
Comparative study of nanohydroxyapatite microspheres for medical applications

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Abstract: This study concerns the preparation, physical, and *in vitro* characterization of two different types of hydroxyapatite (HA) microspheres, which are intended to be used as drug-delivery systems and bone-regeneration matrices. Hydroxyapatite nanoparticles (HA-1 and HA-2) were prepared using the chemical precipitation synthesis with H_3PO_4 , $\text{Ca}(\text{OH})_2$, and a surfactant, SDS (sodium dodecyl sulfate), as starting reagents. The HA powders were dispersed in a sodium alginate solution, and spherical particles were obtained by droplet extrusion coupled with ionotropic gelation in the presence of Ca^{2+} . These were subsequently sintered to produce HA-1 and HA-2 microspheres with a uniform size and interconnected micro-porosity. Both powders and microspheres were characterized using FTIR and X-ray diffraction. Moreover, SEM and mercury intrusion porosimetry were used to analyze the

microspheres, and TEM was used to analyze the powders. Results showed that pure HA and mixtures of HA/ β -TCP in the nanometer range and needlelike shape were obtained for HA-1 and HA-2 powders, respectively. Neutral Red, scanning electron microscopy and confocal microscopy were used to evaluate the behavior of osteoblastic-like MG-63 cells cultured on HA microspheres surfaces for 7 days. Results showed that good adhesion and proliferation of osteoblasts on the HA microspheres surface. Cells built bridges between adjacent microspheres, forming microspheres-cells clusters in both types of materials. © 2007 Wiley Periodicals, Inc. *J Biomed Mater Res 86A*: 483–493, 2008

Key words: hydroxyapatite; nanoparticles; microspheres; osteoblasts; bone regeneration

INTRODUCTION

Many biomaterials have been studied to repair bone defects or restore bone tissue functionality. During the last two decades, extensive research on ceramic materials, like calcium phosphates, has shown that these materials are suitable as bone substitutes due to their biocompatible, bioactive, biodegradable, and osteoconductive characteristics,^{1–5} and when implanted *in vivo*, they are nontoxic and do not induce any antigenic response.⁶

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Bone exhibits natural hydroxyapatite (HA) crystals with needlelike or rodlike shapes, well arranged within the polymeric matrix of collagen type I. These natural nanoparticles formed in physiological environment have a more dynamic response when compared with synthetic material with larger particle size.⁷ To prepare fine HA powders, many chemical processing routes have been employed, including hydrothermal reactions,^{8–10} sol-gel synthesis,^{11–13} pyrolysis of aerosols,^{14,15} and microemulsion,^{16,17} biomimetic process,^{18,19} and chemical precipitation,^{20–24} which is the most used alternative.

Recently, sodium alginate has been presented as an interesting material for biomedical applications, showing capability to be processed under mild conditions.^{25–28} Alginate, seaweed extract composed of chains of alternating α -L-guluronic acid and β -D-mannuronic acid residues, is considered by far the most widely used polymer for immobilization and

microencapsulation technologies. Sodium alginate is water soluble and may polymerize into a solid matrix. Its processing into microspheres is compatible with the presence of proteins and even bacteria or viruses. Alginate microspheres are an inexpensive, reliable and available agent for antigens encapsulation for example in several animal diseases treatment. Besides, alginate microspheres are stable at low pH and can be prepared to a wide variety of sizes.²⁹ The simple and versatile aqueous-based gel formation of sodium alginate in the presence of divalent cations such as calcium has been used to immobilize cells³⁰ for hybrid artificial organs,³¹ hemoglobin carriers,³² macromolecular delivery,^{33–35} and drug-delivery systems.³⁵

This work describes the preparation and initial characterization of novel HA microspheres intended to be used as a biomaterial for bone regeneration or as enzyme-delivery matrices. The purpose of this work is to do the morphological, chemical composition characterization, and *in vitro* study of microspheres based on synthesized nanophased HA obtained by two different procedures.

MATERIALS AND METHODS

Synthesis of nanosized hydroxyapatite

Aqueous solutions of calcium hydroxide ($\text{Ca}(\text{OH})_2$) and ortho-phosphoric acid (H_3PO_4 , 85%), both of analytical grade, were used as reactants for the preparation of two different types of HA nanoparticles. HA powder 1 (HA-1) was synthesized according to the following procedure: first, 1 L of an aqueous suspension of H_3PO_4 (0.6M) was slowly added drop by drop to a 1 L of an aqueous suspension of $\text{Ca}(\text{OH})_2$ (1M) while stirring vigorously for about 2 h at room temperature. Concentrated NaOH was added until a final pH of 10.5 was obtained. The white solution obtained was washed using deionized water and dried in oven at 80°C for 24 h. HA powder 2 (HA-2) was synthesized in the same way, with the difference of the addition of sodium dodecylsulfate addition (10 g) into the calcium solution.

