## STUDY OF N, P, K AND C ON DEGRADATION OF INDIGOSOL BATIK DYE EFFLUENT BY *Aspergillus* sp. GPN

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#### ABSTRACT

Indigosol dye batik effluent is toxic since it contains excess contains decreasing reducing agent sodium hydrosulfite which that is oxidized to alkali and alkaline earth metal sulfate  $(SO_4^{2^\circ})$ , sulfite  $(SO_3^{2^\circ})$  and thiosulfate (S<sub>2</sub>O<sub>3</sub><sup>2-</sup>) that very corrosive. Therefore, the role of environmental biotechnology in the technology for restoring environmental functions is needed efforts to treat batik effluent have been carried out biologically by using microorganism agents. It can be used Aspergillus sp. that has ability to reduce industrial dyes. From the previous research, product of Indigosol Blue batik effluent degradation that produced by Aspergillus sp. GPN did not cause any toxicity in plants even better growth than giving watering with irrigation water. Content in the results of degradation can affect the plants, so the element analysis needs to be performed. ItsIt's must including include substance nitrogen (N) in the form of organic compounds that easily absorbed by plants, does not leave residual organic acids in the soil, has high carbon (C) content and also Phosphorus (P)- and Potassium (K). The aims of this study were to analyse content of N, P, K, organic carbon, and ratio of organic carbon to total nitrogen (C:NC : N) in degradation product of Indigosol Blue dye batik effluent that produced by Aspergillus sp. GPN. The methode that used were spectrofotometry for organic carbon and Pand P2O5, Kjedal for total\_-N ,spectrofotometryN, spectrofotometry and AAS for Kfor K2O. The result of the assay were the product degradation of indigosol blue batik effluent had a total nitrogen content of 0.11%, K%, K2O of 0.01%, organic carbon of 0.24%, C/N ratio of 10.13%, while for phosphorus content of 0.01%. Fermentation of degradation products causes 0% frequency of wilted and dead plants, better when compared to water (40% frequency of wilted plants) and effluent (40% frequency of wilted and dead plants).

Keywords: Aspergillus sp., batik effluent, elements.

### 1. INTRODUCTION

Most of the textile wastewater, especially batik, contains dye residues that have potential to pollute the environment. It is extremely dangerous because it carries chemical substances which are dangerous contains chemicals that are harmful to the environment. The effluent has a detrimental effect on the content of organic matter, suspended solids, oil or fat, and the content of dangerous heavy metals such as Zn, Cd, Cu, Cr and Pb (Nurdalia, 2006). Generally, batik industrial effluent is discharged directly into water bodies or rivers without being treated first (Watini, 2009). One type of synthetic batik dye is Indigosol Blue that one of the synthetic dyes of Anthraquinone, and has molecular bonds of -NH and C=C. This synthetic dye is not easily damaged by chemical or photolytic treatment. This dye

waste has a strong chemical bond structure and is classified as non-biodegradable waste (Herfiani et al., 2017). Indigosol wastewater is toxic because it carries extra since it contains excess ontains excess reducing agent sodium hydrosulfitehydrosulphite which that sulfatesulphate (SO<sub>4</sub><sup>2</sup>), sulfitesulphite— (SO<sub>3</sub><sup>2-</sup>) and thiosulfate (S<sub>2</sub>O<sub>3</sub><sup>2-</sup>) hat very corrosive. Thus, the formed sulfatesulphate deposits can create toxic hydrogen sulfidesulphite(HS) ions (Blackburn et al., 2009).

Efforts to treat batik effluent have been carried out biologically by using microorganism agents. The microorganism agents that can be used is Aspergillus sp. that has ability to reduce industrial waste dyes. Study of Dewi et al. (2018) showed *Aspergillus* sp. 3 strain GPN is the <u>best superior</u> isolate for to decolorizing decolorize the <u>batik</u>effluent. Aspergillus species 3 <u>nearlyalmost</u> completely (99.9%) decolorized the effluent <u>with</u>in the liquid medium after a <u>three\_daythree-day</u> incubation. According to Dewi & Khotimah (2019), *Aspergillus* sp. 3 can remove effluent as a single carbon and nitrogen source dye.

The previous study findfinds that effluent batik dye degradation by *Aspergillus* sp. 3 did now no longer motivet-cause any toxicity in plants. Study of Dewi et al. (2018) confirmedshowed plumule and radicle length of *Zea mays* and *Vigna radiata* grown atom the decolorized effluent being longer than withinside the untreated effluent. The percentage of corn *Z. mays* and mung bean *V. radiata* seed germination on decolorized effluent turned into better was higher-than on untreated effluent. Even, its better growth than giving watering with irrigation water. Content in the results of degradation can affect the plants, so the element analysis needs to be performed.

Elements that can fertilize plants are substance nitrogen (N) in the form of organic compounds that easily absorbed by plants, does not leave residual organic acids in the soil, has high carbon (C) content and alsoand Phosphorus ( $\underline{P}$ ) and  $\underline{P}$ ) and Potassium (K).

Substance N in the form of organic compounds that easily absorbed by plants, does not leave residual organic acids in the soil and has high organic C content such as charcoal hydrates. Nutrients required of plant are not sufficiently available in the soil because the nutrients contained in the soil are relatively little, the quantity of nutrients cannot suffice certain plants, so it is necessary to add nutrients. The role of potassium in plants is to form protein and carbohydrates, harden straw and the bottom of the wood, increase plant retention against disease and improve seed or fruit quality (Effendi, 2004). According to Afandi et al. (2015) carbon is a food supply source for soil microorganisms, so the presence of organic C withinsidein the soil will stimulate the activity of microorganisms. Wahyudi (2009) statedaid that the increase in C-Organic caused by carbon (C) with the application of element on the soil will increase the organic substancematter content in the soil.

