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Synthesis and characterization of β -cyclodextrin/carboxymethyl chitosan/hydroxyapatite fused with date seed extract nanocomposite scaffolds for regenerative bone tissue engineering†

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Herein, for the first time, heterojunctions containing nano-hydroxyapatite/carboxymethyl chitosan/ β -cyclodextrin fused with date seed extract (DSE) were synthesized at different concentrations using co-precipitation to generate BCHD1, BCHD2 and BCHD3 nanocomposite scaffolds with tunable size and amendable surface properties. FTIR, XRD and mechanical analysis studies confirmed the presence of different constituents in the proposed nanocomposite scaffolds displaying strong interactions with an optimal compressive modulus shown by BCHD3 [1533 ± 2.69 MPa], relative to nano-hydroxyapatite/carboxymethyl chitosan (CH) nanocomposite scaffolds. The results of TEM, water contact angle and SEM revealed the homogenous distribution of needle-shaped particles having an average size ranging between 18 and 25 nm with a moderate hydrophilicity [WCA = 42.5°] and an interconnected rough morphology. Furthermore, the BCHD nanocomposite scaffolds were investigated for their biomimetic mineralization ability using simulated body fluid by mimicking the inorganic composition of human blood plasma, which demonstrated superior hydroxyapatite nucleation. The BCHD3 nanocomposite scaffolds showed an advanced protein adsorption [$130 \pm 41 \mu\text{g cm}^{-2}$] and active alkaline phosphatase performance and did not exacerbate lactate dehydrogenase leakage of MG63 osteoblast-like cells relative to CH. The comprehensive *in vivo* study identified better bone regeneration in the repair of critical size calvarial defects [8 mm] in albino rats upon treatment with BCHD3, which is selectively tested as a bare implant in comparison to cerabone, as corroborated by the histopathological and radiological investigations with an average GBD of $94.5 \pm 1.24\%$. Therefore, the above-mentioned results have shed more light on establishing the efficiency of BCHD3 to find applications in osteogenic tissue engineering.

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1. Introduction

Tissue engineering is intended to develop analogues that could sustain, repair, mimic or substitute a biological tissue.¹ The process of bone regeneration through engineering the tissue is an intricate biological event which involves scaffolds and processes such as vascularisation, cellular and signaling cues, along with intracellular interactions. In this milieu, the principle extracellular matrix (ECM) like micro-environment is provided by biocompatible materials which facilitate the intracellular interactions. There are many properties that should be satisfied by a tissue engineered matrix, which include appropriate mechanical strength, optimal porosity and surface chemistry to name a few.² However, as the process of regeneration of bone inside the scaffolds is complex, the synchronized processes such as osteogenesis and angiogenesis which are involved in bone

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regeneration have often been examined piecemeal using several *in vitro* and *in vivo* experimental studies.³ Compared to the existing implants, modern strategies, including nanotechnology,⁴ have opened new horizons to reproduce superior structural properties by fabricating materials which could imitate the nano-scaled architecture of the natural bone. For example, nano-crystalline ceramics such as calcium phosphates⁵ have been shown to fuel the bone cell functions causing rapid degradation relative to microcrystalline calcium phosphate.

Recently, the use of nanohydroxyapatite (NHA) as a bio-material has escalated great attention in the engineering of bone tissues, being the main component of bone. It acts as a template for the mineral phase deposition and triggers new bone formation.⁶ However, the antiresorptive and anabolic ability of NHA are found to be unsatisfactory and its matrices are reported to be brittle and thus, a new strategy is still desired for the development of biocompatible, biodegradable and bioactive scaffolds. To meet this requirement, biopolymers have enthralled researchers because of their inherent properties such as abundance, low cost, extracellular matrix (ECM)-like characteristics appropriate for cell function and survival, biocompatibility, biodegradability and non-toxicity and thus have been the emerging candidates of choice in recent years.⁷ Among the biopolymers, chitosan, the deacetylated form of chitin derived from the exoskeleton of crustaceans, is actually a polyheterosaccharide containing glucosamine and *N*-acetyl glucosamine units bonded by 1–4 glucosidic linkages and has gained much attention in the field of bone tissue engineering.⁸ Nevertheless, it is associated with disadvantages like low solubility, less load bearing potency and high percentage of degradation that have restricted its utilization.⁹ In order to cope with these limitations, carboxymethyl chitosan¹⁰ (CMC), a well warranted chitosan-derived polymer, derived from the carboxylation reaction of amine group substitution by smaller alkyl groups, has engrossed great interest in a wide spectrum of biomedical applications, being biocompatible and cost-effective, and owing to the suitable degradation rate. In addition, CMC is better soluble in water in comparison with chitosan without the aid of acid substances, preventing damage to the pH sensitive antibacterial activity of a drug.^{11,12} At this junction, it is relevant to mention that combining chitosan derivatives with other naturally derived moieties may result in the development of nanocomposite scaffolds with the advantages of enhanced performance in terms of solubilization and permeability. One such structural residue is the class of cyclodextrins,¹³ which are the cyclic oligosaccharides, consisting of 6, 7 or 8 α -D-glucopyranose units, forming toroidal-shaped molecules with a hydrophilic exterior and a hydrophobic interior formed by the carbon backbones of glucopyranose monomers, appropriate for the formation of inclusion complexes. Being oligosaccharides, these molecules bear hydroxyl groups that may possibly be employed for further functionalization.¹⁴ The β -cyclodextrin (β -CD), among cyclodextrins, is acknowledged as a performance accelerator in tissue engineering¹⁵ by various routes such as (1) sequestering growth factors, (2) augmenting oxygen concentrations in the tissue engineered constructs, and (3) regulating collagen self-assembly.^{16–18} Therefore, these co-polymeric

