Structural characterization of sol–gel derived Sr-substituted calcium phosphates with anti-osteoporotic and anti-inflammatory properties†

G. Renaudin,‡a P. Laquerrière,b Y. Filinchuk,c E. Jallotd and J. M. Nedelec*a

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Sol–gel chemistry has been successfully used to prepare un-doped and Sr-doped calcium phosphate ceramics exhibiting a porous structure. The samples composition is very close to the nominal one. All samples present phase mixtures of mainly hydroxyapatite (HAp) and tricalcium phosphate (β-TCP). Doping with Sr2+ ions has a clear effect on the proportions of the different phases, increasing the amount of β-TCP. An amorphous phase is also observed incorporating some 40% of the total amount of strontium. Strontium ions also substitute for calcium both in HAp and β-TCP in specific sites that have been determined from Rietveld refinement on synchrotron powder diffraction data. The soluble amorphous and TCP phases are responsible for a beneficial partial release of strontium ions in solution during interactions between the material and biological fluids. Preliminary in vitro study demonstrates anti-inflammatory effects of strontium for human monocytes cultured in contact with calcium phosphates.

1. Introduction

Due to the global aging of the population, the need for bone substitutes is increasing very quickly. There is also a strong demand for bioceramics with specific properties such as anti-inflammatory, bactericidal or anti-osteoporotic properties. Bone mineral mass is dominated by nano-crystalline non-stoichiometric hydroxyapatite (HAp, Ca5(PO4)3(OH)) and whitlockite (β-TCP, β-Ca3(PO4)2) can be found at many different sites in the human body.1–3 For these reasons, apatite and whitlockite have been widely used as biocompatible materials for bone replacement and for bone prosthesis coatings.4 An interesting alternative is the use of so-called biphasic calcium phosphate (BCP) which is a mixture of HAp and β-TCP. Because of the difference in the solubility of the two phases, the solution behaviour of BCP can be easily tuned. Another interesting possibility is to perform ionic substitution in calcium phosphates to improve their properties and to modulate their bioactivity.

Similarly to the case of Ca, an estimated 99% of the total amount of Sr in the body is confined in bone.1 The total amount of Sr in the human skeleton is small but not insignificant; the Sr/Ca ratio is in the range 0.1–0.3 wt%o.5 Sr is readily taken up in humans.1,7–12 In vitro and in vivo studies have indicated that oral strontium intake not only increases bone formation, the number of bone-forming sites and the bone mineral density, but also reduces bone resorption.1,10–12 Sr is used for treatment of osteoporosis,13,14 and was found to induce osteoblast activity when introduced in biocompatible bone cements.15–19 Better osteointegration has been observed when Sr is present at low concentration level in biomaterials. The role of strontium in human pathology has attracted less attention than the other two important alkaline earths, calcium and magnesium. However there has been increasing awareness of the potential biological role of strontium and its incorporation in calcium phosphate cements has been largely studied in the last decade.15–19 The purpose of this study was to examine the incorporation of Sr2+ ions in hydroxyapatite samples at the atomic level: identification and quantification of the crystallised phases, identification of the Sr-occupied crystallographic sites, identification of an amorphous phase and determination of the chemical composition of the different phases.

2. Experimental

2.1 Sol–gel elaboration of Sr-substituted hydroxyapatite

The sol–gel route previously proposed by the authors31,32 was used. Briefly, to produce 2 g of pure HAp powder, 4.7 g of Ca(NO3)2·4H2O (Aldrich) and 0.84 g of P2O5 (Avocado Research chemicals) were dissolved in ethanol under stirring and refluxed at 85 °C during 24 h. Then, this solution was kept at 55 °C during 24 h, to obtain a white consistent gel and further heated at 80 °C during 10 h to obtain a white powder. Finally, the powder was heated at 1100 °C during 15 h. To prepare Sr-substituted hydroxyapatite, the required amount of Sr(NO3)2 (Aldrich) was added to the solution.

