Delivery of dexamethasone from bioactive nanofiber matrices stimulates odontogenesis of human dental pulp cells through integrin/BMP/mTOR signaling pathways

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Abstract: Therapeutically relevant design of scaffolds is of special importance in the repair and regeneration of tissues including dentin and pulp. Here we exploit nanofiber matrices that incorporate bioactive glass nanoparticles (BGNs) and deliver the odontogenic drug dexamethasone (DEX) to stimulate the odontogenic differentiation of human dental pulp cells (HDPCs). DEX molecules were first loaded onto the BGN, and then the DEX-BGN complex was incorporated within the biopolymer nanofiber matrix through electrospinning. The release of DEX continued over a month, showing a slow releasing profile. HDPCs cultured on the DEX-releasing BGN matrices were viable, proliferating well up to 14 days. The odontogenic differentiation, as assessed by alkaline phosphatase activity, mRNA expression of genes, and mineralization, was significantly stimulated on the matrices incorporating BGN and further on those releasing DEX. The DEX-releasing BGN matrices highly upregulated the expression of the integrin subsets α1, α5, and β3 as well as integrin downstream signaling molecules, including focal adhesion kinase (FAK), Paxillin, and RhoA, and activated bone morphogenetic protein mRNA and phosphorylation of Smad1/5/8. Furthermore, the DEX-releasing BGN-matrices stimulated Akt and mammalian target of rapamycin (mTOR), which was proven by the inhibition study. Collectively, the designed therapeutic nanofiber matrices that incorporate BGN and deliver DEX were demonstrated to promote odontogenesis of HDPCs, and the integrins, bone morphogenetic protein, and mTOR signaling pathways are proposed to be the possible molecular mechanisms. While further in vivo studies are still needed, the DEX-releasing bioactive scaffolds are considered as a potential therapeutic nanomatrix for regenerative endodontics and tissue engineering.

Keywords: drug delivery, therapeutic matrix, dexamethasone, bioactive glass, human dental pulp cells, odontogenic differentiation

Introduction

Biomatrices that have the therapeutic actions can lead to excellent outcome in the repair and regeneration of tissues, including dentin–pulp complexes.1 The controlled and sustained delivery of therapeutic molecules to the target site of injuries and diseases possibly reduces inflammation, enhances angiogenesis, and accelerates odontogenic differentiation of dental pulp stem cells.2,3 Therefore, the design of matrices with therapeutically relevant actions is considered of special importance in improving the repair and regenerative potential of dentin and pulp tissues.

Over the last decades, many different forms of biomatrices have been developed in this area, including porous foams, macrochanneled scaffolds, and nanofibrous meshes.
The latter, among others, have been attractive mainly due to their mimicking the structural feature of the native extracellular matrix at the nanoscale.6-9 Cells recognize the nanofibrous topology, adhere and spread well, and adopt their phenotypes properly depending on the surrounding biochemical cues.5,10 Our previous studies have shown that human dental pulp cells (HDPCs) actively grew on the nanofiber matrix made of biopolymers and differentiated actively into odontoblastic lineage.11 Furthermore, when the nanofiber compositions were tailored to incorporate bioactive inorganic nanoparticles, such as those of calcium phosphates and bioglass, the odontogenic differentiation capacity could be substantially improved.12

In this study, we aimed to utilize the nanofiber matrices and design them with therapeutic capacity through delivering drug molecules relevant for activating odontogenesis of dental stem cells. For this, we incorporated a bioactive nanoparticle composition, namely bioactive glass nanoparticles (BGNs), within biopolymer nanofiber matrices. In fact, BGN, as the bioactive inorganic nanocomponent, exerts excellent bone bioactivity, regulating the genetic profiles of osteoprogenitor/stem cells toward a lineage of bone-forming cells.7,8 Furthermore, as the therapeutic drug, we incorporated dexamethasone (DEX) within the nanofiber matrix.