systems possess the cumulative complimentary properties of their constituent polymers. With regard to date seed added values, *Phoenix dactylifera* L. (date palm), from the Arecaceae family, has recently been studied for its bone regeneration properties.¹⁹ There are various studies that investigated date seed powder as an economical and major source of antioxidants and fibers due to the attendance of valuable dietary components in addition to the presence of flavones, polyphenols and vitamins (particularly vitamins C and E), which make them function as anti-atherogenic.²⁰ In addition, their chemical makeup and phytochemical profile demonstrated *in vitro* bioactivities and showed applications such as expectorant, hypoglycaemic, aphrodisiac, and antidiarrheic. In addition, the recent study represented that the extract of the date seeds plays a cerebroprotective role in male rats.²¹ Therefore, in view of their valuable essential features, we used date seed extracts with other polymeric counterparts in the present work. The reason to recruit these established biocompatible²² components has come from their wide usage in biomedical and pharmaceutical applications due to the associated low cost, biocompatibility and biodegradability.^{23,24}

To our knowledge, heterojunctions containing β -cyclodextrin (B), carboxymethyl chitosan (C), nano-hydroxyapatite (H), and date seed extracts (D) synthesized by co-precipitation to yield the BCHD nanocomposite scaffolds have not been achieved to date. Therefore, the present study was anticipated to provide a favourable combination of chemical, mechanical and biological properties to evaluate the performance of date seed extracts incorporated at different concentrations into a nano-hydroxyapatite/carboxymethyl chitosan/ β -cyclodextrin matrix and its potency was investigated as a competent and efficacious bone implant. The physicochemical characteristics were evaluated in terms of size, shape, crystallinity, morphology and mechanical efficiency. Furthermore, we have also delved into cell viability, ALP, protein adsorption and LDH analysis to study the biocompatible nature of the proposed nanocomposite scaffolds.

2. Materials and methods

2.1. Materials

3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide (MTT), carboxymethyl chitosan (CMC) and β -cyclodextrin (β -CD) were obtained from Sigma-Aldrich (USA). The other chemicals/reagents used in the present study are listed in the ESI† under Section S.2.1.

2.2. Experimental

2.2.1. Preparation of NHA. Hydroxyapatite nanoparticles (NHA) were synthesized by chemical precipitation as reported in our previous studies.^{19,25} The so obtained product was subjected to ball milling to decrease the average size of the particles for the formation of crystalline NHA.

2.2.2. Preparation of date palm seed extract (DSE). The Ajwa date seeds were manually separated from the fruit and kept in water for 4–5 hours. Subsequently, the soaked seeds



were washed repeatedly to completely remove any adhered flesh. Then, the seeds were kept for air-drying followed by drying in an oven for 48 h at 55–60 °C and milled to form a fine powder with the help of a heavy duty grinder.²⁶ To this powder of dried pits, distilled water was added to make the mixture of 50 gm L⁻¹ which was boiled until it turned brownish and then filtered and stored at 4 °C for use.^{7,27}

2.2.3. Synthesis of homogeneous BCHD nanocomposite scaffolds. The BCHD nanocomposite scaffolds were synthesized *via* the co-precipitation approach¹⁹ as described. First, CMC solution was prepared by dissolving 1 g of CMC in 50 mL of DH₂O with continuous stirring at room temperature to obtain a perfectly transparent solution. To this solution, 1 g of NHA solution prepared in 50 mL of DH₂O was added dropwise and the mixture was continuously stirred at room temperature for 4–6 hours. To this mixture, β -CD solution, made by dispersing 1 g of β -CD in 50 mL of DH₂O by sonication for 30 min, is dropped slowly to obtain a homogeneous stock solution. At this point of time, the stock solution is divided into 3 equal parts. To each part, 10 mL, 20 mL and 30 mL of DSE were added dropwise to form BCHD1, BCHD2 and, BCHD3 nanocomposite scaffolds, respectively. Then, the reaction was continued until the contents were thoroughly mixed. The reaction mixture thus obtained was kept under constant stirring at 1200 rpm at room temperature. The pH of the resulting mixture was maintained at ~11 using 0.5 M NaOH to ease the nucleation of NHA followed by stirring at the same rate for another 24 h. The slurry so obtained was aged for 48 h and the precipitate was harvested after repeated washing with DH₂O until the filtrate became neutral. The precipitate so obtained was dried at 50 °C in a vacuum. To compare the properties of the proposed nanocomposite scaffolds, we synthesized the CH (CMC/nano-hydroxyapatite) nanocomposite scaffold adopting the same procedure in a 1 : 1 ratio.

2.3. Characterization

The morphological and chemical characterization are described in the ESI† under Section S.2.2.

2.4. Statistical analysis

Data were obtained from triplicate samples and presented as mean \pm standard deviation. A Student's *t*-test was performed for statistical analysis and the significance was defined at $p < 0.05$ (*).²⁸

3. Animal experiments

3.1. Ethical statement

Ethical consent was obtained for all the animal trials and conducted in conformity to the experiment guidelines permitted by the Institutional Animal Ethics Committee, Jawaharlal Nehru Medical College, and Aligarh Muslim University.²⁹

3.2. Cerabone

In the present study, the commercially available grafting material from Botiss (Zossen, Germany) was employed.³⁰