Two samples were studied: a pure hydroxyapatite sample (named ‘HAp’ in this paper) of nominal composition...
Table 1 Nominal and experimental concentrations (wt%) of Ca, P, Sr in HAp and Sr:HAp samples determined by ICP-AES

<table>
<thead>
<tr>
<th></th>
<th>HAp</th>
<th>Sr:HAp</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ca Nominal</td>
<td>39.88</td>
<td>37.02</td>
</tr>
<tr>
<td>Ca Experimental</td>
<td>38.46</td>
<td>38.79</td>
</tr>
<tr>
<td>P Nominal</td>
<td>18.50</td>
<td>18.07</td>
</tr>
<tr>
<td>P Experimental</td>
<td>17.21</td>
<td>18.42</td>
</tr>
<tr>
<td>Sr Nominal</td>
<td>—</td>
<td>4.26</td>
</tr>
<tr>
<td>Sr Experimental</td>
<td>—</td>
<td>4.23</td>
</tr>
<tr>
<td>% Substitution theoretical</td>
<td>5</td>
<td>5</td>
</tr>
<tr>
<td>% Substitution experimental</td>
<td>—</td>
<td>—</td>
</tr>
</tbody>
</table>

In vitro Sr²⁺ release studies. The interactions between the ceramics and a biological medium were carried out at 37 °C for 1 to 6 days by immersing about 10 mg of powder in a standard Dulbecco’s Modified Eagle Medium (DMEM), the composition of which is almost identical to that of human plasma. The specific surface area to DMEM volume ratio was fixed at 500 cm⁻¹. These in vitro assays were conducted under static conditions: biological fluids were not renewed and therefore contained only limited amounts of P and Ca. Namely, this is a good approach for further development and to compare different doping elements and compositions. ICP-AES was used to analyze the biological fluid before and after bio ceramics interactions.

2.3 X-Ray powder diffraction

Synchrotron measurement. Synchrotron powder diffraction data of both samples were obtained at ESRF (Grenoble, France) on the SNBL. Finely ground powders were introduced into a 0.3 mm diameter glass capillary. Data collections were performed at 295 K with a MAR345 Image Plate detector by using a monochromatic wavelength of λ = 0.710130 Å. The calculated absorption coefficient was $\mu R = 0.68$ (m = powder packing factor and $\mu$ = linear absorption coefficient; $R$ = radius of the capillary, 0.15 mm). Two sample-to-detector distances were used (150 and 350 mm) in order to combine the benefits of high resolution and an extended 2θ range. The detector and the wavelength were calibrated with NIST LaB₆. The highest resolution of the MAR345 detector was used: 3450 × 3450 pixels with a pixel size of 100 μm. The exposure time was 60 s with a rotation of the capillary by 60°. The two-dimensional data (see ESI† Fig. S1) were integrated with the Fit2D program which produced the correct intensity on a relative scale. This 2D detector was used in order to perfectly define the background line, to observe very weak diffraction peaks, and to improve the accuracy of the integrated intensities by achieving a better powder average. Uncertainties of the integrated intensities were calculated at each 2θ point applying Poisson statistics to the intensity data, considering the geometry of the detector. The Instrument Resolutions Functions were determined from the LaB₆ data.

Rietveld refinements. Joint Rietveld refinements based on the simultaneous use of the data measured at 150 mm (large 2θ interval) and 350 mm (high resolution) sample to detector distances were performed for both samples with the program Fullprof.2k Multi-Pattern. Three phases were refined for the HAp sample: hydroxyapatite, whitlockite and lime. For the Sr:HAp sample, portlandite was introduced as the fourth phase. The absence of calcite in both samples was verified. The structural parameters of hydroxyapatite, $\text{Ca}_9(\text{PO}_4)_3\cdot\text{OH}$, were taken from ref. 36: space group $P6_3/m$, $Z = 2$, $a = 9.4218$ Å and $c = 6.8813$ Å, 7 independent atomic positions: two Ca, one P and four O positions. The O4 oxygen position (i.e. the hydroxyl anion) is half occupied, this corresponds to a statistically disordered 4e position around the 2a (0.0,0.14) site. The structural parameters of whitlockite, $\beta$-$\text{Ca}_9(\text{PO}_4)_3\cdot\text{OH}$, were taken from ref. 37: space group $R3c$, $Z = 21$, $a = 10.4352$ Å and $c = 37.4029$ Å, 18 independent atomic positions: five Ca, three P positions and ten O positions. The structural parameters of lime, CaO, were taken...
from ref. 38: space group $Fm\bar{3}m$, $Z = 4$, $a = 4.8071$ Å. The structural parameters of portlandite, Ca(OH)$_2$, were taken from ref. 39: space group $P\overline{3}m1$, $Z = 1$, $a = 3.589$ Å and $c = 4.911$ Å. In the first step all the structural parameters were fixed according to the literature values. Then during the successive refinement cycles they were all refined (first lattice parameters and peak profiles, and next atomic parameters: positional, isotropic temperature and occupancy factors). In the final run, 96 and 115 parameters were refined for the HAp sample (2 zero shifts, 6 scale factors, 20 profile parameters and 68 structural parameters) and for the Sr:HAp sample (2 zero shifts, 8 scale factors, 28 profile parameters and 77 structural parameters) respectively. The determination of the Instrumental Resolution Function (extracted from the powder patterns recorded on NIST LaB$_6$ powder, see ESI† Fig. S2) improved the peak profile modelling while decreasing the number of profile parameters. It allowed also the extraction of the sample intrinsic microstructural parameters: average apparent crystallite size and average maximum strain. The diffraction profiles (both instrumental and sample intrinsic) were used to study the apparent crystallite size and average maximum strain. The use of the monoclinic ordered structural description for the hydroxyapatite phase did not allow increased accuracy of the refinement. The last refinement cycles were performed by introducing the hydrogen atom from the hydroxyl anion. The H4 position was located the site 4e (in accordance with ref. 37), a soft constraint of 0.92 ± 0.05 Å was applied to the distance O4–H4, and the isotropic temperature factors of H4 was constrained to be 1.2 times $B_{iso}$ of the corresponding oxygen atom O4, as is usual for hydroxyl groups. The final refined $d_{O4-H4}$ distances were 0.89(2) Å and 0.92(2) Å for the pure and the Sr-doped hydroxyapatite phase respectively. The previously unconstrained distance converged to 0.67(7) Å in the case of the pure hydroxyapatite phase, and did not converge in the case of the Sr-doped hydroxyapatite phase. The refined structural parameters for hydroxyapatite and whitlockite phases are summarized in Tables 2 and 3 respectively. A Rietveld plot of the Sr:HAp sample for one sample to detector distance is presented in Fig. 2. The other plots can be found in ESI† (Fig. S3 and S4).

