

## **Porous calcium phosphate glass microspheres for orthobiologic applications**

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### **Abstract**

Orthobiologics is a rapidly advancing field utilising cell-based therapies and biomaterials to enable the body to repair and regenerate musculoskeletal tissues. This paper reports on a cost-effective flame spheroidisation process for production of novel porous glass microspheres from calcium phosphate based glasses to encapsulate and deliver stem cells. Careful selection of the glass and pore forming agent, along with a manufacturing method with the required processing window enabled the production of highly porous glass microspheres via a single-stage manufacturing process. The morphological and physical characterisation revealed porous microspheres with tailored surface and interconnected porosity (up to  $76\pm 5\%$ ) with average pore size of  $55\pm 8\ \mu\text{m}$  and surface areas ranging from  $0.34$  to  $0.9\ \text{m}^2\text{g}^{-1}$ . Furthermore, simple alteration of the processing parameters produced microspheres with alternate unique morphologies, such as with solid cores and surface porosity only. The tuneable porosity enabled control over their surface area, degradation profiles and hence ion release rates. Furthermore, biocompatibility of the microspheres was assessed using human mesenchymal stem cells (hMSCs) via direct cell culture experiments and analysis confirmed that they had migrated to within the centre of the microspheres. The novel microspheres developed have huge potential for tissue engineering and regenerative medicine applications.

**Keywords:** *Calcium phosphate glass, porous microspheres, stem cells.*

## 1. Introduction

Orthobiologics is a rapidly advancing field which utilises cell-based therapies and biomaterials to promote healing. The development of orthobiologic solutions for repairing musculoskeletal disorders has provided clinicians alternatives to mechanical stabilisation, which has been the focus of orthopaedic procedures for many years. Orthobiologics have been utilised in a variety of different areas of medicine and their applications continue to expand [1] which is enabling orthobiologics to emerge as the frontline for disease treatment and has the potential to revolutionise medicine [2].

The advantages of using synthetic biomaterials over autograft and allograft bone materials are associated with their scale-up manufacturing and long term preservation in order to supply on-demand [3]. It was discovered that bioactive glasses could be used to stimulate osteogenesis, thereby leading to the concept of tissue regeneration. It has also since been discovered that the dissolution ion products from these glasses could also provide signals to the cells [4].

The three main types of bioactive glasses investigated for biomedical applications include silicates, borates and phosphate based glasses. Silicate based bioactive glasses are based on  $\text{SiO}_2$  (as the glass network former) and include other modifying oxides such as  $\text{Na}_2\text{O}$ ,  $\text{CaO}$  and  $\text{P}_2\text{O}_5$  (and were initially developed by the late Professor Larry Hench and commercialised as 45S5 Bioglass<sup>TM</sup> [5]). They have to a great extent been applied for use in hard tissue repair [5-7] and more recently have been added to toothpaste (Sensodyne®, Novamin<sup>TM</sup>, BioMin). Borate based glasses (where,  $\text{B}_2\text{O}_3$  is the glass network former) degrade faster than silicate-based glasses and have also been investigated for biomedical applications [8]. Huang *et al.* [8] investigated the formulation  $\text{B}_2\text{O}_3$  46.1 -  $\text{CaO}$  26.9 -  $\text{Na}_2\text{O}$  24.4 -  $\text{P}_2\text{O}_5$  2.6 (in mol%) and found that they could convert this glass into apatite following immersion within a dilute (0.02 M)  $\text{K}_2\text{HPO}_4$  solution. Phosphate based glasses are based on  $\text{P}_2\text{O}_5$  (as the glass network former) and have been widely explored using  $\text{Na}_2\text{O}$  and  $\text{CaO}$  including many other modifying oxides such as  $\text{CuO}$  [9],  $\text{Ag}_2\text{O}$  [10],  $\text{TiO}_2$  [11],  $\text{SrO}$  [12],  $\text{MgO}$  [13] to provide enhanced properties or biological responses [14-18]. The main advantage of phosphate glasses over the silicate bioglasses and bioceramic materials, include their fully degradable and controllable resorption profiles (which can range from days/weeks/to many months), by simply tailoring the glass formulations [19, 20].

Microspheres (i.e. spherical particles) exhibit greater advantages over irregular-shaped particles such as improved flow properties, which would be particularly beneficial for biomedical applications (for example, by enabling improved delivery via minimally invasive surgical injection procedures [21]). Microparticles with hollow and/or porous morphologies have become extremely important in many scientific fields due to their advantageous functional properties. For example, hollow microspheres have been exploited as additives for adhesives and as fillers in construction composites due to their durability and light weight [22], whereas porous microparticles have been widely investigated for applications varying from stem cell research [23, 24], drug delivery [25], tissue engineering [26], separation sciences [27], supercapacitors [28] and energy storage [29]. More recently, hollow glass microspheres with porous walls (with porosity on the angstrom level) have been developed and are being investigated to store liquids and gases within their hollow interior with the aim to release them on demand [30]. However, the development of highly porous glass microspheres via a simple manufacturing process, with controlled surface porosity levels and with interconnected porosity (throughout the entire microsphere), could have the potential to totally revolutionise the microsphere industry by enabling many other applications for these materials.

Manufacturing processes for porous materials depend heavily on the type of material used. For example, techniques utilised for making porous glass and/or ceramic scaffolds commonly use methods such as incorporation of a removable space holder [31], polymer foam replication [32], sol-gel [33], gel cast foaming [34] and solid freeform methods (i.e. 3D printing) [35]. These manufacturing processes involve multiple processing steps which can be laborious, time consuming and energy intensive. The dissolution and thermal decomposition of space holders and sacrificial polymer templates can also result in residual contaminants which not only affects further processing but also the mechanical and physical properties of the resultant porous material. In the case of solid freeform fabrication techniques (3D printing), maintaining the viscosity of the ink (glass particles along with the polymer binder) is critical and can be time consuming requiring several steps of printing, drying and sintering [35].

Processes utilised for manufacturing polymer based porous microspheres have mostly been via emulsion-solvent evaporation, spray drying and phase separation techniques [25, 36, 37]. Whilst, ceramic microspheres have commonly been fabricated via gelation, emulsification and precipitation processes [38, 39] and solid (non-porous) and hollow glass microspheres have been fabricated via sol-gel [40] and flame spheroidisation [41] processes. However, when considering the manufacture of porous glass microspheres with fully controlled surface and

fully interconnected porosity, the main challenges have included high production costs alongside lengthy and/or complex manufacturing processes, scale-up limitations and control over porosity size, interconnected porosity and morphology.

