Preparation and characterization of three-dimensional porous collagen/graphene oxide/hydroxyapatite nanocomposite scaffolds for bone tissue engineering

Bhahat Lawlley Zimba\textsuperscript{1,2,*}, Hao Jiang\textsuperscript{1}, Li Chen\textsuperscript{1}, Yaomin Li\textsuperscript{1}, Xunzhi Yu\textsuperscript{1}, Chang Chen\textsuperscript{1}, Jiangling Wan\textsuperscript{1}, Qungzhi Wu\textsuperscript{1}

\textsuperscript{1}Wuhan University of Technology, Wuhan, China
\textsuperscript{2}Malawi University of Science and Technology, Malawi

*Corresponding author: Bhahat Lawlley Zimba: bahat.zimba@yahoo.com

Abstract:

Studies have reported that the incorporation of graphene oxide (GO) and hydroxyapatite (HA) into biocompatible polymers (such as collagen (Col), chitosan, alginate, etc) results in enhanced structural and mechanical properties respectively. The objective of this study was to prepare and characterize three-dimensional (3D) porous Col/GO/HA nanocomposite scaffolds and to investigate cytocompatibility and osteogenic differentiation potential of rat bone marrow mesenchymal stem cells (rBMSCs) on the as-prepared scaffolds. The SEM images revealed that the scaffolds were porous with the pore diameter inversely proportional to the concentration of HA. XRD results were able to depict the characteristic peaks for HA which shows that HA was incorporated into the scaffolds. The rBMSCs which were cultured on the scaffolds were able to attach and proliferate during the 21 days of the experiment which indicates that the as-prepared scaffolds are cytocompatible. The Alizarin red staining demonstrated the presence of calcium deposits as there were orange-red stains on the samples after culturing the cells using the osteogenic differentiation medium. These results demonstrate the promising potential of the 3D porous Col/GO/HA nanocomposite scaffolds for applications in bone tissue engineering.

Keywords: Collagen, Graphene oxide, Hydroxyapatite, Porous scaffolds, Tissue engineering.
Introduction

Scaffolds are used to restore or regenerate damaged tissue in tissue engineering by providing a framework for cells to attach, proliferate, and form extracellular matrix of the targeted tissue [1]. These scaffolds may also serve as carriers for cells, growth factors, and other bimolecular signals. For the scaffolds to effectively aid tissue restoration and regeneration, they also need to exhibit certain characteristics like being biocompatible with the biological systems of the body and they have to be able to undergo biodegradation once implanted [2]. These scaffolds are also supposed to be fabricated in such a way that they should have proper interconnected pores with proper pore size. This will require the scaffolds to have a proper three-dimensional (3D) porous structure when it is being synthesized. There is also need for the scaffolds to have an excellent surface chemistry, topography and adequate mechanical properties [3].

Freeze-drying, which is one of the most common techniques for fabricating scaffolds, was used in this study to prepare collagen (Col)/graphene oxide (GO)/hydroxyapatite (HA) nanocomposite scaffolds for bone tissue engineering. Col was selected to be used due to its great biocompatibility, degradability, and its inherent structural architecture [4]. GO was incorporated due to its distinct and extraordinary physical, chemical, and mechanical properties [5]. Studies have also shown GO-containing biomaterials to be biocompatible, non-cytotoxic, and that they allow effective adhesion and proliferation of human and mammalian cells [6]. Girao et al. [7] reported that the inclusion of GO in Col biomaterials resulted into enhanced structural properties of the materials and this depended both on the pH of the medium used and the weight ratio of GO/Col used during the fabrication. Addition of GO also enhanced osteogenic differentiation of the human mesenchymal stem cells (hMSCs) that were cultured on the GO/Col as compared to those cells cultured on pure Col scaffolds [8].

On the other hand, the incorporation of HA which is mostly calcium phosphate (CaP), has shown considerable increase in the mechanical properties of the biomaterials, a better cellular migration, proliferation and differentiation after stem cells were seeded on the scaffolds [9, 10]. The presence of CaP in Col-based scaffolds has been reported to increase the compressive modulus of up to 1.3 kPa as compared to that of pure Col scaffolds [11]. The compressive modulus has been said to be directly proportional to the concentration of CaP in the scaffolds.

