Porous Polycaprolactone Scaffold Engineered with Naringin Loaded Bovine Serum Albumin Nanoparticles for Bone Tissue Engineering

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A biodegradable three-dimensional scaffolds have gathered attention and are widely studied for bone tissue engineering applications. In the present study, porous polycaprolactone scaffold entrapped with naringin loaded bovine serum albumin nanoparticles (PS-N-BSANP) has been engineered. Further, the prepared nanoparticles and interconnected porous scaffolds were characterized by scanning electron microscopy, X-ray diffraction and fourier transform infrared spectroscopy analysis. X- ray diffraction showed amorphization of naringin in PS-N-BSANP. In addition, sustained naringin release profile was observed from PS-N-BSANP for 12 days which showed a cumulative release of 52.54 micromolar (µM). Furthermore, conditioned medium from PS-N-BSANP showed an increased calcium deposition and collagen matrix formation under osteogenic conditions with C3H10T1/2 cell line. These results suggest that PS-N-BSANP enhanced the osteogenic differentiation potential in bone tissue engineering applications due to the controlled release of naringin.

Keywords: Naringin, Polycaprolactone Porous Scaffold, Bovine Serum Albumin Nanoparticles, Bone Tissue Engineering, C3H10T1/2 cell line.

In recent years, three-dimensional biodegradable scaffolds has gathered more attention in various tissue engineering fields for reconstruction of damaged tissues. In general, cells require an three-dimensional matrix for cell adhesion, proliferation and controlled environment for tissue restoration. Hence, three-dimensional scaffolds are widely used as an extracellular matrix in tissue repair applications. Moreover, three-dimensional scaffolds have been synthesized from many different types of aliphatic polysters and natural polymers for therapeutic applications due to its excellent properties such as biodegradability, biocompatibility and mechanical strength. Among those polymers, polycaprolactone is Food and Drug Administration approved biodegradable material to construct the porous scaffolds for long-term implants.

Regeneration of bone fracture with a bone loss of 3cm is tedious process. Therefore, biodegradable porous scaffolds are extensively evaluated as an alternative for autologous and allogeneic approaches in bone tissue repair. In addition, controlled release of bioactive molecules...
from the porous scaffolds could increase the therapeutic potential in bone regeneration. Naringin is a flavanone glycoside that has shown enormous potential such as anti-inflammatory, antioxidant, anticancer, antidiabetic etc. Also, naringin enhances the proliferation and osteogenic differentiation of human bone marrow-derived mesenchymal stem cells.

We propose controlled release of naringin from a three-dimensional porous scaffold could increase its therapeutic potential in bone regeneration. In this study, three-dimensional porous polycaprolactone scaffold carrying bovine serum albumin nanoparticles loaded with naringin is fabricated and evaluated for its in vitro osteogenic potential using C3H10T1/2 cell line.

**MATERIALS AND METHODS**

Polycaprolactone (average Mn 80,000), Naringin (≥95% (HPLC)), β-glycerophosphate, Dexamethasone, Ascorbic acid, Direct red 80, Hematoxylin were purchased from Sigma-Aldrich (St. Louis, MO). Bovine serum albumin (BSA) fraction V (pH 6-7) purchased from Sisco Research Laboratories Pvt. Ltd., Mumbai, India. Dulbecco’s modified eagle medium-low glucose (DMEM-LG), Antibiotic-antimycotic solution, Dulbecco’s phosphate buffered saline (DPBS), Trypsin-ethylenediaminetetraacetic acid (Trypsin-EDTA) solution, 3-(4,5-Dimethyl-2-thiazolyl)-2,5-diphenyltetrazolium bromide (MTT), Picric acid, Silver nitrate were purchased from Himedia, Mumbai, India. Fetal bovine serum (FBS) was purchased from PAN Biotech, Germany. All other chemicals and solvents used were of analytical grade.

