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Padang, Indonesia, July 25-26, 2018

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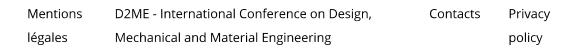
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Preface ICTIS S2B

This issue of Proceedings gathers the papers presented at 2018 the International Conference on Technology, Innovation, Society and Science to Business (ICTIS S2B 2018) held on July 25-26, 2018 in Padang, West Sumatra Indonesia. This special conference is a merger of the 2nd International Conference on Technology, Innovation, Society (ICTIS) and the 15th International Science-to-Business Marketing Conference. Very significant subjects of the future of our societies and economies will be tackled and addressed at the conference and allow the participation from all over the world. ICTIS S2B finally accepted 52 papers after a double blinded peer review process by international reviewers and technical program committee members.

Six keynote speeches were presented from Prof. Thomas Baaken; Dr. Ing Ilham Habibie; Prof. Gideon Maas, Arno Meerman, Arif Musta'in, and Antono Pasha, will be presented during the two-day conference. All the talks were very impressive for the high level of professionalism, and in many cases original ideas and activities have been accomplished or proposed.

The international environment will be a breading place for great discussions, new connections and inspiring talks. The different conference activities, such as keynote presentations, facilitated expert panels, scientific presentations and practical workshops provide a brought range of new knowledge and creative solutions. Participants will gain new ideas, models, instruments and neat pieces of knowledge and innovation to apply in practice, either as an academic or practitioner. We are also indebted to session chairs, international reviewers, conference secretariat who dedicated to make the conference run smoothly and properly and ensure the proceedings quality. Last but not the least, we should express our thanks to all delegates, who showing the high level of international interest in the subject. It is exactly your participation that make the conference to its success.

The Proceedings provide the permanent record of what were presented. It indicated the state of development at the time of writing of all aspects of this important topic and will be invaluable to all researchers in the field for that reason. We truly believe the participants will find the discussion fruitful, and we hope you enjoy and find your engagement with their ideas valuable in sustaining your own professional development.

Final thanks go to all authors and participants at ICTIS S2B 2018 for helping to make it a successful event.

Yours Sincerely Conference Organizing Committee July 24, 2018





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August 25, 2018

Solid-State Fermentation of Banana Peels Potential Study for Feed Additive

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Abstract. Agricultural solid wastes present abundantly on earth as crops harvesting as well as processing are countinuesly run. Banana peels are one of agricultural solid wastes produced anywhere the banana processing presents. The peels present abundantly in tropical countries such as Indonesia. The carbohydrate content of banana peels make it useful for the production of many chemicals, including feed. Meanwhile the large need in feed in farming including fish farming could prevent farmer to obtain substantial profit. This research studied the possibility of banana peel as one of abundant Indonesian agricultural solid waste to be utilized as fish feed which is known requiring certain level of protein content. This was done by fermenting the peels in fixed bed reaction mode using surface aeration and non-aeration. The fermentation was conducted using yeast Saccharomyces cerevisiae Y1536 and Rhizopus Oryzae FNCC 6157. The reaction time was varied for 1, 3, and 5 days. The important parameters studied were protein contents, and amylase activity of the fermented banana peels. Despite aeration indicated more operational cost, it showed significant impact on the fermentation of banana peels. The best condition for fermentation using Saccharomyces cerevisiae Y1536 were 5 day fermentation with surface aeration which result in the increase of protein content up to 4.05%, the decrease of fiber content up to 1.08%, and amylase activity of 9.99 DP. Whilst the fermentation using Rhizopus Oryzae FNCC 6157 obtained its best result at 1 day fermentation with aeration, which are protein content increase up to 4.04% and fiber content decrease up to 0.69%. However, the fermentation using this mold showed its best amylase activity result of 12.75 DP at 5 day surface aerated fermentation.

1 Introduction

Waste produced from domestic, agriculture, and industrial activity has found causing many environemtal and health problems. Agricultural activities produces several waste including solid wastes, which in spite of causing environmental and health problems, they store economical potential to be transformed either chemically or non chemically into more beneficial products. Agricultural wastes which can be liquid, slurries or solids, are byproducts of growing and first processing of raw agricultural products such as fruit and vegetables, crops, meat and poultry as well as dairy products. Manures, fertilizers, and residues from slaughterhouse are also included as agricultural solid wastes [1]. Currently, many researches have attempted to utilise agicultural solid wastes to produce more valuable products and improve economy such as fired clay materials for building manufacturing, green desiccants in desiccantcooling systems, biofuels, bio-oil and char, and animal feed [2-6]. Generally agricultural wastes still contain useful nutrition components which are good for animal consumption. The potential of agricultural solid waste

for animal feed should be encouraged because of its benefit for both environment and the farmers.

In fish farming activities, the productivity of fish farmers is usually constrained by production cost where the feed cost can cover up to 50-80% of the total operational cost [7-9]. Protein is the main ingredient of the commercial diet of fish which constitutes 40-56% of the diets [10-11]. Many kinds of fermented fiberrich agricultural waste had been used for fish or shrimp feed additives such as papaya processing waste, corncob, rice husk and apple pomace [9, 12-14]. This carbohydrate-rich and low protein content agriculture wastes can be low cost fish energy source. However, carbohydrate amylolytic activity in carnivorous fish is lower than herbivorous fish [15]. Therefore, adding amylase activities on the carbohydrate feed will be advantageous.