In fact, DEX has been used as the bioactive molecule for bone tissue. Studies have shown that DEX treatment stimulates mesenchymal stem cell (MSC) proliferation and supported osteogenic lineage differentiation; however, the direct use of DEX has been limited mainly due to its toxic side effects. For this reason, the use of DEX through incorporation into biomaterials is favored. When added to electrosprun polycaprolactone (PCL) nanofibers, DEX was shown to increase the level of alkaline phosphatase (ALP) and mineralized matrix in human bone marrow MSCs.14 Recently, nanofiber scaffolds were also designed to deliver DEX in concert with the bone morphogenetic protein 2 (BMP-2), which was shown to improve differentiation and bone repair.15 Moreover, the proliferation and osteogenic differentiation of stem cells derived from periodontal ligament were stimulated by the long-term delivery of DEX that was incorporated within composite nanofibers.16

Based on these series of studies, we proposed to utilize the DEX delivery system made of nano-biomatrices that have nanofibrous morphology and bioactive nanocomponent within the composition, targeting odontogenic differentiation of dental stem cells. We cultured dental stem cells derived from human pulp (HDPCs) upon the DEX-delivering nano-biomatrices and investigated the odontogenesis of cells; moreover, we addressed the biological signaling pathways behind these events.

Materials and methods
Nano-biomatrices and DEX loading
All experiments were approved by the ethics committee of Kyung Hee University (Seoul, Republic of Korea). First, the nanocomponent (BGN) added within the nanofiber matrix was prepared by the sol–gel approach.16,17 In brief, 5 g polyethylene glycol (PEG) was dissolved in 120 mL methanol with pH adjusted to 12.5, and then 0.299 g Ca(NO3)2⋅4H2O was added and dissolved. Separately, 0.793 g TEOS (tetraethyl orthosilicate) was added to 30 mL methanol, which was then added dropwise to the pH-adjusted solution with ultrasonication. A white precipitate was obtained with 24 hours of stirring, which was then separated and washed with a water/ethanol mixture thoroughly and then dried overnight. To remove the PEG template, the powder was calcined at 600°C for 5 hours to obtain BGNs. The BGN powder was then amine-functionalized with 3-aminopropyl triethoxysilane at 80°C for 12 hours.

For the loading of DEX (C16H28FNa2O5P, a phospate form in diosodium salt), the BGN was dispersed at 10 mg/mL in distilled water containing DEX at different concentrations (5 mg/mL, 10 mg/mL, 15 mg/mL, 20 mg/mL, 25 mg/mL, and 30 mg/mL). The amounts of DEX loaded were determined by the absorbance at 242 nm using Ultraviolet–visible (UV–vis) spectrometry (Libra S22, Biochrom). The release of DEX from the BGN was also recorded by adding 10 mg of DEX-loaded BGN in 50 mL distilled water at 37°C for different time periods. At predetermined time points, 1 mL of test sample was withdrawn and the release of DEX was monitored by the UV–vis measurement.

For the generation of nanofiber matrices made of DEX-loaded BGN and biopolymers, the electrospinning technique was used. The polymer solution was composed of 10 wt% gelatin/10 wt% PCL dissolved in TFE (trifluoroethanol) at 40°C, to which the BGN (or BGN-DEX) was added at 2.5 wt% with respect to the polymer. The composite solutions were then electrospun at an injection rate of 1 mL/h, voltage 10.5 V, and distance 14.5 cm to produce nanofibrous non-woven meshes. The prepared samples were examined for the nanostructure morphologies by scanning electron microscopy (SEM, JSM-6510; JEOL, Tokyo, Japan) and transmission electron microscopy (TEM, JEM-3010; JEOL).

The degradation of the composite nanofiber was examined by immersing the nanofiber sample in 10 mL phosphate-buffered saline (pH = 7.4) at 37°C. After immersion for different time periods (7 days, 14 days, 21 days, and 28 days), the sample was taken out, dried, and the weight change was recorded. The release of DEX from the
biomatries was investigated by high-performance liquid chromatography (ACME 9000 system; Younglin Instrument, Ltd., Republic of Korea). Fifty milligrams of the nanofiber mesh was immersed in 10 mL distilled water and incubated at 37°C. At predetermined time points, 1 mL of the release medium was sampled for the DEX analysis and the medium was refreshed.