Despite this demonstrated potential of GO and HA in enhancing the structural as well as functional properties of Col scaffolds, the preparation and characterisation of Col/GO/HA nanocomposite scaffolds for bone tissue engineering has not yet been thoroughly investigated. It is against this background that this study wants to investigate the preparation and characterisation of Col/GO/HA freeze-dried nanocomposite scaffolds that can have applications in bone tissue engineering. In this work, a series of 3D porous Col/GO/HA nanocomposite scaffolds were prepared through freeze-drying technique. The as-prepared Col/GO/HA porous scaffolds were characterized through scanning electron microscopy (SEM), X-ray diffraction (XRD), Fourier Transform Infra-Red (FT-IR) spectrum, mechanical test. Furthermore, the cytocompatibility and the osteogenesis ability of the rat bone marrow stem cells (rBMSCs) on the as-prepared scaffolds were evaluated in vitro.
Materials and Methods

Reagents

Lyophilised collagen (acid soluble) was purchased from Sichuan Ming Yi Biotechnology Co., Ltd (China). The modified Hummers' method [12] was used to synthesize Graphene oxide. Calcium hydroxide (Ca(OH)\(_2\)) and Sodium tripolyphosphate (Na\(_5\)P\(_3\)O\(_10\)) (Sinopharm. Chemical Reagent Co., Ltd) were of analytical grade. Hydroxyapatite was prepared within the laboratory from Calcium hydroxide and Sodium tripolyphosphate by modifying the method used by Yang et. al [13]. Healthy 12 weeks old specific pathogen free grade Wistar rats were sacrificed and BMSCs were extracted from the femur and tibia. The stem cells were then sub-cultured in complete growth medium.

Preparation of the 3D porous Col/GO/HA nanocomposite scaffolds

An aqueous acetic acid (AA) solution was prepared by diluting 0.642 mL of AA in 85 mL of deionised water. Col was dissolved in the AA aqueous solution until the solution was homogenous, then 10 mL of GO solution (0.001 mg/mL) was added. Finally, HA was added to the solution at different concentrations. The samples were then frozen at a final freezing temperature of -20°C for 24 hrs. Thereafter, the samples were lyophilized in a vacuum at -51°C for 36 hrs (12 hrs per day for 3 days) to obtain the scaffolds. In between the freeze-drying, the samples were kept frozen at -20°C.

Six samples were prepared and the concentration of Col was kept constant (0.5% wt./vol.) in all the samples. Out of the six samples, GO was added to five of them at a constant concentration (0.01% wt./vol.). Only four of the samples contained HA in them. The ratio of HA:Col in the four samples was 0.25:1, 0.5:1, 1:1 and 1.25:1, respectively. One sample only contained Col and GO and the last sample which was used as a control sample only contained Col dissolved in aqueous AA.

Characterizations of the as-prepared porous scaffolds

The microstructure of the as-prepared scaffolds was analysed using the scanning electron microscopy (SEM). The samples were cut into 0.5 cm by 0.3 cm and were fixed to an adhesive carbon stub. Then the samples were coated with gold in vacuum using an ion sputter (HITACHI E-1010) to enhance their conductivity during imaging. The microstructure of the scaffolds was imaged by a Scanning Electron Microscope (HITACHI S-4800) at a voltage of 5.0 kV to examine the surfaces of the scaffolds and the interconnection of the pores in the scaffolds. Fourier Transform Infra-Red (FTIR) spectroscopy was used to compare and analyse the characteristics of the as-prepared scaffolds. The compressive stress of the as-prepared nanocomposite scaffolds was measured using INSTRON (ElectroPuls E1000, America).
In vitro analysis

Cell viability was determined by seeding rat rBMSCs, that were harvested from Wistar rats, on the scaffolds in 24-well culture plates at a density of $2 \times 10^3$ cells/well using Dulbecco’s Modified Eagle’s Medium (DMEM) supplemented with 15% foetal bovine serum (FBS) and 1% penicillin/streptomycin. The cells were then incubated with 100 uL of MTT (0.5 mg/mL, 3-(4, 5-dimethyl-2-thiazolyl)-2,5-diphenyl-2-H-tetrazolium bromide) at 37°C for 4 hrs. Thereafter, the reagent was removed and 500 uL of DMSO was added to each well. 150 uL of DMSO solution was transferred to 96-well culture plate before evaluating the cell number in each well using an enzyme-linked immunosorbent assay.