**Preparation of naringin loaded bovine serum albumin nanoparticles (N-BSANP)**

N-BSANP was synthesized by coacervation process as described earlier. In brief, naringin (2% w/w) was added to the 2% (w/v) of BSA in deionized water followed by one hour incubation at room temperature. After incubation, ethanol was added dropwise until turbidity under constant stirring. Further, coacervates were hardened with 8% glutaraldehyde, incubated for 2 hours under constant stirring. After incubation, nanoparticles were obtained by centrifugation at 18000g, 4°C for 30 minutes. The pellet was suspended in DPBS and washed thrice and freeze-dried. The entrapment efficiency was calculated from the amount of free naringin present in the supernatant using the formula

\[
\text{Entrapment Efficiency} = \left[\frac{T_{\text{Nar}} - F_{\text{Nar}}}{T_{\text{Nar}}}\right] \times 100\%
\]

Where, \(T_{\text{Nar}}\) is the total amount of naringin added and \(F_{\text{Nar}}\) is the free naringin in the supernatant.

**Preparation of porous polycaprolactone scaffold entrapped with naringin loaded bovine serum albumin nanoparticles (PS-N-BSANP)**

PS-N-BSANP was engineered by solvent casting and particulate leaching method. Polycaprolactone (10% w/v) was dissolved in equal volumes of dichloromethane and ethanol. N-BSANP (5% w/w) was added to the polycaprolactone solution and sonicated on ice to achieve the homogeneous mixture. To this, finely ground sucrose (1:5 ratio; polycaprolactone to sucrose) was added and stirred to make an uniform slurry, cast into glass molds and dried at room temperature. Further, sucrose was leached out with deionized water to form a porous structure followed by treatment with 0.5M NaOH for 4 hours to impart hydrophilicity. The scaffolds were neutralized by repeated washing with deionized water and 50% ethanol. Furthermore, scaffolds were stabilized using an absolute ethanol and washed with DPBS, UV sterilized for 30 minutes. Polycaprolactone scaffold prepared the same without N-BSANP was treated as control (PS).

**Characterization of prepared nanoparticles and porous scaffolds**

**Scanning electron microscopy (SEM)**

To determine the size of the prepared nanoparticles, N-BSANP was sputtered with gold and analyzed by SEM (TESCAN VEGA3 SBU) with 25kV applied voltage and 25000x magnifications. The average size of the nanoparticles were calculated manually by measuring 30 particles, selected randomly from SEM image using ImageJ software (ImageJ 1.51j8, National Institutes of Health, USA) and the results were expressed as mean±standard deviation. Similarly, the fabricated scaffolds were cut into small pieces and sputtered with gold, morphology of the scaffolds were observed by SEM with an applied voltage of 10kV and 200μM magnifications.

**X-ray diffraction (XRD)**

The prepared nanoparticles and porous scaffolds.
scaffolds were characterized by X-ray diffraction using the PANalytical X’Pert PRO (Almelo, The Netherlands) with a current of 30mA and a load of 40kV tension. The 2θ scan ranges from 10º–100º with a step size of 0.0167.

**Fourier transform infrared spectroscopy (FTIR)**

Samples (3mg) were mixed with 300 mg of potassium bromide and the pellet was made with mechanical pellet maker. Further, the samples were analyzed in the range of 4000–500cm$^{-1}$ with a resolution of 1cm$^{-1}$ using an Alpha FTIR spectrophotometer (Bruker Optics, Mumbai, India).

**Naringin release from PS-N-BSANP**

PS-N-BSANP was cut into 1cm × 1cm and incubated in 2ml of DPBS under static condition. At fixed time intervals, 700µl was removed and replaced with equal amounts of DPBS, read at 284 nm using UV-Vis spectrophotometer (Systronics, India). The concentration of released naringin was obtained from the naringin standard curve.

**Cell culture**

C3H10T1/2 (Mouse mesenchymal cell line) was purchased from National Centre for Cell Sciences (NCCS, Pune, India) and cultured in undifferentiated state using complete culture medium (DMEM-LG supplemented with 10% FBS and 1% antibiotic-antimycotic solution) under standard culture conditions (37ºC, 5% CO$_2$, humidified incubator). The cells were trypsinized and subcultured using trypsin-EDTA solution at 80% confluence.

**Preparation of conditioned media (CM)**

The sterile scaffolds (PS and PS-N-BSANP) were incubated in 1ml of complete culture medium under standard culture conditions. Then, the media were collected and replaced with fresh complete culture medium every day. The collected media were termed as conditioned media. Furthermore, CM obtained from PS scaffold termed as CM-PS, from PS-N-BSANP scaffold as CM-PS-N-BSANP and used for further studies.

**Evaluation of in vitro osteogenic potential**

Cells (5 x 10$^4$) were seeded in 12 well culture plates and allowed to attach overnight under complete culture medium. The groups were labeled as CM-PS (control) and CM-PS-N-BSANP. Further, the complete culture medium was replaced in respective groups with CM and osteogenic medium (DMEM-LG containing 10% FBS, 1% antibiotic-antimycotic solution, 100µg/ml ascorbic acid, 20nM dexamethasone and 20mM β-glycerophosphate) in equal volumes with a change of media every day. To determine the cytotoxicity and osteogenic potential of CM the following assays were performed.

**MTT assay**

The cytotoxicity of the CM was evaluated by MTT assay. On day 3, the cells were rinsed with sterile DPBS followed by MTT solution (0.5mg/ml in DPBS) was added and then incubated for four hours to form formazan crystals. After incubation, the formazan crystals were solubilized with an appropriate volume of DMSO and read at 590nm using UV-Vis spectrophotometer.

**von Kossa and picrosirius red staining**

von Kossa staining and picrosirius red staining were done to visualize the calcium deposition and collagen matrix formation respectively. On day 12, wells were washed with DPBS and fixed with 10% neutral buffered formalin.

For von Kossa staining, silver nitrate solution (1%) was added and exposed to ultraviolet light for 45 minutes. Further, to stop the reaction 5% sodium thiosulfate solution was added followed by washing with distilled water. Calcium deposits stained as black coloured crystals were observed under microscope at 10x magnification.

For picrosirius red staining, cells were incubated with Weigert’s hematoxylin for 8 minutes and washed with distilled water. Further, picrosirius red was added and incubated for one hour and washed with acidified water to remove the excess stain. Furthermore, red coloured stained collagen matrix formation was observed at 10x magnification under microscope.

**Statistical analysis**

The results of MTT assay were expressed as mean±standard deviation. The statistical analysis was evaluated by unpaired student’s t-test using GraphPad PRISM 5.01 (San Diego, CA) and the significant difference (p < 0.05) was considered as statistically significant.

**RESULTS AND DISCUSSION**

**Characterization of the nanoparticles and porous scaffolds**

The nanoparticles were synthesized by
conservation process and the average size of prepared nanoparticles were found as 203.21±38.99nm for N-BSANP (Fig. 1A). These particles did not form agglomerates and were spherical in shape. BSA is a biodegradable protein which is extensively used to synthesize the nanoparticles for sustained release of bioactive molecules in various drug delivery applications due to its excellent properties such as biocompatible, nontoxic, nonimmunogenic etc.\textsuperscript{14,15}

Further, the SEM images of PS (Fig. 1B) and PS-N-BSANP (Fig. 1C) showed interconnected porous structure, which is a most important aspect for cell penetration into the scaffolds. The scaffolds with highly interconnected porous network are considered advantageous for cell migration, attachment, tissue ingrowth, angiogenesis and also it could supports to transport nutrients and wastes\textsuperscript{16}.

XRD analysis was done to determine the crystallinity of prepared nanoparticles and porous scaffolds, the results are shown in Fig. 2. Naringin diffraction spectra (Fig. 2A) showed many numbers of intense peaks\textsuperscript{17}. Further, the broad diffraction peak was seen in N-BSANP spectra (Fig. 2B) corresponds to amorphous BSA, also absence of naringin peaks implies amorphization of naringin during encapsulation which increases the dissolution of hydrophobic drugs\textsuperscript{18}. Also, two semicrystalline peaks were observed for PS (at 2\textdegree of 21.6° and 24°) (Fig. 2C) and PS-N-BSANP (at 2\textdegree of 21.8° and 24.1°) (Fig. 2E) which corresponds to polycaprolactone\textsuperscript{6}. In addition, PS-N-BSANP unleached spectra (Fig. 2D) showed many intense peaks corresponding to sucrose compared to PS-N-BSANP scaffold, which confirmed that the sucrose has leached out completely in PS-N-BSANP scaffold.

FTIR spectra revealed the characteristic peaks for prepared nanoparticles and porous scaffolds. Naringin spectra (Fig. 3A) showed the characteristic peaks at 1647.7 cm\textsuperscript{-1}, 1519.1 cm\textsuperscript{-1}, 1456.8 cm\textsuperscript{-1}, 1363.4 cm\textsuperscript{-1}, 1264.1 cm\textsuperscript{-1}, 1178.6 cm\textsuperscript{-1},
1137.2 cm\(^{-1}\), 1088.8 cm\(^{-1}\), 888.1 cm\(^{-1}\), 835.7 cm\(^{-1}\) and 639 cm\(^{-1}\). Further, N-BSANP characteristic peaks were showed in Fig. 3B. In N-BSANP, the bands at 1647.6 cm\(^{-1}\), 1270.2 cm\(^{-1}\) and 1173.9 cm\(^{-1}\) indicates the presence of naringin in prepared nanoparticles which confirms that naringin was not modified chemically during entrapment process. As previously reported, a strong band at 1735.33 cm\(^{-1}\) (C=O stretching) and 2863.21 cm\(^{-1}\) (asymmetric CH\(_2\)-stretching) were observed for PS scaffold (Fig. 3C) which corresponds to aliphatic polyester of polycaprolactone\(^{20}\). Similarly, characteristic peaks of polycaprolactone were observed at 1731.62 cm\(^{-1}\), 2944.69 cm\(^{-1}\) (symmetric CH\(_2\)-stretching) and 2865.55 cm\(^{-1}\) in PS-N-BSANP scaffold (Fig. 3D). Also, the band for bovine serum albumin nanoparticles was observed at 1243.05 cm\(^{-1}\) and also the bands at 1469.31 cm\(^{-1}\) and 1367.80 cm\(^{-1}\) confirmed the naringin characteristic peaks in PS-N-BSANP scaffold. These results indicates that there was no chemical modification of naringin nor chemical bonding to bovine serum albumin nanoparticles and polycaprolactone during PS-N-BSANP scaffold fabrication.

**Naringin release from PS-N-BSANP**

The nanoparticles were synthesized by coacervation process and the entrapment efficiency of naringin was found to be 97.38%. Sustained release of naringin was analyzed from porous PS-N-BSANP scaffold and the cumulative release was found to be 52.54 micromolar (µM) at the end of 12 days (Fig. 4). In general, drug delivery systems helps to deliver bioactive agents in a controlled manner at the targeted site\(^{22}\). Incorporation of various drug delivery systems into three-dimensional scaffolds have been gathered more consideration for delivery of bioactive molecules with desired concentrations which can be more advantageous in bone tissue engineering approach\(^{22,23}\). Previous studies have reported that, various drugs, growth factors, antiinflammatories and antibiotics have successfully incorporated into the three-dimensional scaffolds and have demonstrated clinical potentials in the treatment

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**Fig. 3.** Fourier transform infrared spectroscopy analysis of nanoparticles and porous scaffolds. Where, (A) - Naringin; (B) - Naringin loaded bovine serum albumin nanoparticles; (C) - Polycaprolactone scaffold; (D) - Polycaprolactone scaffold entrapped with naringin loaded bovine serum albumin nanoparticles
and management of bone-related pathologies\textsuperscript{22,24,25}. Further, targeted drug delivery improves the drug absorption and intracellular diffusion, sustained release for longer time, enhances drug efficacy and also it could reduce the drug degradation\textsuperscript{26}. The naringin release profile suggests that N-BSANP was successfully entrapped into the porous polycaprolactone scaffold and it releases naringin in a controlled manner.

**In vitro osteogenic potential of CM**

The MTT assay showed that the conditioned media from the fabricated porous scaffolds were not toxic with C3H10T1/2 cell line (Fig. 5). The absorbance values of CM-PS-N-BSANP group showed significant increase (P< 0.05) in cell population compared with control. Previously reported studies has shown proliferative effect of naringin on human bone mesenchymal


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