Rietveld Refinement of the Crystal Structure of Hydroxyapatite Using X-ray Powder Diffraction

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Abstract The stoichiometric hydroxyapatite, Ca₁₀(PO₄)₆(OH)₂ is prepared by an aqueous precipitation method at room temperature, the main reactants were Ca(OH)₂ and H₃PO₄ without addition of ammonia solution. The sample was analyzed by Fourier transformed infrared spectroscopy (FTIR) which reveals the presence of a small amount of carbonate due to absorbance of carbon dioxide from the air during synthesis process. The crystal structure was carried out by X-ray powder diffraction data and the Rietveld method using FullPROF software. We confirmed that this material has a hexagonal structure (space group P6₃/m; Z = 1). Unit-cell parameters with higher precision (a = b = 9.4159(4) Å, c = 6.8819(3) Å, α = β = 90°; and γ = 120°).

Keywords: hydroxyapatite, infrared spectroscopy, X-ray powder diffraction, Rietveld refinement

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1. Introduction

Although single-crystal and powder X-ray diffraction patterns contain essentially the same information, in the single-crystal case this information is distributed in three-dimensional reciprocal space, whereas in the powder case the three-dimensional diffraction data are effectively compressed into one dimension (intensity versus diffraction angle 2θ) as a consequence of the random orientational distribution of the crystallites in the powder sample. Such compression usually results an extensive overlap of peaks in a powder X-ray diffraction pattern. As a result, the process of extracting intensity data from the powder X-ray diffraction pattern can be unreliable and leads to difficulties in attempting to carry out structure solution using such data.

In orthopedic and dental surgeries fields, the most commonly used material for coatings is hydroxyapatite (HAp). The HAp Ca₁₀(PO₄)₆(OH)₂ is found in natural human hard tissue and considered as the main inorganic component of bone and teeth [1,2,3]. It has been proven that synthetic hydroxyapatite exhibits properties such as osteoconductivity and osteoinductivity which stimulate bone regeneration [4]. Due to its excellent biocompatibility, the HAp has been widely used as a coating material for implanted devices [5,6]. Another important characteristic of HAp is its biodegradability. Previous studies [7-12] have shown that hydroxyapatite tends not to trigger a response from the immune system, embedding itself successfully in the surrounding tissue and degrading over time as the host tissue begins to regenerate.

The hydroxyapatite structure was published nearly simultaneously by Náray-Szabó [13] and Mehmel [14] in 1930. It possesses a hexagonal structure with a P6₃/m space group and cell dimensions a=b=9.42Å, and c=6.88Å, where P6₃/m refers to a space group with a six-fold symmetry axis with a threefold helix and a mirror plane [15]. It has an exact stoichiometric Ca/P ratio of 1.67.

The aim of this study was to undertake an X-ray powder diffraction study of a nanometer scale crystalline HAp powder synthesized by a novel method.

2. Materials and Methods

2.1. Simple Preparation

The hydroxyapatite sample is prepared via an aqueous precipitation reaction of Ca(OH)₂ and H₃PO₄ as described in our previous paper [16]. In brief, the pure HAp was prepared by a precipitation method at room temperature (RT). After the total addition of the H₃PO₄ solution, the pH remained greater than 10 (no ammonia was therefore added) and the suspension was continuously agitated for an additional 2h at the same medium temperature and then matured over a period of 48 h at RT. This matured suspension was decanted and subjected to thermal treatment at 90°C for 24 h and ground to a fine powder in an agate mortar. The resulting oven-dried precipitate is calcined at 900°C in a dry air atmosphere in order to improve its crystallinity.

XRD measurements were performed for phase identification and phase purity with the use of a Shimadzu 6100 diffractometer equipped with a diffracted beam...
monochromator and CuKα wavelength. A single HAp phase (card #9-432) was identified in all specimens by using the ICDD (International Center for Diffraction Data) data bank.

The functional groups present in the prepared powder calcined at 900°C were identified by FTIR (Vertex 70 Spectrometer). To perform a IR spectrum, 1% of the powder was mixed and ground with 99% KBr. Tablets of 10 mm diameter for FTIR measurements were prepared by pressing the powder mixture under 5 tons for 2 min and the spectrum was taken in the range of 400 to 4000 cm⁻¹ with resolution 4 and 128 times scanning.

2.2. Rietveld Refinement Theory

A detailed description of the mathematical procedures implemented in the Rietveld analysis has been earlier reported [17,18,19,20]. Here, we give brief details of the analysis of experimental powder diffraction patterns that we’ve done. The crystal structure characterization of the synthesized sample has been made by employing the Rietveld’s whole-profile fitting method using FullPROF software [21].