Preparation of HA microspheres

Microspheres were prepared using a methodology previously described.^{32,34} Briefly, pharmaceutical-grade sodium alginate (Protanal 10/60 LS) with a high α -L-guluronic acid content (65–75%) was kindly donated by Pronova Biopolymers and used without further purification. About 1.5 g of Na-alginate was dissolved in 50 mL of distilled and deionized water until total homogenization, where upon nanophased HA powder was added to the alginate solution in a ratio of 2:8 w/w until thorough dispersion. This paste was extruded using a droplet generator device (Encapsulation Unit Var 1-J1, Nisco) into a 0.1M CaCl_2 crosslinking

solution, wherein spherical-shaped particles were instantaneously formed, and allowed to harden for 30 min. The size of the microspheres was controlled by regulating the extrusion flow rate using a single-syringe infusion pump (EW 47-900-00, Cole Palmer) and by applying a coaxial air stream. The extrusion flow rate for HA-1 was twofold the flow rate for sample HA-2. At the completion of the gelling period, microspheres were recovered and rinsed in water to remove the excess CaCl_2 . Finally, they were dried overnight in a vacuum oven at 30°C and sintered at 1100°C for 1 h.³⁶

Characterization of nanosized HA

Infrared spectroscopy

Chemical characterization was performed using Fourier transformed infrared (FTIR) spectroscopy. For this purpose, HA was reduced to powder and analyzed as KBr pellets using a PerkinElmer System 2000 spectrometer.

Transmission electron microscopy

The size and morphology of the HA powders were determined using a transmission electron microscope (TEM, Zeiss model EM 902A) at an accelerating voltage of 80 kV. A HA solution was prepared by dispersing 0.5 mg of HA in 1 mL of ethanol. This was placed in an ultrasonic bath overnight to ensure thorough dispersion. A drop of the solution, agitated for 1 min in vortex, was placed on a Formvar carbon-coated copper 400 grid and left for 1 min. Excess liquid was removed with filter paper, and the grid was allowed to dry for 10 min. TEM images were acquired and observed with Axion Camera Zeiss and AxionCs 40AC Zeiss, v 4.2.0.0 software.

X-ray diffractometry

X-ray diffraction (XRD) characterization of HA-1 and HA-2 powders was conducted using a Panalytical X'Pert PRO alpha-1 with a RTMS X'Celerator detector. It used Ni-filtered $\text{Cu K}\alpha$ radiation over the 2θ range of 10°–90° at a scan rate of 2.4°/min and with a sampling interval of 0.002° at 40 mA and 45 kV.

Characterization of nano-HA microspheres

Inverted stand optical microscope analysis

The size of the microspheres was estimated using an inverted stage microscope (Olympus PME3-ADL) with an eye piece lens equipped with a micrometer rule, with an accuracy of 10 μm . The average diameter was determined from 20 microspheres for each material.

Infrared spectroscopy

Chemical characterization was performed using FTIR spectroscopy. For this purpose, nanoHA microspheres were reduced to powder and analyzed as KBr pellets using a PerkinElmer System 2000 spectrometer.

Scanning electron microscopy analysis

The microspheres morphology was analyzed by digital imaging and scanning electron microscopy (SEM). Using a JEOL JFC-100 fine coat ion sputter device, samples were sputter coated with gold. Then, they were studied with a JEOL JSM-6301F microscope at an accelerating voltage of 5 kV.

Mercury porosimetry method

Mercury intrusion porosimetry equipment (Quantachrome Poremaster model No. 60) was used to determine the surface area, pore number fraction, and porosity from both types of microspheres. Samples were dried at room temperature; about 1.8 g dried microspheres were used in this determination and Helium was used for gas adsorption. The data reported were obtained using Quantachrome Poremaster for Windows, version 3.0.

X-ray diffractometry

X-ray diffraction (XRD) characterization of HA microspheres was conducted using a Panalytical X'Pert PRO alpha-1 with a RTMS X'Celerator detector. It used Ni-filtered Cu K α radiation over the 2θ range of 10° – 90° at a scan rate of $2.4^\circ/\text{min}$ and with a sampling interval of 0.002° at 40 mA and 45 kV.

Biological assessment

MG-63 cells were cultured at 37°C in a humidified atmosphere of 5% CO_2 in air, α -MEM (GIBCO 22561-021), 10% fetal bovine serum (FBS), 1% fungizone, and 0.5% gentamicin. For the subculture, the cell monolayer was washed twice with phosphate-buffered saline and incubated with trypsin-EDTA solution (0.05% trypsin, 0.25% EDTA) for 5 min at 37°C to detach the cells. Cells were resuspended in culture medium for reseeding. Microspheres were sterilized at 120°C for 20 min, placed on a 96 PS culture plate (14 mg of microspheres in each well, ca. 1 cm^2 of surface area), and incubated with PBS overnight. Cells were then seeded on microspheres preconditioned with α -MEM with 10% FBS, 1% fungizone, and 0.5% gentamicin. MG-63 cells were cultured (2×10^4 cells/ cm^2) up to 7 days under controlled conditions (cells seeded on 96 TCPS plates) and on the surface of both type of microspheres. The control cultures and seeded microspheres were evaluated throughout the incubation time (7 days) for cytotoxicity (Neutral Red) and observed by confocal

microscopy (CSLM) and SEM. Medium was substituted by fresh one every other day.

Cytotoxicity tests (Neutral Red)

Cytotoxicity of HA-1 and HA-2 microspheres was investigated on MG-63 osteoblast-like cells, which express a number of features characteristic of relatively immature osteoblasts³⁴ using Neutral Red assay. A 0.4% w/v solution of neutral red dye tested for cell cultures (Color Index 50040) in distilled water was prepared and stored at 4°C (stock solution). The working solution was prepared immediately before use by diluting the stock solution in complete medium to yield a final concentration of $50\text{ }\mu\text{g/mL}$. Five replicates were performed for each type of microspheres. The negative control was microspheres in culture medium in PS 96 wells culture plates. The positive control was MG-63 in culture medium in 96 wells TCPS culture plates. After incubation for 1, 3, 5, and 7 days at 37°C , the supernatant was discarded, and 0.150 mL of neutral red dye solution was added to the wells for 3 h at 37°C . The supernatant was removed, the wells washed twice with PBS, and 0.1 mL of lysing solution (50% ethanol in 1% acetic acid) was added. The color intensity of each well was read at 540 nm with reference to 620 nm, the optical densities of the replicates were averaged, and the viability read in negative control was discounted.