The aims of this study were to <u>analyse ANALYSE</u> content of N, P, K, organic carbon, and ratio of organic carbon to total nitrogen (C:NC: N) in degradation product of Indigosol Blue dye batik effluent that produced by *Aspergillus* sp. GPN.

## 2. METHODOLOGY

## 2.1. Reculture Aspergillus sp. GPN

Materials and tools are prepared, alcohol sprayed on palms and tables. The Bunsen fire is light. The wrapping paper of the petri dish is removed. The mouth of the petri dish is heated. A hole in the PDA made using a straw. A piece of PDA took using a loop needle. Inoculated on new PDA media, placed in the middle of the PDA. The petri dish is covered, heated then wrapped in plastic wrap. Then incubated at room temperature for 3-7 days.

#### 2.2. Preparation of Aspergillus sp. GPN inoculumGPN inoculum

Materials and tools are prepared, alcohol sprayed on palms and tables. The Bunsen fire is light. The wrap paper on the Erlenmeyer flask areis removed. The mouth of Erlenmeyer flask is heated. A hole in the PDA made using a straw. A piece of PDA took using a loop needle. Inoculated on PDB media. The Erlenmeyer flask is covered, heated then wrapped in plastic wrap. Then incubated on shaking incubator at room temperature for 3-7 days.

### 2.3. Efluent Degradation

Indigosol added on *Aspergillus* sp. 3 preparation then incubated on shaking incubator at room temperature. Observed in each time interval. The degradation result is filtered then used for Macro Nutrient Assay.

#### 2.4. Macro Nutrient Assay

#### 2.4.1. Assay of K Nutrient

The testing used Flame photometer. The 100 mL sample solution added in 100 mL volumetric flask. Diluted to the limit. The standard series of samples were made of 4 mL, 6 mL, and 8 mL. Each sample was diluted into a 100 mL volumetric flask. The absorbance measured using a flamefotometerflame photometer.

#### 2.4.2. Assay of C Nutrient

The C test used the walkey and black methods. A sample of 100 mL added with 5 mL of  $K_2Cr_2O_7 2 N$  solution and then shaked. The  $H_2SO_4$  pa. 98% of 7 mL is added then shaken. The solution is left for 30 minutes then shaken. Standard solution C of 5 ml is pipetted into a 100 ml volume measuring flask.  $H_2SO_4$  of 5 mL and 7 ml of  $K_2Cr_2O_7 2 N$  solution were added and then shaken. The solution is left for 30 minutes then shaken. Blanko as standard 0 ppm C were prepared by diluted the sample with ion-free water. After cool, the volume is adjusted to the mark of 100 ml. Then shaken back until homogeneous and overnight left. Then measured with a spectrophotometer at a wavelength of 651 nm.

## 2.4.3. Assay of P Nutrient

AnalyzAnalysinge the P content by inserting the waste sample in a cuvette before measuring it in a UV-Vis spectrophotometer. Blanks using distilled water were included in the cuvette. Then the absorbance value was measured using a UV-Vis spectrophotometer with a wavelength of 650-750 mn.

#### 2.4.4. Assay of N Nutrient

The sample for the analysis of the N content used the Kjeldahl micro method. There are three stages in the Kjeldahl micro method, namely digestion, distillation and titration which functions to obtain the values of Norganic, N-NH<sub>4</sub>, and N-NO<sub>3</sub>. The N-organic test begins to make titrant A as a sample, and A1 as a blank. The tTitran A and A1 were prepared by adding 0.25 g of selenium mixture to which 3 mL of H<sub>2</sub>SO<sub>4</sub> pa were added in a Kjeldahl flask, then shaken until evenly distributed and then left for 2 hours. Gradual digestion with a temperature of 150°C to a maximum temperature of 350°C for three3 hours and clear liquid. Cool then dilute with a touchlittle distilled water. The solution is transferred to a distillatordistillation boiling flask and then added a little boiling stone and distilled water until it reaches the volume of the flask. Then 10 mL of 40% NaOH were added. The destination container includes 10 mL of 1% boric acid in Erlenmeyer and 3 drops of Conway indicator liquid are added until the Erlenmeyer reaches 75 mL. Continue the distillate titration for A1 with 0.05 N H<sub>2</sub>SO<sub>4</sub> to the end point when the color changes from green to pink.

The N-NH<sub>4</sub> test was initiated using titrant B as the sample and B1 as the blank. Sample 1 g in a boiling distillator flask plus a little boiling rock, 0.5 mL of liquid paraffin, and 10 mL of distilled water for B while 100 mL for B1. Then add 10 mL of 1% boric acid, 3 drops of Conway indicator, and 10 mL of 40% NaOH and then distill it until it reaches 75 mL. The distillate was titrated with 0.05 N H<sub>2</sub>SO<sub>4</sub> until the color of the solution changed from green to pink.