The experimental profiles were fitted with the most suitable pseudo-Voigt analytical function. For both the Kα1 and Kα2 profiles, the line broadening function B (2θ)

\[ pV(x) = \eta L(x) + (1 - \eta) G(x). \]

The L(x) and G(x) are, respectively, the Lorentzian and Gaussian components. Considering the integrated intensity of the peaks as a function of structural parameters only, the Marquardt least-squares procedures were adopted for minimization of the difference between the observed and simulated powder diffraction patterns. The minimization was carried out using the reliability index parameter such as the residuals for the weighted pattern Rwp, the pattern Rp, Braggs factor RBragG, structure factor RF, and goodness of fit χ². All these parameters were used as numerical criteria of the quality of the fit of calculated to experimental diffraction data and are represented by the following relations [22]:

\[ R_{wp} = 100 \left[ \frac{\sum w_i (y_{i(obs)} - y_{i(cal)})^2}{\sum w_i y_{i(obs)}^2} \right]^{1/2} \]

\[ R_p = 100 \left[ \frac{\sum y_{i(obs)} - y_{i(cal)}}{\sum y_{i(ols)}} \right] \]

\[ R_B = 100 \left[ \frac{\sum I_{i(obs)} - I_{i(cal)}}{\sum I_{i(obs)}} \right] \]

\[ R_{exp} = 100 \left[ \frac{(N - P + C)}{\sum w_i y_{i(ols)}^2} \right]^{1/2} \]

\[ \chi^2 = \left[ \frac{R_{wp}}{R_{exp}} \right]^2. \]

where \( y_{i(obs)} \) is the experimental intensities, \( y_{i(cal)} \) is the calculated intensities, \( w_i = (1/y_{i(obs)}) \) is the weight experimental observations, N is the number of experimental observations, P is the number of fitting parameters, C number of constraints and I is the integrated intensity. The goodness of fit indicator (GOF) is established by comparing \( R_{wp} \) with the expected error \( R_{exp} \). This leads to the value of goodness of fit:

\[ \text{GOF} = \frac{R_{wp}}{R_{exp}}. \]

3. Results and Discussions

The X-ray powder diffraction pattern of the HAp sample is shown in Figure 1. The XRD pattern shows the characteristic peaks of hydroxyapatite, according to the International Center for Diffraction Data database, ICDD-PDF 9-0432. The XRD pattern indicates the presence of a majority phase corresponding to the hexagonal singony – P63/m space group associated to pure crystalline nanometer sized HAp. In addition, no other minority phase has been detected, within the X-ray detection limits of ~ 2%.

![Figure 1. X-ray powder diffraction pattern of Hap sample](image1)

![Figure 2. FTIR of as-synthesized hydroxyapatite sample](image2)

The functional groups present in the prepared sample were investigated by FT-IR studies (Figure 2). The spectra registered for the hydroxyapatite showed the apatite characteristic bands. The presence of phosphate group PO₄³⁻ is highlighted by the 474 cm⁻¹ band associated to the v2 double degenerated bending mode and the bands found at 570 and 603 cm⁻¹ characteristic to the ν₄ vibration mode. Furthermore, the band registered at 962 cm⁻¹ is evidence of the ν₁ non-degenerated symmetric stretching mode of the P-O bond and the bands found at 1040 and 1098 cm⁻¹ are associated to the ν₃ vibration mode of the functional group. Sharp bands at 3573 cm⁻¹ and 633 cm⁻¹ correspond to stretching and vibrational modes of hydroxyl vibrations,
respectively. On the other hand, the CO$_3^{2-}$ functional group is highlighted by the presence of several vibration bands. The band found at around 875 cm$^{-1}$ can be characteristic to the ν$_2$ vibration mode and the stretching vibrations of the CO$_3^{2-}$ group are highlighted by the band in the region 1400 and 1500 cm$^{-1}$. The carbonate presence in the specimens is due to absorbance of carbon dioxide from the air during synthesis of the sample and it has been reported by others [23,24].

3.1. Peaks’ Search and the Initial Structure Determination

The refinements were performed using the computer program FullProf which has been mainly developed to perform Rietveld analysis. The average intensity profile Analysis using the peak-search tool in the program WinPLOTR [25] yielded 28 peaks between 20° and 60° in 2θ range. These peak positions were used as input for the program DICVOL [26] which is an automatic program for finding the unit cell. As a result, its basic structure is a Hexagonal system with lattice parameters a = b = 9.4153 Å and c = 6.8822 Å. Determination of the most likely space groups was achieved by the program CHECKGROUP. The best solution resulted in space group is P6$_3$/m.

3.2. Refinement of XRD Data

The inherent parameters determined by the standard data and other experimental parameters were used as input data in the program FULLPROF, the atomic positions reported in the literature [27,28] were used for the initial refinement. The pseudo-Voigt profile function of Thompson, Cox and Hastings [29] was employed to model the peak shape of the Bragg reflections. The refinement procedure was started by refining: the scale factor, zero point of detector, 1$^{st}$ background parameter (polynomial function of sixth order) and lattice parameters; then gradually more parameters were opened; the peak profile (pseudo-Voigt) parameters, atomic coordinates and isotropic displacement parameters B. In the final step the individual anisotropic thermal parameters (ATPs) could be improved. Attempts to refine the ATPs of the hydrogen atom were not successful, then Isotropic displacement parameter is Kept fixed at the value 1.5. The structure refinement was stable at all stages and converged rapidly to the results summarized in Table 1.

