

**KORESPONDENSI UNTUK IOP Conference Series: Earth and Environmental Science Science 741 (2021) 012019**

- 1.Submitted Abstract to Proceeding committee: IOP Conf. Series: Earth and Environmental Science with a title: Growth dynamics of mold-yeast and bacteria during the production process of saga tauco [*Adenanthera pavonina*] (by system 30 -11-2020)
- 2.Letter of Acceptance (14-12-2020)
- 3.The Author submits PPT for presentation (15-12-2020)
4. The Author submits full paper: Growth dynamics of mold-yeast and bacteria during the production process of saga tauco [*Adenanthera pavonina*] (26-12-2020)
5. The Author received abstract which has been corrected by reviewer (11-01-21)
6. The Author sent the corrected full paper (12-1-2021)
7. The editor sent back the manuscript that had to be corrected (18-01-2021)
8. The Author sent back the final full paper (30-01-2021)
9. Paper published (1-06-2021)



PRESENTER\_abu amar <aamar3884biugm@gmail.com>

---

## Letter of Acceptance 2nd ICBEAU 2020

---

**ICBEAU UNAND** <icbeau.unand@gmail.com>  
To: Abu Amar <aamar3884biugm@gmail.com>

Mon, Dec 14, 2020 at 10:24 PM

Dear, Abu Amar  
There is your LoA from 2nd ICBEAU 2020.  
Thank you

Best Regard,  
2nd ICBEAU 2020 committee

---

 **Abu Amar - ID 107.pdf**  
176K



PRESENTER\_abu amar &lt;aamar3884biugm@gmail.com&gt;

---

**PPT: ABU AMAR. ID 107.Growth dynamics of mold-yeast and bacteria during the production process of saga tauco (Adenanthera pavonina)**

---

PRESENTER\_abu amar &lt;aamar3884biugm@gmail.com&gt;

Tue, Dec 15, 2020 at 2:28 PM

To: icbeau.unand@gmail.com

dear International Conference Committee,  
attached my PPT with a title: Growth dynamics of mold-yeast and bacteria during the production process of saga tauco (*Adenanthera pavonina*)  
my ID 107.  
best regard  
Abu Amar

**ID 107.ABUAMAR ICBEAU2020.pptx**

2128K

## Growth dynamics of mold-yeast and bacteria during the production process of Saga Tauco (*Adenanthera pavonina*)

Abu Amar, Syahril makosim, Setiarti Sukotjo, Nafarin Ahadiati, Ezer Weisman  
Agroindustrial Technology Department, Institut Teknologi Indonesia Serpong Tangerang Selatan  
Corresponding author: [abu.amar@iti.ac.id](mailto:abu.amar@iti.ac.id) or [aamar3884biugm@gmail.com](mailto:aamar3884biugm@gmail.com)

**Abstract.** As a plant protein source saga bean has not been used by the society. Several studies regarding saga bean being a food product have been initiated. Saga bean can be used as Tauco. The study was to observe the growth dynamics of microbe in Sagabean Tauco during the production process. The manufacturing of sagabean Tauco was the same as making of soybean Tauco. It was soaking, boiling peeling, soaking overnight, followed by steaming, and inoculating with tempeh starter. Saga tempeh was crushed then mixed with glutinous rice flour that had been roasted. After drying, it was put in brine solution for fermentation. Measurements taken were total microbes, total yeast-mold, and total lactic acid bacteria. Other measurements were total protein, total suspended solids, total acid, pH value, and ash content, this was done to see the relationship between the presence of microorganisms and their metabolic processes during the Tauco production. The result showed that the total number of microbes had increased during the fermentation process until a certain period, then it was constant and decreased according to the growth curve of microorganisms in general. For yeast and molds being relatively fluctuating and tending to increase, this seemed to relate to the presence of halophilic microorganisms in the product. Regarding the pH of the product during fermentation, it had a relevant value, while the ash content experienced a fluctuating value. If it had been related to the presence of microorganisms, the metabolism of the existing microorganisms would have affected the chemical properties of the Saga Tauco.

**Keywords:** Saga Tauco (*Adenanthera pavonina*, L), yeast mold, Lactic acid bacteria.

**Formatted:** Left: 0.98", Right: 0.98", Top: 1.58", Bottom: 1.06", Width: 8.27", Height: 11.69"

**Commented [SE1]:** Follow the IOP format  
<https://drive.google.com/file/d/1rZIDay3mvSgn3QE17SChTQ8J6O1ePhMO/view>  
<https://drive.google.com/file/d/1qYxRGO9S45hONFGpOvuQkFj3twdWh9hc/view>

**Formatted:** Left, Space Before: 79.4 pt, After: 28.35 pt

**Formatted:** Left, Right: 0", Space Before: 79.4 pt, After: 28.35 pt, Adjust space between Latin and Asian text, Adjust space between Asian text and numbers, Tab stops: 0.64", Left + 1.27", Left + 1.91", Left + 2.54", Left + 3.18", Left + 3.82", Left + 4.45", Left + 5.09", Left + 5.73", Left + 6.36", Left + 7", Left + 7.63", Left + 8.27", Left + 8.91", Left + 9.54", Left + 10.18", Left

**Formatted:** Font: 10 pt

**Formatted:** Justified, Indent: Left: 0.98"

**Formatted:** Font: 10 pt

**Formatted:** Font: 10 pt

**Formatted:** Indent: Left: 0.64"

**Commented [SE2]:** Sort by alphabet

# Growth dynamics of mold-yeast and bacteria during the production process of saga tauco (*Adenanthera pavonina*)

A Amar<sup>1</sup>, S Makosim<sup>1</sup>, S Sukotjo<sup>1</sup>, N Ahadiyanti<sup>1</sup>, E Weisman<sup>1</sup>

<sup>1</sup>Agro-Industrial Technology, Institut Teknologi Indonesia Serpong Tangerang Selatan  
Corresponding Author [abu.amar@iti.ac.id](mailto:abu.amar@iti.ac.id) or aamar3884biugm@gmail.com

**Abstract.** As a plant protein source sagabean has not been used by society. Several studies regarding sagabean being a food product have been initiated. Sagabean can be used as tauco. The study was to observe the growth dynamics of microbe in Sagabean tauco during the production process. The manufacturing of sagabean tauco was the same as making of soybean tauco. It was soaking, boiling peeling, soaking overnight, followed by steaming, and inoculating with tempeh starter. Saga tempeh was crushed then mixed with glutinous rice flour that had been roasted. After drying, it was put in a brine solution for fermentation. Measurements taken were total microbes, total yeast-mold, and total lactic acid bacteria. Other measurements were total protein, total dissolved solids, total acid, pH value, and ash content, this was done to see the relationship between the presence of microorganisms and their metabolic processes during the tauco production. The result showed that the total number of microbes had increased during the fermentation process until a certain period, then it was constant and decreased according to the growth curve of microorganisms in general. For yeast and molds being relatively fluctuating and tending to increase, this seemed to relate to the presence of halophilic microorganisms in the product. Regarding the pH of the product during fermentation, it had a relevant value, while the ash content experienced a fluctuating value. If it had been related to the presence of microorganisms, the metabolism of the existing microorganisms would have affected the chemical properties of the saga tauco.

**Keywords:** Lactic Acids Bacteria, Saga Tauco (*Adenanthera pavonina*) Yeast Mold

## 1. Introduction

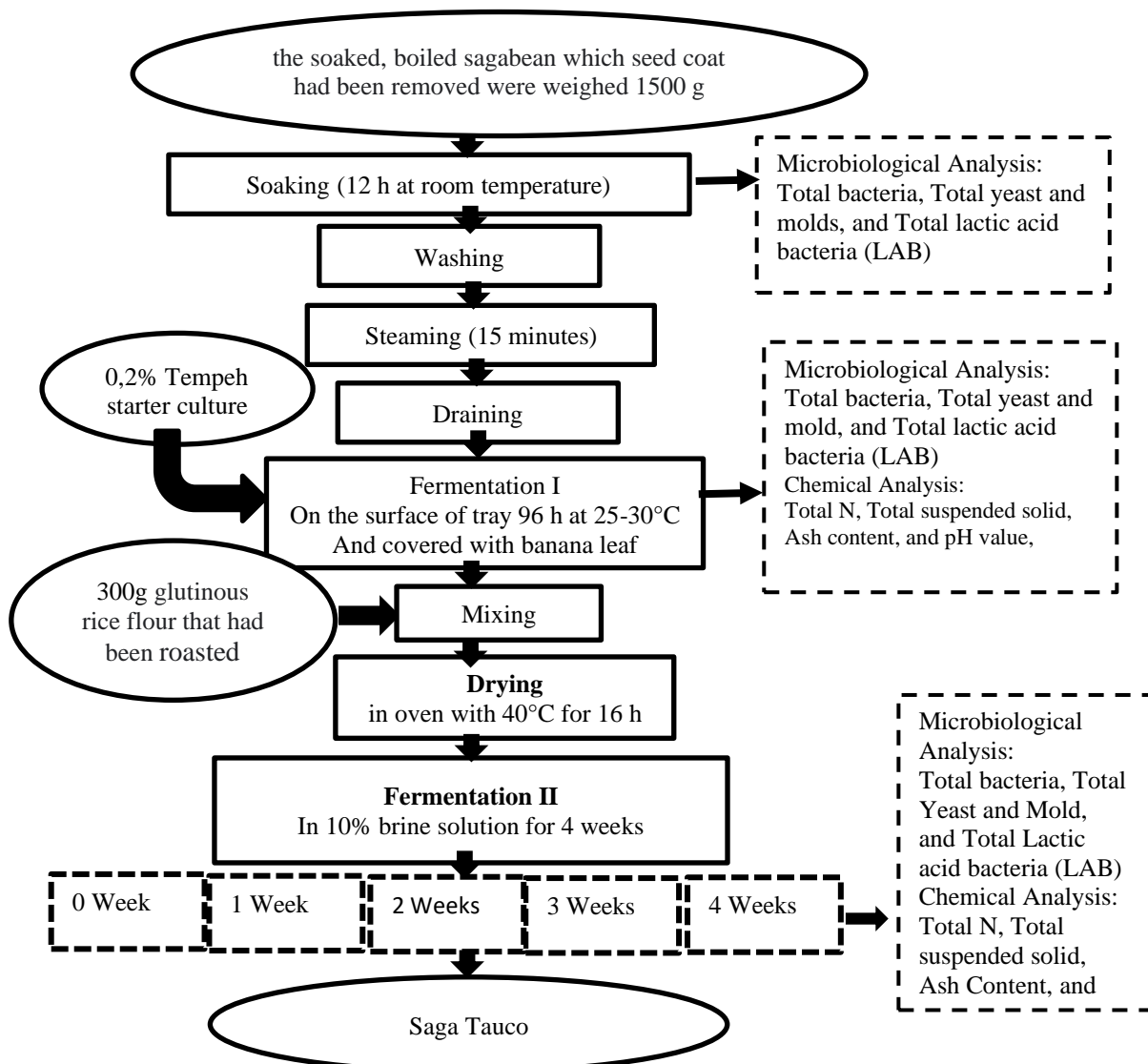
The import of soybeans in each period in Indonesia has increased, as an example from 2013 to 2018 with the total amount of imports in \$ fluctuating but with an upward trend. [1]. The use of soybeans in Indonesia is primarily for tempeh, tofu, soy sauce, soy milk, and tauco. Tauco was originally produced in Cianjur, West Java [2]. Tauco can be used as a seasoning or food flavoring with a distinctive taste and is relatively durable because of its relatively high salt content [3]. Soybean as a raw material for Tauco has similar biological and chemical properties as well as functionally with sagabean (*Adenanthera pavonina*). The economic feasibility of the tauco production process must use at least 60 kg of soybeans once a process [4]. Thus, if sagabeans are to be used as a substitute for soybeans, a lot of sagabean production is needed. This is what is able to move the economy of the farming community to produce sagabean. Sagabean are not only used as raw material for tempeh but they can also be used for fresh cheese. [5] In making tauco, mold and bacteria greatly affect the quality of tauco. Several types of yeast and lactic acid bacteria had been identified in the processing of soybean tauco. In the first and second weeks, two types of yeast, namely *Sacharomyces* and *Phicia*, dominated, until the third and fourth weeks only *Sacharomces* appeared, while the fifth week onwards, lactic acid bacteria, namely *Streptococcus*. dominated [6]. A previous study reported that the dynamics of microbial growth in tempeh that were processed in different ways would provide different microbial profiles [7]. In our

study, the microbiological and chemical reviews of tauco saga were examined during the production process.

## 2. Material and Methods

The materials used in this study were ripe sagabeen (*Adenanthera pavonina*, L) from the ITI Serpong campus, South Tangerang. Tempe starter culture was obtained from Bandung (Raprima), banana leaves which were used as a cover during stage I fermentation. 10% Salt Solution was used during stage II fermentation. The chemicals and media used were NaOH, HCl, TCAA, H<sub>2</sub>SO<sub>4</sub>, (Merck), PDA, NA, and MRS Agar (All media from Difco).

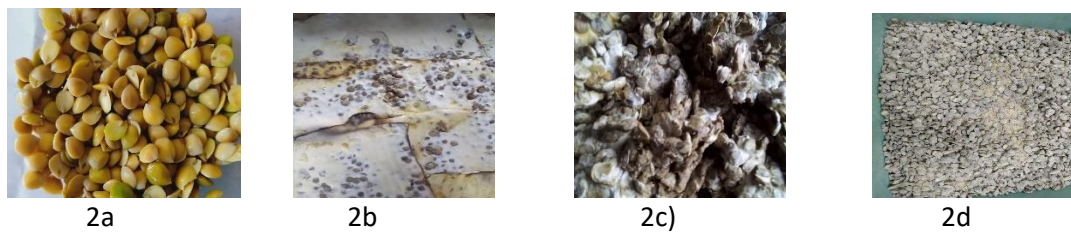
The method in this research was descriptive quantitative. Sagabeen tauco based was produced the same as manufacturing of soybean Tauco (Fig.1). This research was repeated 2 times. The parameters observed were microbiological analysis of the product during the process [8] which included the total yeast mold (using PDA media), the total bacteria (using Nutrient agar), and the total lactic acid bacteria (using MRSa media). To determine the role of the various types of microorganisms mentioned above, the measurement of Total N (Kjeldahl), Total Dissolved Solids [9] pH, and Ash content were measured. [10]. The experimental design for data analysis results of Total microbe, total protein. pH, total acids, ash content, and total suspended solids were in Random Block Design with seven treatments (soaking process, mold fermentation, 0 weeks in a brine solution, 1 week in a brine solution, 2 weeks in a brine solution, 3 weeks in brine solution and 4 weeks in brine solutions) and the experiment was repeated two times.



**Figure 1.** Research Flowchart of Saga Tauco Production Process

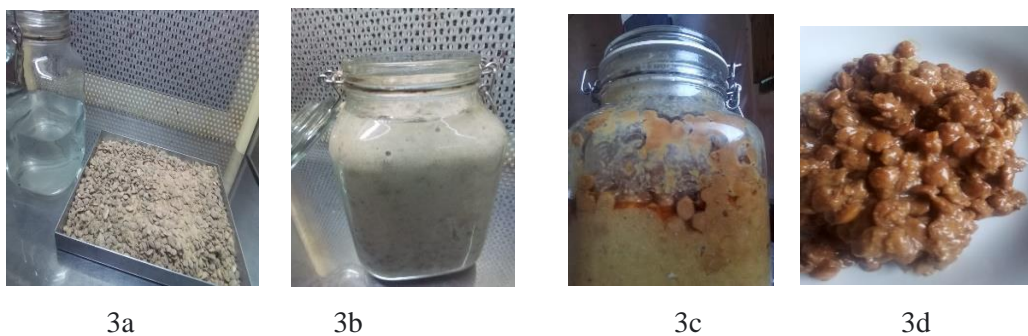
### 3.Results and Discussion

Based on the diagram in Figure 1, to carry out the saga tauco production process, two of good sagabean were prepared then soaked insufficient clean water for 24 hours, after an imbibition event occurred, the sagabeans were washed with water until they were clean then were boiled with boiling water for 1 hour. After 1 hour of boiling was stopped and cold water was added to make it easier to remove, separating the endosperm from the seed coat. This was to facilitate the process of stripping the sagabean, the endosperm was then washed and soaked for 24 hours to provide an opportunity for microbes to carry out their activities. As shown in Figure 1, the microbiological analysis was carried out during the immersion including the total bacteria, the total yeast molds, and the total LAB. Then they have washed again and steamed for 15 minutes and continued with cooling to room temperature. Currently, they were ready to inoculate with a tempeh starter (2a). The tempeh was 96 hours old (2b) Up to this stage, it called the fermentation process I, then they were cut into small pieces and added the glutinous rice flour that had been roasted and finely ground to provide nutrients to microbes later when soaking in a salt solution (2c). up to dry in an oven at 40 ° C for 16 hours (2d).



**Figure 2** Visualization of the endosperm of sagabean up to Fermentation I stage and it had been reduced in size and added with roasted glutinous rice flour ready to be continued in Phase II fermentation

The next step was preparation for the Fermentation II in a previously prepared 10% salt solution. The tempeh that had been cut into small pieces and was dried and had been added with roasted glutinous rice flour was then put in a 10% salt solution with a volume of 1000 ml (3a) while it was mixed in a jar so as the starting point of fermentation II, namely 0 weeks (Figure 3b) and incubated at 25-30 ° C for 4 weeks. Periodically, from 0 to 4 weeks of age, microbiological and chemical analyzes were carried out.



**Figure 3.** Fermentation II started from entering the fermentation product I into a 10% salt solution until Saga tauco was formed

After 2 weeks of age, it could be seen that saga tauco had begun to form with a slightly brownish color change. The oil appeared at the top of the product (Figure 3c), then the mass of the tauco started to concentrate and coalesced into a compact mass, but the saga bean granules were still visible. In line with the fermentation time, the mass of the tauco became more massive and if taken with a spoon, the texture became softer, this showed the metabolic process of microorganisms in the tauco production process to carry out its function. It was due to the enzymes produced by the microorganisms that existed during the process both from fermentation stage I which was dominated by the molds from tempeh starter, and fermentation stage II which came from microorganisms that were resistant to salt solutions.

### 3.1 Profile of microorganisms during the Tauco saga production process

The role of microorganisms during the soaking process certainly affected the saga bean endosperm which would be processed into saga tauco. At least the microorganisms during soaking initiated the implementation of metabolic processes in the saga bean. This was due to the decrease in the pH of the saga bean endosperm from the initial 7.13 at the beginning of soaking and after 12 hours of soaking the pH of the sagabean endosperm fell to 6.42. This indicated the presence of organic acids produced by microorganisms during the soaking process. The number of microorganisms starting from soaking and fermentation I can be seen in table 1.

**Table 1.** Profile of microorganisms during soaking and during fermentation with tempeh starter or fermentation stage I in the age of 96 hours (log CFU / g product)

Batch	Number of colony in Log CFU/g product					
	After soaking			After fermentation I with tempeh Starter 96 h		
	NA media	PDA media	MRSA media	NA Media	PDA media	MRSA media
1	7.477	6.623	-	7.362	8.204	6.491
2	7.146	5.623	-	7.255	8.176	6.380
Mean	7.312	6.123		7.308	8.190	6.436

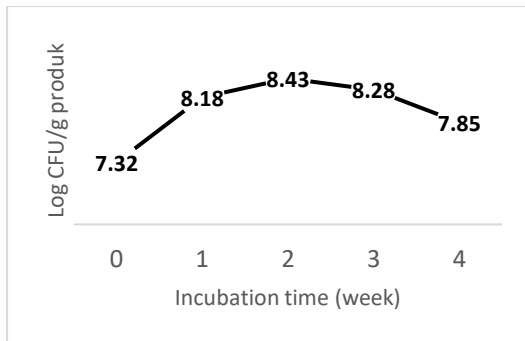
The number of bacteria after soaking was much higher than that of yeast mold colonies. It could be understood that in the soaking water, bacteria and mold competition occurs. However, the LAB was not detected. This was a different result compared to the previous study. The previous study reported that the soaking water of soybean which would be processed into tempeh had enough LAB colonies/g product [7]. The mold and yeast population (PDA media) in Saga tempeh dominated reaching 8.19 log CFU /g of product compared to the total bacteria (NA media) which only reached 7.308 and even for the total LAB (MRSA media) only reached 6.436 log CFU/g product. This made a lot of sense because the tempeh starter contains mainly the mold spores of *Rhizopus oligosporus* and *Rhizopus oryzae*. The mold and yeast present in tempeh starter, hydrolyzed saga bean with the microbial enzymes they produced during fermentation into simple compounds and as a result produced organic acid which could lower the pH. This decrease in pH provided comfort for LAB as evidenced by the growth of LAB in MRSA reaching 6.436 log CFU / g of product.

The bacteria growth curve of on NA media during the incubation process from 0 weeks to 4 weeks shown in Fig 4a. Thus, it could be said that the total growth of bacteria in tauco fermentation followed the usual bacterial growth curve, namely, there was a lag phase or an early phase then an exponential phase in the first and second weeks, and until the peak, then in the third and fourth week, it had decreased. Some of the factors determining this occurrence were the large variety of other microorganisms that contributed to the fermentation process of tauco. The variety of microbes in tauco greatly accelerated the substrate hydrolysis process, this was indicated by the increase in the total dissolved solids during fermentation (Fig. 5a). The fermented soybean food is known for its attractive flavor, texture and superior digestibility [11] Therefore, the saga tauco also had the opportunity to be a product that was easily digested by the human body.

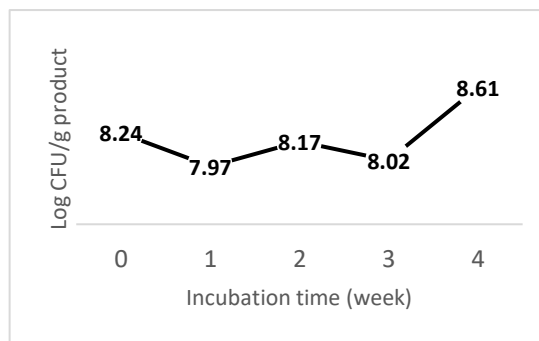
The dynamics of yeast and mold growth were very volatile (Fig 4b). It was assumed that the growth of mold and yeast species alternately at the beginning was dominated by aerobic fungi, then if oxygen availability ran out, yeast was dominated by the anaerobic tolerant and relatively halophilic yeast. And



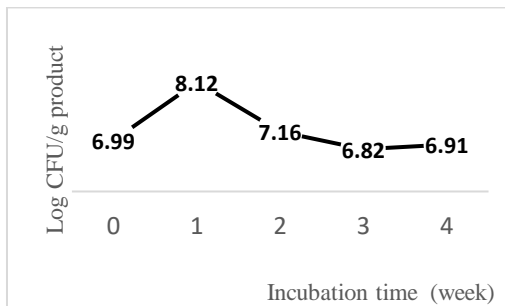
near the end of the fermentation process the number of halophilic microorganisms predominated. Usually, at the end of fermentation, salt-resistant yeast would dominate compared to other microbes.



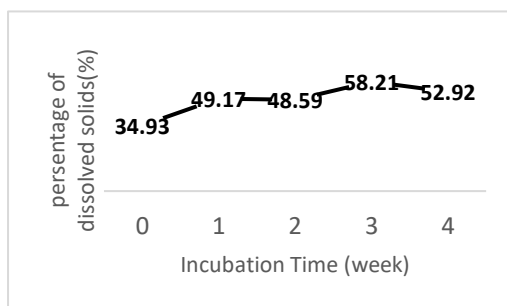
**Figure 4a.** Total growth of bacteria (NA media) in the Saga Tauco during the production process



**Figure 4b.** Total growth of yeast and mold (PDA media) in the Saga Tauco during the production process



**Figure 4c.** Total growth of LAB (MRSA media) in the Saga Tauco during the production process



**Figure 5a.** The percentage of total dissolved solid in the Saga Tauco during the production process

During the fermentation process in a salt solution, all salt-resistant microbes, be they lactic acid bacteria (LAB), Yeast, or even molds, synergized or might compete to hydrolyze the existing substrate so that the total dissolved solids increased as well as the pH value tended to rise slowly. When compared to Figures 4b and 4c there was a clear correlation in the first week of the maximum amount of LAB in contrast to minimal mold and yeast. It might be possible that the LAB produces antimicrobials substance so that mold and yeast life was suppressed. This result was in line with previous research which proved that LAB was able to suppress the growth of mold and yeast at the beginning of tempeh fermentation [7]. In contrast to the research conducted by Feng et al 2005 [12], who reported that the growth of *Rhizopus oligosporus* on grain fermentation was not affected by the presence of LAB. The presence of LAB in saga tauco is very beneficial because it had the potential as a probiotic microorganism. Other researchers reported that traditional fermented food in Indonesia was a potential source of probiotics [13].

### 3.2 Chemical Analysis in Saga Tauco during the production process

The total dissolved solids in tauco during the production process had increased, (Fig 5a) indicating that tauco was a product rich in nutrients, at least a lot of dissolved solids that facilitated the absorption system of the human body. This was due to the many microbial enzymes in tauco that came from yeast, mold, and LAB that were present in tauco during the production process. The total nitrogen in tauco during its production process fluctuated which was not too different, while the pH value tended to increase slightly. The pH value of the saga tauco ranged from 4.5 to 5.03 was due to the action of the LAB. This was caused by protein hydrolysis by the protease enzyme from microorganisms, leading to the breakdown of amino acids into volatile compounds such as ammonia. Likewise, the ash content in

the saga tauco during the production process slightly increased. (Table 2). A previous study reported that LAB produced organic acids which contributed to the sensory value of the fermented product [14]. In addition to the sufficient total dissolved solids, tauco made from soybean also contained sufficient antioxidants and total phenol which have a positive effect on body health. [15].

**Table 2.** Total Nitrogen content, pH value, and Ash content of saga tauco during production process\*

Incubation time (week)	Total N content (%)	Total Ash content (%)	pH value
0	13.80 <sup>a</sup>	6.85 <sup>c</sup>	4.50 <sup>a</sup>
1	13.68 <sup>a</sup>	5.84 <sup>a</sup>	4.80 <sup>b</sup>
2	13.53 <sup>a</sup>	5.94 <sup>a</sup>	4.83 <sup>b</sup>
3	14.05 <sup>a</sup>	6.34 <sup>b</sup>	4.82 <sup>b</sup>
4	14.08 <sup>a</sup>	6.14 <sup>ab</sup>	5.03 <sup>c</sup>

\*average of two replicates. the same letter in the same column shows no significant difference

#### 4. Conclusion

In saga tauco production process, this existing microbe's growth was very dynamic and contributed to the nutritional value of tauco proven by the increasing dissolved solids.

#### Reference

- [1] Biro Pusat Statistik 2019: Impor Kedelai Menurut Negara Asal Utama, 2010-2018. <https://www.bps.go.id/statistictable/2019/02/14/2015/impor-kedelai-menurut-negara-asal-utama-2010-2018.htm>
- [2] Astawan M 2009 *Sehat dengan Hidangan Kacang dan Biji-bijian* Jakarta: Penebar Swadaya
- [3] Harti A S and Kusumawati H N 2013 *Jurnal KESMADASKA* (4) 2 :89-95
- [4] Nandiyanto A B D, Ismiati R, Indrianti J, Abdullah, A G 2017 *IOP Conference series. Materials Science and Engineering* 288. IOP Publishing
- [5] Amar A, Makosim S, Marwati 2017 *Jurnal IPTEK ITI* 1 (2) Doi: <https://doi.org/10.31543/jii.v1i2>
- [6] Umniyatie S and Mariyam S 2002 *Jurnal Chimera* 7 (2) <http://journal.um.ac.id/index.php/chimera/index>
- [7] Nurdini A L, Nuraida L, Suwanto A, Suliantari 2015 *International Food Research Journal* 22(4): 1668-1674
- [8] Fardiaz S 1989 *Petunjuk Laboratorium Analisis Mikrobiologi Pangan*. Pusat Antar Univ Pangan dan Gizi IPB
- [9] Legowo A M and Nurwantoro 2004 *Analisis Pangan*, Prodi Teknologi hasil ternak Fakultas Peternakan Universitas Diponegoro Semarang
- [10] Andarwulan N, Kusnandar F, Hariyadi P 2011 *Analisis Kimia pangan* Universitas Terbuka Jakarta
- [11] Nouts M J R and Kiers J L 2005. *Journal of Applied Microbiology* 98, 789–805 doi:10.1111/j.1365-2672.2004. 02471.x
- [12] Feng X M, Erickson A R B, Schnurer, J 2005. *International Journal of Food Microbiology* 104.249-256
- [13] Nuraida L 2015 *Food Science and Human Wellness* 4 (2015) 47–55
- [14] Rathore S, Salmerón I, Pandiella S 2012 *Food Microbiology* 30: 239–244.
- [15] Larasati N 2017 *Jurnal Pangan dan Agroindustri* Vol.5 No.2:85-95.

#### Acknowledgments

Thanks to Research and Public Service Centre Institut Teknologi Indonesia Serpong for financial support during research with the contract number 023/KP/PRPM-PP/ITI/VII/2020



PRESENTER\_abu amar <aamar3884biugm@gmail.com>

---

## Artikel Review

---

**ICBEAU UNAND** <icbeau.unand@gmail.com>  
To: Abu Amar <aamar3884biugm@gmail.com>

Mon, Jan 11, 2021 at 11:49 AM

Dear author,

Your article has been pre-reviewed. We have corrected your article and have adjusted it to the IOP format.

Please correct it according to IOP template and submit your article in the OJS system


Thank you.

Best regard,

Section Editor

---

### 2 attachments

 **107-Article Text-340-1-18-20201213.docx**  
421K

 **ICBEAU20-Template-JPCSEExampleWordDocument.docx**  
535K



PRESENTER\_abu amar <aamar3884biugm@gmail.com>

---

## Artikel Review


---

**PRESENTER\_abu amar** <aamar3884biugm@gmail.com>  
To: ICBEAU UNAND <icbeau.unand@gmail.com>

Tue, Jan 12, 2021 at 3:22 PM

thank you very much I have already fixed it according to what the reviewer wanted. I expect it to be well received  
best regard  
Abu Amar  
[Quoted text hidden]

---

 **107-Article Text-340-1-18-20201213 final.docx**  
383K

## Growth dynamics of mold-yeast and bacteria during the production process of saga tauco (*Adenanthera pavonina*)

**Abstract.** Sagabeen as a plant protein source has not been used by society. Several studies regarding Sagabeen being a food product have been initiated. Sagabeen can be used as tauco. The study was to observe the growth dynamics of microbe in Sagabeen tauco during the production process. The manufacturing of Sagabeen tauco was the same as making of soybean tauco. It was soaking, boiling peeling, soaking overnight, followed by steaming, and inoculating with tempeh starter. Saga tempeh was crushed then mixed with glutinous rice flour that had been roasted. After drying, it was put in a brine solution for fermentation. The observation were total microbes, total yeast-mold, total lactic acid bacteria, total protein, total dissolved solids, total acid, pH value, and ash content. This aims of this research was to see the relationship between the presence of microorganisms and their metabolic processes during the tauco production. The result showed that the total number of microbes had increased during the fermentation process until a certain period, then it was constant and decreased according to the growth curve of microorganisms in general. For yeast and molds being relatively fluctuating and tending to increase. This seemed to relate to the presence of halophilic microorganisms in the product. Regarding the pH of the product during fermentation, it had a relevant value, while the ash content experienced a fluctuating value. If it had been related to the presence of microorganisms, the metabolism of the existing microorganisms would have affected the chemical properties of the saga tauco.

**Keywords:** Lactic Aaids, bacteria, saga tauco, *Adenantrea pavonina*, yeast mold

### 1. Introduction

The import of soybeans in each period in Indonesia has increased, as an example from 2013 to 2018 with the total amount of imports in \$ fluctuating but with an upward trend. [1]. The use of soybeans in Indonesia is primarily for tempeh, tofu, soy sauce, soy milk, and tauco. Tauco was originally produced in Cianjur, West Java [2]. Tauco can be used as a seasoning or food flavoring with a distinctive taste and is relatively durable because of its relatively high salt content [3]. Soybean as a raw material for Tauco has similar biological and chemical properties as well as functionally with sagabeen (*Adenanthera pavonina*). The economic feasibility of the tauco production process must use at least 60 kg of soybeans once a process [4]. Thus, if sagabeens are to be used as a substitute for soybeans, a lot of sagabeen production is needed. This is what is able to move the economy of the farming community to produce sagabeen. Sagabeen are not only used as raw material for tempeh but they can also be used for fresh cheese. [5] In making tauco, mold and bacteria greatly affect the quality of tauco. Several types of yeast and lactic acid bacteria had been identified in the processing of soybean tauco. In the first and second weeks, two types of yeast, namely *Sacharomyces* and *Phicia*, dominated, until the third and fourth weeks only *Sacharomces* appeared, while the fifth week onwards, lactic acid bacteria, namely *Streptococcus*. dominated [6]. A previous study reported that the dynamics of microbial growth in tempeh that were processed in different ways would provide different microbial profiles [7]. In our study, the microbiological and chemical reviews of tauco saga were examined during the production process.

### 2. Material and Methods

The materials used in this study were ripe Sagabeen (*Adenanthera pavonina*, L) from the ITI Serpong campus, South Tangerang. Tempe starter culture was obtained from Bandung (Raprima),

**Commented [A1]:** Konsisten Sagabeen..S huruf besar. Cek untuk lainnya...

**Commented [A2]:** Tempe or tempeh..?

banana leaves which were used as a cover during first stage of fermentation. The second stage of fermentation used 10% salt solution. The chemicals and media used were NaOH, HCl, TCAA, H<sub>2</sub>SO<sub>4</sub>, (Merck), PDA, NA, and MRS Agar (All media from Difco).

The method in this research was descriptive quantitative. Sagabean tauco based was produced the same as manufacturing of soybean Tauco (Figure1). This research was repeated twice. The parameters observed were microbiological analysis of the product during the process [8] which included the total yeast mold (using PDA media), the total bacteria (using Nutrient agar), and the total lactic acid bacteria (using MRS media). The measurement of total N (Kjeldahl), total dissolved solids [9] pH, and ash content were measured [10] to determine the role of the various types of microorganisms mentioned above. The experimental design for data analysis results of total microbe, total protein, pH, total acids, ash content, and total suspended solids were in Random Block Design with seven treatments (soaking process, mold fermentation, 0 weeks in a brine solution, one week in a brine solution, two weeks in a brine solution, three weeks in a brine solution and four weeks in brine solutions) and the experiment was repeated twice.

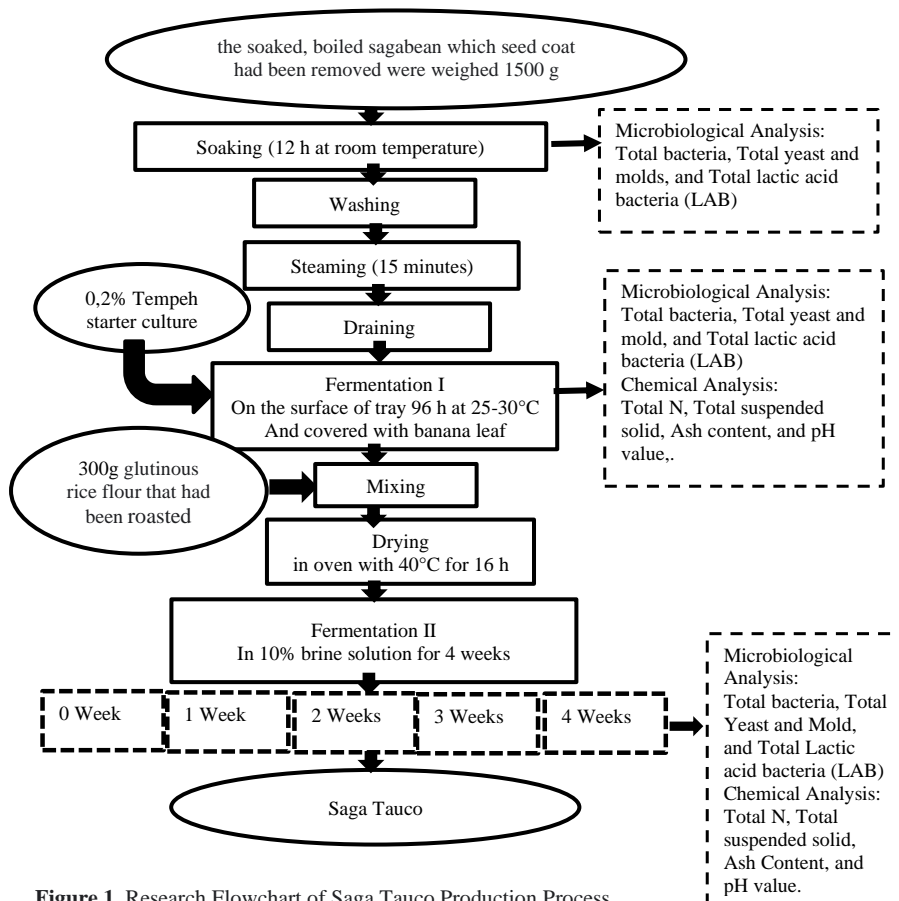
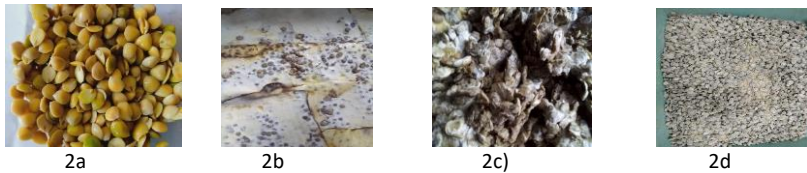


Figure 1. Research Flowchart of Saga Tauco Production Process

### 3.Results and Discussion

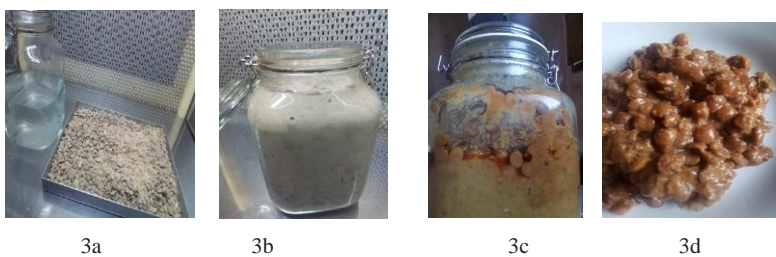
Based on the diagram in Figure 1, to carry out the saga tauco production process, two of good sagabean were prepared then soaked insufficient clean water for 24 hours, after an imbibition event occurred, the Sagabeans were washed with water until they were clean then were boiled with boiling water for 1 hour. After 1 hour of boiling was stopped and cold water was added to make it easier to remove, separating the endosperm from the seed coat. This was to facilitate the process of stripping the sagabean, the endosperm was then washed and soaked for 24 hours to provide an opportunity for microbes to carry out their activities. As shown in Figure 1, the microbiological analysis was carried out during the immersion including the total bacteria, the total yeast molds, and the total LAB. Then they have washed again and steamed for 15 minutes and continued with cooling to room temperature. Currently, they were ready to inoculate with a tempeh starter (2a). The tempeh was 96 hours old (2b) Up to this stage, it called the first fermentation process, then they were cut into small pieces and added the glutinous rice flour that had been roasted and finely ground to provide nutrients to microbes later when soaking in a salt solution (2c) up to dry in an oven at 40 ° C for 16 hours (2d).



**Figure 2** Visualization of the endosperm of sagabean up to first fermentation stage and it had been reduced in size and added with roasted glutinous rice flour ready to be continued in second phase fermentation

**Commented [A3]:** Keterangan Gambar 2a, 2b, 2c, 2d harus dijelaskan gambar apa.

The next step was preparation for the second fermentation in a previously prepared 10% salt solution. The tempeh that had been cut into small pieces and was dried and had been added with roasted glutinous rice flour was then put in a 10% salt solution with a volume of 1000 ml (3a) while it was mixed in a jar so as the starting point of second fermentation, namely 0 weeks (Figure 3b) and incubated at 25-30 °C for 4 weeks. Periodically, from 0 to 4 weeks of age, microbiological and chemical analyzes were carried out.



**Figure 3.** The second fermentation started from entering the first fermentation product into a 10% salt solution until Saga tauco was formed.

**Commented [A4]:** Keterangan Gambar 3a, 3b, 3c, 3d harus dijelaskan gambar apa.

After 2 weeks of age, it could be seen that saga tauco had begun to form with a slightly brownish color change. The oil appeared at the top of the product (Figure 3c), then the mass of the tauco started to concentrate and coalesced into a compact mass, but the saga bean granules were still visible. In line

with the fermentation time, the mass of the tauco became more massive and if taken with a spoon, the texture became softer, this showed that the metabolic process of microorganisms in the tauco production process to carry out its function. It was due to the enzymes produced by the microorganisms that existed during the process both from first fermentation stage which was dominated by the molds from tempeh starter, and second fermentation stage which came from microorganisms that were resistant to salt solutions.

### 3.1 Profile of microorganisms during the Tauco saga production process

The role of microorganisms during the soaking process certainly affected the saga bean endosperm which would be processed into saga tauco. At least the microorganisms during soaking initiated the implementation of metabolic processes in the saga bean. This was due to the decrease in the pH of the saga bean endosperm from the initial 7.13 at the beginning of soaking and after 12 hours of soaking the pH of the sagabean endosperm fell to 6.42. This indicated the presence of organic acids produced by microorganisms during the soaking process. The number of microorganisms starting from soaking and first fermentation can be seen in Table 1.

**Table 1.** Profile of microorganisms during soaking and during fermentation with tempeh starter or first fermentation stage in the age of 96 hours (log CFU / g product)

Batch	Number of colony in Log CFU/g product					
	After soaking			After first fermentation with tempeh starter 96 h		
	NA media	PDA media	MRSA media	NA Media	PDA media	MRSA media
1	7.477	6.623	-	7.362	8.204	6.491
2	7.146	5.623	-	7.255	8.176	6.380
Mean	7.312	6.123		7.308	8.190	6.436

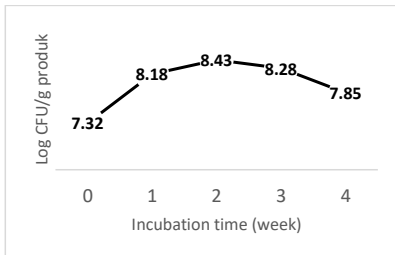
The number of bacteria after soaking was much higher than that of yeast mold colonies. It could be understood that in the soaking water, bacteria and mold competition occurs. However, the LAB was not detected. This was a different result compared to the previous study. The previous study reported that the soaking water of soybean which would be processed into tempeh had enough LAB colonies/g product [7]. The mold and yeast population (PDA media) in Saga tempeh dominated reaching 8.19 log CFU /g of product compared to the total bacteria (NA media) which only reached 7.308 and even for the total LAB (MRSA media) only reached 6.436 log CFU/g product. This made a lot of sense because the tempeh starter contains mainly the mold spores of *Rhizopus oligosporus* and *Rhizopus oryzae*. The mold and yeast present in tempeh starter, hydrolyzed saga bean with the microbial enzymes they produced during fermentation into simple compounds and as a result produced organic acid which could lower the pH. This decrease in pH provided comfort for LAB as evidenced by the growth of LAB in MRSA reaching 6.436 log CFU / g of product.

The bacteria growth curve of on NA media during the incubation process from 0 weeks to 4 weeks shown in Figure 4a. Thus, it could be said that the total growth of bacteria in tauco fermentation followed the usual bacterial growth curve, namely, there was a lag phase or an early phase then an exponential phase in the first and second weeks, and until the peak, then in the third and fourth week, it had decreased. Some of the factors determining this occurrence were the large variety of other microorganisms that contributed to the fermentation process of tauco. The variety of microbes in tauco greatly accelerated the substrate hydrolysis process, this was indicated by the increase in the total dissolved solids during fermentation (Figure 5a). The fermented soybean food is known for its attractive flavor, texture and superior digestibility [11]. Therefore, the saga tauco also had the opportunity to be a product that was easily digested by the human body.

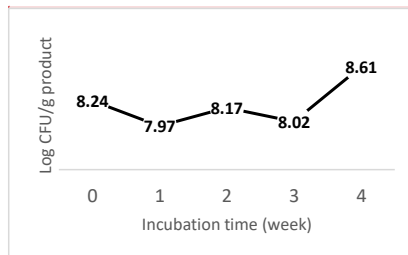
The dynamics of yeast and mold growth were very volatile (Figure 4b). It was assumed that the growth of mold and yeast species alternately at the beginning was dominated by aerobic fungi, then if oxygen availability ran out, yeast was dominated by the anaerobic tolerant and relatively halophilic yeast. And near the end of the fermentation process the number of halophilic microorganisms



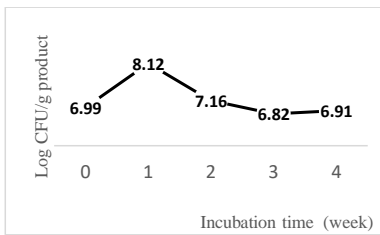
predominated. Usually, at the end of fermentation, salt-resistant yeast would dominate compared to other microbes.



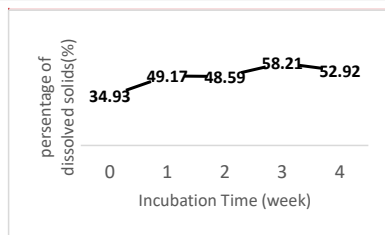
**Figure 4a.** Total growth of bacteria (NA media) in the Saga Tauco during the production process



**Figure 4b.** Total growth of yeast and mold (PDA media) in the Saga Tauco during the production process



**Figure 4c.** Total growth of LAB (MRSA media) in the Saga Tauco during the production process



**Figure 5a.** The percentage of total dissolved solid in the Saga Tauco during the production process

During the fermentation process in a salt solution, all salt-resistant microbes, be they lactic acid bacteria (LAB), Yeast, or even molds, synergized or might compete to hydrolyze the existing substrate so that the total dissolved solids increased as well as the pH value tended to rise slowly. When compared to Figures 4b and 4c there was a clear correlation in the first week of the maximum amount of LAB in contrast to minimal mold and yeast. It might be possible that the LAB produces antimicrobials substance so that mold and yeast life was suppressed. This result was in line with previous research which proved that LAB was able to suppress the growth of mold and yeast at the beginning of tempeh fermentation [7]. In contrast to the research conducted by Feng et al 2005 [12], who reported that the growth of *Rhizopus oligosporus* on grain fermentation was not affected by the presence of LAB. The presence of LAB in saga tauco is very beneficial because it had the potential as a probiotic microorganism. Other researchers reported that traditional fermented food in Indonesia was a potential source of probiotics [13].

### 3.2 Chemical Analysis in Saga Tauco during the production process

The total dissolved solids in tauco during the production process had increased, (Figure 5a) indicating that tauco was a product rich in nutrients, at least a lot of dissolved solids that facilitated the absorption system of the human body. This was due to the many microbial enzymes in tauco that came from yeast, mold, and LAB that were present in tauco during the production process. The total nitrogen in tauco during its production process fluctuated which was not too different, while the pH value tended to increase slightly. The pH value of the saga tauco ranged from 4.5 to 5.03 was due to the action of the LAB. This was caused by protein hydrolysis by the protease enzyme from microorganisms, leading to the breakdown of amino acids into volatile compounds such as ammonia.

**Commented [A5]:** Weeks  
Cek untuk semuanya.

**Commented [A6]:** The percentage...

Likewise, the ash content in the saga tauco during the production process slightly increased (Table 2). A previous study reported that LAB produced organic acids which contributed to the sensory value of the fermented product [14]. In addition to the sufficient total dissolved solids, tauco made from soybean also contained sufficient antioxidants and total phenol which have a positive effect on body health [15].

**Table 2.** Total Nitrogen content, pH value, and Ash content of saga tauco during production process\*

Incubation time (week)	Total N content (%)	Total ash content (%)	pH value
0	13.80 <sup>a</sup>	6.85 <sup>c</sup>	4.50 <sup>a</sup>
1	13.68 <sup>a</sup>	5.84 <sup>a</sup>	4.80 <sup>b</sup>
2	13.53 <sup>a</sup>	5.94 <sup>a</sup>	4.83 <sup>b</sup>
3	14.05 <sup>a</sup>	6.34 <sup>b</sup>	4.82 <sup>b</sup>
4	14.08 <sup>a</sup>	6.14 <sup>ab</sup>	5.03 <sup>c</sup>

\*average of two replicates. the same letter in the same column shows no significant difference

#### 4. Conclusion

In saga tauco production process, this existing microbe's growth was very dynamic and contributed to the nutritional value of tauco proven by the increasing dissolved solids.

#### Reference

- [1] Biro Pusat Statistik 2019: Impor Kedelai Menurut Negara Asal Utama, 2010-2018. <https://www.bps.go.id/statictable/2019/02/14/2015/impor-kedelai-menurut-negara-asal-utama-2010-2018.htm>
- [2] Astawan M 2009 *Sehat dengan Hidangan Kacang dan Biji-bijian* Jakarta: Penebar Swadaya
- [3] Harti A S and Kusumawati H N 2013 *Jurnal KESMADASKA* (4) 2 :89-95
- [4] Nandiyanto A B D, Ismiati R, Indrianti J, Abdullah, A G 2017 *IOP Conference series. Materials Science and Engineering* 288. IOP Publishing
- [5] Amar A, Makosim S, Marwati 2017 *Jurnal IPTEK ITI* 1 (2) Doi: <https://doi.org/10.31543/jii.v1i2>
- [6] Ummiyatie S and Mariyam S 2002 *Jurnal Chimera* 7 (2) <http://journal.um.ac.id/index.php/chimera/index>
- [7] Nurdini A L, Nuraida L, Suwanto A, Suliantari 2015 *International Food Research Journal* 22(4): 1668-1674
- [8] Fardiaz S 1989 *Petunjuk Laboratorium Analisis Mikrobiologi Pangan*. Pusat Antar Univ Pangan dan Gizi IPB
- [9] Legowo A M and Nurwantoro 2004 *Analisis Pangan, Prodi Teknologi hasil ternak* Fakultas Peternakan Universitas Diponegoro Semarang
- [10] Andarwulan N, Kusnandar F, Hariyadi P 2011 *Analisis Kimia pangan* Universitas Terbuka Jakarta
- [11] Nouts M J R and Kiers J L 2005. *Journal of Applied Microbiology* 98, 789–805 doi:10.1111/j.1365-2672.2004.02471.x
- [12] Feng X M, Erickson A R B, Schnurer, J 2005. *International Journal of Food Microbiology* 104.249-256
- [13] Nuraida L 2015 *Food Science and Human Wellness* 4 (2015) 47–55
- [14] Rathore S, Salmerón I, Pandiella S 2012 *Food Microbiology* 30: 239–244.
- [15] Larasati N 2017 *Jurnal Pangan dan Agroindustri* Vol.5 No.2:85-95.

# Growth dynamics of mold-yeast and bacteria during the production process of saga tauco (*Adenanthera pavonina*)

**Abstract.** Sagabean as a plant protein source Sagabean has not been used by the society. Several studies regarding Sagabean being a food product have been initiated. Sagabean can be used as tauco. The study was to observe the growth dynamics of microbe in Sagabean tauco during production process. The manufacturing of Sagabean tauco was the same as making of soybean tauco. It was soaking, boiling peeling, soaking overnight, followed by steaming, and inoculating with tempe starter. Saga tempe was crushed then mixed with glutinous rice flour that had been roasted. After drying, it was put in brine solution for fermentation. The observation were total microbes, total yeast-mold, total lactic acids bacteria, total protein, total dissolved solid, total acid, pH value and ash content. This aims of this reearch was to see the relationship between the presence of microorganisms and their metabolic processes during the tauco production. The result showed that the total number of microbes had increased during the fermentation process until a certain period, then it was constant and decreased according to the growth curve of microorganisms in general. For yeast and molds being relatively fluctuating and tending to increase. This seemed to relate to the presence of halophilic microorganisms in the product. Regarding the pH of the product during fermentation, it had a relevant value, while the ash content experienced a fluctuating value. If it had related to the presence of microorganisms, the metabolism of the existing microorganisms would have affected the chemical properties of the saga tauco.

**Keywords:** Lactic acids bacteria, saga tauco (*Adenanthera pavonina*) yeast mold

## 1. Introduction

The import of soybeans in each period in Indonesia has increased, as an example from 2013 to 2018 with the total amount of imports in \$ fluctuating but with an upward trend. [1]. The use of soybeans in Indonesia is primarily for tempe, tofu, soy sauce, soy milk and tauco. Tauco was originally produced in Cianjur, West Java [2]. Tauco can be used as a seasoning or food flavoring with a distinctive taste and is relatively durable because of its relatively high salt content [3]. Soybean as a raw material for Tauco has similar biological and chemical properties as well as functionally with Sagabean (*Adenanthera pavonina*). The economic feasibility of the tauco production process must use at least 60 kg of soybeans once a process [4]. Thus, if Sagabeans are to be used as a substitute for soybeans, a lot of Sagabean production is needed. This is what is able to move the economy of the farming community to produce Sagabean. Sagabean are not only used as raw material for tempe but they can also be used for fresh cheese. [5] In making tauco, mold and bacteria greatly affect the quality of tauco. Several types of yeast and lactic acid bacteria had been identified in the processing of soybean tauco. In the first and second weeks, two types of yeast, namely *Sacharomyces* and *Phicia*, dominated, until the third and fourth weeks only *Sacharomces* appeared, while the fifth week onwards, lactic acid bacteria, namely *Streptococcus*. dominated [6]. Previous study reported that the dynamics of microbial growth in tempe that were processed in different ways would provide different microbial profiles [7]. In our study, the microbiological and chemical reviews of tauco saga were examined during the production process.

## 2. Material and Methods

The materials used in this study were ripe Sagabean (*Adenanthera pavonina*, L) from the ITI Serpong campus, South Tangerang. Tempe starter culture was obtained from Bandung (Raprima), banana leaves which were used as a cover during first stage fermentation. The second stage fermentation used 10%

salt solution. The chemicals and media used were NaOH, HCl, TCAA, H<sub>2</sub>SO<sub>4</sub>, (Merck), PDA, NA and MRS Agar (All media from Difco).

The method in this research was descriptive quantitative. Sagabeau taucu based was produced the same as manufacturing of soybean Taucu (Figure.1). This research was repeated twice. The parameters observed were microbiological analysis of the product during the process [8] which included the total yeast mold (using PDA media), the total bacteria (using Nutrient agar) and the total lactic acid bacteria (using MRS media). The measurement of Total N (Kjeldahl), total dissolved solids [9] pH and ash content were measured [10] to determine the role of the various types of microorganisms mentioned above. The experimental design for data analysis results of total microbe, total protein. pH, total acids, ash content and total suspended solids were in Random Block Design with seven treatments (soaking process, mold fermentation, 0 week in brine solution, one week in brine solution, two weeks in brine solution, three weeks in brine solution and four weeks in brine solutions) and the experiment was repeated twice.

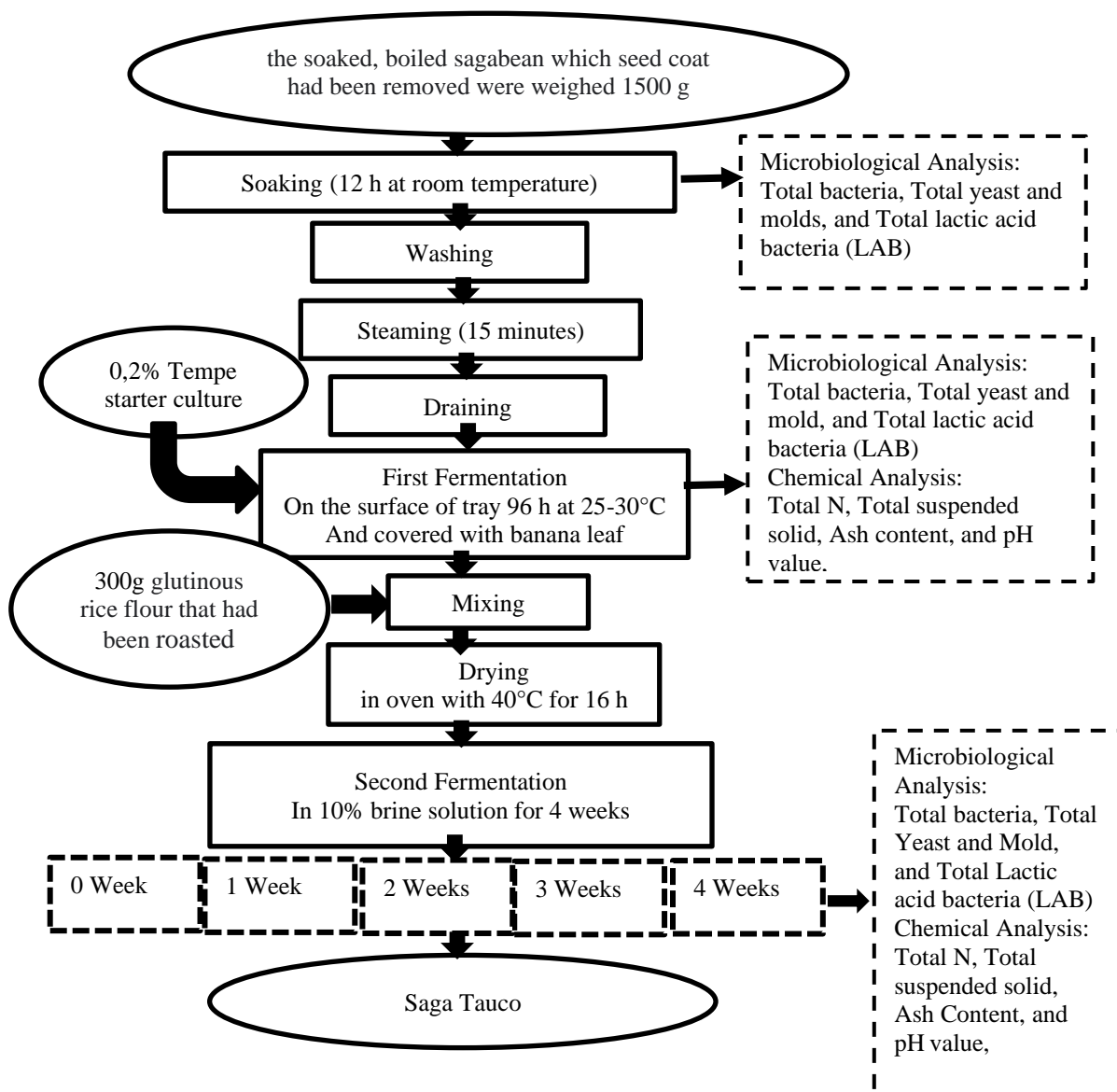


Figure 1: Research Flowchart of Saga Taucu Production Process

### 3.Results and Discussion

Based on the diagram in Figure 1, to carry out the saga tauco production process, two of good Sagabean were prepared then soaked in sufficient clean water for 24 hours, after an imbibition event occurred, the Sagabeans were washed with water until they were clean then were boiled with boiling water for 1 hour. After 1 hour of boiling was stopped and cold water was added to make it easier to remove, separating the endosperm from the seed coat. This was to facilitate the process of stripping the Sagabean, the endosperm was then washed and soaked for 24 hours to provide an opportunity for microbes to carry out their activities. As shown in Figure 1, microbiological analysis was carried out during the immersion including the total bacteria, the total yeast molds and the total LAB. Then they were washed again and steamed for 15 minutes and continued with cooling to room temperature. Currently they were ready to inoculate with tempe starter (2a). The tempe was 96 hours old (2b) Up to this stage, it called the first fermentation process, then they were cut into small pieces and added the glutinous rice flour that had been roasted and finely ground to provide nutrients to microbes later when soaking in salt solution (2c). up to drying in an oven at 40 ° C for 16 hours (2d).

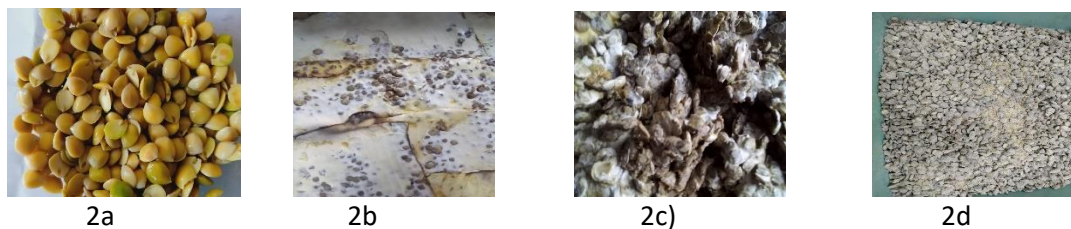


Figure 2: Visualization of the endosperm of Sagabean up to first fermentation and it had been reduced in size and added with roasted glutinous rice flour ready to be continued in second fermentation. Saga seed endosperm that is ready to be inoculated with the tempe starter (2a), saga tempe 96 hours after fermentation (2b), saga tempe pieces mixed with roasted glutinous rice flour (2c), saga tempe pieces that have been mixed with roasted glutinous rice flour and dried in oven 40°C for 16 hours ready to press to the second fermentation with saline solution (2d)

The next step was preparation for the second fermentation in a previously prepared 10% salt solution. The tempe that had been cut into small pieces and was dried and had been added with roasted glutinous rice flour was then put in a 10% salt solution with a volume of 1000 ml (3a) while it was mixed in a jar so as the starting point of second fermentation, namely 0 weeks (Figure 3b) and incubated at 25-30 ° C for 4 weeks. Periodically, from 0 to 4 weeks of age, microbiological and chemical analyzes were carried out.

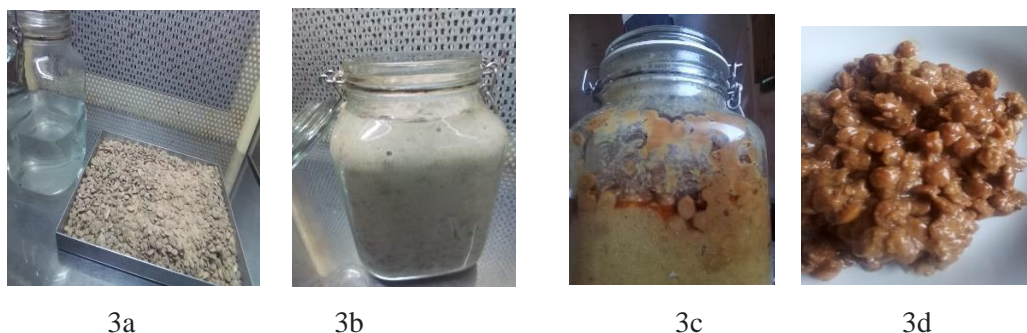


Figure 3: Fermentation II started from entering the fermentation product I into a 10% salt solution until Saga tauco was formed. Saga tempe pieces that have been mixed with roasted glutinous rice flour and dried in oven 40°C for 16 hours ready to press to the second fermentation with saline solution. Glass vessel filled with brine solution and saga tempe pieces that have been mixed with glutinous rice flour,

saga tempe pieces that have been mixed roasted glutinous rice flour and dried in oven 40°C for 16 hours ready to process to the second fermentation with saline solution (3a), the second fermentation was incubated for 0 weeks in a glass vessel containing brine solution (3b), second fermentation was incubated for 4 weeks (3c), tauco saga incubated for 4 weeks is ready for consumption (3d)

After 2 weeks of age, it could be seen that saga tauco had begun to form with a slightly brownish color change. The oil appeared at the top of the product (Figure 3c), then the mass of the tauco started to concentrate and coalesced into a compact mass, but the saga bean granules were still visible. In line with the fermentation time, the mass of the tauco became more massive and if taken with a spoon, the texture became softer, this showed that the metabolic process of microorganisms in the tauco production process to carry out its function. It was due to the enzymes produced by the microorganisms that existed during the process, both from first fermentation stage which was dominated by the molds from tempe starter and second fermentation stage which came from microorganisms that were resistant to salt solutions.

### 3.1 Profile of microorganisms during the Tauco saga production process

The role of microorganisms during the soaking process certainly affected the saga bean endosperm which would be processed into saga tauco. At least the microorganisms during soaking initiated the implementation of metabolic processes in the saga bean. This was due to the decrease in the pH of the saga bean endosperm from the initial 7.13 at the beginning of soaking and after 12 hours of soaking the pH of the saga bean endosperm fell to 6.42. This indicated the presence of organic acids produced by microorganisms during the soaking process. The number of microorganisms starting from soaking and first fermentation can be seen in Table 1.

Table 1. Profile of microorganisms during soaking and during fermentation with tempe starter or first fermentation stage in the age of 96 hours (log CFU / g product)

Batch	Number of colony in Log CFU/g product					
	After soaking			After first fermentation with tempe starter 96 h		
	NA media	PDA media	MRSA media	NA Media	PDA media	MRSA media
1	7.477	6.623	-	7.362	8.204	6.491
2	7.146	5.623	-	7.255	8.176	6.380
Mean	7.312	6.123		7.308	8.190	6.436

The number of bacteria after soaking was much higher than that of yeast mold colonies. It could be understood that in the soaking water, bacteria and mold competition occurs. However, the LAB was not detected. This was different result compared to previous study. Previous study reported that in the soaking water of soybean which would be processed into tempe had an enough LAB colonies/g product [7]. The mold and yeast population (PDA media) in Saga tempe dominated reaching 8.19 log CFU /g of product compared to the total bacteria (NA media) which only reached 7.308 and even for the total LAB (MRSA media) only reached 6.436 log CFU/g product. This made a lot of sense because the tempe starter contains mainly the mold spores of *Rhizopus oligosporus* and *Rhizopus oryzae*. The mold and yeast present in tempe starter, hydrolyzed saga bean with the microbial enzymes they produced during fermentation into simple compounds and as a result produced organic acid which could lower the pH. This decrease in pH provided comfort for LAB as evidenced by the growth of LAB in MRSA reaching 6.436 log CFU / g of product.

The bacteria growth curve of on NA media during the incubation process from 0 week to 4 weeks shown in Fig 4a. Thus, it could be said that the total growth of bacteria in tauco fermentation followed the usual bacterial growth curve, namely there was a lag phase or an early phase then an exponential phase in the first and second weeks, and until the peak, then in the third and fourth week it had decreased. Some of the factors determining this occurrence were the large variety of other microorganisms that contributed to the fermentation process of tauco. The variety of microbes in tauco greatly accelerated

the substrate hydrolysis process, this was indicated by the increase in the total dissolved solids during fermentation (Figure. 5a). The fermented soybean food is known for its attractive flavor, texture and superior digestibility [11] Therefore, the saga tauco also had the opportunity to be a product that was easily digested by the human body.

The dynamics of yeast and mold growth were very volatile (Figure 4b). It was assumed that the growth of mold and yeast species alternately at the beginning was dominated by aerobic fungi, then if oxygen availability ran out, yeast was dominated by the anaerobic tolerant and relatively halophilic yeast. And near the end of fermentation process the number of halophilic microorganisms predominated. Usually at the end of fermentation, salt-resistant yeast would dominate compared to other microbes.

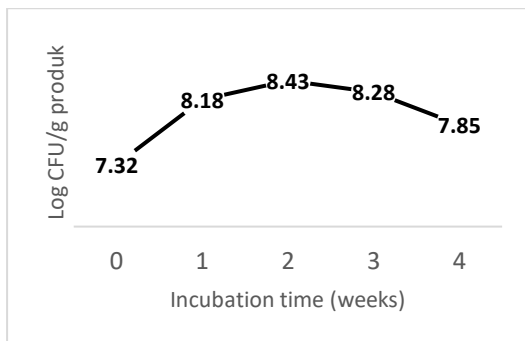


Figure 4a: Total growth of bacteria (NA media) in the Saga Tauco during the production process

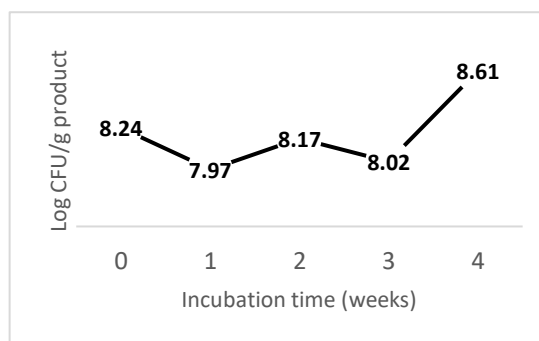


Figure 4b: Total growth of yeast and mold (PDA media) in the Saga Tauco during the production process

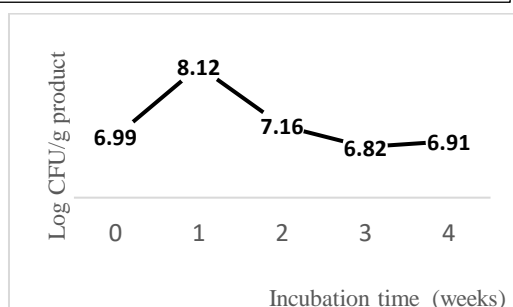


Figure 4c: Total growth of LAB (MRSA media) in the Saga Tauco during the production process

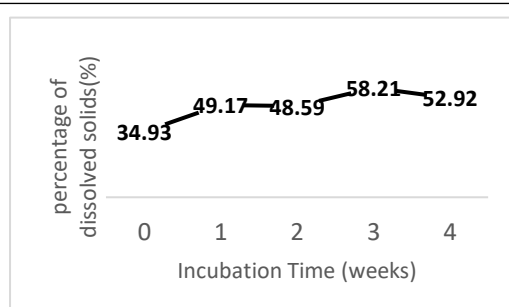


Figure 5a: The percentage of total dissolved solid in the Saga Tauco during the production process

During the fermentation process in a salt solution, all salt-resistant microbes, be they lactic acid bacteria (LAB), Yeast or even molds, synergized or might compete to hydrolyze the existing substrate so that the total dissolved solids increased as well as the pH value tended to rise slowly. When compared to Figures 4b and 4c there was a clear correlation in the first week of the maximum amount of LAB in contrast to minimal mold and yeast. It might be possible that LAB produces antimicrobials substance, so that mold and yeast life was suppressed. This result was in line with previous research which proved that LAB was able to suppress the growth of mold and yeast at the beginning of tempe fermentation [7]. In contrast to the research conducted by Feng et al 2005 [12], who reported that the growth of *Rhizopus oligosporus* on grain fermentation was not affected by the presence of LAB. The presence of LAB in saga tauco is very beneficial because it had the potential as a probiotic microorganism. Other researcher reported that traditional fermented food in Indonesia was a potential source of probiotics [13].

### 3.2 Chemical Analysis in Saga Tauco during the production process

The total dissolved solids in tauco during the production process had increased, (Figure 5a) indicating that tauco was a product rich in nutrients, at least a lot of dissolved solids that facilitated the absorption

system of the human body. This was due to the many microbial enzymes in tauco that came from yeast, mold, and LAB that were present in tauco during the production process. The total nitrogen in tauco during its production process fluctuated which was not too different, while the pH value tended to increase slightly. The pH value of the saga tauco ranged from 4.5 to 5.03 was due to the action of LAB. This was caused by protein hydrolysis by the protease enzyme from microorganisms, leading to the breakdown of amino acids into volatile compounds such as ammonia. Likewise, the ash content in the saga tauco during the production process slightly increased. (Table 2). Previous study reported that LAB produced organic acids which contributed to the sensory value of the fermented product [14]. In addition to the sufficient total dissolved solids, tauco made from soybean also contained sufficient antioxidants and total phenol which have a positive effect on body health. [15].

Table 2. Total Nitrogen content, pH value and Ash content of saga tauco during production process\*

Incubation time (week)	Total N content (%)	Total ash content (%)	pH value
0	13.80 <sup>a</sup>	6.85 <sup>c</sup>	4.50 <sup>a</sup>
1	13.68 <sup>a</sup>	5.84 <sup>a</sup>	4.80 <sup>b</sup>
2	13.53 <sup>a</sup>	5.94 <sup>a</sup>	4.83 <sup>b</sup>
3	14.05 <sup>a</sup>	6.34 <sup>b</sup>	4.82 <sup>b</sup>
4	14.08 <sup>a</sup>	6.14 <sup>ab</sup>	5.03 <sup>c</sup>

\*average of two replicates. the same letter in the same column shows no significant difference

#### 4. CONCLUSION

In saga tauco production process, this existing microbe's growth was very dynamic and contributed to the nutritional value of tauco proven by the increasing dissolved solids.

#### Reference

- [1] Biro Pusat Statistik 2019: Impor Kedelai Menurut Negara Asal Utama, 2010-2018. <https://www.bps.go.id/statictable/2019/02/14/2015/imp-or-kedelai-menurut-negara-asal-utama-2010-2018.htm>
- [2] Astawan M 2009 *Sehat dengan Hidangan Kacang dan Biji-bijian* Jakarta: Penebar Swadaya
- [3] Harti A S and Kusumawati H N 2013 *Jurnal KESMADASKA* (4) 2 :89-95
- [4] Nandiyanto A B D, Ismiati R, Indrianti J, Abdullah, A G 2017 *IOP Conference series. Materials Science and Engineering* 288. IOP Publishing
- [5] Amar A, Makosim S, Marwati 2017 *Jurnal IPTEK ITI* 1 (2) Doi: <https://doi.org/10.31543/jii.v1i2>
- [6] Umniyatie S and Mariyam S 2002 *Jurnal Chimera* 7 (2) <http://journal.um.ac.id/index.php/chimera/index>
- [7] Nurdini A L, Nuraida L, Suwanto A, Suliantari 2015 *International Food Research Journal* 22(4): 1668-1674
- [8] Fardiaz S 1989 *Petunjuk Laboratorium Analisis Mikrobiologi Pangan*. Pusat Antar Univ Pangan dan Gizi IPB
- [9] Legowo A M and Nurwantoro 2004 *Analisis Pangan*, Prodi Teknologi hasil ternak Fakultas Peternakan Universitas Diponegoro Semarang
- [10] Andarwulan N, Kusnandar F, Hariyadi P 2011 *Analisis Kimia pangan* Universitas Terbuka Jakarta
- [11] Nouts M J R and Kiers J L 2005. *Journal of Applied Microbiology* 98, 789–805 doi:10.1111/j.1365-2672.2004.02471.x
- [12] Feng X M, Erickson A R B, Schnurer, J 2005. *International Journal of Food Microbiology* 104.249-256
- [13] Nuraida L 2015 *Food Science and Human Wellness* 4 (2015) 47–55
- [14] Rathore S, Salmerón I, Pandiella S 2012 *Food Microbiology* 30: 239–244.
- [15] Larasati N 2017 *Jurnal Pangan dan Agroindustri* Vol.5 No.2:85-95.

#### Acknowledgements

Thanks to Research and Public Service Centre Institut Teknologi Indonesia Serpong for financial support during research with the contract number 023/KP/PRPM-PP/ITI/VII/2020





PRESENTER\_abu amar &lt;aamar3884biugm@gmail.com&gt;

---

## Proceedings of 2nd International Conference of Bio-Based Economy for Application and Utilization now available online

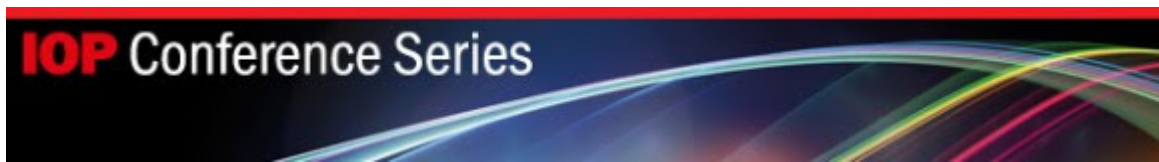
---

IOP Conference Series team <jpcs@iopublishing.org>  
Reply-To: IOP Conference Series team <jpcs@iopublishing.org>  
To: aamar3884biugm@gmail.com

Tue, Jun 1, 2021 at 8:00 PM

Your article [Growth dynamics of mold-yeast and bacteria during the production process of saga tauco \[\*Adenanthera pavonina\*\]](#) is online.

Visit [iopscience.org/jpcs](https://iopscience.org/jpcs) | [View this email online](#) | [Unsubscribe](#)



### Proceedings of 2nd International Conference of Bio-Based Economy for Application and Utilization.

Thank you for publishing your paper '[Growth dynamics of mold-yeast and bacteria during the production process of saga tauco \[\*Adenanthera pavonina\*\]](#)' in the IOP Conference Series: Earth and Environmental Science™. Your article has now been published online.

#### Create an account in ScholarOne

As part of our commitment to provide the best possible publishing service to our authors, we encourage you to [create an account in ScholarOne](#), so you can benefit from the following:

- Invitations to write and referee papers in your research area.
- Stay up-to-date on resources available to help you with getting your work published and promoted.
- Associate your existing ORCID ID with your account or [create a free ORCID ID](#).

[Create a ScholarOne Account →](#)

Thank you, and we hope to work with you again soon.

**Anete Ashton**

Publisher, Conference Series

[Journal of Physics: Conference Series](#)

[jpcs@iopublishing.org](mailto:jpcs@iopublishing.org)

**IOP Publishing**

This email has been sent to you because it is a required legal notice, customer update or other important alert. It is not a marketing or promotional email. This is why you are receiving this email even though you may have unsubscribed from IOP Publishing marketing emails.

For more information, please see our [privacy policy](#).

**IOP Publishing Limited** Registered in England under Registration No 467514. Registered Office: Temple Circus, Temple Way, Bristol BS1 6HG England VAT No GB 461 6000 84. Please consider the environment before printing this e-mail.