Cell morphology was analysed by seeding rBMSCs on the scaffolds in 24-well culture plates at a density of $2 \times 10^3$ cells/cm² using growth medium. After 7 days, cells were fixed in 4% (vol/vol) paraformaldehyde for 30 min at room temperature. The cells were then dehydrated gradually by using ethanol solutions with different concentrations before the scaffolds were freeze-dried. The scaffolds were coated with gold ions in vacuum before being examined. SEM was then used to observe the cell morphology on the scaffolds.

To evaluate the osteogenic differentiation abilities of the cells on the scaffolds, rBMSCs were seeded on the scaffolds at a density of $2 \times 10^3$ cells/well using osteogenic induction medium (DMEM supplemented with 15% FBS, 1% penicillin/streptomycin, 10 nM dexamethasone, 10 mM $\beta$-glycerophosphate and 0.2 mM ascorbic acid). After 14 days, rBMSCs were then fixed with 4% (vol/vol) paraformaldehyde for 30 min at room temperature. The fixed cells were then stained with Alizarin Red Solution (2%) for 30 min at 37°C before being viewed under a light microscope. Alizarin red staining has been used extensively to demonstrate the osteogenic differentiation potential of various stem cells in bone regeneration studies [14-16].

Results

Structural characteristics of the scaffolds

The scaffold porosity was analysed and it was determined using the formula below:

$$\text{Porosity} = \frac{W_t - W_r - W_d}{W_c - W_r} \times 100\%$$

Where, $W_c$ is the weight of container filled with ethanol, $W_d$ is the weight of the dry scaffolds only, $W_t$ is the total weight of the container filled with ethanol and the dry scaffold that has been soaked in the container, and $W_r$ is the weight of the container filled with ethanol after removing the soaked scaffold [17]. Using the above formula, it was found that the porosity of the scaffolds was as follows: Col/GO/HA (HA:Col =0.25:1) had a porosity of 99.9%, Col/GO/HA (HA:Col =0.5:1) was 99.4% porous, Col/GO/HA (HA:Col=1:1) had 99.1%, Col/GO/HA
(HA:Col=1.25:1) had 98.3%, Col/GO had 99.4% and pure Col had 99.9% porosity results.

Figure 1. SEM images of as-prepared porous scaffolds. (a) Col/GO/HA (HA:Col =0.25:1); (b) Col/GO/HA (HA:Col =0.5:1); (c) Col/GO/HA (HA:Col=1:1); (d) Col/GO/HA (HA:Col =1.25:1); (e) Col/GO; and (f) pure Col.

Figure 1 above shows SEM images of as-prepared Col/GO/HA, Col/GO and pure Col scaffolds. The images show the scaffolds having macropores with diameter ranging between 30-350 μm. The scaffolds with a higher concentration of HA have a smaller number of pores with a diameter larger than 100 μm.
Figure 2. XRD patterns of as-prepared porous scaffolds. (a) Col/GO/HA (HA:Col =0.25:1); (b) Col/GO/HA (HA:Col =0.5:1); (c) Col/GO/HA (HA:Col =1:1); (d) Col/GO/HA (HA:Col =1.25:1); (e) Col/GO; and (f) pure Col.

The XRD machines was used to analyse the incorporation of HA and GO materials into the Col scaffolds. Figure 2 above shows the XRD patterns of pure Col, Col/GO, and Col/GO/HA composite scaffolds. Different peaks were recorded in the XRD patterns and some of which are characteristic peaks for the materials which were used in this study. Col/GO/HA composite scaffolds showed the characteristic peaks for HA that appeared at $2\theta = 31.7^\circ$ (for HA:Col =0.25:1), 25.8$^\circ$ & 31.7$^\circ$ (for HA:Col =0.5:1), 32.2$^\circ$ & 32.8$^\circ$ (for HA:Col =1:1), and 31.7$^\circ$ & 32.2$^\circ$ (for HA:Col =1.25:1). The characteristic peak for GO at $2\theta = 11^\circ$ was only reported in Col/GO/HA (HA:Col =1.25:1) while the characteristic peak for Col at $2\theta = 21^\circ$ was only visible in Col/GO/HA.
Figure 3. FTIR spectra of as-prepared porous scaffolds. (a) Col/GO/HA (HA:Col =0.25:1); (b) Col/GO/HA (HA:Col =0.5:1); (c) Col/GO/HA (HA:Col=1:1); (d) Col/GO/HA (HA:Col =1.25:1); (e) Col/GO; and (f) pure Col

