Original Research Paper

Production and Evaluation of Hydroxyapatite (HAp) Properties of Broiler's Composite Bone (BCB) Waste at Different Sintering Temperatures

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Article history Received: 06-10-2017 Revised: 28-03-2018 Accepted: 7-05-2018

Corresponding Author: Muhammad Irfan Said Lab. of Animal By-Products Processing Technology, Faculty of Animal Science, Hasanuddin University, Makassar, Indonesia Tell/Fax: (+62-411) 587217 Email: irfan.said@unhas.ac.id Abstract: Bone is one of the perishable livestock wastes, so serious attention is needed. The wastes are produced by Poultry Slaughterhouse (PSh) primarily in the process of separating meat from bone to produce boneless products. The main constituent of bone is the crystals Ca and P. which are the major elements of Hydroxyapatite (HAp). The purpose of this study was to produce HAp from the Broiler's Composite Bone (BCB) waste as well as to evaluate the properties of BCB at different sintering temperatures. HAp was produced through 3 stages, i.e., calcination, precipitation and sintering. The calcination was carried out at 1000°C for 5 h and the resultant was evaluated by X-Ray Diffraction (XRD) and followed by precipitation with HPO₄ 0.3 M. The next stage was sintering process, using 6 level temperatures of 550°C, 600°C, 650°C, 700°C, 750°C, 800° and followed by XRD analysis. The results showed that calcination at 1000°C for 5 h demonstrated the efficiency of 41.99%. The intensity peak of calcination of the BCB was found at 20: 32.5250°, 19.7200° and 21.1200° (before calcination) and 32.5187°, 33.5626° and 33.0000° (after calcination). Sintering condition demonstrated significant effects on HAp yield. We also found that sintering process at 700°C (97.2%) was the most optimum temperature to produce HAp from BCB.

Keywords: Hydroxyapatite (HAp), Composite Bone, Broiler, Calcination, Sintering

Introduction

The waste products from poultry slaughterhouse (PSh) industry have remained a greater challenge (Enginuity worldwide LLC, 2017; Bodirsky *et al.*, 2014; Khan and Ghouri, 2011) because of their susceptibility to deterioration (Centner, 2004). Bone constitutes is one of the most abundant solid wastes in PSh industry (Arshadi *et al.*, 2015) and may transmit diseases. On the contrary, many beneficial applications of the bone were reported such as catalyst (Corro *et al.*, 2016) and

inexpensive organic fertilizer (Gousterova *et al.*, 2008). Without further use, the bone waste might be a serious pollutant that contaminates air, water and soil, especially in PSh (Kazemi-Bonchenari *et al.*, 2017). Bone contains a valuable component that needs to explore. It was reported to have a high content of calcium and phosphorus (Gopi *et al.*, 2014). Its high mineral content may represent the adequacy of mineral supply contained in the feed (Onyango *et al.*, 2003). As by-product, the bone is not only applicable for fabrication of feed, but also for environmentally friendly packaging material



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(Said *et al.*, 2016; 2011). The broiler's composite bone refers to bone in the whole body except claw and head.

In Indonesia, the broiler's composite bone (BCB) is abundant and needs a further exploration. Based on MA (2015), broiler population reached 1,528.33 million. If assumed 70% of the population is slaughtered, the number of slaughtered broiler per month is 127.26 million (70% ×1,528.33 million/12 months). The bone waste accounts for 20.85% of total body weight. With average weight of 1.5 kg per head, the potential bone waste reaches 39.83 million kg per month (20.85% ×191.04 million kg = 39.83 million kg).