In this paper, the authors report a new and cost-effective process for manufacturing highly porous calcium phosphate (CaP) glass microspheres with tailorable surface and fully interconnected porosity (up to 76%) within each microsphere produced, via a single-stage flame spheroidisation process. Preliminary biological analyses were performed via direct cell culture studies using human mesenchymal stem cells including the suitability of these microspheres to support cell adhesion and growth.

## **2. Materials and methods**

### *2.1 Manufacture of solid and porous calcium phosphate glass microspheres*

Calcium phosphate glass in the system  $40\text{P}_2\text{O}_5 \cdot 16\text{CaO} \cdot 24\text{MgO} \cdot 20\text{Na}_2\text{O}$  (mol% denoted as P40) was prepared using  $\text{NaH}_2\text{PO}_4$ ,  $\text{CaHPO}_4$ ,  $\text{MgHPO}_4$  and  $\text{P}_2\text{O}_5$  (Sigma-Aldrich, UK) as starting materials. These precursors were placed into a 100 ml volume Pt/5% Au crucible and dried at  $350^\circ\text{C}$  for 30 minutes in a furnace. The sample was then melted at  $1150^\circ\text{C}$  for 90 minutes. The molten glass was then quenched onto a steel plate and allowed to cool to room temperature.

Once cooled, the glass was ground into microparticles using a Retsch PM100 milling machine and sieved into varying size ranges spanning  $63\text{--}300\ \mu\text{m}$ . The solid non-porous microspheres were produced using the flame spheroidisation method which utilised an oxy/acetylene flame spray gun (MK 74, Metallisation Ltd, UK). Porous glass microspheres were produced by introducing porogens to the glass feed, which were then fed simultaneously into the thermal spray gun. Post manufacture, the microspheres were collected from cooling trays and were then subjected to an acid wash step by gently stirring them in acetic acid (5M) for 2 min followed by washing using deionised water (for 5 min) and then dried in an oven at  $50^\circ\text{C}$  for 24 h.

### *2.2 Characterisation methods*

#### *2.2.1 Scanning electron microscopy (SEM) and energy dispersive X-ray (EDX) analysis*

The surface and cross-sectional morphology was qualitatively examined using scanning electron microscopy (SEM - Philips XL30, FEI, USA) operated at 20kV. Cross sections were achieved by embedding the microspheres in a cold set epoxy resin and polished with SiC paper

followed by a diamond cloth. Quantitative compositional analysis of the CaP microspheres was carried out using an energy dispersive X-ray (EDX, INCA Oxford Instruments) detector at an accelerating voltage of 20 kV and a working distance of 10 mm. All samples were carbon coated using an evaporation coater (Edwards coating System E306A) in order to avoid image distortion due to charging.

### *2.2.2 Mercury porosimetry and BET surface area analysis*

The porosity of the microspheres was investigated using mercury intrusion porosimetry (Micromeritics Autopore IV 9500). A 5cc powder penetrometer (Micromeritics) with 1cc intrusion volume was used for all samples. An empty penetrometer test was also conducted as a blank before running the samples.

The surface area was determined via gas adsorption using the BET technique (Micromeritics Gemini 2360) where approximately 0.5 g of the microspheres were initially degassed under nitrogen at 110°C for at least 2 h before performing multipoint BET analysis.

### *2.2.3 Degradation and ion release studies*

Degradation analyses of the microspheres was evaluated by means of mass loss measurements in ultra-purified water (Milli-Q) (1% w/v) at 37°C over 28 days (the medium was refreshed every 2 days). The degraded microspheres were removed from the medium at various time points (1, 3, 7, 14, 21, and 28 days) and placed onto tissue paper to blot dry then weighed immediately.

$$\text{Mass Loss (\%)} = \frac{M_0 - M_t}{M_0} \quad (1)$$

Where  $M_0$  is the initial mass and  $M_t$  is the mass at test time point t.

Ion release profiles of the microspheres was conducted in Dulbecco's Modified Eagle Medium (DMEM, Sigma-Aldrich, UK) cell culture media (1% w/v) over 14 days (the media was again refreshed every 2 days) using inductively coupled plasma mass spectrometry (ICP-MS, Thermo-Fisher iCAP-Q model).

### *2.2.4 Cell culture study*

#### *2.2.4.1 Preparation of CaP microspheres extract*

Porous CaP microspheres were sterilised through two 15-min wash steps using 70% EtOH. After complete evaporation of 70% EtOH at room temperature in sterile conditions, porous microspheres were soaked in standard cell culture (SC) medium (low glucose DMEM (Gibco) supplemented with 10% foetal calf serum (Gibco), 1% penicillin and streptomycin, 1% L-Glutamine, 1% of non-essential amino acids) at a final concentration of 100 mg ml<sup>-1</sup> at 37°C and 5% CO<sub>2</sub>. After 24 h, the conditioned medium containing the microsphere ion extracts was collected, filtered using 0.22 µm syringe filters and used to prepare three serial dilutions with 1:10 ratio using SC medium.

#### 2.2.4.2 MTT assay

The evaluation of cytotoxicity was performed using a “Cell Growth Determination Kit, MTT based” (Sigma Adrich, UK); for this purpose, 1x10<sup>4</sup> immortalised human bone marrow-derived mesenchymal stem cells (hMSCs) were seeded in 96-well plate in 100 µl of SC medium. After 24 h, SC medium was replaced with ion extract conditioned medium or the 3 serial dilutions prepared as shown in section 2.2.4.1. Moreover, SC medium alone or supplemented with 5% of DMSO were included as controls. After 24 h, cells were washed once with PBS and incubated during 3 h at 37°C and 5% of CO<sub>2</sub> with a SC medium supplemented with 10% of MTT solution. After incubation, media was aspirated and 100 µl of DMSO was added to each well and incubated for 15 min in gentle shaking to facilitate the formazan salts dissolution. Absorbance was read using a plate reader at 570 nm, setting 650 nm as reference. The cell viability percent was calculated in comparison to SC medium control using the following formula:

$$Viability (\%) = \frac{OD570s}{OD570c} \quad (2)$$

Where, *OD570s* and *OD570c* are the optical densities measured in the samples of interest and in the control, respectively.

#### 2.2.4.3 Direct seeding of cells on porous CaP microspheres

For direct seeding of cells onto the porous CaP microspheres, 2x10<sup>4</sup> immortalized GFP-labelled human bone marrow-derived mesenchymal stem cells (hMSCs) were seeded on 10 mg of sterile microspheres into low-adherent 48-well plates previously coated with 1% (w/v) solution of poly(2-hydroxyethyl methacrylate) (poly-HEMA) and Ethanol 95% in SC medium. Cells were cultured by 14 days at 37°C and 5% CO<sub>2</sub>. The media was refreshed every 48 h.