### 2.4 Preliminary biological study of inflammation reactions

**Cell culture.** Elutriated monocytes were used to evaluate tumor necrosis factor (TNF-α) production. Monocytes from five healthy consenting donors were collected by leukapheresis and purified by counterflow elutriation. Purity, after CD14 staining, was at least 95%, the rest being neutrophils. Cells were maintained in RPMI-1640 medium (Life Technologies, France) supplemented with 10% fetal calf serum (Life Technologies, France), 2 mM glucose, penicillin (5000 U mL$^{-1}$, Life Technologies) and streptomycin (25 μg mL$^{-1}$, Life Technologies) at 37 °C in a saturated 5% CO$_2$–95% air atmosphere. All reagents were certified endotoxin-free. The cells were cultured in petri dishes (Life Technologies). The density was 2 × 10$^6$ cells mL$^{-1}$ in a 5 mL total volume. Cells of each donor were exposed during 6 h (time chosen to have an interaction between cells and particles) to the different particles described above. The negative control consisted of monocytes in culture medium alone (data not shown) and the positive control was monocytes plus lipopolysaccharide (LPS). LPS (Sigma-Aldrich) were chosen because of their inflammatory actions on the target cells.

The viability of cells was evaluated by trypan blue exclusion test. The viability was >95% in all of the experiments (before and after exposure to particles).

**Quantification of TNF-α by ELISA.** After 6 h, TNF-α determinations in cell culture supernatants were performed using commercially available ELISA kits (Quantikine, R&D Systems, Inc., Minneapolis, MN) according to the manufacturer’s