Figure 3 above shows the FTIR spectra of pure Col, Col/GO, and Col/GO/HA composite scaffolds, revealing the absorption peaks that are characteristic to the different components that make up the scaffolds. A wide and strong absorption peak near 3430 cm$^{-1}$ is visible in all the scaffold samples. A number of peaks were also seen between 500 cm$^{-1}$ to 750 cm$^{-1}$ wavelength.

Mechanical properties of as-prepared Col/GO/HA nanocomposite scaffolds

The ability of the scaffold materials to resist compression was measured using INSTRON-5967 and the results are shown in Table 1. The results show that compressive stress at maximum load is almost directly proportional to the concentration of HA. It is worth noting that the compressive stress in Col/GO/HA (HA:Col=1:1) was lower than that of Col/GO/HA (HA:Col=0.5:1). Pure Col scaffolds had the least compressive stress of all at maximum load.

Table 1. Compressive stress of as-prepared porous scaffolds.

<table>
<thead>
<tr>
<th>Samples</th>
<th>Height of pressure plate [mm]</th>
<th>Maximum load [N]</th>
<th>Compressive stress at maximum load [MPa]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Col/GO/HA\textsuperscript{a}</td>
<td>10.00000</td>
<td>3.49194</td>
<td>0.04446</td>
</tr>
<tr>
<td>Col/GO/HA\textsuperscript{b}</td>
<td>10.00000</td>
<td>4.83388</td>
<td>0.06155</td>
</tr>
<tr>
<td>Col/GO/HA\textsuperscript{c}</td>
<td>10.00000</td>
<td>4.41889</td>
<td>0.05626</td>
</tr>
<tr>
<td>Col/GO/HA\textsuperscript{d}</td>
<td>10.00000</td>
<td>6.12309</td>
<td>0.07796</td>
</tr>
<tr>
<td>Col/GO</td>
<td>10.00000</td>
<td>4.37390</td>
<td>0.05569</td>
</tr>
<tr>
<td>Pure Col</td>
<td>10.00000</td>
<td>1.27305</td>
<td>0.01621</td>
</tr>
<tr>
<td><strong>Average value</strong></td>
<td>10.00000</td>
<td><strong>4.08579</strong></td>
<td><strong>0.05202</strong></td>
</tr>
<tr>
<td><strong>Standard variance</strong></td>
<td>0.00000</td>
<td><strong>1.62286</strong></td>
<td><strong>0.02066</strong></td>
</tr>
</tbody>
</table>

Note: a: (HA:Col =0.25:1); b: (HA:Col =0.5:1); c: (HA:Col=1:1); d: (HA:Col =1.25:1).
Figure 4. SEM images of rBMSCs cultured on as-prepared scaffolds. (a) Col/GO/HA (HA:Col =0.25:1); (b) Col/GO/HA (HA:Col =0.5:1); (c) Col/GO/HA (HA:Col =1:1); (d) Col/GO/HA (HA:Col =1.25:1); (e) Col/GO; and (f) pure Col. The magnified areas in the images show the rBMSCs and their morphology on the scaffolds.

**Cell culture**

After Day 7 of culturing rBMSCs on the Col/GO composite scaffolds using growth media, the cells were dehydrated, fixed and freeze-dried while on the scaffolds before taking the scaffolds for imaging. Figure 4 above shows the SEM images of the as-prepared scaffolds with rBMSCs cultured on them. The magnified areas in the SEM images show the rBMSCs and their morphology on the scaffolds. The rBMSCs can be seen covering almost the entire surface of the composite scaffolds.
Figure 5. MTT assay for cell viabilities of rBMSCs cultured on as-prepared porous scaffolds. When compared with pure Col scaffold, a considerable increase on cell viability was observed. The cell viability cultured on Col/GO/HA (1:1) displayed the highest OD value among the tested groups.