#### 2.2.4.4 Presto Blue assay

Metabolic activity of cells seeded on porous CaP glass microspheres was analysed using Presto Blue reagent at day 2, day 5 and day 12 according to the manufacturer's indications. Briefly, cells were incubated with a solution of SC medium supplemented with 10% of Presto Blue at 37°C for 40 min. The fluorescence was then measured at 560 nm and 590 nm as excitation and emission wavelengths using a plate reader (Tecan infinite 200).

#### 2.2.4.5 Cell staining and imaging

Cell imaging was performed at day 14 in paraformaldehyde-fixed cells. For Environmental Scanning Electron Microscopy (ESEM), post fixed cells were washed twice with distilled water and analysed using a FEI Quanta 650 ESEM microscope.

For confocal laser scanning microscopy analysis, cell nuclei were counterstained with 10  $\mu\text{g ml}^{-1}$  of Hoechst 33258 for 10 min and cytoskeletal actin fibres were visualised using Vectashield mounting medium containing TRITC-Phalloidin (Vector Laboratories, UK). Confocal laser scanning microscopy was carried out on a Zeiss LSM 880 microscope using a 10x 0.3 NA or 20x 0.5 NA objectives setting 5.77  $\mu\text{m}$  and 1.77  $\mu\text{m}$  spacing respectively and sequential imaging channels for each fluorophore. The following lasers were assigned to each fluorophore: Hoechst:  $\lambda_{\text{ex}}$ . 405 nm laser,  $\lambda_{\text{em}}$ . 405-437 nm, FITC-GFP:  $\lambda_{\text{ex}}$ . 488nm laser,  $\lambda_{\text{em}}$ . 496-556 nm, TRITC-Phalloidin:  $\lambda_{\text{ex}}$ . 561nm laser,  $\lambda_{\text{em}}$ . 564-600 nm.

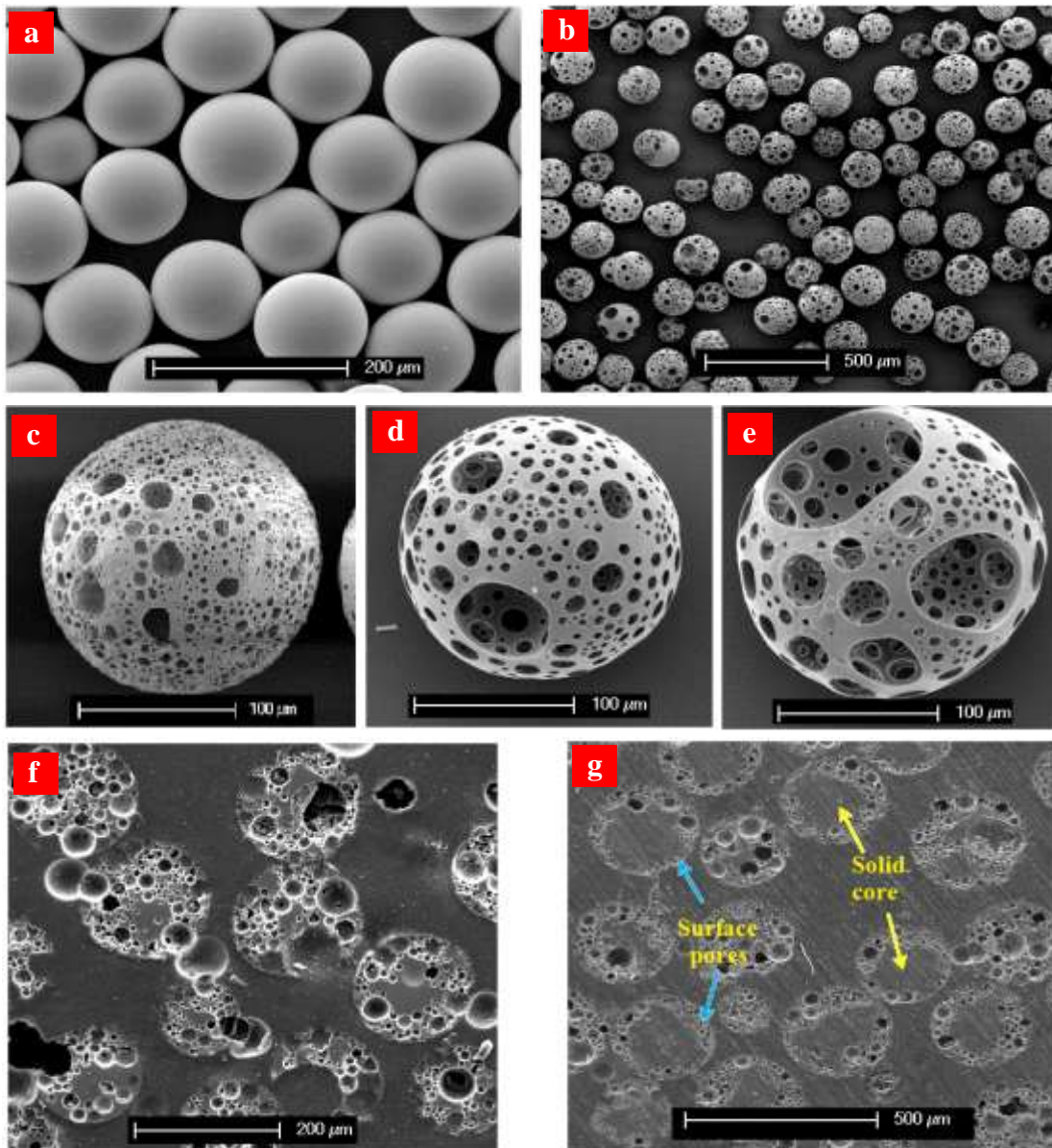
#### 2.2.5 Statistical analysis

For the cell study, one biological replicate was included in each assay. Data is presented as mean  $\pm$  SEM. Statistical analysis was performed using GraphPad PRISM 7.01 software package via the One-way ANOVA with Tukey's multiple comparison post hoc test. A 95% confidence level was considered significant.

### 3. Results

Both solid (non-porous) and highly porous CaP glass microspheres were produced via the flame spheroidisation process as shown in Figure 1. Figure 1a shows uniform non-porous glass microspheres with an average diameter of 108 ( $\pm 10$ )  $\mu\text{m}$ . Figure 1b represents the yield and distribution of pore morphologies achieved utilising the novel manufacturing process

developed herein. Figure 1c shows a SEM image of a porous CaP glass microsphere with a wide distribution of smaller surface pores (ranging from macroporous sized features to 30  $\mu\text{m}$ ). Whilst Figures 1d+e highlights porous CaP glass microspheres with a combination of larger (between 50 - 80  $\mu\text{m}$ ) and smaller (from 5 - 20  $\mu\text{m}$ ) surface pores. Comparatively larger surface pores with similar internal larger interconnected pores were also produced (as shown in Figure 1e).