### Table 2 Refinement results on the hydroxyapatite phase in the two HAp and Sr:HAp samples

<table>
<thead>
<tr>
<th>Phase</th>
<th>Atom</th>
<th>Site</th>
<th>x</th>
<th>y</th>
<th>z</th>
<th>$B_{iso}$/Å$^2$</th>
<th>Occ. $^b$</th>
</tr>
</thead>
<tbody>
<tr>
<td>$Ca_4(PO_4)_3(OH)$</td>
<td>$Ca_1$</td>
<td>4f</td>
<td>1/3</td>
<td>2/3</td>
<td>0.0013(2)</td>
<td>0.29(2)</td>
<td>I(—)</td>
</tr>
<tr>
<td>$P6_3/m$, $Z = 2$</td>
<td>$Ca_2$</td>
<td>6h</td>
<td>0.2462(1)</td>
<td>0.9926(1)</td>
<td>½</td>
<td>$= B_{Ca(1)}$</td>
<td>I(—)</td>
</tr>
<tr>
<td>$a = 9.4239(1)$ Å</td>
<td>$P1$</td>
<td>6h</td>
<td>0.3988(2)</td>
<td>0.3691(2)</td>
<td>½</td>
<td>$= B_{(Ca)^1}$</td>
<td>I(—)</td>
</tr>
<tr>
<td>$c = 6.8841(1)$ Å</td>
<td>$O1$</td>
<td>6h</td>
<td>0.3260(3)</td>
<td>0.4830(3)</td>
<td>½</td>
<td>$= B_{Ca(1)}$</td>
<td>I(—)</td>
</tr>
<tr>
<td>$R_{Bragg} = 0.026; 0.013$</td>
<td>$O2$</td>
<td>6h</td>
<td>0.5875(3)</td>
<td>0.4653(3)</td>
<td>½</td>
<td>$= B_{(Ca)^1}$</td>
<td>I(—)</td>
</tr>
<tr>
<td>$R_b = 0.021; 0.018$</td>
<td>$O3$</td>
<td>12i</td>
<td>0.3407(2)</td>
<td>0.2559(2)</td>
<td>0.2030(9)</td>
<td>$= B_{(Ca)^1}$</td>
<td>I(—)</td>
</tr>
<tr>
<td>$R_{ap} = 0.029; 0.027$</td>
<td>$H4$</td>
<td>4e</td>
<td>0.073(3)</td>
<td>0.191(1)</td>
<td>0.057(3)</td>
<td>$= 1.2 \times B_{(Ca)^1}$</td>
<td>I(—)</td>
</tr>
</tbody>
</table>

* $^a$ Two values represent the measurements performed at 150 and 350 mm respectively. $^b$ Site occupancy factor.
compare the different conditions.

between positive control and stimulated cells.

were presented in relative concentration. Results were presented

as median

and lime). Portlandite was introduced as the fourth phase in

preliminarily determined by X-ray diffraction. This amorphous phase can be expressed as

an amorphous (Ca,Sr)O equivalent. According to the apparent Sr deficit in the crystallised phases (indicated by the

synthesis of a single phase Sr:HAp sample) and about 28% is

inserted in the whitlockite phase (with the Ca2.87(6)Sr0.13(6)(PO4)2

composition). Rietveld refinement for the HAp sample indicated

that a very small amount of CaO (i.e. 0.2 wt%) led to the

introduction of Sr into the apatite. The presence of whitlockite, with a lower Ca/PO4 ratio of

1.66(2) and 1.63(3) refined ratios, respectively). This deficit in the case of the Sr:HAp sample may be explained by the presence of CaO. Nevertheless a deficit in alkaline earth atoms was observed: it was in the range of the standard deviation in the case of the HAp sample, and more pronounced in the case of the Sr:HAp sample (i.e. 1.66(2) and 1.63(3) refined ratios, respectively). This deficit in the case of the Sr:HAp sample may be explained by the presence of an alkaline-earth rich amorphous phase, not accessible by X-ray diffraction. This amorphous phase can be expressed as an amorphous (Ca,Sr)O equivalent. According to the apparent Sr deficit in the crystallised phases (indicated by the refined compositions of Ca4.88(1)Sr0.12(1)(PO4)3(OH) and Ca2.87(6)Sr0.13(6)(PO4)2 leading to a Sr amount of only 2.86(4)%), the amorphous (Ca,Sr)O should mainly be composed of strontium oxide. Calculation was performed, in the case of the Sr:HAp sample, to estimate the amount of amorphous phase by simultaneous comparison of the nominal and refined (Ca + Sr)/PO4 ratios and the nominal and refined Sr amounts (see Table 4). An estimation of 2.2 wt% of amorphous (Ca,Sr)O, which is composed of around 90 wt% strontium oxide, was obtained. These results indicate that a large amount of about 40% of the Sr introduced during the synthesis is present in the amorphous phase, about 32% is inserted in the apatite phase (of composition Ca4.88(1)Sr0.12(1)(PO4)3(OH) relatively far from the theoretical Ca4.75Sr0.25(PO4)3(OH) composition corresponding to the synthesis of a single phase Sr:HAp sample) and about 28% is inserted in the whitlockite phase (with the Ca2.87(6)Sr0.13(6)(PO4)2 composition). Rietveld refinement for the HAp sample indicated that a very small amount of CaO (i.e. 0.2 wt%) led to the

Table 3 Refinement results on the whitlockite phase in the Sr:HAp sample

<table>
<thead>
<tr>
<th>Phase</th>
<th>Atom</th>
<th>Site</th>
<th>x</th>
<th>y</th>
<th>z</th>
<th>Biso/Å²</th>
<th>Occ.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ca2.87(6)Sr0.13(6)(PO4)2</td>
<td>Ca1</td>
<td>18b</td>
<td>0.728(1)</td>
<td>0.861(2)</td>
<td>0.1700(7)</td>
<td>0.30(9)</td>
<td>1(—)</td>
</tr>
<tr>
<td>3.1 Quantitative phases analysis</td>
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</tbody>
</table>

Quantitative phases analysis was performed on the basis of the joint Rietveld refinements. Three phases were observed and quantified in the case of the HAp sample (apatite, whitlockite and lime). Portlandite was introduced as the fourth phase in the Sr:HAp sample. Results of the quantitative phase analysis are summarized in Table 4.