Figure 5 above shows the cell viabilities for rBMSCs on Col/GO/HA, Col/Go and pure Col scaffolds. The bar graphs show that the rBMSCs remained viable on the scaffolds up to Day 21. On average, the results show that the proliferation of the rBMSCs increased as the days went by. The viability only decreased slightly in all the composite scaffolds between Day 1 and Day 6. The viability was measured using the optic density (OD) readings from the graphs. The lowest OD was recorded in Col/GO/HA (0.25:1) composite scaffolds on Day 6 and the highest was recorded in Col/GO/HA (1:1) composite scaffolds on Day 18.

The osteogenic differentiation ability of the rBMSCs was analysed using the Alizarin Red staining test. Figure 6 below shows the Alizarin red staining results for the cells that were harvested from the scaffolds on which they were cultured. Osteogenesis was revealed in the rBMSCs which were differentiated for 14 days using the osteogenic differentiation media. Alizarin red was used to stain for traces of calcium deposits in the samples and the orange-red stains in the samples reveal a positive test for calcium deposits. The arrows in the images show the orange-red stained calcium deposits that indicate the osteogenic capability of the rBMSCs to form bone tissue.
Figure 6. Images of rBMSCs stained using Alizarin red after the cells were cultured for 14 days on as-prepared porous scaffolds. (a) Col/GO/HA (HA:Col =0.25:1); (b) Col/GO/HA (HA:Col =0.5:1); (c) Col/GO/HA (HA:Col=1:1); (d) Col/GO/HA (HA:Col =1.25:1); (e) Col/GO; and (f) pure Col. The arrows show the calcium deposits that indicate the osteogenic capability of the rBMSCs to form bone tissue.

Discussion

**Structural analysis of the scaffolds**

As shown in Figure 1, Col/GO/HA, Col/GO and pure Col scaffolds all exhibited an isotropic structure following freezing and lyophilisation. The application of the scaffolds and the type of cells used determine the optimal pore size for the scaffolds and according to Gibson et al. [18], the optimal size of the pores range from 20–500 µm. A larger pore size has been reported to be associated with higher final freezing temperatures in which there is a slower rate at which heat is extracted from the scaffolds thereby requiring an extended freezing time [19]. As it can also be observed from Figure 1, the number of pores with a diameter larger than 100 µm decreased as the concentration of HA increased in the samples and this can be attributed to the increased total concentration of the materials in Col/GO/HA scaffolds as compared to pure Col scaffolds. These results are consistent with the ones reported by Xia et al. [20] in
which an increase in the apatite content in the collagen-based scaffolds from 0-54 wt.% resulted into a decrease in the number of pores with a diameter of more than 100 μm.

It has been reported that scaffolds with porosity of greater than 90% are better suited for cell migration, nutrient and waste transportation as well as the vascularization throughout the scaffolds [21]. Pure Col and Col/GO/HA (HA:Col=0.25:1) scaffolds exhibited the highest porosity (99.9%) and also largest diameter of the pore sizes as compared to the rest of the scaffolds. As expected, the lowest porosity (98.3%) was recorded in Col/GO/HA (HA:Col=1.25:1) considering that studies have already reported that the addition of HA to Col-based scaffolds and increasing the total concentration of the materials used reduces the porosity of the scaffolds. Col/GO scaffolds had a porosity of 99.4% but the diameter of the pores was noticeably much smaller than that of pure Col scaffolds. On the contrary, Norahan et al. [22] found that the addition of GO had no bearing on the pore size and porosity of Col scaffolds. This difference may be due to the variations in the addition methods of GO in these two studies.

The incorporation of HA and GO into the scaffolds was analysed using XRD. From the XRD patterns in Figure 2, it can be seen that the each of the Col/GO/HA composite scaffolds showed at least one of the characteristic peaks for HA that appeared at 2θ = 25.8°, 31.7°, 32.2°, or 32.8°. This is in line with what other studies have demonstrated that HA has typical 2θ indices at 25.8°, 31.7°, 32.2°, 32.8°, and 40° [23]. This, therefore, indicates that HA was duly incorporated in the Col scaffolds. The characteristic peaks that appear broader due to the presence of HA in the samples indicate a low crystallinity and this suggests a similarity with the minerals present in natural bones [24]. Only Col/GO/HA (HA:Col =1.25:1) composite scaffolds showed the characteristic peak for GO at 2θ = 11° and also only the Col/GO/HA composite scaffolds (samples a, b, c & d) showed the characteristic peaks for Col at 2θ = 21°. The homogenous distribution of GO in the collagen solution and the formation of fully exfoliated structures may be associated with the nonappearance of regular characteristic peaks for GO [25].