Testing for N-NO<sub>2</sub> begins with titrant C as the sample and C1 as a blank with the same procedure. The remainder of the N-NH<sub>4</sub> determination of the sample could cool then added with distilled water to the original volume. The distillate is entered by adding 10 mL of 1% boric acid in 100 mL Erlenmeyer plus 3 drops of Conway indicator. Then distill it by adding 2 g of devarda alloy. Distillation is started indefinitely so that the foam does not overflow after the foam is almost exhausted starting from low temperature after boiling the temperature is increased to a normal. Distillation to 75 mL of liquid. Then titrated with 0.05 N H<sub>2</sub>SO<sub>4</sub> until the color of the solution changes from green to pink. The results of N-organic, N-NH<sub>4</sub>, and N-NO<sub>3</sub>. Were calculated using the formula then totaled.

## 2.5. Plant Assay

The degradation products obtained were then tested on plants. This experiment was compared with the degradation product which was mixed with a small amount of goat urine and then fermented using EM 4 (Effective Microorganism-4). Then the spinach plants were watered and compared with irrigation water and batik waste. Observations were made on the frequency of wilting and the frequency of death.

## **3. RESULTS**

In this research assay of N,PN, K, K, and C element content derived from the degradation process of Indigosol Blue batik effluent was carried out. The test results for assay of nutrient were presented in Table 1.

Table 1 showed results of total nitrogen content in degradation product which has a value of 0.11%, while for phosphorus it has a value of 0.01%. because it is possible to fertilize plants, it is compared with the content in-organic liquid <u>organic fertilizers (LOF)</u>. According to Minister of Agriculture Regulation No. 70/Permentan/SR.140/10/2011, the minimum quality standard for nitrogen and phosphorus content of <u>LOFliquid organic fertilizer</u> is 3-6%. The results degradation product of this research werewas not reach quality standards for nitrogen and phosphorus content has a value of <1%.

 Table
 1.
 The Results for Assay nutrient from

 Degradation Indigosol Blue Batik Effluent

No	Parame-	Unit	Method	Result •
	ter			
1	N-Total	%	Kjeidahl	0.11
2	$P_2O_2$	%	Spekrofotometri	0.01
3	C-organic	%	Spectrophotometric	0,24
4	K <sub>2</sub> O	%	Spectrophotometric,	0,01
			AAS	
5	C/N	%	-	10.13
	Ratio			

Minister of Agriculture Regulation No.70/Permentan/SR.140/10/2011 contains standards for LOFHquid organic fertilizers, but types of samples obtained in this research cannot be categorized as Hquid organic fertilizersLOF. Nitrogen content obtained is 0.11% and Phosphorus is 0.01%. According to its regulation, the minimum quality standard for nitrogen and phosphorus content of LOFHquid organic fertilizer is 3-6%. Based on the research of Ni'am et al. (2015), stated that the fermented cassava wastewater with addition of cow urine, and foot of cow washing water contained 5.11% Nitrogen and 3.08% Phosphorus.

Plants need one of the macro nutrients Nitrogen for protein formation and vegetative growth Nitrogen is a macro nutrient needed by plants for vegetative growth and protein formation (Sarief, 1986). Nitrogen

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deficiency will cause plants grow stunted, leaves turn yellow, and reducing growth activity (Gardner et al., 1991). The following is a nitrogen formation reaction According to Novizan (2005):

Protein Organic Ingredients Aminization Reaction Amino acid Amino acid Amino acid Amino acid Amino acid Ammonification Reaction - Ammonia (NH<sub>3</sub> &) & Ammonium NH<sub>4</sub>\*

Amonia Nitrification -Reaction of nitrification by Nitroccus bacteria Nitrosomonas- and Nit

The nitrogen contained in degradation product with indigosol blue effluent does not reach quality standards to Minister of Agriculture Regulation No.70/Permentan/SR.140/10/2011, the minimum quality standard for nitrogen content of liquid organic fertilizer is 3-6%. Based on research Wulandari (2015), that factor causing the decrease in Nitrogen (N) content is due to limited oxygen (O<sub>2</sub>) causing \_-ammonia (NH<sub>3</sub>) cannot be converted into nitrate (NO<sub>3</sub>) and nitrogen (N), causes it to evaporate into the air in the form of ammonia (NH<sub>3</sub>).

Phosphorus is a macro nutrient that is needed by plants, there are no other elements that can replace its function in plants. Plants must get or contain enough P for normal growth (Winarso, 2005). Phosphorus in plants has an important function, namely in the process of cell enlargement photosynthesis, respiration, division, respiration, and photosynthesis cell enlargement. Phosphorus can improve the quality of fruits, vegetables, grains, and important in the formation of seeds. Phosphorus can help in the process of accelerating root development and germination, can increase water use efficiency, increase resistance to disease which ultimately improves crop quality. If there a lack of phosphorus, plant will show symptoms of growth such as slow growth, stunted growth, and stunted root development. Symptoms that will arise on leaves are diverse, some plants show an abnormal glossy dark green color, fruit maturation is inhibited. development of fruit shape, color, and seeds develop abnormally (Kusuma, 2014).

The Phosphorus contained in degradation product with indigosol blue effluent does not reach quality standards to Minister of Agriculture Regulation No.70/Permentan/SR.140/10/2011, the minimum quality standard for phosphorus content of liquid organic fertilizer\_LOF is 3-6%. According to Stoffella & Khan (2001) stated, that\_In the substrate, there is a relationship between phosphorus and nitrogen content. the greater the nitrogen contained, the greater the number of microorganisms that break down phosphorus. Phosphorus on the substrate will be used by certain microorganisms for cell growth and development.phosphorus content is related to nitrogen content in the substrate. The greater nitrogen contained, more microorganisms that remodel phosphorus. The phosphorus content in the substrate will be used by some microorganisms to build their cells.