3.3. Experimental design: surgery and implantation procedure

A total of eighteen male adult albino rats (300–350 g) were employed for the study. Animals were randomly assigned to three groups ($n = 6$ per group). The critical size 8 mm defect was created in the cranial bone of all animals in each group. In group (1), the defect was kept blank and treated as the control group; in group (2), cerabone (CB) was filled in the defect; and in group (3), the defect was filled with the BCHD3 nanocomposite scaffold. The intended surgical area was prepared by shaving skin and disinfection with 7.5% povidone-iodine solution (Win-Medicare, India). The surgeries were performed following the experimental procedure described in our previous studies,^{19,29} which involved general anaesthesia, skin incision, periosteum incision, creation of a critical defect in the parietal bone, implantation and closure of the defect that took around 30–45 minutes. The operative procedure was uneventful and postoperatively, the animals also received antibiotic cover (gentamycin) for 5 days. The rats were allowed to move freely in their cages without external support. The wounds healed normally and no abnormal behavior was noticed during the 8 week duration of the study. The animals were sacrificed after the completion of the experimental period and calvarial samples were harvested, cleaned and subjected to radiovisiographic analysis. Subsequently, these samples were decalcified for histopathological observations.

3.4. Radiographic analyses

For radiographic samples, the imaging system (Satelac-Acteon, X-Mind, Italy; X-ray tube-Toshiba, Japan) was used. The radiographs were recorded using a radiographic sensor (Suni Plus-0543, USA). Image Analysis toolbox (MATLAB 6.1, The MathWorks Inc., Natick, MA, 2000) was used to quantify the gain in bone density (GBD) as reported previously.^{19,29} The results were expressed as mean \pm standard deviation (SD). The *in vivo* study analysis was performed using Tukey–Kramer multiple comparison. All results were considered to be significant at the 5% critical level.¹⁹

3.5. Preparation of calvarial samples

The harvested tissue of calvarial samples was fixed in formalin (10%) for 48 h and kept for decalcification [in ethylene diamine tetra acetic acid (EDTA), 10%] after the RVG analyses. The decalcified tissue samples were processed for paraffin sections. The sections were stained with hematoxylin–eosin (HE) and Sirius Red (SR) stains and observations were made using a light microscope (Trinocular Olympus RX40, Japan) as described in our previous studies.²⁹

4. Results and discussion

4.1. Fourier transform infrared (FTIR)

For the elucidation of the chemical interactions between the various components present in the nanocomposite samples, FTIR analyses were carried out [Fig. 1(i)]. The FTIR scan of the carboxymethyl chitosan/nano-hydroxyapatite nanocomposite





Fig. 1 FTIR spectra (i), XRD diffractograms (ii), mechanical strength analysis: compressive modulus (iii), and compressive stress–strain (iv) curves of BCHD1, BCHD2, BCHD3 and CH nanocomposite scaffolds where the fracture point has been shown by yellow color, * $p < 0.05$ showing the statistical difference compared with the CH nanocomposite scaffold.

scaffold (CH) showed the presence of the representative bands of NHA in the range of 536–566 and 603–608 cm^{-1} , respectively,³¹ corresponding to phosphate bending modes of vibrations of NHA. Also, the band characteristics of the symmetric and asymmetric deformations of COO^- signifying the presence of carboxymethyl groups in CMC are shifted to 1634 and 1400 cm^{-1} , respectively, indicating the coordinative interaction between calcium ions and CMC.³² The stretching mode of vibration of the phosphate group in NHA overlapped with the $-\text{C}-\text{O}-$ band of CMC, resulting in a sharp peak³³ at 1030 cm^{-1} . The FTIR spectra of BCHD1, BCHD2 and BCHD3 nanocomposite scaffolds exhibited the presence of characteristic peaks of CMC and NHA; however, the stretching mode of vibration of the phosphate group in NHA overlapped with the $-\text{C}-\text{O}-$ band of β -CD, $-\text{C}-\text{O}-$ band of CMC³⁴ and $\text{C}-\text{O}$ stretching vibration of cellulose and hemicellulose present in DSE,¹⁵ resulting in a broad band^{15,19,35} in the region around 1020–1042 cm^{-1} . The band representing the more hydrophilic character of CMC is overlapped with the OH stretching band of NHA, resulting in the appearance of a broad band centered at 3268 cm^{-1} in the CH nanocomposite scaffold; however the band became broader and shifted slightly to higher wavenumber and appeared in the region of 3402–3416 cm^{-1} with the addition of β -CD and DSE in comparison to CH. This is possibly due to the overlapping of the OH stretching band of NHA with the OH stretching bands of CMC, β -CD and DSE, signifying the possibility of intermolecular hydrogen bonding among the components.^{15,35} The peak observed at 2942 cm^{-1} was due to the asymmetric C–H stretching vibration from CH_2 groups in CMC present in all the nanocomposite scaffolds.^{36–38} The appearance of new bands in the range of 2318–2352 cm^{-1} in BCHD1, BCHD2,

cellulose present in DSE,¹⁵ resulting in a broad band^{15,19,35} in the region around 1020–1042 cm^{-1} . The band representing the more hydrophilic character of CMC is overlapped with the OH stretching band of NHA, resulting in the appearance of a broad band centered at 3268 cm^{-1} in the CH nanocomposite scaffold; however the band became broader and shifted slightly to higher wavenumber and appeared in the region of 3402–3416 cm^{-1} with the addition of β -CD and DSE in comparison to CH. This is possibly due to the overlapping of the OH stretching band of NHA with the OH stretching bands of CMC, β -CD and DSE, signifying the possibility of intermolecular hydrogen bonding among the components.^{15,35} The peak observed at 2942 cm^{-1} was due to the asymmetric C–H stretching vibration from CH_2 groups in CMC present in all the nanocomposite scaffolds.^{36–38} The appearance of new bands in the range of 2318–2352 cm^{-1} in BCHD1, BCHD2,



and BCHD3 nanocomposite scaffolds may be attributed to the coordinative interactions between Ca^{2+} of NHA and OH of $\beta\text{-CD}^{15}$ that is responsible to support the nucleation of apatite crystals. Thus, the analysis of the FTIR spectra confirmed that all the components used during synthesis, *i.e.* NHA, CMC, $\beta\text{-CD}$ and DSE, have been incorporated in the final system BCHD and exhibited the expected interactions among different components present in the nanocomposite scaffolds.