A wide and strong absorption peak that corresponds to the vibrational peak of -OH group can be seen near 3430 cm⁻¹ in all the samples. The -OH functional group can either be from the HA crystals or from the water that was absorbed by the composite scaffolds. For all the Col/GO/HA scaffolds, except sample d (HA:Col =1.25:1), the peaks near 988 cm⁻¹, 1063 cm⁻¹ and 1136 cm⁻¹ are indicative of the various vibration modes of the phosphate group that is present in HA. The characteristic peaks that are associated with the carbonate group were also visible in all the scaffolds near 1449 cm⁻¹ (a), 1450 cm⁻¹ (b), 1450 cm⁻¹ (c), 1449 cm⁻¹ (d), 1452 cm⁻¹ (e) and 1451 cm⁻¹ (f). These characteristic peaks for phosphate and carbonate groups are consistent with what was reported previously [26]. Since all the scaffolds contained Col, the typical amide peaks were observed on all the scaffolds with amide I near 1649 cm⁻¹ (a), 1650 cm⁻¹ (b), 1651 cm⁻¹ (c), 1659 cm⁻¹ (d and e) and 1655 cm⁻¹ (f), amide II near 1557 cm⁻¹ (a and e), 1561 cm⁻¹ (b, c, and f) and 155 cm⁻¹ (d), amide III near 1237 cm⁻¹ (a), 1229 cm⁻¹ (b), 1235 cm⁻¹ (c), 1240 cm⁻¹ (d, e, and f). Similarly, a different study stated that FTIR spectra of collagen-based scaffolds showed characteristic amide I, II and III peaks at 1645 cm⁻¹, 1549 cm⁻¹ and 1243 cm⁻¹, respectively [27].
Mechanical properties of as-prepared Col/GO/HA nanocomposite scaffolds

From the results of compressive stress at maximum load in Table 1, it is clear that the addition of GO and HA to Col scaffolds greatly enhanced the compressive stress of the scaffolds. The compressive stress was recorded with the highest value at a ratio of HA:Col = 1.25:1 in Col/GO/HA scaffolds. This means that the compressive stress increased by almost 480% after adding HA to Col scaffolds. The deposits of HA that are tangled between the Col fibres result into increased resistance of the collagenous scaffolds to deformation [28] and also the addition of GO to Col enhances the strength of the scaffolds [22]. The addition of HA and GO to Col enhances the mechanical and degradation properties of the scaffolds since Col on its own lacks such essential properties in bone tissue engineering.

In Vitro Analysis

From The attachment, proliferation and the morphological changes of the rBMSCs that were cultured on the scaffolds were observed using SEM. Figure 4 shows the SEM images of the cells on the scaffolds. From the images, it is evident that all the scaffolds were able to support cell adhesion and proliferation. In contrast to the SEM images in Figure 1, the images in Figure 4 show rBMSCs covering almost the entire surface of the scaffolds as they had proliferated successfully. Norahan et al. [22] also reported in their study that the cells that were cultured on collagen-based scaffolds had successfully attached and covered the surface of the materials.

The viability of the rBMSCs on the as-prepared scaffolds was assessed using an MTT assay after culturing the cells for 21 days using growth media. It can be seen from the results in Figure 5 that the materials are not cytotoxic in nature. This is evident by the continued proliferation of the rBMSCs for the entire duration they were on the scaffolds. At Day 1 of incubation, the cells were viable on all the scaffolds with no much difference amongst the scaffolds except for Col/GO/HA (1:1) which showed a slightly increased cell proliferation than the other scaffolds. There was a reduction in cell proliferation on all the scaffolds at Day 6 except for pure Col which was used as the control. This suggests that the presence of GO and HA was responsible for the slight suppression of the proliferation of the rBMSCs that were seeded on the test materials. This claim is supported by several preceding studies which reported that the presence of numerous surface functional groups in GO increases intracellular reactive oxygen species levels which may somehow induce cell apoptosis [29-31]. HA has also been considered to demonstrate some levels of cytotoxicity mainly due to its intrinsic characteristics like alkalinity [14, 32].