Nutrient enrichment had been attempted by fermentation process which can increase protein content, required metabolic enzymes, and pathogenic bacteria inhibitory substance. Several fungi had shown to have either amylolytic or cellulolytic activities and therefore can be used to ferment carbohydrate-rich wastes while

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after fermentation they can provide protein sources needed for feed component. Some fungi such as Saccharomyces cerevisiae, Candida utilis, Aspergillus niger, and Rhizopus oligosporus have been known to be able to increase the protein content of biomass through fermentation [9]. Fermented corn-cob by mold consortium consisting of Trichoderma viridae, Trichoderma reesei, Aspergillus oryzae, and Rhizopus oligosporus had been used in feed formulation of Java barb fish and had been shown to increase the growth of Java barb [13]. Yeast protein is easily digestible and widely used for fish and shrimp feed supplement which in spite of providing protein source, it can enhance immune system and stimulating bacterial disease resistance [12]. When used as feed additive, the increased protein content of apple pomace after fermentation using G. butleri had been able to increase 44% in body mass of fish [16].

This research attempted to investigate the effect of solid-state fermentation of abundantly available waste banana peels by fungi *Saccharomyces cerevisiae* and *Rhizopus oryzae* on the protein and fiber content as well as the amylase activity of the peels in order to see its potential for fish feed additive. Solid-state fermentation itself has many applications including production of a large number of enzymes used in feed industry [17] The expected results were the increase of protein content of the banana peels and the availability of amylolitic activities of fermented banana peels.

2 Materials and Methods

2.1 Materials and Preparation Stage

The banana peels used in this research is from the species of *Musa acuminate* or *Musa balbisiana* or in Indonesian name, Pisang Kepok. The peels were collected locally from market and banana processing food seller. Immediately after collection, the peels were washed and cut into small size (1x 0.2 cm). The protein and fiber content of the peels were analyzed before fermentation.

There were two types of fermentation microorganism used in this research. The first type is yeast *Saccharomyces cerevisiae* Y1536 obtained from Institute Pertanian Bogor culture collection (IPBCC) while the second type is mold *Rhizopus oryzae* FNCC 6157 obtained from Food and Nutrition Development and Research Center (FANDARC), Gadjah Mada University (UGM). The fungi cultures were cultivated using potato dextrose agar (PDA).

2.2 Fermentation Stage

Prior to fermentation, the fungi cultures being used in the fermentation stage were transfered into 25 mL potato dextrose broth (PDB). The broths were shaken for 24 hours at speed of 150 rpm using incubator shaker.

The fermentations were conducted using tray solidstate bioreactors set up from Erlenmeyer flask in which the banana peels were stacked. As much as 225 grams of banana peel was firstly steamed for 30 minutes inside Erlenmeyer flask. It is then cooled and inoculated with 24 hour broth innoculum in biological safety cabinet to prevent contamination. The fermentation system set up is shown on Fig 1. This fermentation was conducted with and without aeration. To prevent contamination from outside air, membrane filters were mounted on the top of the Erlenmeyer flasks. The reaction time was varied 1, 3, and 5 days. At the end of fermentation, the peels were analyzed for the protein and fiber content as well as amylase activity.



Fig. 1. Solid-state fermentation set up

2.3 Chemical and Biological Analysis

The protein contents of the fermented banana peels were analyzed using total nitrogen (Kjelhdal method). The fiber contents were analyzed using fiber anaylisis by SNI 01-2891-1992. The amylase activities of the fermented banana peels were assayed using method by Fuwa (1954) [18]. The method was colorimeterically and the activity was expressed as dextrinising power (DP). Principally the substrate for dextrinising power determination used was commercial cassava amylose which was prepared according to Fuwa method. The values of DP were measured by the change of blue values of amylose-iodine complexes. Amylase unit (1 DP) was defined as the amount of amylase which would produce 10 percent fall in blue colour intensity of amylase-iodine complexes.

3 Results and Discussions

The locally collected banana peels were analysed for the total carbohydrate, fiber, lipid, protein, water, and ash contents. The result showed that the banana peels studied contained 79.59-85.66 % water. By dry basis, the peels also contained 55.05-58% total carbohydrate, 6.5-8.18% fiber, 7.36-12.23% lipid, 2.08-3.18% protein, and 13.97-20.24% ash. The result showed that the peels were rich in carbohydrate and were very suitable for energy sources. However, in order to be used as fish feed additives, effort must be done to improve their protein content as well to decrease the fiber contents. These can be achieved through fermentation. The process of fermentation by amylase producing fungi can also produce amylase enzyme which may be needed for increasing feed digestion rate of fish.

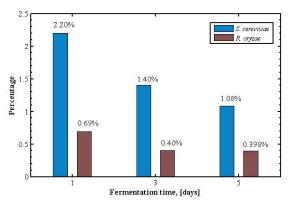


Fig. 2. Fiber content with aeration

The fiber contents resulted after fermentation is shown on Fig. 2 for aerated fermentation and Fig. 4 for non-aerated fermentation. While the protein content after fermentation is shown on Fig. 3 for aerated fermentation and Fig. 5 for non-aerated fermentation. From Fig.2, it is shown that the *Rhizopus oryzae* FNCC 6157 can decrease fiber faster and higher than *Saccharomyces cerevisiae* Y1536. It is understood that mould produces hydrolytic enzymes very well and better than yeast. By comparing Fig. 2 and Fig. 4, it can be said that aeration affected fiber decrease in the fermentation of banana peels by *Saccharomyces cerevisiae* and *Rhizopus oryzae*. This aeration increased the growth rate of those fungi and therefore increased the production of cellulase, fiber hydrolyzing enzyme.

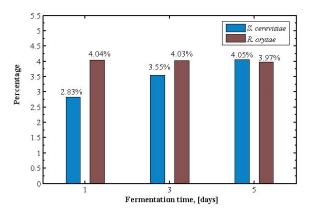


Fig. 3. Protein content with aeration

Fig.3 and Fig. 5 presents the protein content of the banana peels after fermentation with aeration, and without aeration, respectively. The protein content increases with fermentation time. The figures also show that *Rhizopus oryzae* fermentation caused protein increase higher than *Saccharomyces cerevisiae*. This protein content increased because the fungi grew and used the banana peels as substrate, broke down the complex carbohydrate of the peels and synthesized protein as the component of fungi cell. The increase of protein content of banana peels made them potential to be used as animal feed additive, including fish feed additive.

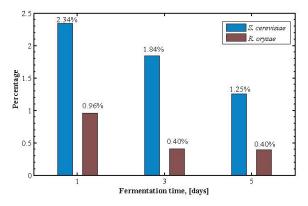


Fig. 4. Fiber content without aeration

The protein content increase was higher in fermentation with aeration compared to without aeration for each of the fungi because aeration favored the fungi growth. In the result of this research, it also can be seen that the highest protein content can be achieved up to 4% for 3 day aerated fermentation by *R. oryzae*. Higher protein content improvement can be achieved by lengthening the fementation time by *S. cereviae*.

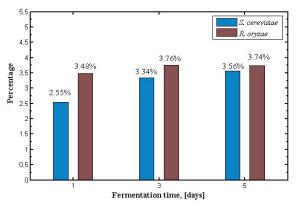


Fig. 5. Protein content without aeration

The amylase activity profiles after fermentation with and without aeration are shown in Fig. 6 and Fig. 7. The activity increased with fermentation time for each of the fungi because this enzyme was needed for their growth on banana peels which are composed mostly by starch.

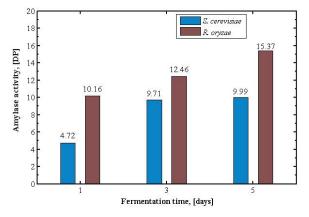


Fig. 6. Amylase activity result with aeration

The amylase activity produced by *Rhizopus oryzae* was higher than that produced by *Saccharomyces cerevisiae*. That is the reason for the higher growth rate of *R. oryzae* which cause the higher protein content and the lower fiber content resulted from *Rhizopus* banana peel fermentation as depicted from previous figures. The highest amylase activity was achieved by 5 day aerated fermentation by *Rhizopus oryzae* which resulted as high as 15.37 DP. As Comparison, 1 mL Taka amylase resulted in 6.5 DP [18].

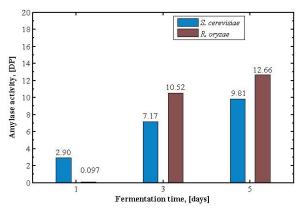


Fig. 7. Amylase activity result without aeration

The results has shown the potential usage of fermented banana peel as fish feed additive according to the increasing protein content, decreasing fiber content amylolitic enzyme to increase feed digestion. Previous publications have also shown that mixture of fungi such as T. viridae, T. reseei, A. oryzae, and Rhizopus oligosporus can ferment corn-cob to produce organic acid (citric, pantoteni, acetic and lactic), fatty acids, glucose, vitamin, mineral and various enzymes (lipase, protease and cellulase) [13]. Living cultures of Zygomycetes including Rhizopus oryzae have been used as probiotic organism to prevent fish infectious disease [19]. In spite of that, Saccharomyces cerevisiae is included as authorized probiotic in feeding stuffs under Council Directive 70/524/EEC [20]. Higher protein content from banana peel fermentation than obtained in this research could be achieved combining with other molds or with S. cerevisiae and other probiotic microorganisms in the future.

4 Conclusions and Recommendations

Banana peels fermentations by *Saccharomyces cerevisiae* Y1536 and *Rhizopus oryzae* FNCC 6157 had shown their potential as fish feed additive because of the increasing protein content and decreasing fiber content after fermentation. The fermentation also showed the presence of amylase activity needed for fish feed digestion. Banana peels fermentation by *Rhizopus oryzae* FNCC 6157 resulted in better protein increase, fiber decrease as well as amylase activity.

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