Physicochemical properties of glucan in differently processed oat foods influence glycemic response. J. Agric. Food Chem. 2009, 57, 8831–8838. [CrossRef]
31. Wang, Q.; Ellis, P.R. Oat β-glucan: Physico-chemical characteristics in relation to its blood-glucose and cholesterol-lowering properties. Br. J. Nutr. 2014, 112, S4–S13. [CrossRef] [PubMed]
32. Li, K.; Chen, W.; Wang, W.; Tan, H.; Li, S.; Yin, H. Effective degradation of curdlan powder by a novel endo-β-1→3-glucanase. Carbohydr. Polym. 2018, 201, 122–130. [CrossRef] [PubMed]
33. Chungho, N.; Suntaro, A.; Wittisuwannakul, R. β-1,3-glucanase isozymes from the latex of Hevea brasiliensis. Phytochemistry 1995, 39, 505–509. [CrossRef]
34. Freimund, S.; Sauter, M.; Kappeli, O.; Butler, H.A. New non-Degrading Isolation Process for 1,3-β-D-glucan of high purity from baker’s yeastSaccharomyces cerevisiae. Carbohydr. Polym. 2003, 54, 159–171. [CrossRef]
35. Synytsya, A.; Mic’ková, K.; Synytsya, A.; Jablonsky’, I.; Spěváček, J.; Erban, V.; Kováříková, E.; Čopíková, J. Glucans from fruit bodies of cultivated mushrooms Pleurotus ostreatus and Pleurotus eryngii: Structure and potential prebiotic activity. Carbohydr. Polym. 2009, 76, 548–556. [CrossRef]
36. McClean, B.; Glennie-Holmes, M. Enzymic quantification of (1→3) (1→4)-β-D-glucan in barley and malt. J. Inst. Brew. 1985, 91, 285–295. [CrossRef]
37. Yodit ung, T.; Kedjarune-Leggat, U.; Smythe, C.; Wittisuwannakul, R.; Pitakpornpreecha, T. L-quebrachitol promotes the proliferation, differentiation, and mineralization of MC3T3-E1 cells: Involvement of the BMP-2/Runx2/SPARC signaling pathway. Molecules 2018, 23, 3086. [CrossRef]
38. Javed, A.; Chen, H.; Ghor i, F.Y. Genetic and transcriptional control of bone formation. Oral Maxillofac. Surg. Clin. N. Am. 2010, 22, 283–293. [CrossRef]
39. Prins, H.J.; Braat, A.K.; Gawil lita, D.; Dhert, W.J.A.; Egan, D.A.; Tijssen-Slump, E.; Yuan, H.; Coffer, P.J.; Rozemuller, H.; Martens, A.C. In vitro induction of alkaline phosphatase levels predicts in vivo bone forming capacity of human bone marrow stromal cells. Stem Cell Res. 2014, 12, 428–440. [CrossRef]
40. Yazid, M.D.; A riff in, S.H.Z.; Senafi, S.; Razak, M.A.; Wahab, R.M.A. Determination of the differentiation capacities of murine’s primary mononucleated cells. Cancer Cell Int. 2010, 28. [CrossRef]
41. Golub, E.E.; Boesze-battaglia, K. The role of alkaline phosphatase in mineralization. Curr. Opin. Orthop. 2007, 18, 444–448. [CrossRef]
42. Li, L.J.; Kim, S.N.; Cho, S.A. Comparison of alkaline phosphatase activity of MC3T3-E1 cells cultured on different Ti surfaces: Modified sandblasted with large grit and acid-etched (MSLA), laser-treated, and laser and acid-treated Ti Surfaces. J. Adv. Prosthodont. 2016, 8, 235–240. [CrossRef] [PubMed]
43. Beck, G.R.; Sullivan, E.C.; Moran, E.; Zer ler, B. Relationship between alkaline phosphatase levels, osteopontin expression, and mineralization in differentiating MC3T3-E1 osteoblasts. J. Cell. Biochem. 1998, 728, 269–280. [CrossRef]
44. Sun, L.Y.; Hsieh, D.K.; Lin, P.C.; Chiu, H.T.; Chiou, T.W. Pulsed Electromagnetic Fields Accelerate Proliferation and Osteogenic Gene Expression in Human Bone Marrow Mesenchymal Stem Cells During Osteogenic Differentiation. Bioelectromagnetics 2010, 31, 209–219. [CrossRef]
45. Hu, L.; Yin, C.; Zhao, F.; Ali, A.; Ma, J.; Qian, A. Mesenchymal stem cells: Cell fate decision to osteoblast or adipocyte and application in osteoporosis treatment. Int. J. Mol. Sci. 2018, 19, 360. [CrossRef]
46. Chen, G.; Deng, C.; Li, Y.P. TGF-β and BMP signaling in osteoblast differentiation and bone formation. Int. J. Biol. Sci. 2012, 8, 272–288. [CrossRef]
47. Javed, A.; Bae, J.S.; Afza, F.; Gutierrez, S.; Prapat, J.; Zaidi, S.K.; Lou, Y.; Van Wijnen, A.J.; Stein, J.L.; Stein, G.S.; et al. Structural coupling of Smad and Runx2 for execution of the BMP2 osteogenic signal. J. Biol. Chem. 2008, 283, 8412–8422. [CrossRef]
48. Xuan, D.; Li, S.; Zhang, X.; Hu, F.; Lin, L.; Wang, C.; Zhang, J. Mutations in the RUNX2 gene in Chinese patients with cleidocranial dysplasia. Ann. Clin. Lab. Sci. 2008, 38, 15–24.
49. Adhami, M.D.; Rashid, H.; Chen, H.; Clarke, J.C.; Yang, Y.; Javed, A. Loss of Runx2 in committed osteoblasts impairs postnatal skeletogenesis. J. Bone Min. Res. 2015, 30, 71–82. [CrossRef]
50. Zhang, R.; Oyajobi, B.O.; Harris, S.E.; Chen, D.; Tsao, C.; Deng, H.W.; Zhao, M. Wnt/β-catenin signaling activates bone morphogenetic protein 2 expression in osteoblasts. Bone 2013, 52, 145–156. [CrossRef]
51. MacDonald, B.T.; Tamai, K.; He, X. Wnt/β-Catenin signaling: Components, mechanisms, and diseases. Dev. Cell. 2009, 17, 9–26. [CrossRef] [PubMed]
52. Gong, Y.; Slee, R.B.; Fukai, N.; Rawadi, G.; Roman-roman, S.; Regnato, A.M.; Wang, H.; Cundy, T.; Glorieux, F.H.; Lev, D.; et al. LDL receptor-related protein 5 (LRP5) affects bone accrual and eye development. *Cell* **2001**, *107*, 513–523. [CrossRef]