The synthesis conditions correspond to hydroxyapatite single phase samples; i.e. a (Ca + Sr)/PO4 ratio of 1.67. However both syntheses yielded multiphase samples. The HAp sample contained 93 wt% of apatite but Sr:HAp contained only 65% apatite. The presence of whitlockite, with a lower Ca/PO4 ratio of 1.50, was compensated by the presence of CaO. Nevertheless a deficit in alkaline earth atoms was observed: it was in the range of the standard deviation in the case of the HAp sample, and more pronounced in the case of the Sr:HAp sample (i.e.1.66(2) and 1.63(3) refined ratios, respectively). This deficit in the case of the Sr:HAp sample may be explained by the presence of an alkaline-earth rich amorphous phase, not accessible by X-ray diffraction. This amorphous phase can be expressed as an amorphous (Ca,Sr)O equivalent. According to the apparent Sr deficit in the crystallised phases (indicated by the refined compositions of Ca4.88(1)Sr0.12(1)(PO4)3(OH) and Ca2.87(6)Sr0.13(6)(PO4)2 leading to a Sr amount of only 2.86(4)%), the amorphous (Ca,Sr)O should mainly be composed of strontium oxide. Calculation was performed, in the case of the Sr:HAp sample, to estimate the amount of amorphous phase by simultaneous comparison of the nominal and refined (Ca + Sr)/PO4 ratios and the nominal and refined Sr amounts (see Table 4). An estimation of 2.2 wt% of amorphous (Ca,Sr)O, which is composed of around 90 wt% strontium oxide, was obtained. These results indicate that a large amount of about 40% of the Sr introduced during the synthesis is present in the amorphous phase, about 32% is inserted in the apatite phase (of composition Ca4.88(1)Sr0.12(1)(PO4)3(OH) relatively far from the theoretical Ca4.75Sr0.25(PO4)3(OH) composition corresponding to the synthesis of a single phase Sr:HAp sample) and about 28% is inserted in the whitlockite phase (with the Ca2.87(6)Sr0.13(6)(PO4)2 composition). Rietveld refinement for the HAp sample indicated that a very small amount of CaO (i.e. 0.2 wt%) led to the

instructions. The sensitivity of each kit was 7.8 pg mL−1. Results were presented in relative concentration. Results were presented as median ± interquartile space (IQS) of relative variation between positive control and stimulated cells.

A Mann & Whitney non-parametric test was performed to compare the different conditions.

3. Results and discussion

3.1 Quantitative phases analysis

Quantitative phases analysis was performed on the basis of the joint Rietveld refinements. Three phases were observed and quantified in the case of the HAp sample (apatite, whitlockite and lime). Portlandite was introduced as the fourth phase in the Sr:HAp sample. Results of the quantitative phase analysis are summarized in Table 4.

The synthesis conditions correspond to hydroxyapatite single phase samples; i.e. a (Ca + Sr)/PO4 ratio of 1.67. However both syntheses yielded multiphase samples. The HAp sample contained 93 wt% of apatite but Sr:HAp contained only 65% apatite. The presence of whitlockite, with a lower Ca/PO4 ratio of 1.50, was compensated by the presence of CaO. Nevertheless a deficit in alkaline earth atoms was observed: it was in the range of the standard deviation in the case of the HAp sample, and more pronounced in the case of the Sr:HAp sample (i.e.1.66(2) and 1.63(3) refined ratios, respectively). This deficit in the case of the Sr:HAp sample may be explained by the presence of an alkaline-earth rich amorphous phase, not accessible by X-ray diffraction. This amorphous phase can be expressed as an amorphous (Ca,Sr)O equivalent. According to the apparent Sr deficit in the crystallised phases (indicated by the refined compositions of Ca4.88(1)Sr0.12(1)(PO4)3(OH) and Ca2.87(6)Sr0.13(6)(PO4)2 leading to a Sr amount of only 2.86(4)%), the amorphous (Ca,Sr)O should mainly be composed of strontium oxide. Calculation was performed, in the case of the Sr:HAp sample, to estimate the amount of amorphous phase by simultaneous comparison of the nominal and refined (Ca + Sr)/PO4 ratios and the nominal and refined Sr amounts (see Table 4). An estimation of 2.2 wt% of amorphous (Ca,Sr)O, which is composed of around 90 wt% strontium oxide, was obtained. These results indicate that a large amount of about 40% of the Sr introduced during the synthesis is present in the amorphous phase, about 32% is inserted in the apatite phase (of composition Ca4.88(1)Sr0.12(1)(PO4)3(OH) relatively far from the theoretical Ca4.75Sr0.25(PO4)3(OH) composition corresponding to the synthesis of a single phase Sr:HAp sample) and about 28% is inserted in the whitlockite phase (with the Ca2.87(6)Sr0.13(6)(PO4)2 composition). Rietveld refinement for the HAp sample indicated that a very small amount of CaO (i.e. 0.2 wt%) led to the
formation of a noticeable amount of whitlockite (i.e., 7.0 wt%) due to the similar Ca/PO₄ ratios of hydroxyapatite (1.67) and whitlockite (1.50) phases. Introduction of strontium during the synthesis increased the formation of the alkaline earth oxide with poor crystallinity (from 0.2wt% in the HAp sample to about 3 wt% in the Sr:HAp sample by considering lime, portlandite and the amorphous equivalent) which is likely located at the surface of the crystallised grains of hydroxyapatite and whitlockite in the form of a thin amorphous surface layer. The appearance of whitlockite in the Sr containing apatite samples has been previously reported. Nevertheless the presence of a small amount (less than 1 wt% evaluated from high quality X-ray powder patterns) of CaO, as well as Ca(OH)₂ and the amorphous equivalent (less than 1 wt% evaluated from high quality X-ray powder patterns) of CaO, as well as Ca(OH)₂ and the amorphous equivalent (less than 1 wt%) which is likely located at the surface of the crystallised grains of hydroxyapatite and whitlockite is not attributed to the stoichiometry of the apatite structure. At high temperature, the Sr-doped whitlockite phase is more stable than the Sr-doped hydroxyapatite phase and its formation is compensated by the combined extraction of alkaline earth oxide in order to keep the Sr:HAp sample by considering lime, portlandite and the amorphous equivalent which is likely located at the surface of the crystallised grains of hydroxyapatite and whitlockite in the form of a thin amorphous surface layer). The appearance of whitlockite in the Sr containing apatite samples has been previously reported. Nevertheless the presence of a small amount (less than 1 wt% evaluated from high quality X-ray powder patterns) of CaO, as well as Ca(OH)₂ and the amorphous equivalent, simultaneously with the whitlockite phase, indicated that the whitlockite formation is not attributed to the non-stoichiometry of the apatite structure. At high temperature, the Sr-doped whitlockite phase is more stable than the Sr-doped hydroxyapatite phase and its formation is compensated by the combined extraction of alkaline earth oxide in order to keep the ratio (Ca + Sr)/PO₄ = 1.67. This observation correlates with the fact that single phase poorly crystalline Sr:HAp samples are obtained by moderate heat treatment. Such hydroxyapatite inhibiting formation, favouring whitlockite formation, has already been described in the case of smaller Mg²⁺ ion insertion.

3.2 Srhydroxyapatite solid solution study
This study investigated the Ca-rich side of the Sr-apatite solid solution, i.e. Ca₅₋ₓSrₓ(PO₄)₃H₂O with x = 0 and 0.12. The solid solution extends from x = 0 (i.e. undoped hydroxyapatite) to x = 5 (i.e. belovite Sr₅(PO₄)₃(OH)⁴⁻). Fig. 3 reports the lattice parameters and unit cell volume variations in the solid solution according to the present study and from the literature for x = 0, 0.09, 0.12, 0.51, 1.158, 3.192 and 5.45. The compound with x = 0.09 has to be carefully considered here because of its non-

Table 4 Results of the quantitative phase analyses using the Rietveld method. An amorphous (Ca,Sr)₂O₃ amount was estimated assuming the refined values for the crystalline phases and the synthesis conditions: (Ca + Sr)/PO₄ = 1.67 with 5 atom% of Sr

<table>
<thead>
<tr>
<th>Phase Formula</th>
<th>Wt%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Apatite</td>
<td>92.8(6)%</td>
</tr>
</tbody>
</table>
| F
c ironic     | 0.4% |
| 2.86(4)%      |       |

Refined:

<table>
<thead>
<tr>
<th>Phase Formula</th>
<th>Wt%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Apatite</td>
<td>66.5(6)%</td>
</tr>
</tbody>
</table>
| F
c ironic     | 0.4% |
| 2.86(4)%      |       |

Estimated assuming the used synthesis ratios:

<table>
<thead>
<tr>
<th>Phase Formula</th>
<th>Wt%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Apatite</td>
<td>65.0%</td>
</tr>
</tbody>
</table>
| F
c ironic     | 0.4% |
| 2.86(4)%      |       |

Ref. 44:

formation of a noticeable amount of whitlockite (i.e., 7.0 wt%) due to the similar Ca/PO₄ ratios of hydroxyapatite (1.67) and whitlockite (1.50) phases. Introduction of strontium during the synthesis increased the formation of the alkaline earth oxide with poor crystallinity (from 0.2wt% in the HAp sample to about 3 wt% in the Sr:HAp sample by considering lime, portlandite and the amorphous equivalent which is likely located at the surface of the crystallised grains of hydroxyapatite and whitlockite in the form of a thin amorphous surface layer). The appearance of whitlockite in the Sr containing apatite samples has been previously reported. Nevertheless the presence of a small amount (less than 1 wt% evaluated from high quality X-ray powder patterns) of CaO, as well as Ca(OH)₂ and the amorphous equivalent, simultaneously with the whitlockite phase, indicated that the whitlockite formation is not attributed to the non-stoichiometry of the apatite structure. At high temperature, the Sr-doped whitlockite phase is more stable than the Sr-doped hydroxyapatite phase and its formation is compensated by the combined extraction of alkaline earth oxide in order to keep the ratio (Ca + Sr)/PO₄ = 1.67. This observation correlates with the fact that single phase poorly crystalline Sr:HAp samples are obtained by moderate heat treatment. Such hydroxyapatite inhibiting formation, favouring whitlockite formation, has already been described in the case of smaller Mg²⁺ ion insertion.
deviation from the linear Vegard’s law at the Ca-rich side of the solid solution).

3.3 Sr-doped whitlockite study

This study investigated the insertion of a small amount of Sr in the whitlockite structure. Table 3 indicates that Sr is preferentially inserted in the Ca4 site. More generally Sr was inserted in the low density column described and named the A column by Yashima et al.37 (see Fig. 2 in ref. 37). 75% of inserted Sr atoms are located in this A column (composed of the Ca4 and Ca5 sites), compared to only 25% of inserted Sr atoms in the dense B column (made of the Ca1, Ca2 and Ca3 sites). Fig. 5 represents the five independent Ca sites in Sr-free whitlockite. The Ca4 site presents an unusually low coordination number (CN = 3) with a planar polyhedron in pure whitlockite.37 If we take into account the three following neighbouring oxygen atoms, a coordination number of 6 can be assumed, with three shorter distances $d_{Ca4-O1} = 2.53 \, Å$ and three larger distances $d_{Ca4-O9} = 3.12 \, Å$ (the latter are represented by broken lines in Fig. 5d). Although such a coordination polyhedron is more appropriate, a deficiency in the calculated bond valence remains for Ca4: the bond valence sum (BVS) equals 0.68 or 0.79 by taking into account 3 or 6 neighbouring oxygens respectively. In our sample, the half-occupied Ca4 site is 25%–25% occupied by Ca and Sr atoms. This large insertion of strontium in the Ca4 site modifies its coordination polyhedron. In the Sr-doped whitlockite structure, the Ca4 site has six neighbouring oxygen atoms with quite equivalent (Ca, Sr)–O bond distances: three distances $d_{Ca4-O1} = 2.63 \, Å$ and three distances $d_{Ca4-O9} = 2.79 \, Å$. These six bonds give the calculated BVS of 1.03 (by considering Ca4 occupied by 25% Ca and 25% Sr) which is still too low for a divalent cation. The insertion of Sr in the Ca4 site has clearly a stabilising effect on the whitlockite structure. Not only is the geometrical shape of the Ca4 polyhedron considerably improved, but also those for the other cation sites. Fig. 5 represents the environments of all the cationic sites as refined in the Sr-doped whitlockite. Ca1, Ca2 and Ca3 compose the dense B column37 and have CN = 8 with a more or less distorted monocapped pentagonal bipyramidal coordination polyhedra.

![Fig. 3](image1.png) Lattice parameters (top) and unit cell volume (bottom) as a function of the $x$ value for the solid solution $Ca_{5-x}Sr_x(PO_4)_3(OH)$. Details of the Ca-rich side are shown in the insets.

![Fig. 4](image2.png) Calcium environments of the Ca1 site (a) and Ca2 site (b) in the hydroxyapatite structure.

![Fig. 5](image3.png) Calcium environments of Ca1 site (a), Ca2 site (b), Ca3 site (c), Ca4 site (d) and Ca5 site (e) in the whitlockite structure.
whitlockite structure explains the fact that hydroxyapatite formation was inhibited when strontium was introduced during the synthesis process. The stabilizing role of Mg in whitlockite has also been reported.47 Like strontium, Mg2+ cation is located in the A column with a large preference for the Ca5 site in agreement with its small ionic radius.

3.4 In vitro Sr release and preliminary results on inflammation reactions in contact with Sr:HAp

The release of strontium after interaction in Sr-free culture medium (DMEM) is presented for HAp and Sr:HAp samples in Fig. 6. The detection level (around 0.04 ppm) is represented by the horizontal line. Whereas no significant amount of Sr is released from the undoped sample, a measurable quantity of strontium ions is detected for the Sr:HAp sample. The plateau concentration of ionic products (0.2 ppm) is reached after only a few days. This is an interesting amount for osteoporosis therapy: indeed it is in the same order of magnitude as that one measured in vivo in plasmas of animals treated with Sr-based drugs which were found to have obvious anti-osteoporotic effects.47 The easy delivery of Sr2+ ions in biological conditions can be related to the observed substitution of Ca for Sr in the whitlockite phase present in Sr-doped ceramics; according to the high solubility product of whitlockite compared to hydroxyapatite.47 Indeed, some substituted strontium in the hydroxyapatite phase is still present in the materials after this interaction time and could serve as a reservoir for future osteointegration of implants upon bone re-modelling. Optimization of the preparation could allow the ceramic composition (HAp/β-TCP) to be finely tuned and thus allow fine tuning of the delivery rate on different time scales.

3.5 Inflammatory process study: TNF-α production

In order to investigate how strontium-substituted HAp modulates TNF-α synthesis, monocyte cells were incubated with HAp and Sr:HAp powder for 6 h. After 6 h, no TNF-α protein was not detected in control cells. Due to the low production of TNF-α induced by the HAp powder, we stimulated cells with LPS to investigate if the strontium had an influence on TNF-α production (Fig. 7). The production of TNF-α significantly decreases for Sr-doped HAp powder (values normalized to LPS stimulated cells). Thus strontium has a positive effect on cytokine production possibly decreasing the inflammatory reactions of human monocytes in contact with HAp powder. Further work is under way to study the effect of Sr concentration and the production of other inflammatory and anti-inflammatory cytokines. Nevertheless, this is a first promising result allowing the use of Sr-doped HAp to limit the inflammatory process during HAp implantation.48,49

4. Conclusions

The sol–gel process has been successfully used to prepare Sr-doped calcium phosphates. All samples are porous and exhibit phase mixtures. With respect to the undoped sample, the Sr-doped sample exhibits a higher proportion of β-TCP phase and a significant amount of amorphous phase. As far as the substitution is concerned, Sr2+ ions are found in three phases, HAp, β-TCP and an amorphous phase defined as (Ca,Sr)O. These strontium ions are preferentially found in the crystalline phases (60%) both in HAp (32%) and β-TCP (28%) but a large amount (40%) are found in the amorphous phase. From these results, it is clear that doping can be efficiently used to modulate the phase proportions and thus the solution behaviour of the material. The effective partial substitution of Sr in HAp makes this material very interesting for orthopaedic applications. The Sr-HAp cell parameters agree fairly well with Vegard’s law and Sr2+ ions substitute for Ca2+ only in the Ca2 site at low levels of substitution. Sr substitution in whitlockite occurs preferentially in the Ca4 site which presents an unusually low 3-fold coordination in the pure Ca3(PO4)2 phase.

Furthermore, the presence of the Sr-containing amorphous phase and the Sr-doped soluble β-TCP phase are responsible for a partial release of Sr2+ ions in solution upon interaction in physiological fluids. The level of Sr2+ release in solution is very promising from the viewpoint of a possible anti-osteoporotic effect demonstrated by other commercial drugs. For the first time, to our knowledge, a significant anti-inflammatory effect of Sr-doped HAp has been demonstrated by measuring the production of TNF-α by human monocytes in contact with the materials. Considering all these results, these materials appear to
be good candidates for bone substitutes or bone tissue engineering applications combining good bioactivity, expected antiosteoporotic properties because of the presence of strontium and anti-inflammatory properties.

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