In term of chemical components, the bone is rich in collagen and calcium (Ockerman and Hansen, 1999). Collagen extract is now widely used as an antiosteoporosis and anti-aging food supplement (English, 2011; English and Cass, 2001). Furthermore, duck beak bone was reported to be a promising source of of Hydroxyapatite HAp (Son et al., 2014). The collagen extract and HAp could be used as a composite material (Imanieh and Aghahosseini, 2013). Chest skin of broiler was also a good source of collagen, especially collagen type I (Bilgen et al., 1999). The extraction of collagen from bone produced HAp $[Ca_{10}(PO_4)_6(OH)_2]$ by $\pm 93\%$ as by-product (Niakan et al., 2013; Gabriela et al., 2008). Previous studies reported that HAp could contribute to the bone regeneration (Kattimani et al., 2016) and serve as biomaterial in medicine science by means of HAp-collagen engineering (Shibata et al., 2005; Khiri et al., 2016; Heinz et al., 2010; Acharya et al., 2016). In dentistry, HAp was useful material for repairing and treating teeth and dental parts (Pepla et al., 2014; Görken et al., 2013).

The properties of HAp are highly depending on source and preparation conditions (such as heating temperature). The use of cattle's bone, shellfish and rock for fabrication of HAp has been studied (Khiri *et al.*, 2016), whereas study pertaining broiler' bone composites for HAp production is rather scarce. This research was very important to increase the added value of BCB waste and reduce the pollution effect of PSh; thus, our study aimed to produce and evaluate the properties of HAp produced from broiler's composite bone at different sintering temperatures.

Materials and Methods

Materials

The BCB waste was obtained from PSh of Daya village, Makassar, Indonesia. The other materials included aquadest, $(NH_4)_2HPO_4$ 0.3 M, Whatman filter paper No.42 and aluminum foil. The instruments were X-Ray Diffraction (XRD) (*Shimadzu 6000*), furnace (*Furnace 6000*), oven (*Memmert*), analytical scale (*Sartorius TE 214S*), magnetic stirrer (*C-MAG HS 7*),

thermometers, desiccators, stop watch (*General*), beaker glass (*Pyrex*), Erlenmeyer (*Pyrex*), measuring cylinders (*Pyrex*) and volume pipettes (*Pyrex*).

Methods

BCB waste (180 g) was cleaned from meat and fat, washed and then weighed. The HAp was produced by using three following stages, i.e., calcination, precipitation and sintering. The method was carried out according to Khiri *et al.* (2016).

Stage 1. Calcination

BCB was heated at 1000°C for 5 h using furnace, leading to removal of organic materials and conversion of CaCO₃ into CaO through elimination of CO₂ gasses (Equation 1). At this stage, the potential amount of waste was also determined. BCB (60 g) was transferred into 6 porcelain containers (10 g BCB per container). The Calcination Efficiency (CE) of the BCB was calculated as follow: CE = A/B ×100%, where A = weight (g) of BCB after calcination; B = weight (g) of BCB before calcination. XRD analysis was also performed to evaluate the resultant of calcination (using CaO as a standard) according to Joint Committee on Powder Diffraction Standard (JCPDS) 82-1691:

$$CaCO_3 \to CaO + CO_2 \tag{1}$$

Stage 2. Precipitation

The calcined BCB was powdered using electrical grinder to obtain a particle size of 45 μ m. The powder was used to prepare 1.0 M of calcium hydroxide (CaOH)₂ solution by mixing with aquadest (100 mL) at 40°C, as previously determined using a stoichiometric equation. The solution was stirred using a magnetic stirrer for 2 h to obtain a uniform mixture of Ca(OH)₂. The reaction process occurs according to Equation 2:

$$CaO + H_2O \to Ca(OH), \tag{2}$$

To produce HAp powder, $Ca(OH)_2$ suspension was added with 100 mL of phosphate acid (HPO₄) solution at a rate of 15-20 drops/min. Next, the solution was stirred with a magnetic stirrer and maintained at pH 8 by adding ammonium hydroxide (NH₄OH) solution to complete the reaction then stored for 24 h at room temperature. The ratio of Ca:P was 1.67: 2. The white precipitate formed was then filtered using Whatman no 42 paper and dried for 5 h at 110°C. The results were weighed using analytic scales. The reaction process was presented in Equation 3:

$$10Ca(OH)_{2} + 6H_{3}PO_{4} \rightarrow Ca_{10}(PO_{4})_{6}(OH)_{2} + 18H_{2}O$$
 (3)