53. Kato, M.; Patel, M.S.; Levasseur, R.; Lobov, I.; Chang, B.H.; Li, D.A.G.; Hartmann, C.; Li, L.; Hwang, T.; Brayton, C.F.; et al. Cbfa1-independent decrease in osteoblast proliferation, osteopenia, and persistent embryonic eye vascularization in mice deficient in Lrp5, a Wnt Coreceptor. *J. Cell Biol.* **2002**, *15*, 303–314. [CrossRef] [PubMed]

54. Okamoto, M.; Udagawa, N.; Uehara, S.; Maeda, K.; Yamashita, T.; Nakamichi, Y.; Kato, H.; Saito, N.; Minami, Y.; Takahashi, N.; et al. Noncanonical Wnt5a enhances Wnt/β-Catenin signaling during osteoblastogenesis. *Sci. Rep.* **2014**, *4*, 1–8. [CrossRef] [PubMed]

55. Rodríguez-carballo, E.; Gámez, B.; Ventura, F. P38 MAPK Signaling in osteoblast differentiation. *Front. Cell Dev. Biol.* **2016**, *4*, 1–20. [CrossRef] [PubMed]

56. Matsuguchi, T.; Chiba, N.; Bandow, K.; Kakimoto, K.; Masuda, A.; Ohnishi, T. JNK activity is essential for Atf4 expression and late-Stage osteoblast differentiation. *J. Bone Miner. Res.* **2009**, *24*, 398–410. [CrossRef]

57. Xu, R.; Zhang, C.; Shin, D.Y.; Kim, J.M.; Lalani, S.; Li, N.; Yang, Y.S.; Liu, Y.; Eiseman, M.; Davis, R.J.; et al. C-Jun N-terminal kinases (JNKs) are critical mediators of osteoblast activity in vivo. *J. Bone Miner. Res.* **2017**, *32*, 1811–1815. [CrossRef]

58. Xiao, G.; Jiang, D.; Thomas, P.; Benson, M.D.; Guan, K.; Karsenty, G.; Franceschi, R.T. MAPK pathways activate and phosphorylate the osteoblast-specific transcription factor, Cbfa1. *J. Biol. Chem.* **2000**, *275*, 4453–4459. [CrossRef]

59. Matsushita, T.; Chan, Y.Y.; Kawanami, A.; Balmes, G.; Landreth, G.E.; Murakami, S. Extracellular signal-regulated kinase 1 (ERK1) and ERK2 play essential roles in osteoblast differentiation and in supporting osteoclastogenesis. *Mol. Cell. Biol.* **2009**, *29*, 5843–5857. [CrossRef]

60. Hu, Y.; Chan, E.; Wang, S.X.; Li, B. Activation of P38 mitogen-activated protein kinase is required for osteoblast differentiation. *Endocrinology* **2003**, *144*, 2068–2074. [CrossRef]