| Table 1. Lattice parameters and agreement factors for HA |
| --- | --- | --- | --- | --- | --- |
| **Unit cell parameters** | a=b=9.4159 Å; c=6.8819 Å; α=β=90°; γ=120° |
| Number of space group | 176 |
| Hermann-Maguin symbol | P6$_3$/m |
| Volume (Å$^3$) | 528,400 |
| Bragg R-factor | 1.82 |
| R$_f$-factor | 1.50 |
| R factors | Rp=9.99; R$_{wp}$=12.8; R$_{exp}$=9.43 |
| Chi squared $\chi^2$ | 1.84 |
| GOF | 1.35 |

![Figure 3. Final Rietveld refinement of the HAp sample: calculated (black solid line), experimental (red dots mark) X-ray powder diffraction pattern and difference curve (blue)](image)

Table 2. Atomic coordinates and occupancies for sample HAp

| Atom | Wyckoff position | x | y | z | Occ | Biso |
| --- | --- | --- | --- | --- | --- | --- |
| O1 | 6h | 0.3259 | 0.48244 | 0.25000 | 5.9154 | 2.4421 |
| O2 | 6h | 0.5839 | 0.46367 | 0.25000 | 5.8946 | 1.3273 |
| O3 | 12i | 0.3381 | 0.25338 | 0.06712 | 12.1170 | 1.6715 |
| O(H) | 4e | 0.0000 | 0.00000 | 0.22352 | 2.0015 | 1.0004 |
| H | 4e | 0.0000 | 0.00000 | 0.16450 | 1.9402 | 1.0004 |
| Ca1 | 4f | 0.3333 | 0.66667 | 0.00112 | 4.0060 | 1.8642 |
| Ca2 | 6h | 0.2484 | 0.99304 | 0.25000 | 5.9143 | 1.2372 |
| P | 6h | 0.3980 | 0.36911 | 0.25000 | 5.9393 | 1.0943 |

Table 3. Anisotropic displacement parameters for HAp sample

| Atom | U$_{11}$ | U$_{12}$ | U$_{13}$ | U$_{22}$ | U$_{23}$ | U$_{33}$ |
| --- | --- | --- | --- | --- | --- | --- |
| O1 | 0.00879 | 0.01597 | 0.00391 | 0.00391 | 0.01477 | 0.00000 |
| O2 | 0.01400 | 0.00091 | 0.00008 | 0.00008 | 0.00501 | 0.00000 |
| O3 | 0.01300 | 0.00530 | 0.00078 | 0.00078 | 0.01042 | 0.00116 |
| O(H) | 0.00545 | 0.00545 | 0.03762 | 0.03762 | 0.00273 | 0.00000 |
| Ca1 | 0.00763 | 0.00763 | 0.00809 | 0.00809 | 0.00381 | 0.00000 |
| Ca2 | 0.00482 | 0.00151 | 0.01071 | 0.01071 | 0.00103 | 0.00000 |
| P | 0.00367 | 0.00325 | 0.00762 | 0.00762 | 0.00215 | 0.00000 |
The calculated and experimental diffraction profiles are shown in Figure 3 together with the difference curve obtained after the final refinement, the dots mark is the experimental pattern and the solid line is the calculated diffraction pattern. The lower trace is the difference between observed and calculated patterns, and the vertical lines mark the positions of the calculated Bragg peaks. It is worth noting the good agreement between both of them. Atomic coordinates, occupancies, lattice parameters and agreement factors for HAp are collected in Table 2. The anisotropic displacement parameters for Hap are listed in Table 3.

The maximum theoretical occupancy for O and H at the 2a position is 2. Sample HAp shows 2 and 1.94 for O and H, respectively. Considering that the H atoms are forming the hydroxyls, our results show that there are 1.93 H atoms per unit cell combined with 1.93 O atoms which will result in the same amount of OH. The composition at the 2a Wyckoff position in HAp was calculated to be: \((\text{OH})_{0.07}\)

The fact that the sites of the O1, O2 and P atoms show some vacancies, whereas the sites of O3 maintain significantly high occupancy, undoubtedly show evidence for replacement of \(\text{PO}_4^{3-}\) by \(\text{CO}_3^{2-}\) ions in the apatite lattice. Figure 3 shows the projection of the HAp structure on (001) plane. It reveals two types of tunnels (Type I and type II) which give it an ionic exchange property (substitution) either on calcium site or on the site of phosphorus and hydroxyl.

4. Conclusion

A single-phase of HAp was prepared following an aqueous precipitation method. The XRD pattern suggests that the prepared sample has the hydroxyapatite characteristic structure, the FTIR spectra suggests the presence of HAp characteristic bands associated mainly to phosphate, hydroxyl and carbonate groups. The crystal structure was refined by the Rietveld analysis and revealed that HAp obtained is not pure hydroxyapatite but oxy-hydroxyapatite slightly carbonated \(\text{Ca}_{10}(\text{PO}_4)_{6-x}(\text{CO}_3)_x(\text{OH})_2\).

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