Stage 3. Sintering

Sintering refers to heating process below melting point to form a new crystal phase as desired. This process aims to assist chemical reactions of materials making up both ceramic and metal. The HAp powder was sintered at a temperature level of 550°C, 600°C, 650°C, 700°C, 750°C, 800°C (50°C intervals) using a furnace for 1 h. After that, the HAp powder was cooled in a furnace at room temperature. The results were weighed to determine the efficiency. The Sintering Efficiency (SE) of BCB was calculated as follow: CE = A/B $\times 100\%$, where A = weight (g) of BCB after sintering, B = weight (g) of BCB before sintering. Furthermore, to know the characteristics of $Ca_{10}(PO_4)_6(OH)_2$ produced, analysis using XRD (Shimadzu 6000) was carried out. The results of sintering were also analyzed using a diffractometer at the diffraction angle (2θ) with scanning distance of 5°. The standard for characterization of HAp (Ca₁₀(PO₄)₆(OH)₂) product followed the Joint Committee Powder Diffraction Standard (JCPDS) 18-0272.

Parameters of Study

The study parameters were (1) Calcination Efficiency (CE), (2) X-Ray diffraction of calcination, (3) Sintering Efficiency (SE) and (4) X-Ray diffraction of sintering.

Experimental Design and Statistical Analysis

The pattern of X-Ray diffraction (diffractogram) was descriptively analyzed, while CE and SE parameters. The data were evaluated using Analysis of Variance (ANOVA). Significant difference between means was compared using Duncan's Multiple Range Test (DMRT) at 5% (Steel and Torrie, 1991).

Results and Discussion

Calcination Efficiency (CE)

The CE value is important to evaluate the effectiveness of the calcination process. High efficiency was expected in a production process. The CE of BCB at calcination condition of 1000°C for 5 h was presented in Table 1.

HAp, a type of bioceramic, has been widely used as an alternative material of artificial bone. It is selected for several biological reasons such as biocompatibility, bio-affinity, bioactivity, osteoconduction (Dubok, 2000), osteo-integration (Hench and Thompson, 2010) and osteo-induction (under certain conditions) (Weiner and Wagner, 1998). In composition, HAp was composed of calcium and phosphate ions, so there has been no reported adverse local or systemic toxicity. When HAp was applied to a new bone, it directly binds HAp through the apatite layer. This occurs because of the lack of calcium carbonate in bone implants (Dorozhkin, 2012; LeGeros, 2008).

The results showed that CE of BCB at heating temperature of 1000°C for 5 h to produce CaO was 41.99% (Atirah, 2017). This CE was lower than that reported by Putri (2012) using different sources of bone, i.e., chicken egg shell of (54.04 ± 1.00) , duck egg shell (55.03 ± 1.30) and quail egg shell (53.19 ± 3.00) . The low CE can be caused by the insufficient temperature and processing time. The heating process eliminates all organic materials. Chicken bone shows a high crystallinity with single phase pattern. High crystallinity can increase the mechanical strength of the HAp. However, in excessive level of crystallinity, this may also complicate the absorption process in the body compared with other materials (Suzuki *et al.*, 2005).

In recent years, the heating process was recommended to remove organic compounds (Lin et al., 1999). The crystalline phase composition of the sintered bone was similar to that of the natural bone of 93% HAp (Ca₁₀(PO₄)₆(OH)₂) of bone weight and about 7% βtricalcium phosphate (Ca₃(PO4)₂, β -TCP) (Martin, 2000). The heat treatment yields an interconnected porous structure (up to 70% porous). Therefore, it allows for faster bone growth (Lin et al., 1999). The ratio of Ca/P ratio also affects the mechanical properties of HAp. The greater the ratio of Ca/P results in the higher strength, but the maximum ratio is 1.67. The mechanical properties of HAp decreased at Ca/P ratio of >1.67 or <1.67. In recent studies, many researchers have tried to extract HAp from natural materials such as coral, shell, eggshell, cuttlefish shells, natural gypsum and natural calcite (Herliansyah et al., 2012). The raw materials were CaCO₃ or CaSO₄. Some chemical reaction processes were performed to produce HAp. In our study, since BCB is from living beings, the production of HAp from BCS has certain advantages, including the reduction of polluted waste. Additionally, the BCB from Psh and meat processing industry is abundant and cheap. Thus, this suggests that the BCB may substitute the role of other natural ingredients.

X-Ray Diffraction of Calcined BCB

The diffraction profile of BCB was important to evaluate the potential content of HAp present in the sample. The standard used in this evaluation was Joint Committee on Powder Diffraction Standard (JCPDS) 82-1691 special for CaO. The diffractogram of BCB before and after calcination at 1000°C for 5 h was obtained by using X-Ray Diffraction (XRD). As presented in Fig. 1, diffractogram shows a similarity pattern to JCPDS 82-1691. This indicates a process of converting CaCO₃ into CaO by eliminating CO₂ gasses Fig. 1. Demonstrates that BCB can be used as a candidate of bone replacement material according to its corresponding crystallinity properties. The BCB crystallization can be controlled by using different

levels of time and temperature when the preparation process begins (Ooi *et al.*, 2007).

Different intensity of X-Ray absorption in BCB before and after calcination is presented in Table 2.

Table 1	CE of of BCB at calcination condition o	of 1000°C for 5 h

Repetition	Before calcination (g)	After calcination (g)	CE (%)
1	15.3255	6.2423	40.73
2	12.7125	5.2624	41.40
3	12.3155	5.1060	41.46
4	10.5020	4.6418	44.20
5	12.4025	5.1237	41.31
6	7.4982	3.2124	42.84
	Average		41.99±1.29

Table 2: Intensity of X-Ray absorption of BCB before and after calcination at 1000°C for 5 h

No	Peak No.	2 Theta (θ) (deg)	D(A)	I/I1	FWHM (deg)	Intensity (Counts)	Integrated Int (Counts)
a. Before calcination	s Strongest 3 pe	aks				. ,	` <i>`</i> _
1	14	32.5250	2.75069	100	1.73000	181	14456
2	5	19.7200	4.49833	48	0.0000	87	0
3	6	21.1200	4.20320	36	0.0000	66	0
b. After calcination S	Strongest 3 peak	S					
1	19	32.5187	2.75121	100	0.51750	143	3637
2	21	33.5626	2.66799	62	0.53870	88	2908
3	20	33.0000	2.71218	44	0.00000	63	0

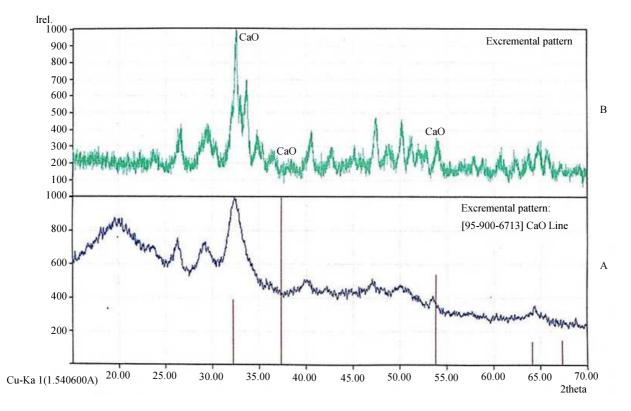


Fig. 1: Diffractogram of CaO crystal formation using X-Ray on BCB before (A) and after (B) calcination at 1000°C for 5 h

Based on Table 2 and Fig. 1, we found the 3 highest peaks of CaO crystallization present before and after calcination. The crystallinity degree of CaO as the main HAp constituent can be observed from its diffraction patterns at the peaks. The maximum intensity peak of BCB was found at 2θ : 32.5250° ; 19,7200° and 21,1200° (before calcination) and $32,5187^{\circ}$; $33,5626^{\circ}$ and $33,0000^{\circ}$ (after calcination) (Atirah, 2017). The molarity of Na₃PO₄ affects content of CaO. Higher level of CaO was responsible

for higher purity degree of HAp. At molarity >5 M, this may damage the crystal structure of the material, leading to reduction of CaO concentration.

Sintering Efficiency (SE)

The formation of HAp crystal involves sintering process after precipitation. The standard for characterizing HAp was the Joint Committee on Powder Diffraction Standard (JCPDS) 18-0272. The SE value was presented in Fig. 2.

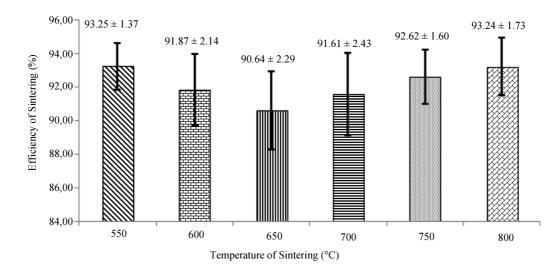


Fig. 2: SE of BCB after sintering process at various temperatures of 550-800°C for 1 h (temperature interval of 50°C)

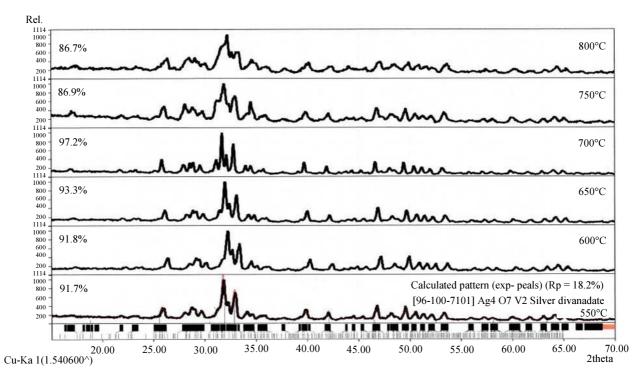


Fig. 3: Diffractogram of HAp formation using X-Ray of sintered BCB

As depicted in Fig. 2, different sintering temperatures showed no significant effects on the amount of dried HAp precipitate (p>0.05). This suggests that sintering temperatures (550-800°C) could be performed to produce HAp precipitates. The amount of dry sintering precipitates did not affect the levels of HAp in the precipitates. Therefore, to know the amount of crystal content of HAp, characterization using XRD was required (Fig. 3).

X-Ray Diffraction of Sintered BCB

Figure 3 exhibits the diffractogram of sintered BCB at temperature variations of 550-800°C for 1 h. The results showed that sintering temperature of 700°C resulted in peak with the most optimum intensity, suggesting that this sintering condition could produce HAp crystallinity at optimum level. Powdered HAp has a great potential as the bone replacement. This is because it was easily absorbed by the bones and can be received by the body. There were two kinds of HAp powder, i.e., micro and nano sized HAp. Micro HAp has a small surface area and has a strong crystal bond, so that easily absorbed by the body. In addition, it can also increase biostability and strength. However, HAp with nanostructures has better density, strength and bioactive properties (Pepla *et al.*, 2014).

Conclusion

The calcination process to produce CaO at 1000°C for 5 h resulted in CE of 41.99% ± 1.29 (n = 6). Diffractogram of maximum intensity of BCB before calcination was at 20: 32,5250°; 19,7200° and 21,1200°, whereas after calcination was at 20: 32,5187°; 33,5626° and 33,0000°. The sintering temperature (550-800°C) showed no effect on a number of dry precipitates of HAp. The sintering temperature at 700°C (97.2%) was the most optimum sintering temperature to produce HAp from BCB.

Acknowledgment

The authors would like thank to the Ministry of Research, Technology and Higher Education, Republic of Indonesia, the Rector of Hasanuddin University (Unhas) and the Institute for Research and Community Service of Hasanuddin University for providing financial support under Research Grant Program of Riset Unggulan Unhas (Runas).

Author Contribution

Muhammad Irfan Said: Participating in all experiments, data analyzing and manuscript writing preparation.

Effendi Abustam: Participating in all experiments, collecting the field and performing statistical analysis.

Asdar Gani: Composing the experimental design and assiting data analysis.

Paulina Taba: Preparing the manuscript and analyzing data.

Atirah: Collecting the field data and contributing to the writing of the manuscript.

Ethics

This article is original and has not been published or presented elsewhere. All the authors have approved the manuscript and agree with submission to this journal. There is no conflict interest to be declared.

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