Cell morphology (SEM and CSLM)

For SEM, microspheres–cells constructs were washed twice with PBS and fixed with 1.5% glutaraldehyde in 0.14% cacodylate solution for 30 min at room temperature. They were dehydrated by suspending in successive ethanol solutions (50, 70, 80, 90, and 100%) for 10 min in each solution. After dehydration, the samples were transferred to hexamethyldisilazane and air dried at room temperature overnight. Finally, the microspheres–cells constructs were sputter coated with gold and examined under a JEOL JSM-6301F microscope at an accelerating voltage of 5 kV.

For CSLM, samples were washed twice with PBS, fixed in 4% v/v formaldehyde (methanol-free; Polyscience) for 15 min, permeabilized with 0.1% v/v Triton X-100 for 5 min, and incubated in 10 mg/mL bovine serum albumin and 100 $\mu\text{g/mL}$ RNase for 1 h at room temperature. F-actin filaments were stained with Alexafluor-conjugated phalloidin (Molecular Probes) for 20 min at RT in dark, and nuclei were counterstained with 10 $\mu\text{g/mL}$ propidium iodide (Sigma) for 15 min. Finally, samples were washed with PBS, mounted in Vectashield[®], and stored at 4°C in dark. CSLM images were acquired on a BioRad MRC 600 microscope.

RESULTS

The FTIR spectra of powders and microspheres using HA-1 and HA-2 materials are represented in Figure 1. It may be seen that the crystallinity

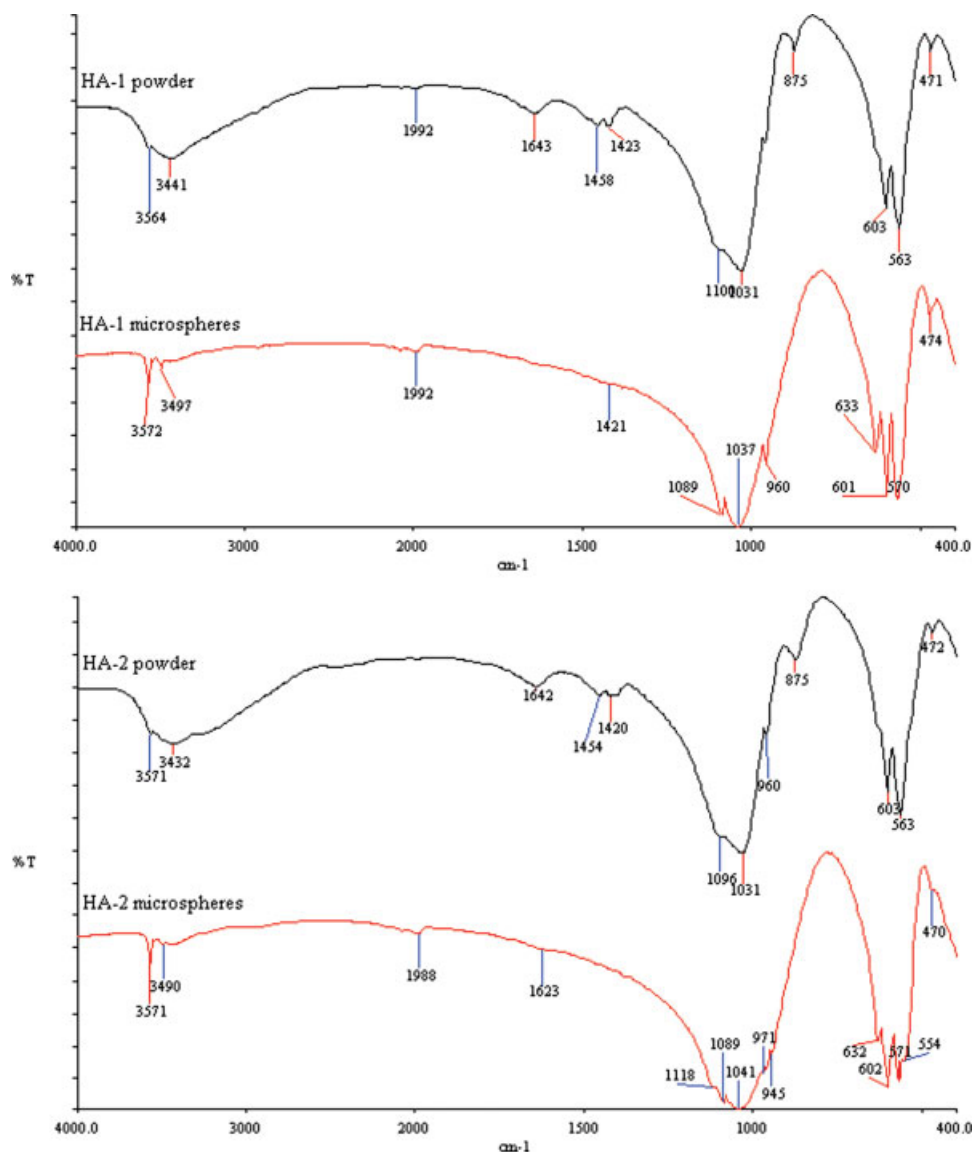


Figure 1. FTIR spectra of HA-1 and HA-2 samples (powders and microspheres). [Color figure can be viewed in the online issue, which is available at www.interscience.wiley.com.]

increased from HA-1 powder to HA-1 microspheres, especially on the phosphate bands at $560\text{--}633\text{ cm}^{-1}$ and around 1030 cm^{-1} . Occluded water gives rise to a band at $1420\text{--}1650\text{ cm}^{-1}$ observed in the HA-1 powder and vanished after the sintering process for HA-1 microspheres. The presence of the CO_3^{2-} group is observed in the HA-1 powder and lost in the spectra of microspheres, which suggests that CO_3^{2-} could be not into the crystalline network but on the surface of the material. This last sample presented higher crystallinity than the original powder material. Both HA-1 and HA-2 materials can be easily recognized as nonstoichiometric HAs. In the spectra of the HA-2 powder, a bands appears at $472\text{ (v}_2\text{)}$, $563\text{ (v}_4\text{)}$, 605 , and $960\text{ (v}_1\text{)}\text{ cm}^{-1}$ that correspond to the PO_4^{3-} group. On the other hand, bands are present at 1642 , $1454\text{ (v}_3\text{)}$, 1420 , and 875 cm^{-1} due to the presence of

carbonate group, which is missing after the sintering process similarly to HA-1 materials. The O—H stretching OH— is observed in the band at 3571 cm^{-1} , and the O—H stretch H_2O is shown in the band 3432 cm^{-1} .

HA-2 microspheres are characterized by sharp, well-resolved bands at 3571 , 602 , and 571 cm^{-1} . The most intense feature in the IR spectra is the 1041 cm^{-1} band with some shoulders at 1118 , 971 , and 945 cm^{-1} , suggesting the presence of two phases; HA and β -TCP. The band at 1089 cm^{-1} corresponds to P—O stretching vibrations. The broad bands at 3497 and 1992 cm^{-1} is due to the presence of both water and structural hydroxyl groups. The bands at 3572 and 633 cm^{-1} have been assigned to stretching and bending vibrations of the structural hydroxyl group.

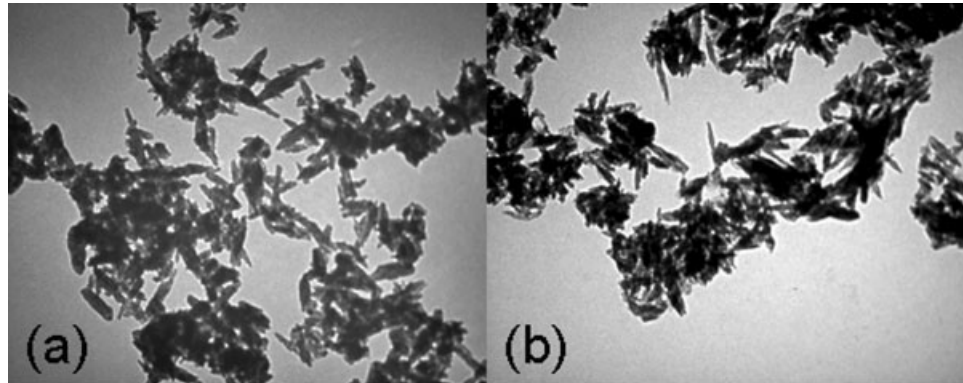


Figure 2. TEM images of HA-1 (a) $\times 20,000$ and HA-2 (b) $\times 20,000$ powders.

Figure 2 shows the morphology of the nanoparticles after the sintering process, of both materials, that generates the microspheres. From TEM images in Figure 2(a,b), some agglomerates of HA and the needlelike shape of the particles in both materials may be seen. HA-1 and HA-2 samples present smaller agglomerates wherein the nanometer size (less than 100 nm in length) of the HA particles is observed.

The alginate/HA beads were formed by dropping the paste into an aqueous solution of calcium ions. During the sintering process, the alginate was burned off and the HA particles became associated, maintaining the original spherical shape. The aver-

age diameter of HA microspheres varied depending on the type of HA used, thus, for HA-1 samples, it was $471 \pm 23 \mu\text{m}$ ($n = 20$); whereas for the HA-2 samples, it was $407 \pm 42 \mu\text{m}$ ($n = 20$). HA-2 microspheres shrank to about 19% of their original size after the sintering process, whereas the shrinkage for HA-1 microspheres was of about 28%.

SEM images of the microspheres are presented in Figure 3. After the sintering process, their size decreases due to alginate burn-off, even though their original shape is maintained. HA microspheres are depicted in Figure 3(a,b) illustrating their spherical shape as well as their uniform size. Higher surface roughness is evidenced in HA-2 microspheres when

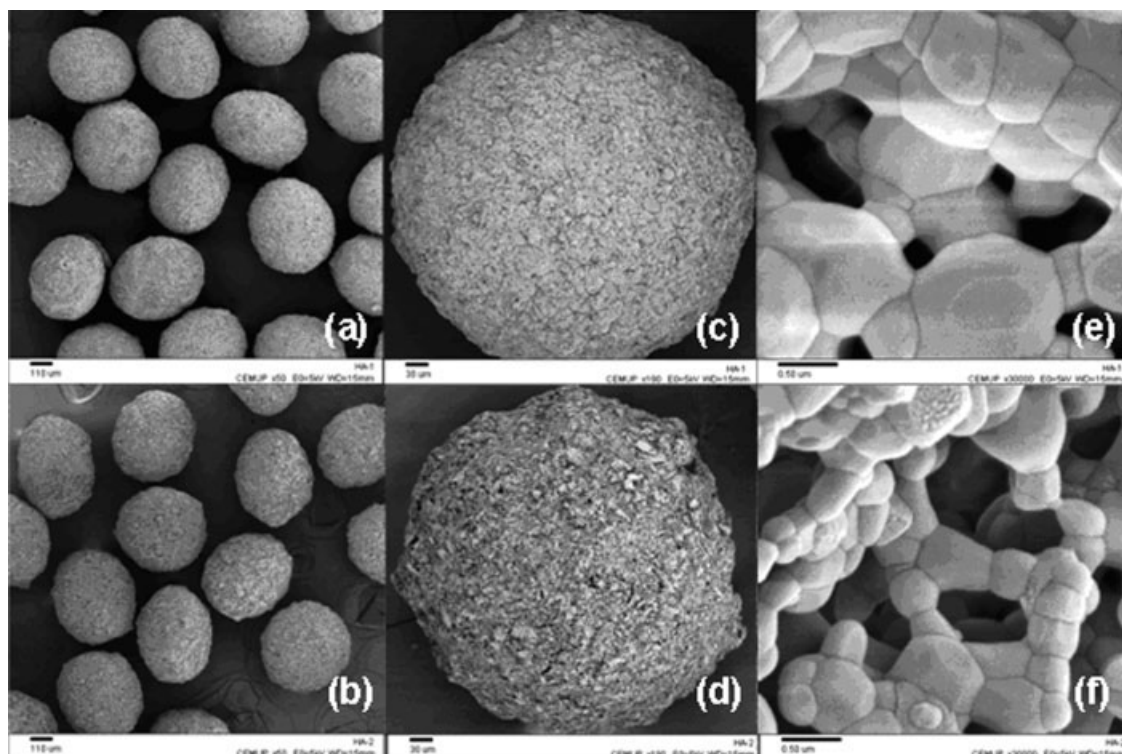


Figure 3. SEM micrographs of HA-1 (a,c,e) and HA-2 (b,d,f) microspheres: (a,b) $\times 50$; (c,d) $\times 190$; and (e,f) $\times 30,000$.

TABLE I
Mercury Porosimetry

Determinations/Samples	HA-1	HA-2
Diameter average (μm)	0.5311	0.8211
Pore volume (cc/g)	0.1507	0.1876
Pore surface area (m^2/g)	0.4958	0.5444
Pore number fraction	0.00196	0.00336
Total surface area (m^2/g)	1.9308	1.4023
Apparent density (g/cc)	1.7653	1.6726
Total intraparticle porosity (%)	45.063	47.571
Theoretical porosity (%)	45.052	45.871

compared with HA-1 [Fig. 3(c,d)]. Interconnectivity seems to be higher in HA-2 than in HA-1 microspheres. This can be due to the thickness of the walls that surround pores [Fig. 3(e,f)].

Data obtained from mercury intrusion porosimetry analysis, which are in line with the above results (Table I), do not show significant differences between both types of microspheres related to the pore surface area, apparent density, or total intraparticle porosity. Porosity does not appear to be a significant HA-2 feature in comparison with HA-1 microspheres, however, evident differences about diameter, pore volume, and pore fraction number are found, all being higher for HA-2.

XRD patterns of all samples are shown in Figure 4. HA-1 powder is represented by the pattern in Figure 4(a) and some undefined humps can be seen, because of the nonsintered state. Figure 4(a,c)

presents many similarities even though after the sintering process they revealed the presence of different phases. The pattern in Figure 4(b) shows clearly the peaks corresponding to pure monoclinic HA, however, Figure 4(d) shows a mixture of monoclinic HA and β -TCP.

In this work, the cytotoxicity tests of HA microspheres were performed using human MG-3 osteoblast-like cells line. Results presented in Figure 5 showed that there is continued growth during the culture time using both HA-1 and HA-2 microspheres. Cells seeded on TCPS, used as a control, showed the same trend, but with a higher growth rate than both microspheres. The control presented less number of viable cells in the first day probably due to the less available surface area compared to the microspheres. The HA-1 microspheres presented a slight increment in the number of viable cells in comparison with HA-2 microspheres at the end of the culture, and higher growth rate as well.

SEM images of MG-63 osteoblast cells cultured for 7 days on the surface of HA-1 and HA-2 microspheres based on nanosized HA are presented in Figure 6. The surface of both HA-1 and HA-2 microspheres is almost completely covered by layers of cuboidal cells, which became denser along the entire time of culture, forming confluent monolayers. Cells built bridges between neighboring microspheres (HA-1 and HA-2 microspheres) forming microspheres–cells clusters (see Fig. 7). CLSM images in Figure 8 show HA-1 and HA-2 microspheres uni-

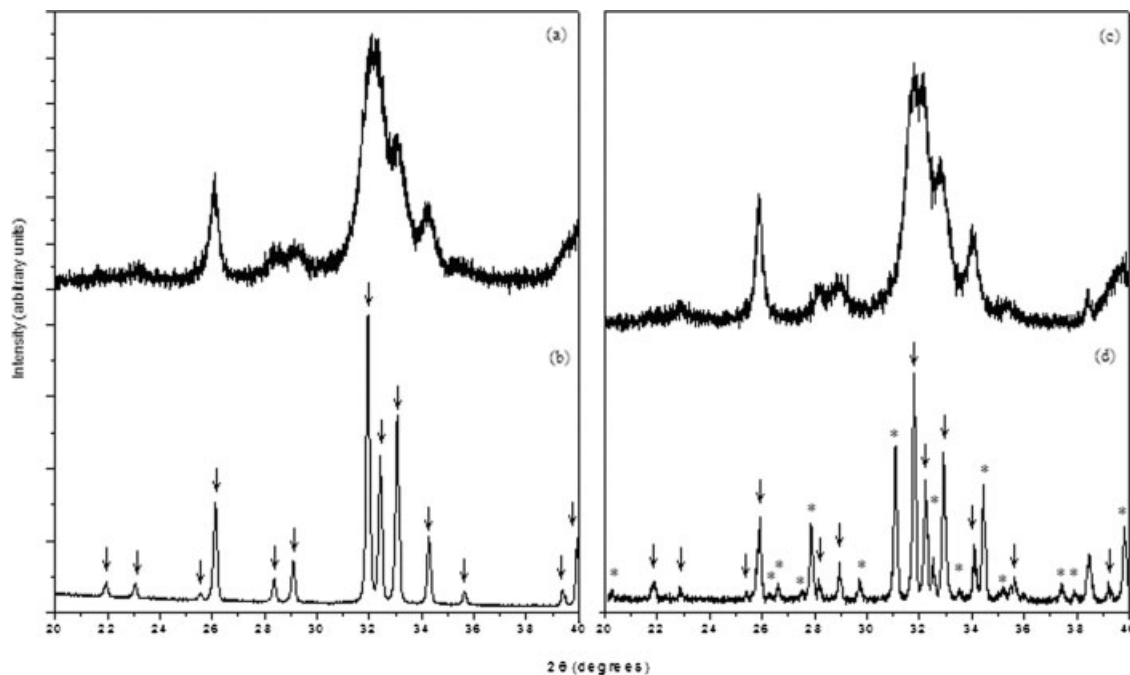


Figure 4. X-ray diffraction of powders and microspheres of HA-1 and HA-2. (a) HA-1 powder, (b) HA-1 microspheres, (c) HA-2 powder, and (d) HA-2 microspheres. 6: monoclinic HA, *: β -TCP.

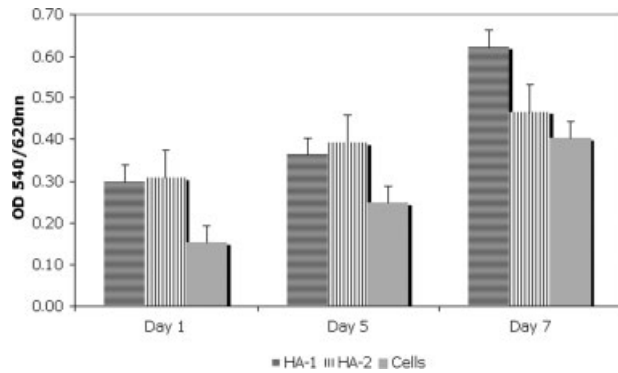


Figure 5. MG-63 cells culture on microspheres based on hydroxyapatite nanosized.

formly colonized by osteoblasts cells after 5 days of culture.

Statistical analysis

Data are presented as mean $\pm$ standard deviation ($n = 5$), and they were analyzed using the Student's t test for independent samples. Differences between groups were considered statistically different when $p < 0.05$.

DISCUSSION

In this work, nanoHA microspheres synthesis was carried out, using two different types of nanosized HA, and a subsequent characterization at morphological and physical level was performed. HA microspheres were prepared using the droplet extrusion process combined with ionotropic gel formation in the presence of calcium ions. The methodology used to prepare HA microspheres has a remarkable advantage as it allows the preparation of spherical-shaped particles with an adequate and uniform size.^{34,36–39}

The microspheres described in this work are composed of two types of HA nanoparticles, both processed by chemical precipitation, and differing in the SDS addition. The synthesis of HA can take place through a variety of processes. One such method is the wet chemical route, wherein HA is precipitated from either an acid-based reaction or a reaction between various salts. Among the different synthesis methods, the wet or precipitation method of synthesizing HA has proved to be popular and for this reason this preparation method has been extensively published. The chemical precipitation method involves the addition of phosphate ions to a solution