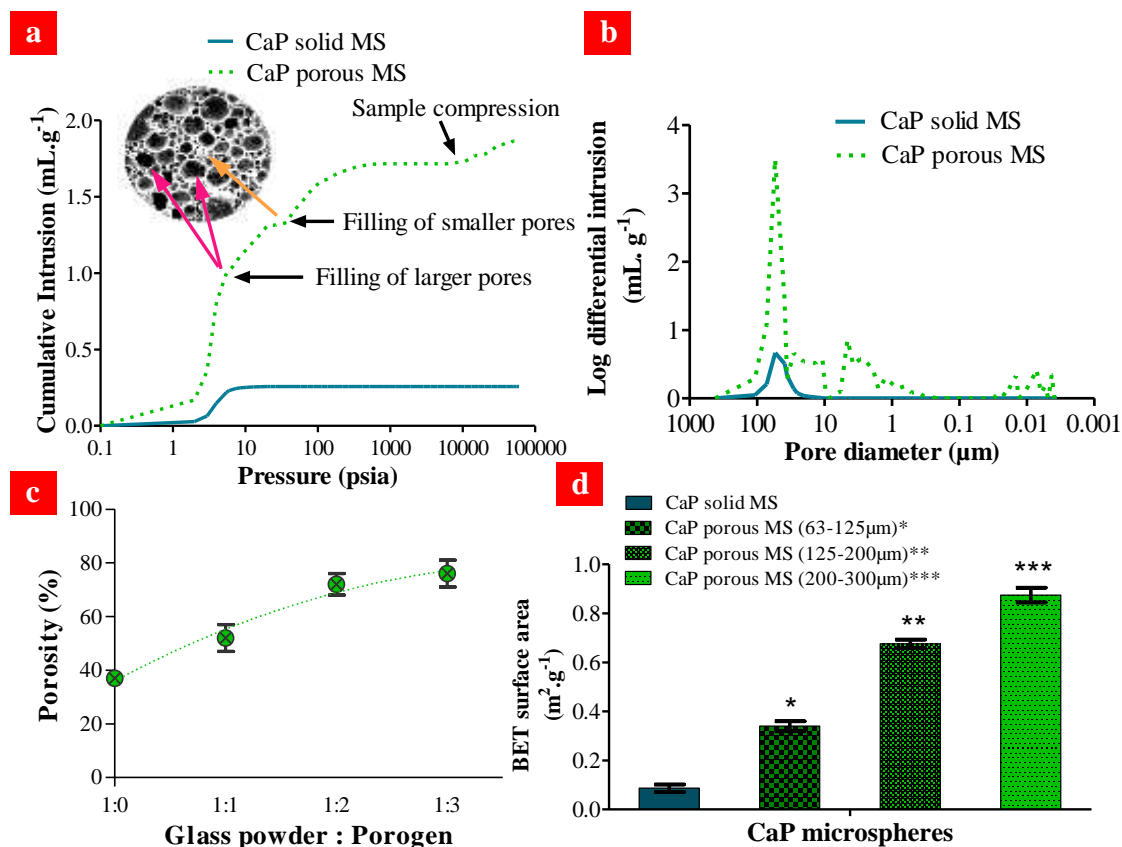
**Figure 1:** Representative SEM images of solid (non-porous) and porous CaP glass microspheres: **a)** Shows the distribution and yield of solid microspheres, **b)** Scale-up production of porous glass microspheres, **c)** Surface morphology of porous glass microsphere with smaller surface pores, **d)** Porous glass microsphere with a mixture of small and large



*surface pores, e) Porous glass microsphere with comparatively larger surface pores, f) Cross-sectional SEM images of resin embedded porous CaP glass microspheres with interconnected open porosity, and g) Microspheres with surface pores (indicated with blue arrows) and solid core (indicated by yellow arrows).*

The cross-sectional SEM images of the porous microspheres (as shown in Figure 1f) revealed the inner porosity and interconnectivity of the pores, which showed that the microspheres consisted mainly of smaller pores embedded within larger pores. Microspheres with surface porosity and a solid core were also manufactured as can be seen in Figure 1g when altering manufacturing parameters i.e. size of initial particles.

Characterisation of the CaP microspheres revealed porosity levels of up to and over 76 ( $\pm 5$ )% could be achieved depending on the process parameters. Mercury porosimetry analysis of porous and solid CaP glass microspheres is shown in Figure 2a. The mean pore size distribution for the porous microspheres was determined to be approximately 55 ( $\pm 8$ )  $\mu\text{m}$  and also revealed inner pore features at the submicron and nano porosity levels (as seen in Figure 2b). Figure 2c highlights the percentage porosity change with increasing glass powder/porogen ratio content. The porosity of the compact solid microspheres (with a size range of between 100-125  $\mu\text{m}$ ) was found to be approximately 37 ( $\pm 4$ ) % where the interparticulate gaps between the solid microspheres were characterised as pores. However, for the porous glass microspheres approximately 76 ( $\pm 5$ ) % porosity levels were obtained. The surface area (ascertained via BET analysis) for the non-porous CaP microspheres was 0.09 ( $\pm 0.01$ )  $\text{m}^2\text{g}^{-1}$  and the porous CaP microspheres were found to have an order of magnitude higher surface area which ranged from between 0.34 ( $\pm 0.02$ ) to 0.90 ( $\pm 0.03$ )  $\text{m}^2\text{g}^{-1}$  dependant on the microsphere size range selected (see Figure 2d). For example, the lower diameter size range porous microspheres (i.e between 63-125  $\mu\text{m}$ ) revealed a significant number of larger surface pores (hence a lower surface area of 0.34  $\text{m}^2\text{g}^{-1}$ ). Whilst, a comparatively higher surface area (0.9  $\text{m}^2\text{g}^{-1}$ ) was obtained for the larger diameter size range porous microspheres (i.e. between 200-300  $\mu\text{m}$ ) which revealed a larger number of smaller surface pores.

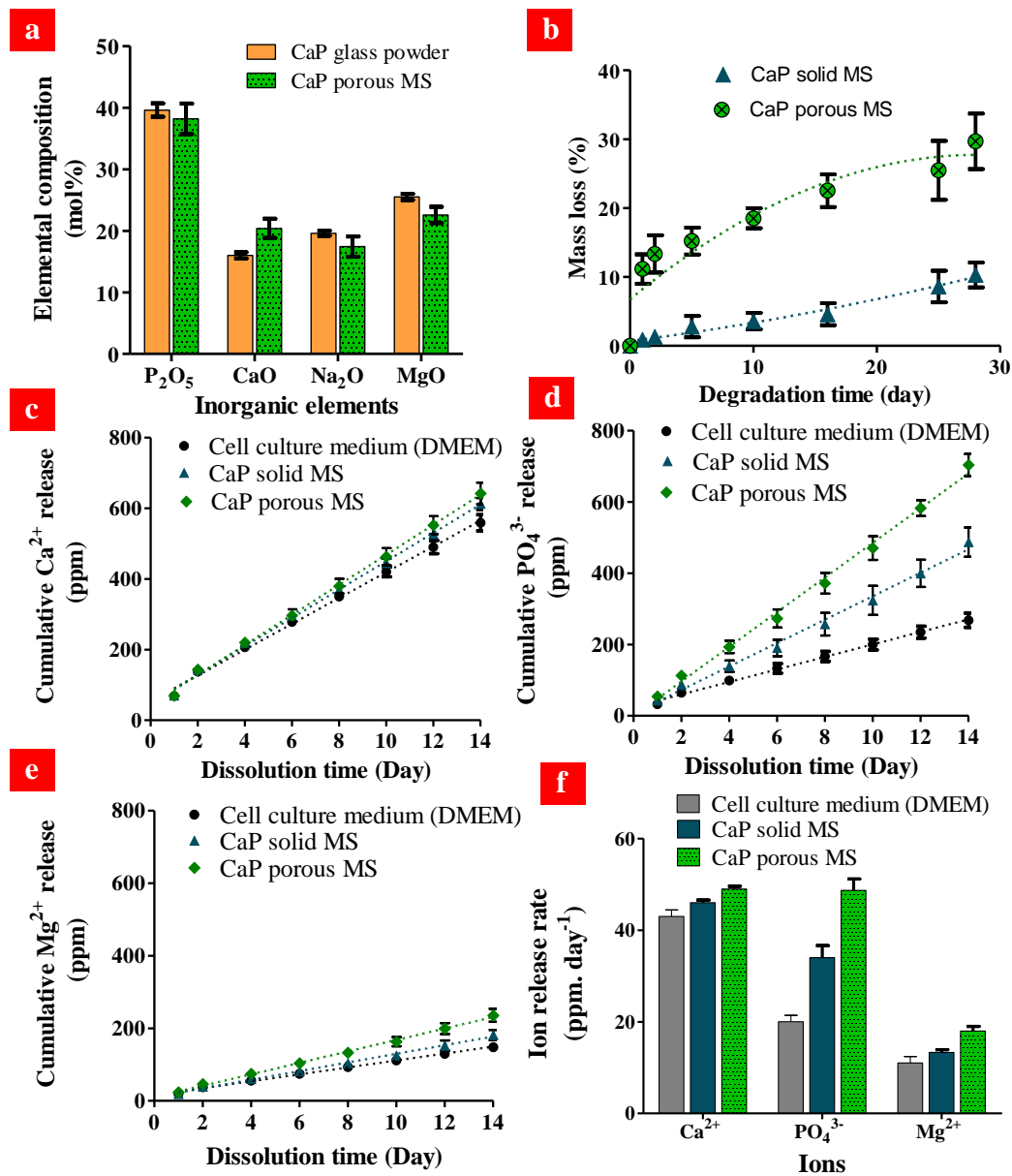


**Figure 2:** Porosity and surface area properties of CaP glass microspheres: **a)** Mercury intrusion curves of the solid and porous CaP glass microspheres, **b)** log differential intrusion curves of solid and porous CaP glass microspheres highlighting the range of pore sizes achieved, **c)** Porosity (%) of microspheres obtained with increasing glass powder/porogen ratio content and **d)** BET surface area analysis of the solid and porous CaP glass microspheres (between 63 to 300 μm diameter ranges).

EDX analysis confirmed negligible changes in glass compositions before and after spheroidisation (see Figure 3a), with only slight localised increases in Ca content which were mainly observed around the pores created.

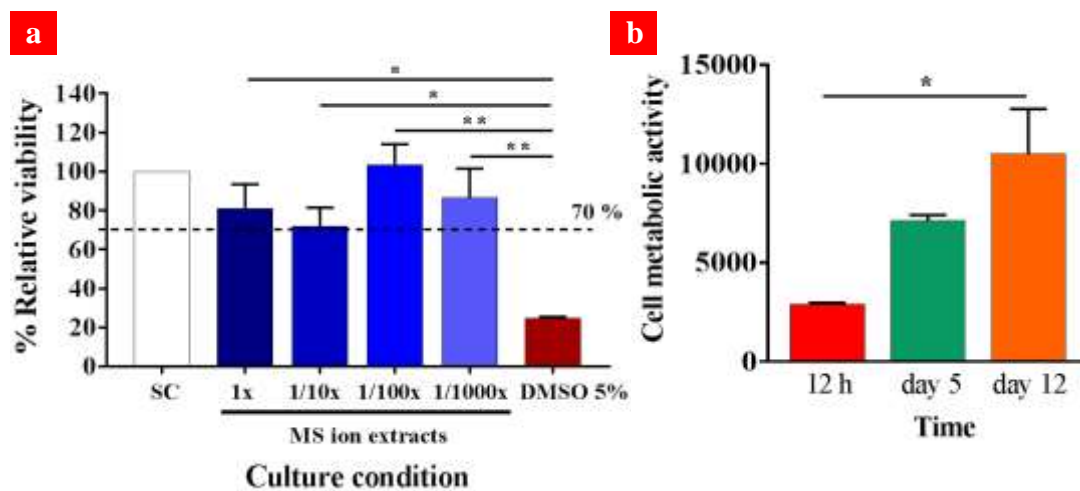
Porous CaP microspheres revealed an approximate 30% mass loss compared to 9% mass loss for the solid non-porous microspheres over a 28 day degradation period in ultra-purified water at 37°C (see Figure 3b). Figures 3c-f show increased ion release profiles (for example, Ca<sup>2+</sup> ~47 (±3) ppm, PO<sub>4</sub><sup>3-</sup> ~49 (±3) ppm, and Mg<sup>2+</sup> ~20 (±3) ppm per day) for the porous CaP glass microspheres (size range 125-200 μm) compared to the solid non-porous microspheres (Ca<sup>2+</sup> ~45 (±1) ppm, PO<sub>4</sub><sup>3-</sup> ~34 (±2) ppm, and Mg<sup>2+</sup> ~13 (±1) ppm per day) when degraded in

standard cell culture medium (DMEM) at 37°C for 14 days, which was attributed to their higher surface area.



**Figure 3:** Chemical composition, degradation and ion release profiles of CaP glass microspheres: a) Chemical composition of the glass (ascertained via EDX analysis) before and after spheroidisation, b) Mass loss profiles of solid and porous CaP glass microspheres during degradation in ultra purified over 28 days at 37°C, and c-e) Cumulative ion release profile (Ca<sup>2+</sup>, PO<sub>4</sub><sup>3-</sup>, and Mg<sup>2+</sup>) of solid and porous CaP microspheres in DMEM cell culture medium over 14 days at 37°C, and f) ion release rates (ppm per day) of solid and porous CaP microspheres calculated from the linear cumulative ion release profiles (observed in c-e).

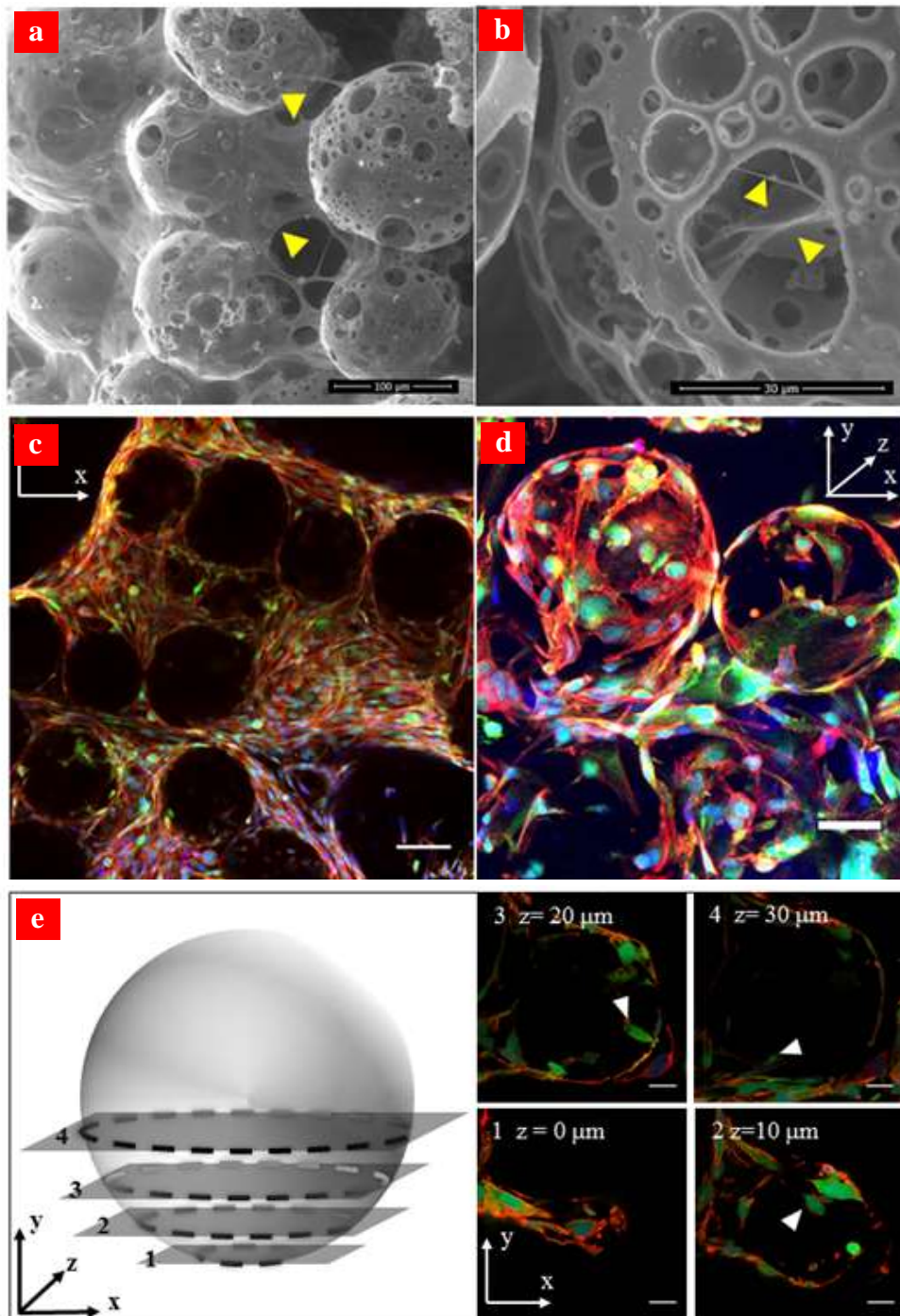
The biocompatibility evaluation of the porous CaP glass microspheres was performed through MTT test according to the ISO standard 10993-5. Results revealed that the proliferation rate of hMSCs cultured with the material ion extracts or with 1:10 serial dilutions was higher than 70 % of the SC medium control (1x: 81% ± 21; 1/10x: 72% ± 16; 1/100x: 103% ± 19; 1/1000x: 87% ± 26), as showed in Figure 4a. The suitability of the microspheres to support cell adhesion and growth was also evaluated by culturing cells directly onto the CaP glass microspheres for up to 14 days in SC medium. Their metabolic activity was analysed at 12 h, day 5 and day 12 which showed a progressive increase of the signal over time, suggesting an increase in cell numbers (see Figure 4b).



**Figure 4:** Biocompatibility evaluation of porous CaP glass microspheres according to ISO 10993. **a)** Results of MTT test showing the relative viability of cells cultured with ion extract conditioned medium (1x) and for the following three serial dilutions (1/10x, 1/100x, 1/1000x). The SC medium only and SC supplemented with 5 % DMSO were included as control. **b)** Shows the results of cell metabolic activity for porous CaP microspheres cultured at 12h, day 5 and day 12 after seeding (\* $p < 0.05$ , \*\* $p < 0.01$ ).

Initial observations confirmed through ESEM imaging showed that the cells were indeed attached and had spread on the microspheres and could also be observed in their inner pore structures (indicated by yellow arrows) where they acquired a 3D conformation (Figure 5a, b).

Further detailed analysis using scanning laser confocal microscopy was performed at day 14, where it was clearly observed that cells were attached to the microspheres and grew on and around them forming macro-aggregates (Figure 5c, d). Moreover, to ascertain how far the stem cells had migrated inside the porous microspheres, z-stack images were taken which confirmed that the cells were present up to half-way inside the porous microspheres (see Figure 5e).



**Figure 5:** *a)* ESEM image of stem cell-microsphere aggregates; the yellow arrows indicate hMSCs attached and bridging across several microspheres, *b)* ESEM image shows colonisation of hMSCs within the pores of the microspheres (yellow arrows), *c, d)* Representative confocal images (in single plane and z-stacked mode) of GFP-labelled hMSCs seeded on porous CaP microspheres at day 14, and *e)* cross sectioned representation of a microsphere and its respective single plane sectioned images; white arrows indicate cells

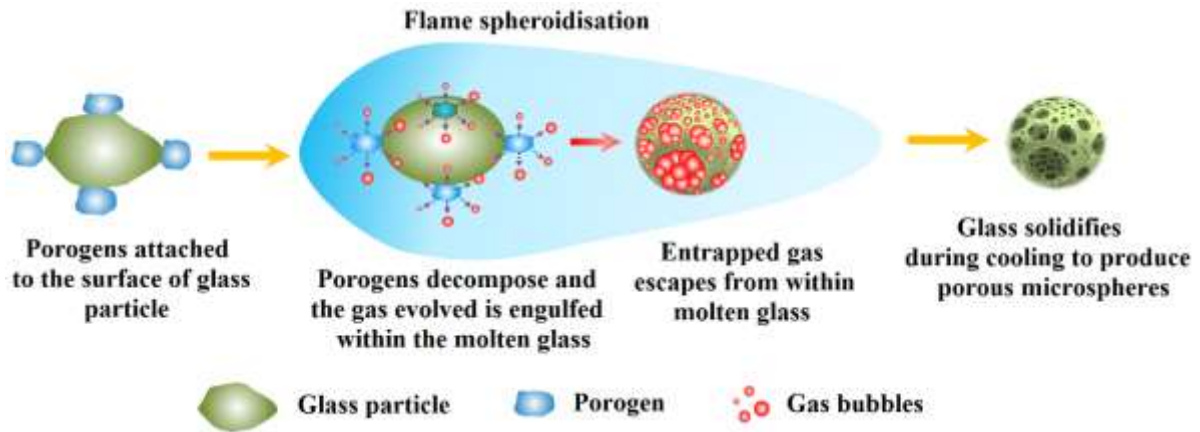
*which have migrated internally within the microsphere. hMSCs are seen in green (labelled using GFP), the nuclei are stained in blue (Hoechst 33259) and the cytoskeleton is seen in red (Phalloidin), respectively. Scale bars: 100  $\mu\text{m}$  (c), 50  $\mu\text{m}$  (d) and 25  $\mu\text{m}$  (e).*

#### **4. Discussion**

Orthobiologic therapies are being developed to help improve the long-term health of patients suffering from disabling musculoskeletal disorders and is a rapidly advancing field which utilises cell-based therapies and biomaterials to promote healing and offers exciting alternatives to traditional orthopaedic options.

The results above demonstrate (for the first time) a highly successful production process to manufacture highly porous CaP glass microspheres. The manufacturing route developed relies on a number of key parameters including the chemical and physical interaction of the porogen with the glass the glass viscosity, along with both being delivered within a suitable thermal processing window.

Figure 6 highlights a schematic representation of the morphological changes that occur and the proposed underlying mechanisms to produce the porous glass microspheres. The manufacturing process involved selection of a glass formulation that will not only melt but then allows it to reach a temperature where the viscosity was sufficiently low such that entrapment of the porogen gas could occur within the molten particle, which was itself driven to produce a spherical morphology due to surface tension from being ejected from the flame. The kinetics of decomposition must ideally be sufficiently slow for gas entrapment and adequately fast for pore formation to occur.



**Figure 6:** Schematic representation of the proposed mechanism for manufacturing porous glass microspheres via the flame spheroidisation process.

Critical to this process was not only the choice of pore forming agent but also the size of the porogens and in some cases how they were applied to affect the pore sizes and levels produced. As shown in Figure 1, a range of pores, pore sizes and quantities are achievable from this process. Furthermore, it was also found that varying the initial particle size and porogen quantities, enabled control over the size of the surface and interconnected pores.

Examining the CaP porous glass microspheres reported here the porosity (as measured via mercury intrusion) showed an increase from approximately 37% for loosely packed non-porous microspheres, which is typical for packed spheres, to over 76% for the porous microspheres (see Figures 2a-c). These results revealed that the inter-particulate gaps also contributed to the overall porosity levels measured for both the solid (non-porous) and the porous microspheres.

This study also showed that both porogen quantity and microparticles size also effected pore size and surface area, as higher quantities of porogen resulted in increased porosity (see Figures 2c) and smaller (63 -125 $\mu\text{m}$ ) particle sizes resulted in larger pore sizes. BET analysis confirmed that surface area could also be controlled up to approximately 0.9  $\text{m}^2 \text{g}^{-1}$  (see Figures 2d). These values appeared to be low, however the resulting porous microspheres resemble a shell-like structure with fully interconnected pores, which was reflected in the values obtained.

Addition of porogen during the spheroidisation process also resulted in slightly localised increase in Ca content which were mainly observed around the pores (see Figures 3a) and were attributed to the use of Ca-based porogens utilised. The introduction of porosity also resulted in an increase in mass loss during degradation (see Fig 3b) and increased the release of ions as expected due to the increased surface area achieved. As such, by simple selection of glass



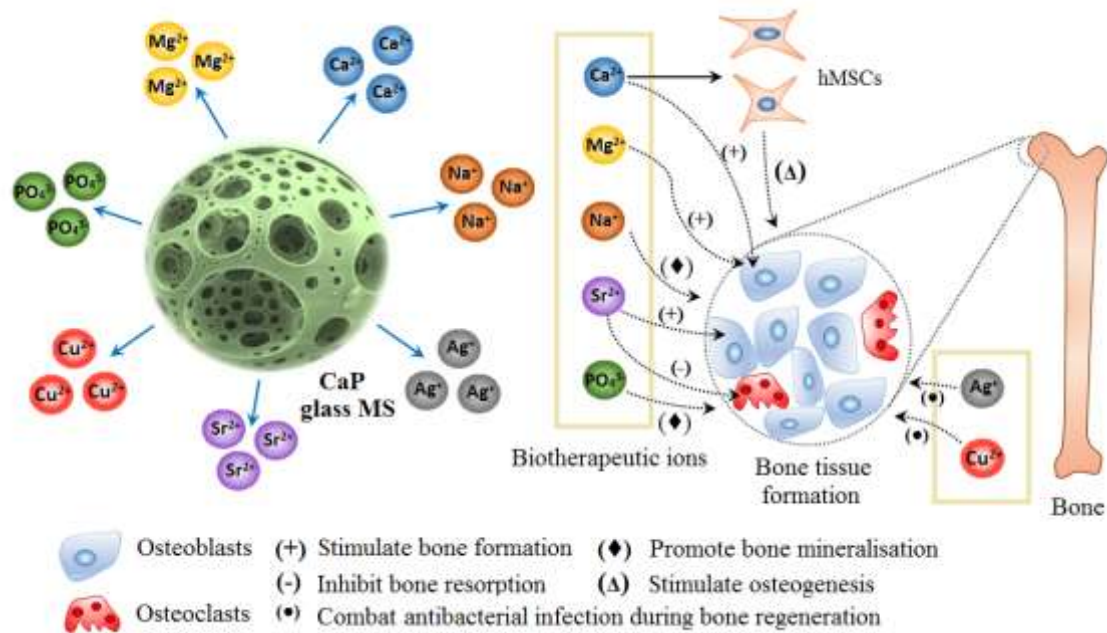
formulations and particle size, the microsphere porosity levels (which in turn relates to their surface area) could easily be manipulated to control ion release rates (as seen in Figures 3c-e) of potentially biotherapeutic ions for specific end applications.

The advantages of larger surface pores are that they could be exploited for applications which require faster ingress of fluids or media through the internal microsphere structure and smaller pores within could be utilised for the entrapment of particulate materials, for example in separation sciences. In particular for biomedical applications, larger pores would be beneficial to accommodate particular cell types, in order to promote tissue engineering or regeneration activities, whereas smaller surface pores could be beneficial for controlling release of drugs and/or other small biological components or biomolecules, entrapped within. Furthermore, where biological cell and material interactions are concerned, the inter-particulate gaps between the microspheres along with the enhanced levels of interconnected porosity created using this simple, single stage manufacturing process, could enable permeation of nutrients and waste products throughout the entire cell/material constructs [42-44].

Recently it has been shown that cell fate, in particular stem cell phenotype, could be influenced by controlled release of specific ions from inorganic materials [45]. Through simple compositional tailoring, the formulation of CaP glasses can easily be modified to release ions of specific interest. The solid and porous microspheres manufactured in this study revealed release of  $\text{Ca}^{2+}$ ,  $\text{PO}_4^{3-}$  and  $\text{Mg}^{2+}$  ions ranging from 45-47, 34-49 and 13-20 ppm per day (see Figure 3f). Furthermore, studies by Ahmed *et al* [46], showed that phosphate anionic species (ranging from ortho, pyro, meta and polyphosphates) could also be controlled, by simple tailoring of the starting glass formulations.

Figure 7 highlights some of the ions which could be released from CaP glasses and their potential biotherapeutic roles in bone repair and regeneration.  $\text{Ca}^{2+}$  ions are known to stimulate proliferation and differentiation of osteoblasts as well as extracellular matrix mineralisation [47] whilst  $\text{Mg}^{2+}$  ions can promote new bone formation [48]. Moreover,  $\text{PO}_4^{3-}$  ions are required for calcium phosphate crystal deposition [49] and extracellular matrix mineralisation while  $\text{Na}^+$  ions are mainly found in extracellular fluid [50].





**Figure 7:** Biotherapeutic ions which can be released from CaP glass and their role in bone tissue formation.

Other biotherapeutic ions can also be released from the CaP microspheres such as  $\text{Sr}^{2+}$  [12],  $\text{Cu}^{2+}$  [9] and  $\text{Ag}^{+}$  [10] by simply doping the glass composition with the desired metal oxide. For instance,  $\text{Sr}^{2+}$  ions are known to inhibit osteoclast activity whilst promoting osteogenesis of mesenchymal stem cells *in vitro* and *in vivo* [51],  $\text{Ag}^{2+}$  and  $\text{Cu}^{2+}$  ions have both demonstrated antimicrobial properties while  $\text{Cu}^{2+}$  ions have been also shown to promote the proliferation of endothelial growth factors and fibroblast growth factor 2 (FGF-2), which are associated with the angiogenesis process [9].

This study also demonstrated that the porous microspheres produced were biocompatible (see Figure 4a) according to the standard ISO 10993-5 (i.e. used for the biological evaluation of biomedical devices). Moreover, when the cells were seeded and cultured directly onto the microspheres, they were not only able to attach but had also migrated to within the centre of the microspheres, revealing colonisation of the inner pore structures (see Figure 5e). The pores formed presented the cells with a high surface area and a niche environment for the cells to grow and proliferate, acting as sites for cell culture [52]. This niche pore environment could also reduce any potential shear stresses experienced by the cells and provide protection for cells from getting damaged via inter-particle abrasion. Furthermore, the pores also closely resemble the native 3D conformation of bone tissue [53]. It has also been proposed that a porous

environment would favour the capture and utilisation of cell secreted factors close to where the cells reside in order to further influence cell phenotype [45].

The porous microspheres developed in this study constitute a versatile and highly promising technology for wider applications in tissue engineering and regenerative medicine (with a particular focus in bone regeneration) as initial examples. Indeed, doping the starting glass formulations with specific biotherapeutic ions could also address other key aspects of bone regeneration to include angiogenesis, osteogenesis and control of osteoclast activity, all potentially also coupled with antimicrobial properties. Further studies are on-going aimed at investigating more in-depth the bone tissue regenerative potential of these microspheres as well as exploring their potential use in alternate biomedical applications.

The porous glass microsphere production process developed herein offers a new cost-effective single stage manufacturing process, with demonstrated scale-up potential and with production levels in the range of kg's per hour at lab scale, which could easily be further scaled-out to commercial scales. Follow-on investigations are on-going to explore utility of this manufacturing process for other glass types (i.e. borates and silicates including glass-ceramics).

## **5. Conclusions**

This study highlights a novel flame spheroidisation process to manufacture highly porous CaP glass microspheres. Microspheres with a range of surface and fully interconnected porosity from 52 ( $\pm 5$ ) to 76 ( $\pm 5$ ) % were manufactured by simply manipulating process parameters such as the ratios of glass particles to porogen used (from 1:1 to 1:3) including the oxy-acetylene gas ratios.

The increase in porosity levels of the microspheres was shown to accelerate degradation over a 28 day degradation period (i.e. a 30% mass loss was observed for porous microspheres, compared to a 9% mass loss for the solid non-porous microspheres). The ion release profiles of the porous microspheres also revealed increasing rates as expected, compared to the non-porous microspheres, which was attributed to their increased surface area profiles.

The cell culture study revealed that the ion extracts from the CaP microspheres (utilising indirect cell culture method according to ISO 10993) did not exert any negative effect on the stem cells (hBM-MSCs). In addition, the direct cell culture study revealed that the porous

microspheres supported cell attachment as well as proliferation and showed that the hBM-MSCs had migrated into and towards the centre of the microspheres. The pore sizes and porosity levels achieved opens up the potential for use of these porous CaP microspheres for other biomedical applications in addition to the proposed orthobiologic applications.

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## Graphical abstract