4.2. X-Ray diffraction (XRD) analysis

The XRD patterns of CH, BCHD1, BCHD2, and BCHD3 nanocomposite scaffolds are shown in Fig. 1(ii). The XRD spectrum of CH exhibited the characteristic peaks of NHA at $2\theta = 25.5^\circ$, 31.9° , *etc.*, in accordance with the standard diffraction pattern of NHA (PDF# 09-0432),³⁷ while the presence of CMC was evident by the appearance of its XRD diffraction peaks³⁸ at $2\theta = 28.9^\circ$ and 46.4° . In the XRD spectra of BCHD1, BCHD2 and BCHD3 nanocomposite scaffolds, the characteristic peaks of NHA and CMC were observed. In addition, the presence of DSE was evident by the appearance of the peak at $2\theta = 16.2^\circ$ ascribed to its native cellulose and hemicellulosic components, while the other peak was overlapped with the CMC and $\beta\text{-CD}$ peaks^{19,39–41} and appeared at $2\theta = 20.4^\circ$. The appearance of a crystalline peak⁴⁰ at $2\theta = 15.9^\circ$ also marked the presence of $\beta\text{-CD}$ in BCHD1, BCHD2 and BCHD3 nanocomposite scaffolds. The broad peaks of the BCHD nanocomposite scaffolds represented low crystallinity of biological apatite, thus mimicking natural bone.⁴²

4.3. Mechanical strength

In the present study, the compressive strength of the CH, BCHD1, BCHD2 and BCHD3 nanocomposite scaffolds was tested following the guidelines for the preparation of the samples [ASTM F 451-95].⁴³ The results of the average

compressive modulus [Fig. 1(iii)] revealed that BCHD3 exhibited a compressive modulus [1533 ± 2.69 MPa] which falls in the range observed for human cortical bone and significantly higher ($p < 0.05$) than CH, suggesting that the addition of $\beta\text{-CD}$ together with DSE which is a rich source of fibers⁴⁴ in the CH matrix has noticeably improved the mechanical parameters. The compressive stress–strain curves of the CH, BCHD1, BCHD2 and BCHD3 nanocomposite scaffolds are shown in Fig. 1(iv), where it can be observed that in the initial phase of compression, there was a sharp increase in stress with subsequent decrease until the point of fracture was attained. The increased intermolecular hydrogen bonding among nano-hydroxyapatite, cellulose and hemicellulose present in DSE, CMC and $\beta\text{-CD}$ moieties may possibly be the reason for enhanced mechanical parameters displayed by the BCHD nanocomposite scaffolds, which might create a rigid framework with suitable mechanical properties, thus facilitating their clinical applications.⁴⁰

4.4. Transmission electron microscopy (TEM) and contact angle measurements

The morphologies of BCHD1, BCHD2, BCHD3 and CH nanocomposite scaffolds were characterized by TEM. Based on the results as displayed in Fig. 2(A–D), needle-like particles were homogeneously distributed in the case of BCHD1, BCHD2 and BCHD3 having an average particle size in the range of 18–25 nm which was found to decrease with an increase in the concentration of DSE in BCHD3 as shown in Fig. 2C. In the case of CH, the average particle size varied in the range of 30–35 nm and the particles were observed to be comparatively crowded resulting in the formation of the aggregated structure.^{32,45} The consistent dispersion of particles in the case of the proposed nanocomposite scaffolds may be due to the interactions of DSE with CMC, NHA and $\beta\text{-CD}$. As the size of the particles was found to be a function of the concentration of DSE, hence, the DSE appeared to play a key



Fig. 2 Representative TEM micrographs of (A) BCHD1, (B) BCHD2, (C) BCHD3, and (D) CH nanocomposite scaffolds [scale bar: 100 nm] with the inset showing water contact angles and their respective SEM images [A'–D'] [scale bar: 10 μm , magnification: $\times 3500$].





Fig. 3 Characteristic SEM micrographs illustrating the bioactivity profile by the SBF study of [A] BCHD1, [B] BCHD2, [C] BCHD3, and [D] CH nanocomposite scaffolds at 4 weeks [magnification: $\times 3500$, scale bar: 5 μm] and at 8 weeks [A'–D'] [magnification: $\times 2500$, scale bar: 10 μm], respectively, with their respective EDX micrographs recorded after 8 weeks.

role in controlling the size of the particles.^{46,47} These results were found to be in line with the previous literature indicating the role of DSE as a size and shape modifying agent.³⁵ The water contact angle studies conducted on BCHD1, BCHD2, BCHD3 and CH nanocomposite scaffolds [Fig. 2(A–D) inset] showed that the contact angle was smaller in the case of BCHD nanocomposite scaffolds with a WCA of 32.6° for BCHD1 and 38.9° for BCHD2, while BCHD3 showed a moderately hydrophilic surface with a WCA of 42.5° and was thus found to be most suitable for the adhesion and proliferation of the cells in comparison to relatively hydrophobic CH with a contact angle of 69.4° in view of the synergic addition of the high degree of $-\text{OH}$ groups present in NHA, CMC, β -CD and DSE.⁴⁸

4.5. SEM

SEM analysis was conducted in order to investigate the microstructure and surface morphologies of BCHD1, BCHD2, BCHD3 and CH nanocomposite scaffolds. Fig. 2(A'–D') represents the porous and rougher surface morphologies of BCHD1, BCHD2, and BCHD3, in which BCHD3 [Fig. 2C'] displayed maximum roughness compared with the CH nanocomposite scaffold where CMC imparted smoothness to the surface [Fig. 2D'].^{38,49} However, the introduction of DSE and β -CD significantly enhanced the roughness in the nanocomposite scaffolds^{15,19}

owing to the presence of intermolecular H-bonding. Therefore, the synergism among NHA, CMC, β -CD and DSE in BCHD nanocomposite scaffolds caused an increased surface area and an interconnected structure with a sufficiently higher porosity appropriate for the transportation of extracellular fluids, nutrients, and hormones to cells⁵⁰ and thus making it a suitable platform for applications in BTE.

4.6. *In vitro* calcification assay and EDS

The representative SEM images at 4 weeks of *in vitro* calcification conducted on BCHD1, BCHD2, BCHD3 and CH nanocomposite scaffolds are shown in Fig. 3(A–D). Having plenty of carboxyl and OH groups, BCHD nanocomposite scaffolds showed flattened and fused flaky apatite growth (shown by red arrows), resulting from the combination of calcium ions with more number of carboxyl and hydroxyl groups present on their surfaces in comparison to the CH nanocomposite scaffold. In accordance with earlier studies,^{51,52} this observation indicated the catalytic effect of the hydroxyl and carboxyl groups in inducing apatite formation. At 8 weeks [Fig. 3(A'–D')], though the carboxyl groups of CMC present in CH acted as the template for subsequent biomimetic deposition,^{43,53} the mineralization got highly increased in all BCHD nanocomposite scaffolds (shown in circular regions) relative to CH. These observations may be



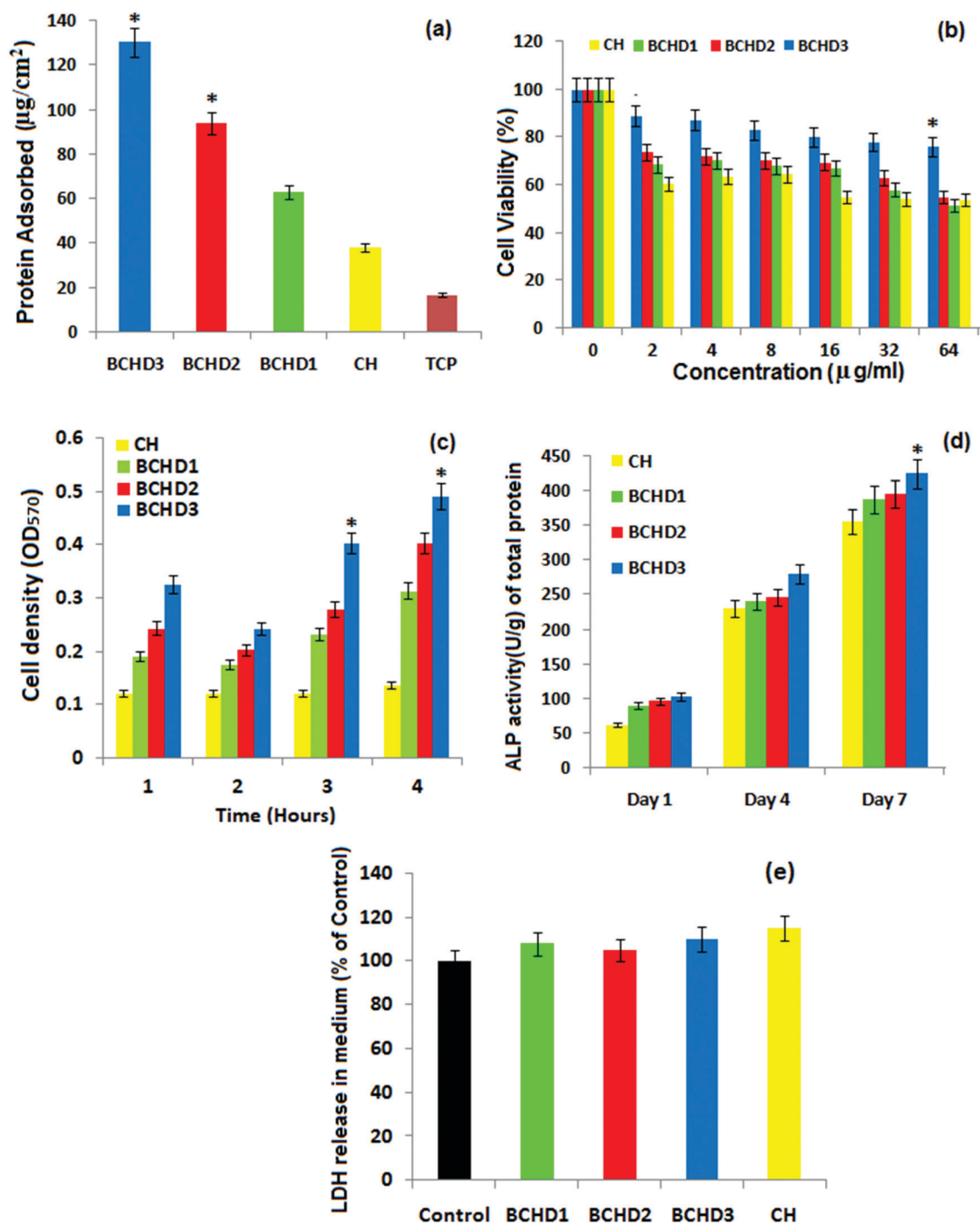


Fig. 4 Biocompatibility studies: (a) protein adsorption assay, (b) cell viability assay, (c) MG-63 cell density measured at 1–4 hours, (d) alkaline phosphatase activity, (e) LDH assays of BCHD1, BCHD2, BCHD3, and CH nanocomposite scaffolds. * $p < 0.05$ shows a statistical difference compared to the CH nanocomposite scaffold.

understood in terms of the relatively rougher surface exhibited by the BCHD nanocomposite scaffolds, leading to enhanced osteoconductivity and bioactivity. Also, it was noticed that the apatite formation occurred as irregular granular depositions comprising mainly spherical and rectangular granules in the case of BCHD2 which appeared to get denser in the case of BCHD3, as shown in Fig. 3C', inferring that with an increase in

the concentration of DSE in BCHD3, more apatite nucleation occurred by the consumption of calcium and phosphate ion precursors present in the surrounding fluid.⁵⁴ These results possibly indicate the ability of β -CD and DSE as a mineralization inducer where DSE is a rich source of minerals essential for bone formation.⁴⁴ These results were found to be in agreement with the former results.^{15,19}

The EDS results of the SBF treated BCHD1, BCHD2, BCHD3 and CH nanocomposite scaffolds after 8 weeks incubation [Fig. 3] clearly showed that the composition and phase resembled the mineral component of the bone. The apatite formed in the case of BCHD nanocomposite scaffolds exhibited⁵⁵ a semiquantitative ratio of Ca/P in the close proximity of 1.6–2.1 which is the characteristic range found in natural bone⁴¹ in comparison to the ratio (1.3) obtained in the case of the CH nanocomposite scaffold.

These results provided a fundamental desirable characteristic of a bioactive material that facilitates osteointegration *in vivo*.

4.7. Protein adsorption assay

Protein adsorption is a crucial index to investigate the biocompatible nature of a scaffold for bone engineering applications. Nanocomposite scaffolds with varying amount of DSE were incubated in FBS solution and compared with CH to investigate their protein adsorption ability. The adsorption of protein on the surface of CH, BCHD1, BCHD2, and BCHD3 nanocomposite scaffolds and a tissue culture plate (TCP) is shown in Fig. 4a. It was observed that all the BCHD nanocomposite scaffolds exhibited significantly higher protein adsorption compared with CH with a maximum average value of $130 \pm 41 \mu\text{g cm}^{-2}$ shown by BCHD3. The addition of β -CD and date seed extracts to the CH matrix enhanced the surface roughness, thus increasing protein-binding channels. These observations suggested the favorable bioaffinity of the BCHD3 nanocomposite scaffold to a range of proteins present in the extracellular matrix such as fibronectin, glycoprotein and the different growth factors that could assist adhesion, proliferation, and differentiation of cells both *in vitro* and *in vivo*.⁵⁶

4.8. Cell proliferation: MTT assay

The cell proliferation was quantitatively evaluated by MTT assay by culturing human osteoblast like MG-63 cells on CH, BCHD1, BCHD2 and BCHD3 nanocomposite scaffolds at various concentrations ($0\text{--}64 \mu\text{g mL}^{-1}$), as shown in Fig. 4b. The BCHD1, BCHD2 and BCHD3 nanocomposite scaffolds inferred higher metabolic activity of the cells compared to CH; however, BCHD3 showed significant cell viability even at higher concentration validating its proficiency in increasing biocompatibility. The optical density value was also detected at a series of time points of 1–4 hours to evaluate cell proliferation (Fig. 4c). Clearly, the results showed higher cell proliferation in the case of the BCHD nanocomposite scaffolds compared with the CH nanocomposite scaffold with BCHD3 displaying significantly higher ($p < 0.05$) cell density at 3 and 4 hours compared with CH. The synergic combination of CMC,³⁶ NHA, β -CD and DSE may possibly lead to a well-interconnected rougher and porous surface in the case of BCHD nanocomposite scaffolds, which is responsible for better structural stability that plays a crucial function in cell attachment and proliferation.⁵⁷

4.9. Alkaline phosphatase (ALP) assay

Generally, ALP assists in the calcium deposition in addition to initial bone mineralization. Therefore, as part of our *in vitro*

investigation, the ALP activity of MG63 cells subjected to BCHD1, BCHD2, BCHD3 and CH samples was monitored and is shown in Fig. 4c. The results illustrated that the BCHD1, BCHD2 and BCHD3 nanocomposite scaffolds significantly increased the expression of ALP ($p < 0.05$) in comparison to the CH at all the tested time points of 1, 4 and 7 days. The superior activity might be attributed to the presence of β -CD and DSE in the CH matrix, which impart porosity and larger surface area concentrating more proteins and growth factors from the culture, hence providing an effective substrate for cellular differentiation stimulating the process of osteogenesis.^{15,19}

4.10. LDH assay

To further understand the biocompatible characteristics, LDH leakage assays of the BCHD1, BCHD2, BCHD3 and CH nanocomposite scaffolds were conducted and the results are shown in Fig. 4d after 24 h of co-culture with MG63 cells. The extent of damage to the cell membrane is indicated by evaluating the percentage of LDH leakage since LDH is a stable cytoplasmic enzyme, which gets released if the plasma membrane integrity is compromised.⁵⁶ It is speculated that in comparison to the control, no significant increase ($p > 0.05$) was observed in the LDH releasing level, further confirming the cytocompatibility of the nanocomposite scaffolds. However the LDH level was found to be slightly lower in the case of quaternary systems relative to CH, warranting the enhanced biocompatibility of the proposed nanocomposite scaffolds⁵⁸ with the addition of β -cyclodextrin and date seed extracts to the CH matrix, resulting in enhanced intermolecular interactions.^{15,19} These observations run parallel to the previous reports and further confirmed the biocompatibility of the BCHD nanocomposite scaffolds.

5. *In vivo* study analysis

5.1. Clinical observations

The visual inspection immediately after the creation of calvarial defects affirmed that the dura mater remains intact in all the groups and there were no complications emerged from the craniotomy. From the clinical aspect, all rats healed well and were observed to gain weight throughout the study period.

5.2. Radiographic and histological assessment

Following *in vitro* characterizations, preliminary *in vivo* investigation was carried out. The histological analysis of hematoxylin and eosin (H&E) and Sirius red (SR) stained sections in the control (blank) group after 4 weeks [Fig. 5 control panel] showed no sign of new bone formation at the defect site. However, active osteoblasts were seen lined up at the margin of the defect and the defect itself was occupied with the fibrous tissue primarily made up of the collagen fibre.^{19,29} The control group at 8 weeks also did not reveal the bone formation at the site of the defect; however, the fibrous tissue which was filling the defect showed increased vascularisation by invading newly formed blood vessels [Fig. 6 control panel]. The RVG results of GBD (gain in bone density) for the control



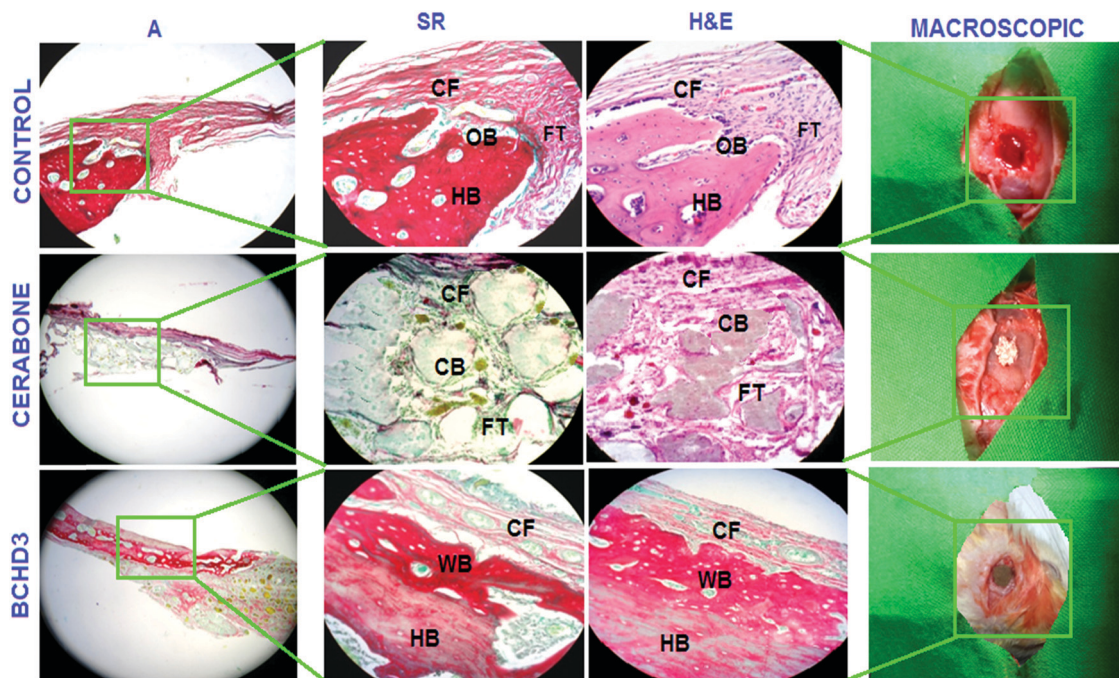


Fig. 5 Histopathology (H&E and SR stains) after 4 weeks post-implantation of the control group, cerabone group and BCHD3 group and their macroscopic view; initial magnification: $\times 100$ (shown by A) and $\times 200$. [OB: osteoblasts, CF: collagen fibre, CB: cerabone, HB: host bone, FT: fibrous tissue, and WB: woven bone].

group at 4 and 8 weeks were found to be $26.60 \pm 0.18\%$ and $37.40 \pm 1.54\%$, respectively [Fig. 7(i, ii) control panels], representing negligible gain which was also in conformity with the histological findings of these groups. The cerabone group at 4 weeks [Fig. 5 cerabone panel] revealed the absence of new bone formation and the defect was seen to be filled with

collagen fibre and cerabone with a GBD of $37.71 \pm 1.12\%$ [Fig. 7(i)]. At 8 weeks, the cerabone group showed a mixed picture having fibrous tissue, some bone and clumps of cerabone as well, thereby suggesting the partial utilization of cerabone for bone formation²⁹ [Fig. 6 cerabone panel] and the GBD was found to be $86.0 \pm 1.4\%$ [Fig. 7(ii)] which supported

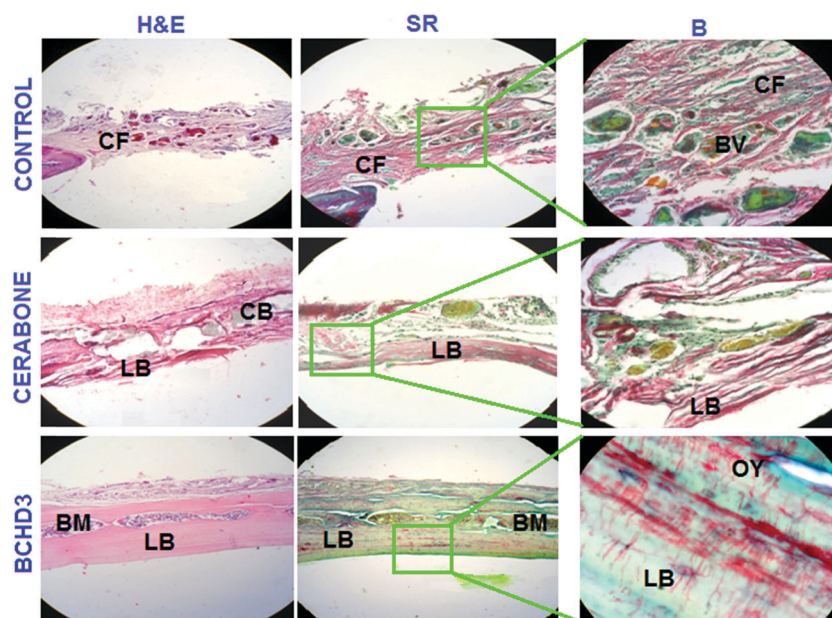


Fig. 6 Histopathology (H&E and SR stains) after 8 weeks post-implantation in the control, cerabone, and BCHD3 experimental groups. Initial magnification: $\times 100$, and B represents magnified view at $\times 200$. [CF: collagen fibre, BV: blood vessels, CB: cerabone, LB: lamellar bone, BM: bone marrow, and OY: osteocytes with multiple processes].





Fig. 7 Gray value profiles of the control, cerabone, and BCHD3 experimental groups after the time period of 4 and 8 weeks. Asterisk indicates the average gray values in the host bone region (indicated by ▲) and defect regions.

the histological findings of partial bone formation. In the case of the BCHD3 group, at 4 weeks, the RVG analysis revealed a GBD of $59\% \pm 1.13\%$, [Fig. 7(i)]; thus, it is really startling and gratifying to note that the BCHD3 group is superior in promoting the bone formation to the extent that even at 4 weeks, there is distinct evidence of woven bone formation. Moreover, by 8 weeks, when there is no bone formation in the control and moderate bone formation in CB, the BCHD3 group showed not only newly formed bone but also remodelled lamellar bone with a GBD of

94.5 ± 1.81 [Fig. 7(ii)]. Notably, in the case of the BCHD3 group at 8 weeks, there was the appearance of bone marrow that is rich in mesenchymal stem cells and osteoblasts and thus marks the critical aspect for the regeneration of bone tissue. From these observations, it may be concluded that BCHD3 not only promoted osteoregeneration, osteoinduction, osteointegration and advanced calcification, but also successfully bridged the critical size defect within 8 weeks even without the use of any growth stimulators and prior cell seeding.



The possible reason for the exceptional performance of BCHD3 may be accounted in terms of synergism among its various constituents. NHA has an ion-inducing effect and releases PO_4^{3-} and Ca^{2+} in the physiological environment, which constitutes a key duo for proper bone mineral acquisition and maintenance, thus participating in the process of osteogenic differentiation leading to new bone formation.³⁶ Next, the component β -CD, being biologically inert, has been approved by the US FDA as the pharmaceutical excipient for numerous drug formulations. Recently, it was reported to show a robust bone anabolic effect *in vivo*.⁵⁹ Another component CMC, being a derivative of carboxymethyl chitin, has also been found to be efficient in causing nucleation of apatite and osteogenesis due to the presence of carboxyl groups.⁶⁰ This causes the induction of inflammatory cytokines and growth factors and thus accounted for restoring cartilage.⁴³ On the other hand, besides being a rich source of proteins, fibres, antioxidants and minerals, DSE has been reported to include several important chemical constituents such as alkaloids, carbohydrates, tannins, *etc.*, which mainly comprise nitrogen and oxygen heteroatoms, that play a crucial role in activating the process of bone formation as reported previously.^{40,61} Furthermore, plant extracts have also been documented to increase the antioxidant defenses and decrease the tissue inflammation, along with accelerating tissue vascularisation and proliferation of the bone cells. They were also reported to stimulate bone repair, promote osteogenesis, speed up calcification, and endorse bone callus formation and mineralization.⁶² Hence, the higher concentration of DSE in the BCHD3 nanocomposite scaffold exhibited a combinatorial effect with other constituents, leading to improvement in the osteoinductivity by enhancing roughness, surface area and topography, therefore influencing the differentiation of stem cells and accelerating the process of new bone formation on the fractured region.^{63–65} Subsequently, these results will thus serve as a cornerstone for future research focused on the design of bioinspired conductive nanocomposite scaffolds incorporating plant extracts for bone tissue regeneration.

6. Conclusion

The present work was intended to invest an agricultural waste product into the pharmaceutical domain, which can offer therapeutic value with a socioeconomic cost. Valorization of these natural origin materials could condense the utilization of synthetic materials for scaffold fabrication, thus avoiding the release of harmful byproducts during fabrication.⁴⁰ The BCHD nanocomposite scaffolds were efficaciously developed by the co-precipitation approach for bone regeneration applications. To summarize, the nanocomposite scaffolds showed a porous and interconnected structure essential for the bone regeneration process. Among various concentrations taken into account, BCHD3, with the maximum DSE content, has stood out as a promising alternative showing excellent bioactivity and cytocompatibility with MG-63 cell lines. Also, the BCHD nanocomposite scaffolds were found to be successful in advancing cell adhesion

and ALP activity besides causing calcium ion deposition, which are regarded as the key parameters for active bone formation. These results demonstrated the portentous role of natural-based quaternary nanocomposite scaffolds with minimum side effects and reduced costs, and they may prove efficacious in treating bone defects.

Conflicts of interest

There are no conflicts of interest to declare.

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