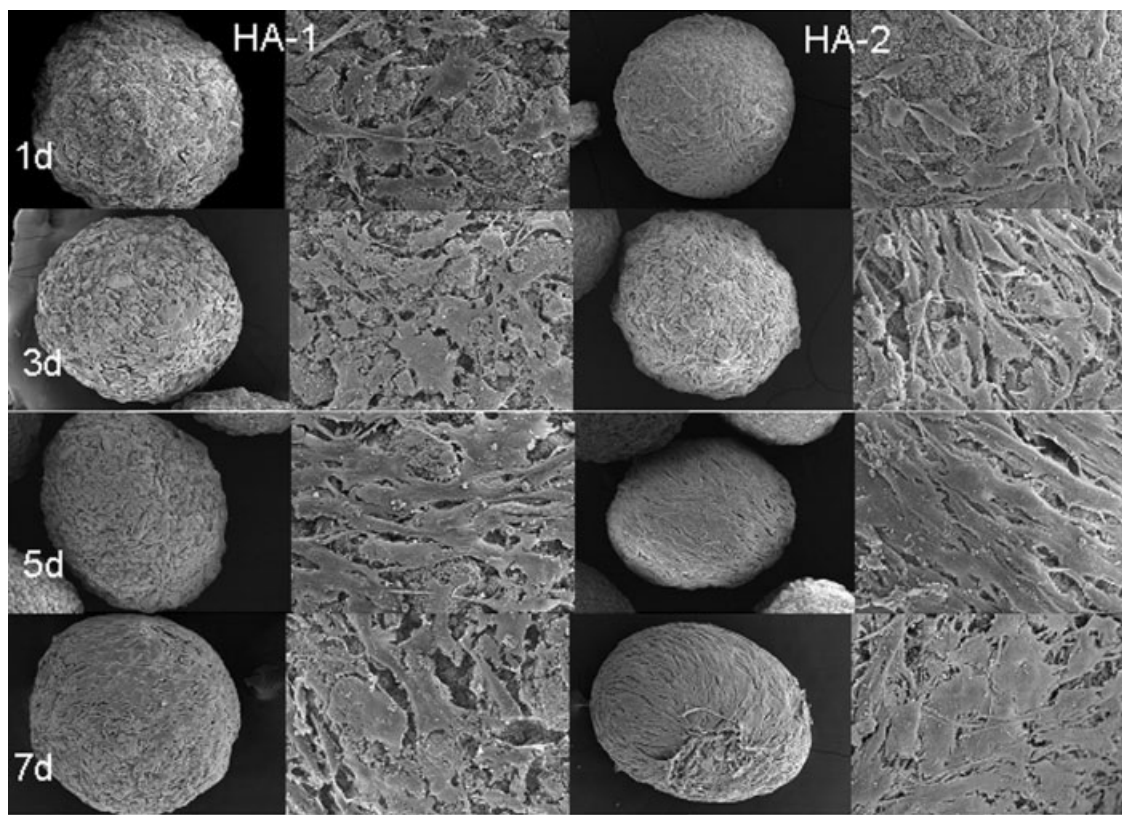


Figure 6. SEM micrographs of 1 to 7 days of MG-63 cells culture on microsphere HA-1 and HA-2 ($\times 180$ and $\times 1,000$).

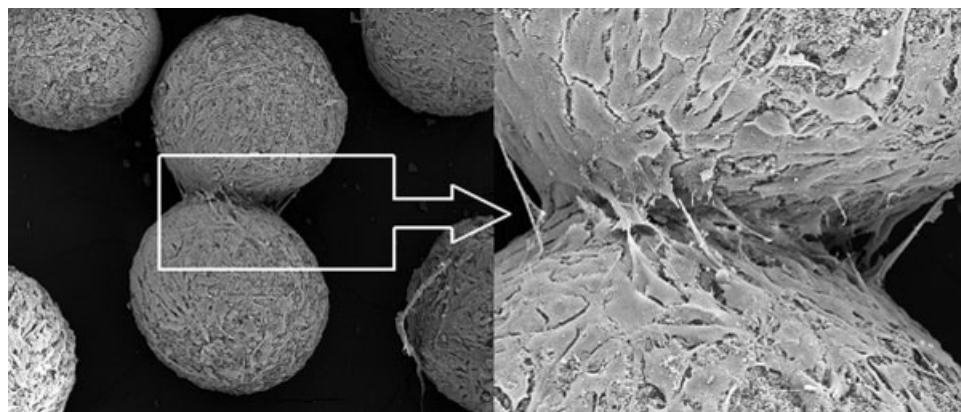


Figure 7. SEM micrographs (left: $\times 110$ and right: $\times 1,000$). Notice that osteoblast cells built bridges between neighbor microspheres. This image corresponds to HA-1 construct after 3 days.

of calcium ions, thereby precipitating HA as well as other calcium phosphates such as β -tricalcium phosphate (β -TCP).

Many chemical processing routes have been employed to prepare fine and agglomerate-free ceramic HA. For that reason, several dispersants have been successfully used to prepare HA particles^{40,41} with smaller size and higher surface area.⁴² In this work, a surfactant (SDS) assisting the chemical precipitation method is found to be able to promote the formation of nanosized HA crystal and at the same time control the agglomeration of particles.

Sodium dodecylsulfate (SDS- $C_{12}H_{25}NaO_4S$) is an ionic surfactant and probably the most researched anionic surfactant compound. The molecule has a tail of 12 carbon atoms that attached to a sulfate group, giving the molecule amphiphilic properties. In this study, SDS could have worked by disrupting bonds interparticles, thereby deagglomerating them.

Also, anions of SDS could effectively impart a negative charge on the particles and create an electrostatic repulsion creating more isolate particles. This same fact could explain the presence of two different phases in the HA-2 material based on SDS: the electrostatic repulsion could have limited the complete reaction between reagents and favor the production of $C_{10}(PO_4)_6(OH)_2$ and β - $Ca_3(PO_4)_2$. Furthermore, the presence of SDS into the calcium phosphates preparation could have filled some spaces that after the burn-out process were empty, creating higher porosity and interconnectivity in HA-2 more than in HA-1 microspheres [Fig. 3(e,f)].

Even though the roll played by SDS in the production of HA-2 material is still unknown, it is not part of the aim of this work and it will need further studies.

The characterization analyses included FTIR, TEM, SEM, mercury porosimetry, and XRD and *in vitro*

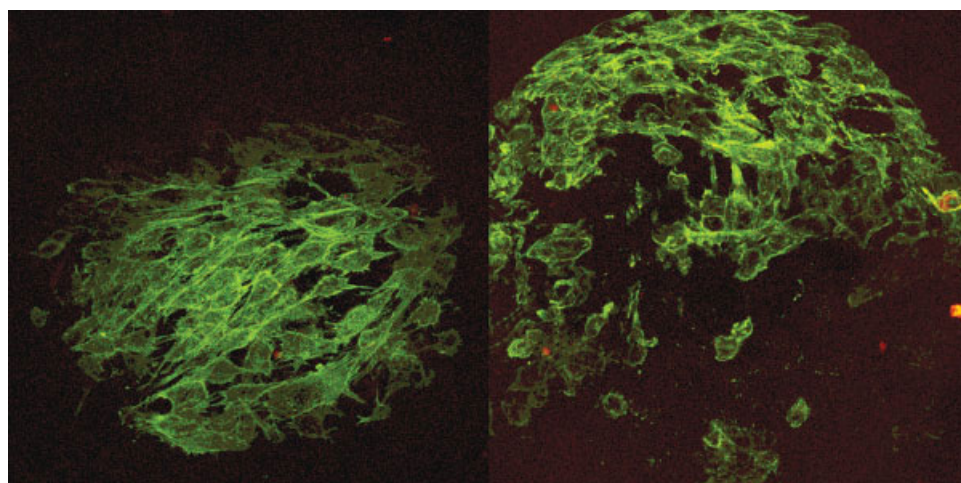


Figure 8. CLSM images of HA-1 (left: $\times 100$) and HA-2 (right: $\times 100$) microspheres after 5 days of MG-63. [Color figure can be viewed in the online issue, which is available at www.interscience.wiley.com.]

tests. FTIR analysis indicated that the characteristic spectra bands of both ceramics, HA-1 and HA-2 powders, are maintained in the microspheres, suggesting the sodium alginate did not induce subsequent modifications in the ceramics structures, for instance, the addition of CO_3^{2-} group, may be due to alginate, in the HA lattice, probably due to the fast sintering cycle, which provides higher porosity in comparison with longer cycles, or it may also derive from the contact with the atmosphere during HA processing. Therefore, alginate may have played two roles in the microspheres preparation: as a vehicle to form spherical-shape particles and as pores inducing agent in the sintering process, all of this without any modification on HA. It is important to point out the multiple-binding sites on the microsphere surface that could allow the attachment of certain macromolecules such as enzymes or antibiotics, among others. This feature may enhance its role as a drug-delivery system component to be used in biomedical applications.

Attractive advantages such as avoiding the use of organic solvents of difficult removal, the manufacturing spherical particles with uniform size, and the possibility of producing particles with different porosities^{28,36} are some characteristics related to microspheres prepared using the method referred here. Besides, in this work, nanophasic HA was used to prepare microspheres what caused a diminution of diameter and therefore a lower surface area; in contrast to previous studies where micrometric commercial HA and calcium-titanium-phosphate (CTP) were used as raw materials.³⁴

The use of nanostructured HA in the microspheres introduced a new question mark about how the bone cells would behave when in contact with this new type of microspheres, since the existing literature related to microspheres based on nanosized particles is scarce. Even though, recent works have indicated the benefits of using nanoHA in several biomaterials different to microspheres. To follow some few examples, adhesion of osteoblasts on non-functionalized with RGD nanoparticulate calcium phosphate was similar to the conventional HA functionalized with RGD⁴³; fibrous scaffolds fabricated by electrospinning from polycaprolactone (PCL) solutions containing nanoparticles of HA promoted much better adhesion and proliferation of cells than the corresponding TCPS⁴⁴; biocompatibility testing of nanoparticles dispersed within a viscous polysaccharide (dextran sulfate) porous sponge showed retention of human bone marrow stromal cells viability, production of extracellular matrix, and alkaline phosphatase expression⁴⁵; investigations about the effect of nanometer HA on the proliferation and osteogenic differentiation of periodontal ligament cells showed that nanoHA has ability to promote the

activity of osteogenetic differentiation for periodontal ligament cells when compared with the dense HA.⁴⁶

The cell culture of fMG-63 tests carried out in this work aimed at analyzing possible differences in behavior resulting from the significant changes in HA crystallites size, and at comparing the two types of material, HA-1 and HA-2 as substrates for cells.

The present studies suggest that the features such as composition and roughness can modulate osteoblast responses to these scaffolds *in vitro*, including cellular proliferation, attachment, and three-dimensional structure of the forming osteoblast multilayer network. Differences in composition (nanophasic HA and HA/ β TCP) and evident roughness (see Fig. 3) higher in HA-2 than HA-1 are the most relevant characteristics, since pore diameter or porosity do not present significant differences.

After 7 days of culturing, the distribution of cells over the HA microspheres appeared similar, suggesting that the cells on both HA-1 and HA-2 were proliferating with cuboidal and well-attached morphology colonizing uniformly both type of microspheres, even though HA-1 microspheres presented a slight increment in the number of viable cells in comparison with HA-2 microspheres at the end of the culture as well as higher growth rate.

Albeit, several authors^{47–49} have shown that surface roughness is an important parameter in the basic biologic responses improving cell attachment and proliferation; in this study, HA-1 microspheres presented slight higher cell viability than HA-1, whereas HA-2 microspheres presented higher roughness than HA-1. This fact could be related to the composition (HA but HA/ β -TCP) revealing that this, the chemical composition, would have higher influence than the roughness on the cellular response.

CONCLUSIONS

Preparation and initial characterization of two different types of HA nanosized microspheres (HA-1 and HA-2) intended to be used as drug-delivery systems, and bone regeneration matrixes were studied in this work. SEM images showed that MG-63 cells, cultured for 7 days, attached and spread through the available surface of these two types of porous microspheres with some orientation, exhibiting adequate osteoblasts morphology. Neutral Red assay results revealed higher cell viability in the HA-1 than in HA-2 microspheres.

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