As it can be seen from Figure 5, the suppression effect of GO and HA on the proliferation of rBMSCs was almost negligible after Day 6. From Day 6 to Day 18, cell viability increased tremendously with an optic density (OD) value of >0.5 for Col/GO/HA (HA:Col =0.5:1) and Col/GO/HA (HA:Col=1:1) at Day 14. The OD value for Col/GO/HA (HA:Col=1.25:1) was also noticeably higher at Day 14 though it was slightly below 0.5 (OD value). The highest OD value
was recorded at Day 18 where Col/GO/HA (HA:Col=1:1) scaffolds had an OD value of about 0.8. These results indicate that the addition of GO to Col scaffolds resulted into an increased cell viability at Day 18 and 21. Furthermore, these results show that HA increased the cell viability even more as compared to GO. This is evident in Figure 5 where the scaffolds that contained HA exhibited a higher OD value than the Col/GO scaffolds. In fact, the scaffolds with a higher HA concentration exhibited a better cell viability than those with a lower HA concentration. It can therefore be concluded that the increased cell viability in Col/GO/HA (HA:Col=1:1) was as a result of the presence of HA in the scaffolds rather than GO. Generally, the MTT assay results shows that the cell viability on the as-prepared scaffolds increased as the days went by.

The osteogenic differentiation results in Figure 6 indicate that the cultured rBMSCs that were harvested from the as-prepared composite scaffolds were able to form calcified tissue which was able to be stained orange-red by the alizarin red staining. These results suggest that the rBMSCs that were cultured on pure Col, Col/GO and Col/GO/HA composite scaffolds were able to successfully differentiate into calcified tissue that resembles bone tissue. In a different study, it was reported that the presence of ascorbic acid, dexamethasone and Beta-Glycerophosphate as growth factors in the osteogenic differentiation media resulted in the rBMSCs forming calcified tissue which is an indication for osteogenesis [15]. There wasn’t much difference in the calcification of the cultures in pure Col and Col/GO scaffolds which may be attributed to the very low concentration of GO that was added to the samples. On the other hand, the addition of HA resulted into an increased levels of calcification with Col/GO/HA (HA:Col =0.5:1) showing the highest levels of calcification. Other studies have reported that GO and HA synergistically increases calcium deposition in stem cells thereby accelerating the osteogenesis of these cells [14, 16, 33].

Conclusions

In summary, a series of 3D porous Col/GO/HA nanocomposite scaffolds were successfully prepared using the freeze-drying technique. The incorporation of GO and HA greatly improved the mechanical as well as the functional properties of the Col-based scaffolds. The rBMSCs were able to successfully attach and proliferate on the as-prepared scaffolds which demonstrates the biocompatibility of the scaffolds. Cell viability was found to be higher on scaffolds with a higher concentration of HA than on the other scaffolds with a lower HA concentration or even without HA at all. In vitro experiments also indicated the presence of calcium deposits after the cells were seeded on the scaffolds and cultured for 14 days using the osteogenic differentiation medium. Out of all the scaffold samples, this study demonstrated that Col/GO/HA (HA:Col =0.5:1) porous composite scaffolds exhibited better structural as well as functional characteristics to be used in bone tissue engineering applications. This, therefore, points to the need for further studies on the hemocompatibilities of these scaffolds as well as conducting animal experiments to establish the extent to which Col/GO/HA (HA:Col=0.5:1) porous composite scaffolds would be able to restore damaged bone tissue.
Abbreviations

AA: acetic acid; CaP: calcium phosphate; Col: collagen; DMEM: Dulbecco’s Modified Eagle’s Medium; DMSO: dimethyl sulfoxide; FBS: foetal bovine serum; FT-IR: Fourier Transform Infra-Red; GO: graphene oxide; HA: hydroxyapatite; hMSCs: the human mesenchymal stem cells; MTT: 3-(4, 5-dimethyl-2-thiazolyl)-2,5-diphenyl-2-H-tetrazolium bromide; OD: optic density; rBMSCs: rat bone marrow mesenchymal stem cells; SEM: scanning electron microscopy; XRD: X-ray diffraction.

References: