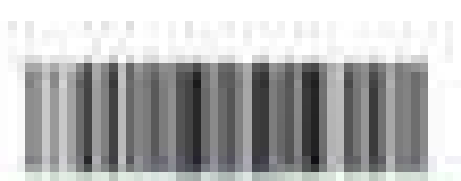


Official Journal of the International Association of Agricultural Engineers (IAAE)  
ISSN 2254-2754 (Print) / ISSN 2254-2762 (Online)  
Volume 11, Number 1, 2017, 1-10  
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DOI: 10.1016/j.cjast.2017.01.001

# Current Journal of Applied Science and Technology

Volume 11, Number 1, 2017  
ISSN 2254-2754 (Print)  
ISSN 2254-2762 (Online)



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# Editorial Policy

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## Journal Information:

**New name:** Current Journal of Applied Science and Technology

**ISSN:** 2457-1024

**Past name:** British Journal of Applied Science & Technology

**ISSN:** 2231-0843 (old)

**NLM ID:** 101664541

**Frequency:** 15 Days

**Journal DOI:** <http://dx.doi.org/10.9734/cjast> (<http://dx.doi.org/10.9734/cjast>)

**Peer-review model:** Advanced OPEN peer review

**NAAS Score (2019):** 5.32 (JrnID: B148)

**Digital Archiving:** Journal Repository (JR), LOCKSS, CLOCKSS

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### **Peer Review Mechanism**

(Up to 6th August 2012, all journals followed strict double blindfold review policy to ensure neutral evaluation. During this review process identity of both the authors and reviewers are kept hidden to ensure an unbiased evaluation.)

### **Advanced OPEN peer review:**

We have migrated to transparent and toughest 'Advanced OPEN peer review' system (Detailed general information is available in this link). High-quality manuscripts are peer-reviewed by minimum two peers in the same field. OPEN peer review system provides the provision to reveal the identities of the authors and reviewers to each other during the review process. In order to add transparency further, details of all reviewers and academic editors are published in the first page of every published paper (in the Article Information section: see example). As a final step to provide highest level transparency in the process, all review comments, authors' feedbacks, all versions of the manuscript and editorial comments are published (along with date) with the paper in 'Review History' link (See example 1, example 2, example 3, etc). This transparent process will help to eradicate any possible malicious/purposeful interference by any person (publishing staff, reviewer, editor, author, etc) during peer review. As a result of this unique system, all reviewers will get their due recognition and respect, once their names are published in the papers (Example Link). If reviewers do not want to reveal their identities, we will honour that request. In that case, only the review reports will be published as 'anonymous reviewer report'.

Additionally 'Advanced OPEN peer review' greatly helps in 'continuity and advancement of science'. We strongly believe that all the files related to peer review of a manuscript are valuable and hold an important place in the continuity and advancement of science. If publishers publish the peer review reports along with published papers, this process can result in savings of thousands of hours of future authors during experiments, manuscript preparation, etc. by minimizing the common errors after reading these previously published peer review reports. Therefore, as per our new official policy update, if the manuscript is published, all peer review reports will be available to the readers. All files (like the original manuscript, comments of the reviewers, revised manuscript, and editorial comment (if any)) related to the peer review, will be available in "Review history" link along with the published paper (Example Link).

Additionally, we believe that one of the main objectives of peer review system is 'to improve the quality of a candidate manuscript'. Normally we try to publish the 'average marks (out of 10)' a manuscript received at initial peer review stage and at final publication stage to record its history of improvement during peer

review. This process further increases the transparency. It is more important to record honestly the 'strength and weakness of a manuscript' than claiming that 'our peer review system is perfect'. Therefore, these transparent processes (i.e. publication of review history files and scores of a particular manuscript) additionally give a clear idea of the strength and weakness of a published paper to the readers, which enhances the chances of proper use of the result of a research (and or reduces the chances of misuse of the weakness of the findings of the paper). Thus this transparent process may prove to be highly beneficial for the society in long run.

We strongly discourage any attempt by the authors to contact the reviewer directly to influence the review process. We also strongly discourage any attempt by the reviewers to contact the authors directly. General guidelines for Peer-review Process are available below.

### **Reviewer suggestion**

It is a prerequisite to submit, with the manuscript, the names, addresses and e-mail addresses of 4 potential reviewers (When suggesting peer reviewers, please follow these guidelines to avoid any probable conflict of interest. Suggested reviewers should not: i) be from the same department or division as one of the authors (the same university, state, country should also be avoided); ii) have been a research guide or student of one of the authors within the past 10 years; iii) have collaborated with one of the authors within the past 10 years; iv) be employees of non-academic organizations with which one of the authors has collaborated within the past 10 years). It is the sole right of the editorial team to decide whether suggested reviewers to be used or not.

### **Reviewer selection**

Reviewer selection is a critical parameter to maintain the high peer review standard of any journal. Many factors are considered during peer reviewer selection like: proof of expertise in terms of published papers in the same area in reputed journals, affiliation, and reputation, specific suggestion, etc. We try to avoid reviewers who are slow, careless or do not provide sufficient justification for their decision (positive or negative). Authors can also identify peers that they want not to review their paper. As far as possible, the editorial team respects requests by authors to exclude reviewers whom they consider to be unsuitable. We also, as much as possible, try to rule out those reviewers who may have an obvious competing interest.

The main force behind our fast, efficient and quality Peer review system is the tremendous hard work of our Peer Reviewers & Editors. We are extremely grateful to the peer reviewers and editors for their great service. A combined list of contributing Peer Reviewers for all journals is published on our official website. From 2012 we have migrated to OPEN peer review system and after the migration, we have stopped updating this list.

### ***Review process flow***

The reviewers' comments are generally sent to authors within 3 weeks after submission. With the help of the reviewers' comments, FINAL decision (accepted or accepted with minor revision or accepted with major revision or rejected) will be sent to the corresponding author. Reviewers are asked if they would like to review a revised version of the manuscript. The editorial office may request a re-review regardless of a reviewer's response in order to ensure a thorough and fair evaluation. Reviewers who may have offered an opinion not in accordance with the FINAL decision should not feel that their recommendation was not duly considered and their service not properly appreciated. Experts often disagree, and it is the job of the editorial team to make a FINAL decision.

Authors are encouraged to submit the revised manuscript within 7-15 days of receipt of reviewer's comment (in case of minor corrections). But at any case, the revised manuscript submission should not go beyond 8 weeks (only for the cases of major revision which involves additional experiment, analysis etc.), in order to maintain this journal's mission of the fast publication. Along with corrected manuscript authors need to submit filled 'review comment form', any rebuttal to any point raised by reviewers. The Editor of the journal will have exclusive power to take the final decision regarding acceptance or rejection of a manuscript during peer review process.

One of the main policies of this journal is 'fast spreading of scientific findings' by publishing suitable manuscripts within 6 weeks after submission (except some abnormal cases). Under special circumstance, if the review process takes more time, author(s) will be informed accordingly. The editorial board or referees may re-review manuscripts that are accepted pending revision. Manuscripts with latest and significant findings will be handled with the highest priority so that it could be published within a very short time. The journal is determined to promote integrity in research publication. In case of any suspected misconduct, journal management will reserve the right to re-review any manuscript at any stages before final publication.

### **General guidelines for Peer Review Process**

- This journal strongly opposes the practice of duplicate publication or any type of plagiarism. If you suspect any unethical practice in this manuscript, kindly write it in the report with some proof/web links.
- Studies which are carried out to reconfirm / replicate the results of any previously published paper with new data-set, may be considered for publication. But these types of studies should have a 'clear declaration' of this matter.
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- Results & discussion (Kindly comment on: 1. Are the data well controlled and robust? 2. Authors should provide relevant and current references during discussion. 3. Discussion and conclusions should be based on actual facts and figures. Biased claims should be pointed out. 4. Are statistical analyses must for this paper? If yes, have sufficient and appropriate statistical analyses been carried out?)
- Conclusion (Is the conclusion supported by the data, discussed inside the manuscript? Conclusions should not be biased and should be based on the data, presented inside the manuscript only. Authors should provide adequate proof for their claims without overselling them)
- Are all the references cited relevant and adequate? Are there any other suitable current references authors need to cite?
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true comments with unfair criticism.

- We are very much reluctant to go against suggestions (particularly on technical areas) of the reviewers. Therefore, authors are requested to treat the suggestions of reviewers with utmost importance.

- Appeal: Rejected papers are given the opportunity for a formal appeal. Appeal requests should be made in writing, not by telephone, and should be addressed to [contact@sciencedomain.org](mailto:contact@sciencedomain.org) with the word "appeal" in the subject line. If an author remains unsatisfied, he or she can write to the Editorial Office, citing the manuscript reference number. In all these cases, it is likely that some time will elapse before journal can respond, and the paper must not be submitted for publication elsewhere during this time. Authors should provide detailed reasons for the appeal and point-by-point responses to the reviewers' and/or Academic Editor's comments. Authors should also be aware that priority is given to new submissions to the journal and so the processing of the appeal may well take longer than the processing of the original submission. If an appeal is rejected, further appeals of the decision will not be considered and the paper may not be resubmitted.

### **Post-publication peer review:**

1. Journal Web sites provide the ability for users to comment on articles to facilitate community evaluation and discourse around published articles. The comment section is mainly dedicated to promote "Post-publication peer review". Therefore, all journals strictly follow 'pre-publication OPEN peer review' and strongly encourage "Post-publication peer review". As a result of this "Post-publication peer review", if authors agree and or journal Editors agree (and or journal agrees) that any correction is necessary, then it will be published FREE of cost by following Correction and retraction policy (<http://www.journalcjust.com/index.php/CJAST/correction-and-retraction-policy>).

2. Users, who want to comment, are encouraged to register on website. But if anybody doesn't want to register, we'll respect the decision. In order to honour 'free flow of thoughts' unregistered user are also welcome to comment. Social login is also encouraged.

3. At the end of every comment, the user must identify himself/herself by providing the following information 1. Full Name 2. Name of the Department, University, institute, etc. (This two information will be displayed publicly). We don't like 'anonymous' comments. Comments with 'forged identity' will be deleted.

Note: Users must see and agree to our complete Comment Policy

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Plagiarized manuscripts would not be considered for publication. If plagiarism is found in any published paper after an internal investigation, a letter would be immediately sent to all the authors, their affiliated institutes and funding agency, if applied and subsequently the paper will be retracted.

Plagiarism policy of this journal is mainly inspired by the plagiarism policy of The Nature. Plagiarism policy of this journal is described below:

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2. Plagiarism can be said to have clearly occurred when large chunks of text have been cut-and-pasted. Such manuscripts would not be considered for publication in this journal. Papers with confirmed plagiarisms are rejected immediately.
3. But minor plagiarism without dishonest intent is relatively frequent, for example, when an author reuses parts of an introduction from an earlier paper.
4. Duplicate publication, sometimes called self-plagiarism, occurs when an author reuses substantial parts of his or her own published work without providing the appropriate references. This can range from getting an identical paper published in multiple journals, to 'salami-slicing', where authors add small amounts of new data to a previous paper. Self-plagiarism, also referred to as 'text recycling', is a topical issue and is currently generating much discussion among editors. Opinions are divided as to how much text overlap with an author's own previous publications is acceptable. We normally follow the guidelines given in COPE website. Editors, reviewers, and authors are also requested to strictly follow this excellent guideline (Reference: Text Recycling Guidelines: <http://publicationethics.org/text-recycling-guidelines>).
5. In case of 'suspected minor plagiarism', authors are contacted for clarification. Depending on all these reports, reviewers and editors decide final fate of the manuscript. If the manuscript is finally accepted and published, then to maintain transparency, all these reports are published in 'publication history' of the paper by following Advanced OPEN peer review system. The journal editors judge any case of which they become aware (either by their own knowledge of and reading about the literature, or when alerted by referees) on its own merits.
6. Use of the automated software is helpful to detect the 'copy-paste' problem. All submitted manuscripts are checked by the help of different databases, eTBLAST, Plagiarism Detection tools, etc. At the same time scientific implication of the case ('suspected minor plagiarism'), also judged by reviewers and editors. Plagiarism Detection tools are useful, but they should be used in tandem with human judgment and discretion for the final conclusion. Therefore, suspected cases of plagiarisms are judged by editors on 'case-to-case basis'.
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Some useful information is available at the following links:

1. <http://www3.imperial.ac.uk/library/researchers/plagiarismdetection>
2. <http://www.nature.com/authors/policies/plagiarism.html>
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As a part of restructuring all journals, we are closing all the manuscripts, where manuscripts are pending dormant for more than 4 weeks after the final acceptance mail. Due to the restructuring of our editorial policy and regulations, we have closed all the files of these types of manuscripts. Files of these types of manuscripts can be kept alive if authors agree for a fresh round of peer review by at least two peer reviewers or re-approved by the present editorial board.

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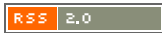
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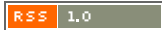
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# Effect of Temperature and Long Storage of Fatty Acid Profile, Value Peroxide and Value TBA Fillet of Snapper (*Lutjanus sp*)

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**Published** Jul 25, 2019

**DOI** <https://doi.org/10.9734/cjast/2019/v36i530249> (<https://doi.org/10.9734/cjast/2019/v36i530249>)

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## Abstract

This research was conducted at the Fisheries Technology Laboratorium Department of Fisheries and Marine Sciences, Gorontalo State University, at the Mey 2018 for three months in a month. Fish has a high nutritional value and is a major food source in many countries. Lipid fish has a high content of polyunsaturated acids (PUFA), especially eicosapentaenoic acid (EPA;20:5n-3) and docosahexenoate acid (DHA; 22: 6n-3). This study aims to the impact effect of temperature and storage time of the fatty acid snapper fillet (*Lutjanus sp*) and damage caused by the storage process. The sample selection stage is Snapper (*Lutjanus sp*) fresh weeds, and the meat is released by gills, bones, then washed and rinsed with ice water to remove blood and other debris. Continue does the cleavage from head to tail without causing his back to be disturbed. Filleting is done by slicing the meat ribs longitudinally to produce boneless meat. Stored at temperatures of 0, 10, 20, 30 and 40°C for 45 days. The results of fatty acid analysis snapper fillet (*Lutjanus sp*) has a saturated fatty acid lauric acid, acid tridecanoic, myristate acid, pentadecanoic acid, palmitic acid, stearic acid, heneicosanoic acid, behenic acid, lignoceric acid, acid and acid heptadecanoate arachidic. Monounsaturated fatty acids oleic acid, nervonic acid, palmitoleic acid, acid and acid erucat eicosanoic, while polyunsaturated fatty acids eicosatetraenoate acid, dokosadinoat



acid, eicosapentaenoic acid, eikosatrioanot acid, arachidonic acid and linolenic acid are susceptible to oxidative damage. Increased primary peroxide snapper fillet (*Lutjanus sp*) the higher the storage temperature increases from 0°C to 40°C. TBA value increases from 0° to 40°C with increased storage time of one day to 45 days storage time. From the results of this study found a way for Snapper fish fillets (*Lutjanus sp*) to have a longer expiration in storage, from 1 day to 45 days without reducing the Omega and DHA content.

Keywords:

Fillet snapper (*Lutjanus sp*), fatty acids, peroxide value and TBA value.

How to Cite

Husain, R., & Sulistijowati, R. (2019). Effect of Temperature and Long Storage of Fatty Acid Profile, Value Peroxide and Value TBA Fillet of Snapper (*Lutjanus sp*). *Current Journal of Applied Science and Technology*, 36(5), 1-9. <https://doi.org/10.9734/cjast/2019/v36i530249>

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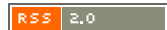
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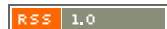
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## **Effect of Temperature and Long Storage of Fatty Acid Profile, Value Peroxide and Value TBA Fillet of Snapper (*Lutjanus sp*)**

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### **Authors' contributions**

*This work was carried out in collaboration between both authors. Both authors read and approved the final manuscript.*

### **Article Information**

DOI: 10.9734/CJAST/2019/v36i530249

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Complete Peer review History: <http://www.sdiarticle3.com/review-history/50066>

**Original Research Article**

**Received 02 May 2019**

**Accepted 08 July 2019**

**Published 25 July 2019**

### **ABSTRACT**

This research was conducted at the Fisheries Technology Laboratorium Department of Fisheries and Marine Sciences, Gorontalo State University, at the Mey 2018 for three months in a month. Fish has a high nutritional value and is a major food source in many countries. Lipid fish has a high content of polyunsaturated acids (PUFA), especially eicosapentaenoic acid (EPA;20:5n-3) and docosahexenoate acid (DHA; 22: 6n-3). This study aims to the impact effect of temperature and storage time of the fatty acid snapper fillet (*Lutjanus sp*) and damage caused by the storage process. The sample selection stage is Snapper (*Lutjanus sp*) fresh weeds, and the meat is released by gills, bones, then washed and rinsed with ice water to remove blood and other debris. Continue does the cleavage from head to tail without causing his back to be disturbed. Filleting is done by slicing the meat ribs longitudinally to produce boneless meat. Stored at temperatures of 0, 10, 20, 30 and 40°C for 45 days. The results of fatty acid analysis snapper fillet (*Lutjanus sp*) has a saturated fatty acid lauric acid, acid tridecanoic, myristate acid, pentadecanoic acid, palmitic acid, stearic acid, heneicosanoic acid, behenic acid, lignoceric acid, acid and acid heptadecanoate

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arachidic. Monounsaturated fatty acids oleic acid, nervonic acid, palmitoleic acid, acid and acid erucat eicosanoic, while polyunsaturated fatty acids eicosatetraenoate acid, dokosadinoat acid, eicosapentaenoic acid, eikosatrioanat acid, arachidonic acid and linolenic acid are susceptible to oxidative damage. Increased primary peroxide snapper fillet (*Lutjanus sp*) the higher the storage temperature increases from 0°C to 40°C. TBA value increases from 0° to 40°C with increased storage time of one day to 45 days storage time. From the results of this study found a way for Snapper fish fillets (*Lutjanus sp*) to have a longer expiration in storage, from 1 day to 45 days without reducing the Omega and DHA content.

**Keywords:** Fillet snapper (*Lutjanus sp*); fatty acids; peroxide value and TBA value.

## 1. INTRODUCTION

In general, the fish has no nutritional value and is a major food source in many countries. Lipid fish has a high content of polyunsaturated acids (PUFA), especially eicosapentaenoic acid (EPA; 20: 5n-3) and docosahexenoate acid (DHA; 22: 6n-3) [1,2]. Processing by freezing the fish has been used for thousands of years because of the quality and high product [3]. The concept of storage by freezing depends on the product temperature decrease to slow decay so that when the fish melted freshness can be maintained [4].

However, fish and fishery products may undergo undesirable changes during storage and can damage the storage time limit. Unwanted changes resulting from the oxidation of proteins [5,6] and lipid oxidation [7,8]. Fish protein experienced a number of changes (cause cannot form aggregates) that modify the structural and functional properties of fish muscle [9].

Degradation of polyunsaturated fatty acids (PUFA) by lipid oxidation during storage led to the formation of volatiles associated with rancidity [1]. The high level of unsaturated fats makes the fish tissue is very susceptible to peroxidation and easily damaged. Oxidative changes primarily related to the taste and texture of the fish. In later stages of the process of lipid peroxidation, changes in color and nutritional value will be observed or secondary products of lipid [10]. This study aims to determine the effect of temperature and storage time of the fatty acid snapper fillet (*Lutjanus sp*) and damages resulting from the storage process.

## 2. MATERIALS AND METHODS

### 2.1 Raw Material

Raw materials snapper (*Lutjanus sp*) obtained from the fish auction place (TPI) kobong, Village Kaligawe, Semarang, Central Java.

### 2.2 Sample Preparation Fish Fillets

The sample selection stage is Snapper (*Lutjanus sp*) fresh weeds, and the meat is released by gills, bones, then washed and rinsed with ice water to remove blood and other debris. Continues does the cleavage from head to tail without causing his back to be disturbed. Filleting is done by slicing the meat ribs longitudinally to produce boneless meat. Stored at temperatures of 0, 10, 20, 30 and 40°C for 45 days.

### 2.3 Proximate Analysis

Chemical analysis: Proximate moisture content, ash content, protein and fat by AOAC [11].

#### Fatty Acid Analysis Procedure

##### a. Hydrolysis

After selecting the sample, then the sample is cleaned with water, then mashed and weighed  $\pm$  10 g of the sample. Hydrolysis with 10 ml HCl. then the water is heated at a temperature of 70°C, boiling. Cold. Extract with 15 ml of diethyl ether and 15 ml of oil, take the top layer. Extract again with 15 ml of diethyl ether and 15 ml of oil, take the top layer, make one with the previous results. Steamed in a water bath with the help of N<sub>2</sub>.

##### b. Analysis of saturated fatty acids and unsaturated

Taken 0.5 ml sample, add 1.5 ml of sodium methanolic solution, cover and heat at 70°C for 5-10 minutes while being shaken. Chill. Add 2 ml Boron trifluoride ethanoic, heat at 70°C for 5-10 minutes. Chill. Extract with 1 ml of heptane and 1 ml saturated NaCl. Take the top layer and put in Eppendorf. Injected into the GC. Injected as much 1 $\mu$  samples on GC - 2010 SHIMADZU.

## 2.4 Analysis Value Peroxide

Samples (0.5 g) put in a test tube, then added 0.1 ml of a solution of ammonium thiocyanate and 0.1 ml of solution feroklorida. The test tube is shaken for 5 seconds and heated at 50°C for 2 minutes, then cooled to a temperature of 25°C. Use absorbance using a spectrophotometer at a wavelength of 510 nm. A blank solution prepared using all solvent without the sample (Hills and Thiel, 1946 modified Adnan [12]).

## 2.5 Analysis Value TBA (Expand)

Analysis of figures TBA performed according to the method Tokur and Korkmaz [13]. Oil added 0.5 g in 50 ml of distilled water, then add another 2.5 mL N HCl then distilled. Absorb distilled to 50 ml, 5 ml of distilled download then added with 5 ml TBA. Thereafter, heated for 30 minutes and cooled. Absorbance at a wavelength of 528 nm. Value TBA = mg malonaldehyde / kg of oil.

## 3. RESULTS AND DISCUSSION

### 3.1 Proximate Fish Snapper (*Lutjanus sp*)

Based on Table 1 fat content snapper (*Lutjanus sp*) in the study was 1.96% compared to that reported by de Castro et al. [14] on tilapia (*Oreochromis niloticus*), i.e 0.79% and tambagui (*Colossoma macropomum*) i.e 1.30%. The fat content rough on snapper (1.96%) is still higher, it indicates that the snapper (*Lutjanus sp*) belong to the fish that contains high fat enough.

**Table 1. Proximate fish snapper (*Lutjanus sp*) and Fe content**

No	Chemical composition	Amount (%)
1.	Moisture	78,39
2.	Ash	1,58
3.	Protein	18,77
4.	Fat	1,96
5.	Carbohydrate by difference	0,30
6.	Fe (ppm)	121,7

Description: Data is derived from repeat 3x

Differences in fish fat content are strongly influenced by the type of fish, fish size, fishing

season and the environment in which the fish live. According to Shaviklo [15] demersal fish higher in fat than pelagic fish that lives in surface waters. Demersal fish usually live in the bottom waters and rarely engage in the activity. In Table 2 also shows the snapper (*Lutjanus sp*) containing Fe total was 121.47 ppm. Okada report [16] describes the color of the flesh of fish due to the Fe content in the meat is very high because it is rich hemoprotein (80%) mainly myoglobin and hemoglobin. Based Okada [16], the content of white meat hemoprotein red snapper low so the meat is white.

### 3.2 Composition Fatty Acid

Based on the analysis of fatty acids snapper fillet (*Lutjanus sp*) are listed in Table 4, snapper fillet (*Lutjanus sp*) has saturated fatty acids (Saturated Fatty Acid, SFAs), which consists of lauric acid, acid tridecanoic, acid myristate, acid pentadecanoic, palmitic acid, stearic acid, heneicosanoic acid, behenic acid, lignoceric acid, acid and acid heptadecanoatearachidic. Acids monounsaturated fats (MUFA) consisting of: oleic acid, acid nervonate, palmitoleic acid, acid erucat and sour eicosanoic, while the unsaturated fatty acid compound (Poly Unsaturated Fatty Acids, PUFAs) consisting of acid eicosatetraenoate, dokosadinoat acid, eicosapentaenoic acid, eikosatrioanate acid, arachidonic acid and linolenic acid are susceptible to oxidative damage.

According Nazemroaya et al. [17]; Karami et al. [18] that when the fish are kept in a long time then, the amount of saturated fatty acids (Saturated Fatty Acid, SFA) will increase as well as fatty acids monounsaturated (Mono Unsaturated Fatty Acid, MUFA) will be slightly experience improved although not too significant. While polyunsaturated fatty acids (Poly Unsaturated Fatty Acids, PUFAs) experienced a reduction process. This is similar to the research conducted at the snapper fillet (*Lutjanus sp*) during storage.

Aranda et al. [19] states that the oxidation of PUFAs begins with the production of hydrogen peroxide with three different ways: (1) autooksidasi, (2) enzymatic oxidation, and (3) photooxidation, which can also occur simultaneously. Product peroxide decomposes by different mechanisms, forming a secondary oxidation products; both the primary and secondary oxidation products can react with

amino groups of proteins, producing compounds that modify the interaction of taste, smell and functional properties of proteins.

Furthermore, the fraction of volatile compounds is indicated as the cause of rancidity in fatty fish [19]. Sebcenic and Beutelspecher [20] states that the oxidation is the process of fat breakdown and lead to the formation of off-flavor compounds and condition is called rancid (rancid). Processed food products were rancid, may experience discoloration and loss of nutritional value for the oxidation of unsaturated fatty acids (PUFA) that impact on the quality. The compound oxide such as peroxides, aldehydes, and ketones harmful to human health. While McClement and Decker [21] states the factors that affect the speed of oxidation include the number and type of oxygen, the chemical structure of lipids, antioxidant and prooxidant form: ferrous metals, sensitiser, storage temperature and properties of packaging materials.

### 3.3 Effect of Temperature and Time against Value Peroxide

Value peroxide as primary products of oxidation snapper fillet (*Lutjanus sp*) during storage at different temperatures and times can be seen in Table 3.

In the treatment of frozen storage temperature (0°C) shows the formation of peroxides to 45 days of storage. Speed peroxide formation increased 9.6 times at a temperature of 0°C with a storage temperature increases, while at a temperature of 10, 20, 30, and 40°C peroxide value increased respectively 14.33, 16.12, 19.19, and 23.05 times meq/kg sample with increased storage temperature. The peroxide values increased from 0° to 40°C with increased storage time of one day to 45 days storage time. Increased primary peroxide snapper fillet (*Lutjanus sp*) the higher the storage temperature increases from 0°C to 40°C.

**Table 2. Composition fatty acids of snapper fillet (*Lutjanus sp*) during storage**

No	Profil asam lemak (%)	Storage temperature (°C)				
		0°C	10°C	20°C	30°C	40°C
1	Lauric acid	0,703	-	-	0,99	-
2	Tridecanoic acid	0,001	-	-	0,86	-
3	Myristic acid	4,687	0,74	4,44	2,69	0,51
4	Pentadecanoic acid	0,001	0,90	1,03	2,46	0,38
5	Palmetic acid	2,283	1,67	4,39	3,58	12,94
6	Stearic acid	0,708	-	9,19	15,46	12,14
7	Heneicosanoic acid	13,462	-	0,78	2,03	-
8	Behenac acid	-	1,04	0,65	0,68	1,54
9	Lignoserat acid	0,762	-	0,74	-	0,55
10	Heptadecanoic acid	17,811	-	1,96	-	-
11	Arachidic acid	2,945	-	-	-	-
I	Saturated Fatty Acid (SFA)	25,552	4,35	32,73	31,75	28,06
1	Oleic acid	8,807	12,63	16,37	22,70	10,82
2	Nervonac acid	1,64	1,34	1,03	1,27	-
3	Palmitoleic acid	13,239	2,59	-	-	-
4	Erucic acid	-	5,43	5,26	5,43	3,58
5	Eicosanoic acid	0,001	8,34	-	-	-
II	Mono Unsaturated Fatty Acid(MUFA)	23,687	23,72	22,66	29,4	14,4
1	Eicosatetraenoic acid	-	-	-	5,60	1,05
2	Docosadinoic acid	0,827	-	0,43	-	0,70
3	Eicosapentanoic acid	6,055	3,52	2,29	0,43	-
4	Eicosatrienoic acid	1,557	2,6,23	9,82	-	-
5	Arachidonac acid	0,692	-	-	-	-
6	Docosaheksanoic acid	5,31	-	0,17	0,25	-
7	Linoleic acid	10, 612	1,52	3,27	2,56	3,57
III	Poly Unsaturated Fatty Acid (PUFA)	25,055	14,99	15,98	8,84	10,32
Total		74,294	103,07	101,37	99,99	82,78

Description: Data is derived from repeat 2x

**Table 3. Date analysis peroxide value snapper fillet (*Lutjanus sp*)**

<b>Temperature 0°C</b>		
<b>N0</b>	<b>Storage temperature (day)</b>	<b>Peroxide Value (meq/kg)</b>
1	0	1.7411
2	5	2.9437
3	10	4.06055
4	15	5.296
5	20	6.7797
6	25	6.29125
7	30	10.905
8	35	11.4923
9	40	12.98225
10	45	13.21835
<b>Temperature 10°C</b>		
<b>N0</b>	<b>Storage temperature (day)</b>	<b>Peroxide Value (meq/kg)</b>
1	0	1.7411
2	3	3.26525
3	6	5.63635
4	9	6.90375
5	12	6.3911
6	15	9.60175
7	18	10.78685
8	21	10.4857
9	24	14.5174
10	27	15.9977
<b>Temperature 20°C</b>		
<b>N0</b>	<b>Storage temperature (day)</b>	<b>Peroxide Value (meq/kg)</b>
1	0	1.7411
2	1	4.21665
3	2	6.96335
4	3	8.3699
5	4	8.5134
6	5	11.22005
7	6	11.963
8	7	15.59645
9	8	15.9638
10	9	18.0701
<b>Temperature 30°C</b>		
<b>N0</b>	<b>Storage temperature (day)</b>	<b>Peroxide Value (meq/kg)</b>
1	0	1.7411
2	0.5	4.5663
3	1	7.06935
4	1.5	9.2808
5	2	10.422
6	2.5	12.1345
7	3	11.83165
8	3.5	16.974
9	4	18.703
10	4.5	20.4086



<b>Temperature 40°C</b>		
<b>NO</b>	<b>Storage temperature (day)</b>	<b>Peroxide Value (meq/kg)</b>
1	0	1.7411
2	0.25	4.67575
3	0.5	7.7808
4	0.75	10.7164
5	1	12.06535
6	1.25	12.6637
7	1.5	18.4146
8	1.75	18.42355
9	2	22.9266
10	2.25	24.62795

Description: Data is derived from repeat 3x

**Table 4. Data analysis value TBA snapper fillet (*Lutjanus sp*)**

<b>Temperature 0°C</b>		
<b>No</b>	<b>Storage temperature (day)</b>	<b>Value TBA (mg MDA/kg)</b>
1	0	2.307378
2	5	3.504316
3	10	5.277034
4	15	6.573477
5	20	6.315832
6	25	9.565049
7	30	11.91097
8	35	11.75207
9	40	13.71696
10	45	14.62961

<b>Temperature 10°C</b>		
<b>No</b>	<b>Storage temperature (day)</b>	<b>Value TBA (mg MDA/kg)</b>
1	0	2.307378
2	3	4.299369
3	6	6.748229
4	9	6.105547
5	12	10.76352
6	15	13.41159
7	18	13.2912
8	21	15.61061
9	24	16.80039
10	27	17.21704

<b>Temperature 20°C</b>		
<b>No</b>	<b>Storage temperature (day)</b>	<b>Value TBA (mg MDA/kg)</b>
1	0	2.307378
2	1	4.424119
3	2	6.195105
4	3	6.932098
5	4	10.79865
6	5	13.33775
7	6	13.45672
8	7	17.74074
9	8	18.62311
10	9	20.31495

Temperature 30°C		
No	Storage temperature (day)	Value TBA (mg MDA/kg)
1	0	2.307378
2	0.5	5.887911
3	1	7.589422
4	1.5	8.931236
5	2	8.773049
6	2.5	13.85214
7	3	14.5131
8	3.5	14.5922
9	4	20.7325
10	4.5	22.03099
Temperature 40°C		
No	Storage temperature (day)	Value TBA (mg MDA/kg)
1	0	2.307378
2	0.25	5.975746
3	0.5	7.946435
4	0.75	8.185896
5	1	8.54802
6	1.25	10.79128
7	1.5	15.43709
8	1.75	15.59018
9	2	20.52694
10	2.25	23.57116

Description: Data is derived from repeat 3x

According to Pak et al. [22], peroxide value is an indicator of the stability of the oil against oxidation, the oxidation products of primary lipid parameters, namely hydroperoxide. Oxidation of lipids/oils naturally easily occurs, because tuna fish oil rich in PUFAs (6 double bonds), while oil containing many double bonds susceptible to lipid oxidation reactions. Thus, the oxygen molecules that are attached to the double bond susceptible to oxidation.

### 3.4 Effect of Temperature to Value TBA

Value TBA is used to measure the secondary products of lipid oxidation, especially coming from PUFA (hide no, 2012) and indicates the level of rancidity, especially in the high PUFA-containing oil [23]. TBA formation as a secondary oxidation products snapper fillet (*Lutjanus sp*) during storage can be seen in Table 4.

Value TBA increased 7.21 times at a temperature of 0°C with an increase in storage temperature, whereas at a temperature of 10, 20, 30, and 40°C peroxide value increased respectively to 9.19, 9.89, 10.63, and 12.59 times the MDA mg/kg of the sample with increased storage temperature. Value TBA increases from 0° to 40° with increased storage time of one day to 45 days storage time. In the treatment of

storage temperature of 0°C-40°C showed the formation of value TBA during storage [23].

## 4. CONCLUSION

Differences in fat content of the fish are strongly influenced by the freshness of the fish is used as a critical factor in resulted isolates, methods of isolation/extraction is used, homogenization of meat processing, the ratio of fish and solvents (viscosity) used, length of extraction, time and temperature processing and dissolution protein. Snapper fillet (*Lutjanus sp*) has a saturated fatty acid comprising: lauric acid, acid tridecanoic, myristate acid, pentadecanoic acid, palmitic acid, stearic acid, heneicosanoic acid, behenic acid, lignoceric acid, acid and acid heptadecanoatearachidic. Fatty acids are monounsaturated consisting of: oleic acid, acid nervonate, palmitoleic acid, acid erucat and sour eicosanoate, while the unsaturated fatty acid compound consisting of: acid eicosatetraenoate, acid dokosadinoate acid, eicosapentaenoic acid eikosatrioanate, arachidonic acid and linolenic acid susceptible to oxidative damage.

Speed formation of peroxide value increased 9.6 times at a temperature of 0°C with a storage temperature increases, while at a temperature of 10, 20, 30, and 40°C peroxide value increased

respectively 14.33, 16.12, 19.19 and 23.05 times meq/kg sample with increased storage temperature. Value TBA increased 7.21 times at a temperature of 0°C with an increase in storage temperature, whereas at a temperature of 10, 20, 30, and [10] 40°C peroxide value increased respectively to 9.19, 9.89, 10.63, and 12.59 times the MDA mg/kg of the sample with increased storage temperature.

## COMPETING INTERESTS

Authors have declared that no competing interests exist.

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