**Cell culture and proliferation**

Immortalized HDPCs, transfected with human telomerase catalytic component, were kindly provided by Professor Takashi Takata (Hiroshima University, Japan). Cells were cultured in α-minimum essential medium supplemented with 10% fetal bovine serum, 100 U/mL penicillin, and 100 mg/mL streptomycin in a humidified atmosphere of 5% CO2 at 37°C. In order to induce odontogenesis and to study the effects of DEX release from the nanofiber matrices, the cell culture medium was composed of either without DEX (negative control) or with DEX (positive control), ie, α-minimum essential medium supplemented with 10% fetal bovine serum, 50 mg/mL ascorbic acid, and 10 mM glycerophosphate, with or without 0.1 μM DEX. The culture medium was replaced every 2 days during the incubation period.

During culture for up to 14 days, the cell viability was assayed by means of a 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTS) assay kit (Cell Titer 96 Aq Non-Radioactive Cell Assay; Promega Corporation, Fitchburg, WI, USA). Briefly, MTS was added to the wells and incubated for 3 hours. The blank used was the culture medium with MTS. The absorbance was measured at 492 nm in an enzyme-linked immunosorbent assay reader.

**Odontogenic stimulation by dexametasonerleasing nanofibers**

For the mineralization assay, the Alizarin Red S (ARS) staining method was used. For this, the treated cells were washed thoroughly with distilled water and then fixed in 10% formalin for 30 minutes at 4°C. Cells were incubated in 40 mM ARS regent (Sigma-Aldrich Co., St Louis, MO, USA) for 20 minutes at room temperature followed by complete washes with distilled water, and then observed under a light microscope (Olympus Corporation, Tokyo, Japan).

**Mechanistic study on signaling pathways**

The signaling mechanisms behind the odontogenic differentiation were investigated in terms of integrins/downstream signals, MAPK pathways, BMP signaling, and Akt pathways. For integrin signaling, the mRNA levels of some integrin subsets representative for HDPCs (α5, α1, β1, and β3) were first analyzed. Furthermore, the integrin downstream signals were identified by the expressions of the phosphorylated form of intracellular adhesive molecules, including focal adhesion kinase (FAK), Paxillin, and RhoA, by means of Western blotting analysis. For BMP signaling, the mRNA levels of BMP-2 and BMP receptors (BMPRa-1, BMPRa-2, and BMPR2) were examined, and the protein expression of p-Smad1/5/8 was also analyzed. In particular, the p-Smad1/5/8 expression was examined by means of the Noggin blocking test. For the Akt/mammalian target of rapamycin (mTOR) signaling study, the expressions of p-Akt, p-70S6K, and p-4EBP1 were analyzed by Western blot, and the blocking study using rapamycin (RAPA) was also carried out.

The conditions for RT-PCR analysis were similar to those used for the odontogenic study, and the primer sequences and PCR conditions were as detailed in Table 1. For Western blotting, the cell extracts were prepared by solubilizing cells in the protein lysis buffer (5 mM ethylenediaminetetraacetic acid [EDTA], 1 mM MgCl2, 50 mM Tris–HCl [pH 7.5], 0.5% Triton X-100, 2 mM phenylmethylsulfonyl fluoride, and 1 mM N-ethylmaleimide) for 30 minutes on ice. Fifty micrograms of the protein was separated by sodium dodecyl sulfate polyacrylamide electrophoresis and electro-transferred to polyvinylidene difluoride membranes. The membranes were treated with...
blocking solution for 1 hour and were incubated with 1:1,000 dilution primary antibody against phospho-focal adhesion kinase (p-FAK; Santa Cruz Biotechnology Inc., Dallas, TX, USA), p-Paxillin (Santa Cruz Biotechnology Inc.), actin (Santa Cruz Biotechnology Inc.), Rho-A (Cell Signaling Technology, Danvers, MA, USA), p-Smad1/5/8 (Cell Signaling Technology), p-p38 (Cell Signaling Technology), and p-Akt (Cell Signaling Technology), p-p-Smad1/5/8 (Cell Signaling Technology), p-p38 (Cell Signaling Technology), and p-Akt (Cell Signaling Technology). The blots were followed by incubation for 1 hour with the appropriate horseradish peroxidase-conjugated secondary antibody. The proteins were then visualized using an enhanced chemiluminescence system (Amersham, Piscataway, NJ, USA).

Statistical analysis
Data are presented as mean and SD. Statistical analysis was performed by one-way analysis of variance followed by multiple-comparison Tukey’s test using SPSS Version 22.0 (IBM Corporation, Armonk, NY, USA). Statistical significance was considered at P < 0.05.

Results and discussion
Nano-biomatrices incorporate bioactive nanoparticles and release DEX drug sustainably
The nano-morphology of the BGN used for the delivery of DEX and also for a nano-additive to polymer matrices was examined by TEM. Spherical nanoparticles with a uniform size of 63 nm ± 12.3 were developed well, and a closer examination revealed the mesopores to be evenly distributed in the nanoparticles (Figure 1A). After the loading with DEX, the TEM image of samples was also revealed (Figure 1B). The mesoporous image appeared diffused due to the pore-filling of DEX molecules. The loading amount of DEX in BGN samples was then monitored when 10 mg/mL of BGN was used while DEX concentration varied up to 40 mg/mL (Figure 1C). DEX loading amount increased gradually with increasing DEX concentration used initially, and then saturated to 630 μg of DEX which gave as high as 63% of loading.

### Table 1

Reverse transcription polymerase chain reaction (RT-PCR) primers and conditions

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<tr>
<th>Genes</th>
<th>Primer sequence (5'-3')</th>
<th>Annealing temperature (°C)</th>
<th>Cycle number</th>
<th>Product size (bp)</th>
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<td>ALP</td>
<td>F: 5'-ACGTGCTGAAGGATGTCATC-3'</td>
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<td>30</td>
<td>476</td>
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<td>R: 5'-CTGTTAAGGCCGATTCACT-3'</td>
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<td>DMP-1</td>
<td>F: 5'-AGAGAAGAGATGGGAGACTGGC-3'</td>
<td>61</td>
<td>35</td>
<td>202</td>
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<td>R: 5'-TGAGCTTCACATGCTGGTGTT-3'</td>
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<tr>
<td>DSPP</td>
<td>F: AGAAGACCTGCGCAGAAAGAAT-3'</td>
<td>60</td>
<td>35</td>
<td>280</td>
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<td>R: TCTGGTGACATGGTCGTT-3'</td>
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<td>Integrin α1</td>
<td>F: 5'-TGTCGCTGGGCTCTGGTCGTTT-3'</td>
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<td>Integrin β1</td>
<td>F: 5'-GAGCCCGCGCGAGAAAAGT-3'</td>
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<td>GAPDH</td>
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<td>306</td>
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<td>R: 5'-AGCCTTCTACAGTGTTGGAAGA-3'</td>
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Abbreviations: ALP, alkaline phosphatase; DMP-1, dentin matrix protein-1; DSPP, dentin sialophosphoprotein; BMPR, bone morphogenetic protein receptor; BMP, bone morphogenetic protein; GAPDH, glyceraldehyde 3-phosphate dehydrogenase.

The significance was considered at P < 0.05.
efficiency. The release of DEX from the BGN samples was then recorded (Figure 1D). The DEX release was highly abrupt, showing ~80% release within 24 hours followed by an almost complete release within 3 days.

Next, the DEX-loaded BGNs were incorporated within the PCL–gelatin nanofiber matrix to produce a nanofibrous mesh through electrospinning. 2.5 wt% of the DEX-BGN complex was loaded within the nanofiber, and the DEX loading amount used was below the saturation point (0.286 mg DEX per 1 mg BGN). The SEM images of the composite nanofibers showed the production of nonwoven mesh of fibrous morphology (Figure 1E). On the TEM image, the internal morphology of the nanofiber was revealed, where the nanospherical BGNs were dispersed well within the biopolymer nanofiber matrix (Figure 1F). The degradation of the composite nanofiber was then examined. The weight loss of the nanofiber during immersion in a saline solution was ~15% for the first 7 days and then ~23% for up to 28 days (Figure 1G).

We further investigated the release of DEX from the produced DEX-BGN-incorporated nanofiber matrices. The DEX amount released at each time point was recorded for up to 35 days (Figure 1H). Interestingly, the released DEX amount was almost similar up to the period tested, except the initial rapid release (10 hours). This initial burst release was mainly due to the DEX molecules present in the biopolymer matrix that might have been pre-released during the preparation of the composite solution and nanofibers. More importantly,
the continued release of DEX over a couple of weeks (and even more possibly) at almost constant dose is considered of great merit in long-term drug delivery systems. From a practical aspect, this DEX-releasing system would be highly beneficial for providing a constant-dosed odontogenic drug to cells during the long-term culture period, without the supplement of the drug within a culture medium; further, this sustained and controlled DEX-releasing system will be potentially useful for in vivo implantable scaffolds where the DEX drug released at a constant dose can favorably act in the surrounding cells for tissue (dentin–pulp) regeneration. This release profile of DEX might be the result of combined actions of the diffusion of DEX and the degradation of biopolymer matrices. If the DEX molecules were released from nondegradable matrix, the release pattern might follow a certain power-law pattern; however, the hydrolytic degradation of the biopolymer matrix might accelerate the DEX release, ultimately leading to an almost linear release pattern. After confirming the release of DEX, we next sought to investigate the effects of DEX-releasing nanofiber matrices on the behavior of HDPCs.

**DEX-delivering bioactive nano-biomatrices stimulate odontogenesis of HDPCs**

Before addressing the odontogenic differentiation of cells, we examined the viability of cells upon the DEX-releasing nanofiber matrices. The culture medium without DEX (--), or with DEX(+) was used as negative and positive control, respectively; moreover, nanofiber matrices without BGN (0BGN), or with BGN only (BGN), or DEX-loaded BGN (DEX-BGN) were used as experimental groups involving scaffolds. The cell viability was measured at day 7 and day 14 of culture, using an MTT assay, as presented in Figure 2. Cells were highly proliferative on day 7 on the nanofiber matrices, with the highest level on 0BGN and the lowest level on DEX-BGN; however, both were still higher than that in DEX(--) control. At day 14, the cell viability increased substantially on the 0BGN matrix; but on other matrices the cell proliferation was not as much, with the DEX(+) control showing the lowest level. While the cell viability on the nano-biomatrices were up- and downregulated with time, this was closely related to the time-dependent cell differentiation as well, making the phenomenon quite complicated.

We next investigated the odontogenic differentiation behaviors of HDPCs. First, the mRNA expressions of odontoblastic marker genes (*ALP*, *DSPP*, and *DMP-1*) were assessed by RT-PCR (Figure 3A). In particular, the *DSPP* and *DMP-1* genes are known to be identifying markers of odontoblasts, and play crucial roles in the early odontoblast differentiation and late dentin mineralization of dental pulp cells. From the RT-PCR bands, the DEX-BGN group, compared to others, showed higher expressions of genes, and particularly at day 14 for *DSPP* gene. Further, we analyzed the transcription factors Runx2 and Osterix, which are the major genes highly expressed during osteogenic/osteogenic induction in various cell types. There were substantially higher expressions of the genes (particularly Osterix) when cultured on the DEX-BGN matrix (Figure 3B).

We further analyzed the odontogenic differentiation of HDPCs by the ALP (Figure 4A). The ALP level increased with the culture time, implying the time-dependent increase of ALP during odontogenic differentiation, a characteristic phenomenon widely reported elsewhere. Compared to DEX(--) control, all the matrix groups showed significantly higher ALP levels. Moreover, the DEX-BGN group had significantly higher ALP level than BGN only, demonstrating the role of DEX released from the matrix. ARS staining was then performed to identify cellular mineralization. Cells cultured for 7 days and 14 days were treated with ARS dye, and the reaction products were gathered and visualized under optical microscopy (Figure 4B). Red stained images became much clearer with increasing time, and, furthermore, the matrix groups showed more stained images than DEX(--), while DEX(+) control exhibited the most. Compared to BGN only, the DEX-BGN group appeared to stain more.

Collectively, the series of assays on odontogenic differentiation of HDPCs clearly demonstrated the effective role of DEX incorporated within the nanofiber matrix. In particular, the DEX-BGN group showed a level almost comparable to that of DEX(+) control, ie, the cells cultured in situ DEX-releasing matrix, which contained DEX-free medium,
behaved similar to those cultured with DEX-maintaining medium in dish. Although this in vitro culture condition can signify the similar effectiveness of the two systems, the DEX-releasing matrix system has the unique merit for the in vivo implantations where additional DEX cannot be supplemented externally. Interestingly, all matrix groups (even BGN- and DEX-free) stimulated the odontogenesis of HDPCs when the results were compared with DEX(−) control; in other words, the nanofiber matrix culture itself had some effective role in odontogenic differentiation under the DEX-free culture medium. In fact, the nanotopological cues have often been shown to activate osteoblast activity of MSCs.26 For the case of HDPCs, nanotopological substrates including biopolymer nanofibers and calcium phosphate nano-grained cements have recently been demonstrated to have similar effects on odontogenic differentiation.27–29 Furthermore, our odontogenesis results showed that the BGN-incorporated nanomatrix was even better than BGN-free nanomatrix, signifying the bioactive nanocomponent role of BGN. Our previous studies have shown that the BGN added to biopolymeric matrices stimulates odontogenesis of HDPCs.11,12 While the results of this study clearly demonstrate the beneficial action of DEX molecules released from the nanofiber matrices in the stimulation of odontogenesis, possible side effects of the DEX drug might also be of concern because DEX, as a powerful glucocorticoid, has been shown to exert adverse effects when used at high doses and for long-term therapy. However, in this study, we designed a system that allows local delivery of small doses of DEX over months, which is considered to relieve the concern that has been raised primarily in the systemic delivery systems. In fact, many local delivery systems of DEX have been reported to stimulate cellular responses in vitro and/or to improve tissue healing in vivo with little side effects.30–33 Even so, future in vivo studies may be needed to confirm this aspect of the current delivery system.
Integrin, BMP, and Akt/mTOR signaling pathways are involved in the events
After confirming the stimulating role of the DEX-releasing bioactive nanofiber matrices in odontogenic differentiation, we next sought to address the signal transduction mechanisms involved in the events. First, we focused on integrin and its downstream pathway, because this has been considered as one of the most important pathways in the matrix-related cell differentiation behaviors.\textsuperscript{34,35} The mRNA expressions of the integrin subsets $\alpha_1$, $\alpha_5$, $\beta_1$, and $\beta_3$ were analyzed via RT-PCR (Figure 5A). The mRNA levels of integrins $\alpha_1$, $\alpha_5$, and $\beta_3$ were upregulated mainly by the BGN and DEX-BGN groups, whereas integrin $\beta_1$ appeared not to be affected. Integrins are the major adhesion receptors that mediate cell adhesion and attachment,\textsuperscript{34} and different combinations of integrin subsets are expressed depending on the underlying substrates and ligands. The integrin subsets activated in our study ($\alpha_2$, $\alpha_5$, $\beta_1$) are known to be associated with collagen binding, which are thus considered to be present at greater levels over the matrices used herein, particularly BGN-containing ones. Our previous studies on bioactive scaffolds incorporating BGN have also shown similar results on the integrin expressions.\textsuperscript{11,12}

Next, the integrin downstream signaling molecules, including FAK, Paxillin, and RhoA, were analyzed by Western blotting (Figure 5B). The expressions of phosphorylated-Paxillin and p-FAK and total RhoA were highly activated with the culture on the nanofiber matrices, and more substantial activation of proteins was revealed on BGN and DEX-BGN. Engagement of cell surface integrins is associated with rapid tyrosine phosphorylation of several intracellular adhesive molecules, such as FAK and Paxillin, which are two focal adhesion-associated proteins that function in transmitting signals downstream of integrins.\textsuperscript{35} Moreover, RhoA, the most extensively studied Rho family member, is known to be activated by the integrins.\textsuperscript{36} Therefore, the significant stimulation of those intracellular adhesive molecules by the BGN and DEX-BGN suggests that integrin and its outside-in signaling pathways are heavily involved in the matrices-based HDPC differentiation behaviors.

We next focused on the BMP signaling pathway. The mRNA expressions of BMP-2 and key BMP receptors (BMPR1a, BMPR1b, and BMPR2) were upregulated by the DEX-BGN group. Furthermore, analysis of the phosphorylation of Smad1/5/8 revealed substantial activation on the DEX-BGN (Figure 6A). To verify the matrix effects on the

![Image](Figure 5 Integrin signaling pathway analyses. Notes: (A) mRNA expressions of integrin subsets ($\alpha_1$, $\alpha_5$, $\beta_1$, and $\beta_3$), by RT-PCR. GAPDH was used as the housekeeping gene. (B) Protein expressions of intracellular adhesive molecules (p-Paxillin, p-FAK, and RhoA), by Western blot. $\beta$-actin was used as the control. Abbreviations: RT-PCR, reverse transcription polymerase chain reaction; p-FAK, phospho-focal adhesion kinase; BGN, bioactive glass nanoparticle; DEX, dexamethasone; GAPDH, glyceraldehyde 3-phosphate dehydrogenase.)

![Image](Figure 6 BMP-2 signaling pathway analyses. Notes: (A) mRNA levels of BMP receptors (R1a, R1b, and R2) and BMP-2, as well as the protein expression of p-Smad1/5/8. (B) Inhibition study to verify the signaling pathway, where a BMP inhibitor, Noggin (10 $\mu$g/mL) was used to treat the cells for 1 hour, and then cultured for 2 days before Western blot analysis. Data are representative of three independent experiments. Abbreviations: BMP, bone morphogenetic protein; BGN, bioactive glass nanoparticle; DEX, dexamethasone; GAPDH, glyceraldehyde 3-phosphate dehydrogenase; Noggin, noggin.)
BMP pathway, the BMP inhibition study was further carried out. For this, cells were pretreated with a culture medium with or without the BMP inhibitor Noggin. The highly activated phosphorylation of Smad1/5/8 by DEX-BGN was significantly inhibited by the treatment with Noggin (Figure 6B). The results clearly demonstrated that the BMP signaling pathway is critically involved in the behavior of HDPCs upon the DEX-delivering bioactive nanofiber matrices.

In fact, the BMP signaling pathway is known to play a vital role in osteogenic differentiation as well as in bone development and regeneration. More recently, it was reported that integrin overlapping with BMP-2 receptors plays an essential role in the BMP-2 induction of osteoblast differentiation. Moreover, the activation of surface integrin of MSCs, which is influenced by the extracellular matrix’s elasticity, in turn, can modulate BMP receptor internalization, thereby regulating the lineage specification of MSCs. Our results also demonstrated that the DEX-releasing nano-biomatrix cues activated the expressions of integrins in HDPCs, which could in turn possibly stimulate the BMP signal transduction mechanism.

We next focused on the mTOR pathway, as this is known to be one of the principal signaling cascades regulating osteoblastic/odontoblastic differentiation. The phosphorylated form of Akt and mTOR targeting p70S6 kinase (p70S6K) and eukaryotic initiation factor 4E-binding protein 1 (4E-BP1) were shown to be highly upregulated in the matrix groups, particularly in the DEX-BGN matrix (Figure 7A). mTOR, a major downstream target of PI3K/Akt, is known to regulate various cellular functions required for cell growth and proliferation by phosphorylating a series of substrates, including p70S6K and 4E-BP1. To clarify the effects in the Akt/mTOR signaling, we tested again the mTOR-inhibition experiment. The cells were pretreated with RAPA, a selective mTOR inhibitor. RAPA, a well-known inhibitor of mTOR, first forms a complex with a 12 kDa cytosolic protein designated FK-506-binding protein 12, and this complex binds mTOR and inhibits its function. After the RAPA treatment, there was significant attenuation of p-p70S6K and p-4E-BP1, which was more noticeable in the DEX-BGN matrix (Figure 7B).

It is known that BMP signaling is linked to the mTOR pathway. BMP expression has been shown to activate P70S6K, a downstream effector of mTOR, suggesting that BMP-induced osteogenesis is mediated by mTOR activation. Furthermore, integrin signaling is also closely related with the Akt/mTOR intracellular signaling.

Suppression of PI3K and mTOR signaling using inhibitors such as LY294002, wortmannin, and RAPA has been shown to promote osteogenesis in vitro. Therefore, the integrin and BMP stimulations, which are influenced by the DEX-releasing bioactive substrates, might trigger the Akt/mTOR transduction signaling pathway. Figure 8 is a schematic diagram showing the signaling pathways of integrins, BMP, and Akt/mTOR, which are possibly involved in the odontogenic stimulation of HDPCs, upon the culture with DEX-releasing bioactive nanofiber matrices.

**Conclusion**

Bioactive nanofiber matrices developed to release DEX and incorporate BGN were demonstrated to stimulate odontogenic differentiation of HDPCs. The activation of integrin, BMP/Smad, and Akt/mTOR signaling pathways are the possible key mechanisms involved in the events. While more in-depth studies on in vivo applicability are necessary, the developed nanomaterials can be a potential therapeutic platform for the regeneration of dentin–pulp complex tissues.

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

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