The nitrogen content and phosphorus content were<u>content</u> was obtained from the structure of Indigosol Blue which contained anthraquinones and had -NH and C=C molecular bonds. This dye waste has a strong chemical bond structure and is classified as nonbiodegradable waste (Herfiani et al., 2017). Indigosol Blue degradation products were identified by Dewi et al. (2021) as simple compounds called aliphatic compounds containing C=C, C-O, and -OH bond.

The results research use of degradation product from tofu industrial waste gave results that affected changes in weight of cucumber plants. In treatment of changes in fruit weight of cucumber plants with a-dose of liquid organic fertilizer-LOF dose from tofu waste of 25 ml showed the best results. That application of liquid organic fertilizer-LOF from tofu industrial waste gives results that have less effect on plant length, number of leaves, leaf area, fruit length, number of fruits, and fruit circumference on cucumber plants (Simajuntak, 2021).

In this research use of degradation product derived from degradation process of indigosol blue batik effluent. The results showed a total nitrogen content of 0.11% and phosphorus 0.01%. According to research Indrayani et al. (2019), states that indigo pasta batik natural dye waste for the manufacture of liquid organic fertilizer <u>LOF</u> with bioactivator EM-4 has nitrogen elements with an anaerobic process of 0.11% and aerobic process is 8.47% and aerobic is 8.49%.

The results of the degradation of batik effluent obtained potassium and carbon content. The carbon (C-organic) content obtained from the structure of Indigosol Blue. This accordance to Herfiani et al. (2017), Indigosol Blue containing Anthraquinone (C<sub>14</sub>H<sub>8</sub>O<sub>2</sub>) and has molecular bonds of -NH and C=C. After degradation by *Aspergillus* sp. 3, the Indigosol Blue changed to a simple-simpler structure which is an aliphatic compound contained bonds of C=C, C-O, C=C, and -OH bonds-that indentified by Dewi et al. (2021).

Potassium (K<sub>2</sub>O) content result was fewer than carbon content. It is accordance to Herfiani et al. (2017), Indigosol Blue not containdoes not contain K<sub>2</sub>Opotassium. The potassium found because the potatoes contained in PBD media. Potatoes has high K<sub>2</sub>Opotassium. The value of the potassium content of potatoes according to UMHS Patient Food and Nutrition Services (2016) is 510 milligrams in a cup medium.

Previous research showshows the degradation of indigosol blue effluent result did not cause toxicity in plants. Degradation results can be applied to plants. This is accordance with research by Dewi et al. (2018) showed plumule and radicle length of *Zea mays* and *Vigna radiata* grown on the decolorized effluent being longer than in the untreated effluent. The percentage of *Z*, mays and V, radiata seed germination on decolorized effluent was higher than on untreated effluent. It takes a further assay on the content of the degradation results:

The result of assay potassium (K<sub>2</sub>O) K2O and Corganic content in degradation indigosol effluent on by spectrophotometry and AAS methods were 0.01%. The Carbon (C-organic) by spectrophotometric methods obtained 0.24%. The results were not reach the minimum standard of Minister of Agriculture Regulation in the previously mentioned Agriculture Minister RegulationNo.261/KPTS/SR.310/M/4/2019, namely the minimum quality requirement for liquid organic fertilizer LOF is 3-6% of K<sub>2</sub>O and minimum 6% C-organic. Beside the degradation of indigosol blue effluent, other batik dyes effluent also havehas the potential as liquid organic fertilizerLOF. Industrial waste for making batik natural dyes that are no longer used in the process, in the form of indigo paste (Strobilanthes cusia), can be reused as raw materials for other industries, such as for LOF with the composting process. The utilization of waste from the industrial process of making natural dyes indigo paste batik (Strobilanthes cusia) in the indigo paste industry, for the manufacture of organic liquid fertilizer through composting by adding EM 4It was studied by Indrayani et al. (2019). The potassium content value were 8.49 were 8.49 ppm, and 2137 mg per liter for aerobic processes. Nitrogen (N) content anaerobically and aerobically has almost the same value, namely 0.11 percent and 0.16 percent, while the value of Phosphorus (P2O2) anaerobically and aerobically also has almost the same value, which is 4.87 ppm and 4.86 ppm, and Potassium (K2O) in liquid organic fertilizer (compost) 2137.53 mg per kg for the aerobic process and 556.30 mg per kg for the anaerobic process.

The degradation result may be used as liquid organic fertilizer\_LOF\_if added with have high nutritional content by others fermentation. Nutrients that can be added obtained from fermenting banana peels or fermenting cow urine. Research of Widyabudiningsih et al. (2021), fermented content of banana peel waste added EM4 has K<sub>2</sub>O 5.80% and C-organic 13.40%. The content of potassium and carbon were reach the minimum standard of liquid organic fertilizerLOF production. While fermented cow urine of Desa Dhusun Thekelan added EM4 in 6 days studied by Kun (2011) contains 7.80% organic C-content and 1.18% potassium. The content of potassium werewas low but the earbon C-content was reachreaching the minimum standard of liquid organic fertilizerLOF production.

Using wastewater as fertilizer by Fraunhofer-Gesellschaft (2012) develop environmentally friendly, chemical-free processes that allow the recovered salt to be converted directly into organic nutrients for food crops. wastewater, sewage sludge, and liquid manure are sources of fertilizer that can be used for food production. The process that occurs is electrochemical which does the precipitation of struvite (magnesium ammonium phosphate), through electrolysis of a solution containing phosphorus and nitrogen. This process does not require the addition of a synthetic base or a salt. In this electrolysis event, water molecules break down into negatively charged hydroxyl ions at the cathode. oxidation then occurs at the anode to form struvite, as magnesium ions migrate through the water and react with the phosphate and ammonium molecules in solution. The struvite is then precipitated from the process water in the form of small crystals that can be directly used as fertilizer, without further processing. This process benefits because it saves energy and is free from chemical processes.) developed a chemical free. eco-friendly process that enables the recovered salts to be converted directly into organic food for crop plants. Sewage sludge, wastewater and liquid manure are valuable sources of fertilizer for food production. The process is electrochemical that precipitates magnesiumammonium phosphate, known as struvite, by means of electrolysis from a solution containing nitrogen and phosphorus. This method does not require the addition of synthetic salts or bases. The electrolytic process splits the water molecules into negatively charged hydroxyl ions at the cathode. At the anode an oxidation takes place which the magnesium ions migrate through the water and react with the phosphate and ammonium molecules in the solution to form struvite. Struvite is precipitated from the process water in the form of tiny erystals that can be used directly as fertilizer, without any further processing. This method savesaves energy and free chemical process. The growing plants were up to four times higher with struvite than with commercially available mineral fertilizers.

Nevertheless, it has the potential as LOF, therefore the test was carried out on plants. The plant used is spinach. Spinach was watered regularly with degradation products, fermented degradation products and then observed the frequency of wilting and death (Table 2).

Fertilizer Manufacturing Effluent Guidelines and Standards initiated by EPA in 1974 and 1975. The regulation amended between 1975 and 1987. The Formatted: Highlight

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regulation covers direct and indirect dischargers. The Effluent Guidelines and Standards are incorporated into NPDES permits for direct dischargers and permits or other control mechanisms for indirect dischargers. The industry manufactures five basic fertilizer chemicals, there are phosphate, ammonia, urea, ammonium nitrate and nitric acid. Phosphate fertilizer manufacturing comprises two principal units: production of sulfurie acid, derived from elemental sulfursulphur, and wet process phosphoric acid, derived from phosphate rock. Ammonia is manufactured using atmospheric nitrogen and hydrogen derived from natural gas or petroleum refinery byproductsby products. Ammonia is sold as a straight fertilizer, and is used to manufacture urea. ammonium nitrate and nitric acid products. Fertilizer manufacturing is included within the following NAICS groups are Nitrogenous Fertilizer Manufacturing (325311), Phosphatic Fertilizer Manufacturing (325312), and Fertilizer (Mixing Only) Manufacturing (325314)

Table 2. The Results of Treatment in Plants

Treatment	Frequency of wilted plants (%)	Frequency of dead plants (%)
Water	40%	0%
Degradation product	0%	0%
Degradation Fermentation	0%	0%
Effluent	0%	40%

Nevertheless, Iti<u>i</u> have<u>has</u> the potential as liquid organic fertilizer, therefore the test was carried out on plants. The plant used is spinach. Spinach was watered regularly with degradation products, fermented degradation products and then observed the frequency of wilting and death (TabelTable 2).

The results of fermentation of degradation products caused 0% wilting frequency and plant death as well as degradation products when compared with treatments using water (40% plant wilt frequency) and effluent (40% plant mortality frequency). This shows that the degradation product of batik liquid waste has the potential as a liquid organic fertilizerLOF but must be redeveloped with certain treatments so as toto achieve the required composition as fertilizer.

## 4. CONCLUSION

The product degradation of indigosol blue batik effluent had a total nitrogen content of 0.11%, K<sub>2</sub>, K<sub>2</sub>O of 0.01%, organic carbon of 0.24%, C/N ratio of 10.13%, while for phosphorus content of 0.01%. Fermentation of degradation products causes 0% frequency of wilted and dead plants, better when compared to water (40% frequency of wilted plants) and

efluentand effluent (40% frequency of wilted and dead plants).

## **AUTHORS' CONTRIBUTIONS**

RSD came up of the presented idea and supervised the findings of this work. MK-verified the analytical methods, helped supervise the project. All authors discussed the results, wrote the manuscript<u>manuscript</u>, and contributed to the final manuscript<u>MK</u> assists in project work and makes analytical methods. All authors contributed in processing the research data, writing the manuscript, and playing a role in working on the final <u>manuscript</u>.

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## Study of N, P, K, and C on Degradation of Indigosol Batik Dye Effluent by *Aspergillus* sp. GPN

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#### ABSTRACT

Indigosol dye batik effluent is toxic since it excesses contains contain decreasing agent sodium hydrosulfite that is oxidized to alkali and alkaline earth metal sulfate (SO42-), sulfite (SO32-), and thiosulfate (S2O32-) that very corrosive. Therefore, the role of environmental biotechnology in the technology for restoring environmental functions is needed efforts to treat batik effluent have been carried out biologically by using microorganism agents. It can be used Aspergillus sp. that can reduce industrial dyes. From the previous research, the product of Indigosol Blue batik effluent degradation that produced by Aspergillus sp. GPN did not cause any toxicity in plants even better grew than giving watering with irrigation water. Content in the results of degradation can affect the plants, so the element analysis needs to be performed. It's must include the substance nitrogen (N) in the form of organic compounds that easily absorbed by plants, do not leave residual organic acids in the soil, has high carbon (C) content, Phosphorus (P), and Potassium (K). This study aimed to analyze the content of N, P, K, organic carbon, and ratio of organic carbon to total nitrogen (C: N) in degradation product of Indigosol Blue dye batik effluent produced by Aspergillus sp. GPN. The methods used were spectrophotometry for organic carbon and P2O5, Kjeldahl for total N, spectrophotometry, and AAS for K2O. The result of the assay was the product degradation of indigosol indigo sol blue batik effluent had a total nitrogen content of 0.11%, K<sub>2</sub>O of 0.01%, organic carbon of 0.24%, C/N ratio of 10.13%, while for phosphorus content of 0.01%. Fermentation of degradation products causes 0% frequency of wilted and dead plants, better when compared to water (40% frequency of wilted plants) and effluent (40% frequency of wilted and dead plants).

Keywords: Aspergillus sp., batik effluent, elements.

## 1. INTRODUCTION

Most of the textile wastewater, especially batik, contains dye residues that have the potential to pollute the environment. It is extremely dangerous because it carries chemical substances which are dangerous to the environment. The effluent has a detrimental effect on the content of organic matter, suspended solids, oil, or fat, and the content of dangerous heavy metals such as Zn, Cd, Cu, Cr, and Pb\_11\_(Nurdalia, 2006). Generally, batik industrial effluent is discharged directly into water bodies or rivers without being treated first [2](Watini, 2009). One type of synthetic batik dye is Indigosol Blue that one of the synthetic dyes of Anthraquinone, and has molecular bonds of -NH and C=C. This synthetic dye is

not easily damaged by chemical or photolytic treatment. This dye waste has a strong chemical bond structure and is classified as non-biodegradable waste [3] (Herfiani et al., 2017). Indigosol wastewater is toxic because it carries extra reducing agent sodium hydrosulfite that is oxidized to alkali and alkaline earth metal sulfate  $(SO_4^2)$ , sulfite  $(SO_3^{2-})$ , and thiosulfate  $(S_2O_3^{2-})$  hat very corrosive. Thus, the formed sulfate deposits can create toxic hydrogen sulfide (HS) ions [4](Blackburn et al., 2009).

Efforts to treat batik effluent have been carried out biologically by using microorganism agents. The microorganism agent that can be used is Aspergillus sp. that can reduce industrial waste dyes. A study by Dewi et al. (2018) showed Aspergillus sp. 3 strain GPN is the Formatted: Font: 20 pt

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superior isolate to decolorize the batik effluent [5]<u>Dewi</u> <u>et-al. (2018)</u>. Aspergillus species 3 nearly completely (99.9%) decolorized the effluent within the liquid medium after a three-day incubation. According to Dewi & Khotimah (2019), Aspergillus sp. 3 can remove effluent as a single carbon and nitrogen source dye [6].

The previous study finds that effluent batik dye degradation by *Aspergillus* sp. 3 did now no longer motive any toxicity in plants. A study by Dewi et al. (2018) confirmed \_pPlumule and radicle length of *Zea mays* and *Vigna radiata* has grown at the decolorized effluent being longer than withinside the untreated effluent  $\frac{A \text{ study by } [5]\text{Dewi et al. (2018)}}{\text{ orn and mung bean seed germination on decolorized effluent turned better than on untreated effluent. Even, it's better growth than giving watering with irrigation water. Content in the results of degradation can affect the plants, so the element analysis needs to be performed.$ 

Elements that can fertilize plants are substance nitrogen (N) in the form of organic compounds that are easily absorbed by plants, do not leave residual organic acids in the soil, has high carbon (C) content, and have Phosphorus (P) and Potassium (K).

Substance N in the form of organic compounds that are easily absorbed by plants does not leave residual organic acids in the soil and has high organic C content such as charcoal hydrates. Nutrients required of the plant are not sufficiently available in the soil because the nutrients contained in the soil are relatively little, the number of nutrients cannot suffice certain plants, so it is necessary to add nutrients. The role of potassium in plants is to form protein and carbohydrates, harden straw and the bottom of the wood, increase plant retention against disease and improve seed or fruit quality [7](Effendi, 2004). According to Afandi et al. (2015) eCarbon is a food supply for soil microorganisms, so the presence of organic C withinside the soil will stimulate the activity of microorganisms [8] Afandi et al. (2015). Wahyudi (2009) said that tThe increase in C-Organic caused by carbon (C) with the application of elements on the soil will increase the organic substance contained in the soil [9].

This study aimed to analyze the content of N, P, K, organic carbon, and ratio of organic carbon to total nitrogen (C: N) in degradation product of Indigosol Blue dye batik effluent produced by *Aspergillus* sp. GPN.

## 2. METHODOLOGY

#### 2.1. Reculture Aspergillus sp. GPN

Materials and tools are prepared, alcohol sprayed on palms and tables. The Bunsen fire is light. The wrapping paper of the petri dish is removed. The mouth of the petri dish is heated. A hole in the PDA was made using a straw. A piece of PDA took using a loop needle. Inoculated on new PDA media, placed in the middle of the PDA. The petri dish is covered, heated then wrapped in plastic wrap. Then incubated at room temperature for 3-7 days.

## 2.2. Preparation of Aspergillus sp. GPN inoculum

Materials and tools are prepared, alcohol sprayed onpalms and tables. The Bunsen fire is light. The wrapping paper on the Erlenmeyer flask is removed. The mouth of the Erlenmeyer flask is heated. A hole in the PDA was made using a straw. A piece of PDA took using a loop needle. Inoculated on PDB media. The Erlenmeyer flask is covered, heated then wrapped in plastic wrap. Then incubated on shaking incubator at room temperature for 3-7 days.

## 2.3. Efluent Degradation

Indigosol was added on *Aspergillus* sp. 3 preparationthen incubated on a shaking incubator at room temperature. Observed in each time interval. The degradation result is filtered then used for Macro Nutrient Assay.

## 2.4. Macro Nutrient Assay

#### 2.4.1. Assay of K Nutrient

The testing used a Flame photometer. The 100 mL sample solution was added in a 100 mL volumetric flask. Diluted to the limit. The standard series of samples were made of 4 mL, 6 mL, and 8 mL. Each sample was diluted into a 100 mL volumetric flask. The absorbance was measured using a flame photometer.

#### 2.4.2. Assay of C Nutrient

The C test used the walker and black methods. A sample of 100 mL was added with 5 mL of  $K_2Cr_2O_7 2 N$  solution and then shake. The  $H_2SO_4$  pa. 98% of 7 mL is added then shaken. The solution is left for 30 minutes then shakenshake. Standard solution C of 5 ml is pipetted into a 100 ml volume measuring flask.  $H_2SO_4$  of 5 mL and 7 ml of  $K_2Cr_2O_7 2 N$  solution were added and then shaken. The solution is left for 30 minutes then shakenshake. Blanko as standard 0 ppm C were prepared by diluting the sample with ion-free water. After cooling, the volume is adjusted to the mark of 100 ml. Then shaken back until homogeneous and overnight left. Then measured with a spectrophotometer at a wavelength of 651 nm.

#### 2.4.3. Assay of P Nutrient

Analyzing the P content by inserting the waste sample in a cuvette before measuring it in a UV-Vis spectrophotometer. Blanks using distilled water were

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Formatted: Line spacing: At least 12,05 pt, Pattern: Clear (Background 1) Formatted: Pattern: Clear (Background 1) included in the cuvette. Then the absorbance value was measured using a UV-Vis spectrophotometer with a wavelength of 650-750 mn.

## 2.4.4. Assay of N Nutrient

The sample for the analysis of the N content used the Kjeldahl micro method. There are three stages in the Kieldahl micro method, namely digestion, distillation, and titration which function to obtain the values of Norganic, N-NH<sub>4</sub>, and N-NO<sub>3</sub>. The N-organic test begins to make titrant A as a sample, and A1 as a blank. The titran A and A1 were prepared by adding 0.25 g of selenium mixture to which 3 mL of H<sub>2</sub>SO<sub>4</sub> pa were added in a Kieldahl flask, then shaken until evenly distributed. and then left for 2 hours. Gradual digestion with a temperature of 150°C to a maximum temperature of 350°C for three hours and clear liquid. Cool then dilute with a touch of distilled water. The solution is transferred to a distillation boiling flask and then added a little boiling stone and distilled water until it reaches the volume of the flask. Then 10 mL of 40% NaOH were added. The destination container includes 10 mL of 1% boric acid in Erlenmeyer and 3 drops of Conway indicator liquid are added until the Erlenmeyer reaches 75 mL. Continue the distillate titration for A1 with 0.05  $N\ H_2SO_4$  to the endpoint when the color changes from green to pink.

The N-NH<sub>4</sub> test was initiated using titrant B as the sample and B1 as the blank. Sample 1 g in a boiling distillation flask plus a little boiling rock, 0.5 mL of liquid paraffin, and 10 mL of distilled water for B while 100 mL for B1. Then add 10 mL of 1% boric acid, 3 drops of Conway indicator, and 10 mL of 40% NaOH and then distill it until it reaches 75 mL. The distillate was titrated with 0.05 N H<sub>2</sub>SO<sub>4</sub> until the color of the solution changed from green to pink.

Testing for N-NO<sub>3</sub> begins with titrant C as the sample and C1 as a blank with the same procedure. The remainder of the N-NH<sub>4</sub> determination of the sample could cool then be added with distilled water to the original volume. The distillate is entered by adding 10 mL of 1% boric acid in 100 mL Erlenmeyer plus 3 drops of Conway indicator. Then distill it by adding 2 g of devarda alloy. Distillation is started indefinitely so that the foam does not overflow after the foam is almost exhausted starting from low temperature after boiling the temperature is increased to a normal. Distillation to 75 mL of liquid. Then titrated with 0.05 N H<sub>2</sub>SO<sub>4</sub> until the color of the solution changes from green to pink. The results of N-organic, N-NH<sub>4</sub>, and N-NO<sub>3</sub>. Were calculated using the formula then totaled.

### 2.5. Plant Assay

The degradation products obtained were then tested on plants. This experiment was compared with the degradation product which was mixed with a small amount of goat urine and then fermented using EM 4 (Effective Microorganism-4). Then the spinach plants were watered and compared with irrigation water and batik waste. Observations were made on the frequency of wilting and the frequency of death.

## 3. RESULTS

In this research assay of N, P, K, and C element content derived from the degradation process of Indigosol Blue batik effluent was carried out. The test results for the assay of nutrients were presented in Table 1.

Table 1 showed results of total nitrogen content in degradation product which has a value of 0.11%, while for phosphorus it has a value of 0.01%. because it is possible to fertilize plants, it is compared with the content in liquid organic fertilizers (LOF). According to Minister of Agriculture Regulation No. 70/Permentan/SR.140/10/2011\_[10], the minimum quality standard for nitrogen and phosphorus content of LOF is 3-6%. The results degradation product of this research was not reached quality standards for nitrogen and phosphorus content has a value of <1%.

Table 1. The Results for Assay nutrient from Degradation Indigosol Blue Batik Effluent

No	Parameter	Method	%
1	N-Total	Kjeldahl	0.11
2	P <sub>2</sub> O <sub>2</sub>	Spekrofotometri	0.01
3	C-organic	Spectrophotometric	0,24
4	K <sub>2</sub> O	Spectrophotometric, AAS	0.,01
5	C/N Ratio	-	10.13

Minister of Agriculture Regulation No.70/Permentan/SR.140/[10/2011] contains standards for LOF, but types of samples obtained in this research cannot be categorized as LOF. The nitrogen content obtained is 0.11% and Phosphorus is 0.01%. According to its regulation, the minimum quality standard for nitrogen and phosphorus content of LOF is 3-6%. Based on the research of Ni'am et al. (2015), stated that tThe fermented cassava wastewater with the addition of cow urine, and foot of cow washing water contained 5.11% Nitrogen and 3.08% Phosphorus [11].

Plants need one of the macronutrients Nitrogen for protein formation and vegetative growth (Sarief, 1986). Nitrogen deficiency will cause plants to grow stunted, leaves turn yellow, and reduce growth activity [12](Gardner et al., 1991). The following is a nitrogen<sup>4</sup> formation reaction [13] According to Novizan (2005):

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Protein Organic Ingredients Aminization Reaction Admino acid Amino acid Ammoniafication Reaction Ammonia (NH<sub>3</sub>) & Ammonium NH<sub>4</sub><sup>+</sup> Ammonia Reaction of nitrification by Nitrococcus bacteria and Nitrosomonas Nitrate (NO<sub>3</sub>)

The nitrogen contained in degradation product with indigosol-indigo sol blue effluent does not reach quality standards to Minister of Agriculture Regulation [10], the minimum quality standard for nitrogen is 3-6%. Based on research by Wulandari (2015), tThat factor causing the decrease in N content is due to limited  $O_2$  causing NH<sub>3</sub> causing to evaporate into the air in the form of NH<sub>3</sub>[14].

Phosphorus is a macronutrient that is needed by plants, there are no other elements that can replace its function in plants. Plants must get or contain enough P for normal growth [15] (Winarso, 2005). Phosphorus in plants has an important function, namely in the process cell enlargement, division, respiration, and of photosynthesis. Phosphorus can improve the quality of fruits, vegetables, grains, and is important in the formation of seeds. Phosphorus can help in the process of accelerating root development and germination, can increase water use efficiency, increase resistance to disease which ultimately improves crop quality. If there is a lack of phosphorus, the plant will show symptoms of growth such as slow growth, stunted growth, and stunted root development. Symptoms that will arise on leaves are diverse, some plants show an abnormal glossy dark green color, fruit maturation is inhibited, development of fruit shape, color, and seeds develop abnormally [16](Kusuma, 2014).

The Phosphorus contained in <u>the</u> degradation product with <u>indigosol\_indigo sol</u> blue effluent does not reach quality standards to Minister of Agriculture Regulation No.70/Permentan/SR.140/[10/2011], the minimum quality standard for phosphorus content of LOF is 3-6%. Stoffella & Khan (2001) stated, In the substrate, there is a relationship between phosphorus and nitrogen content. the greater the nitrogen contained, the greater the number of microorganisms that break down phosphorus\_[17]. Phosphorus on the substrate will be used by certain microorganisms for cell growth and development.

The nitrogen content and phosphorus content were obtained from the structure of Indigosol Blue which contained anthraquinones and had ——NH and C==C molecular bonds. This dye waste has a strong chemical bond structure and is classified as non-biodegradable waste [3] (Herfiani et al., 2017). Indigosol Blue degradation products were identified by Dewi et al. (2021) as simple compounds called aliphatic compounds containing C==C, C-O, and -OH bonds [18].-

The results research use of degradation product from tofu industrial waste gave results that affected changes in weight of cucumber plants. The treatment of changes in fruit weight of cucumber plants with a LOF dose from tofu waste of 25 ml showed the best results. That application of LOF from tofu industrial waste gives results that have less effect on plant length, number of leaves, leaf area, fruit length, number of fruits, and fruit circumference on cucumber plants [19].(Simajuntak, 2021).

In this research use of degradation products is derived from the degradation process of indigosol indigo sol blue batik effluent. The results showed a total nitrogen content of 0.11% and phosphorus of 0.01%. According to research Indrayani et al. (2019), states (That indigo pasta batik natural dye waste for the manufacture of LOF with bio activator EM-4 has nitrogen elements with an anaerobic process of 0.11% and an\_aerobic process of 0.16%. Phosphorus element with the anaerobic process is 8.47% and aerobic is 8.49% [20].Indrayani et al. (2019),.

The results of the degradation of batik effluent obtained potassium and carbon content. The carbon (C-organic) content was obtained from the structure of Indigosol Blue. This accordance to Herfiani et al. (2017), Indigosol Blue contains Anthraquinone ( $C_{14}H_8O_2$ ) and has molecular bonds of -NH and C=C. After degradation by *Aspergillus* sp. 3, the Indigosol Blue changed to a simpler structure which is an aliphatic compound <u>This accordance to [3]Herfiani et al. (2017)</u>. The aliphatic compound contained bonds of C-O, C=C, and -OH that were identified by [18].Dewi et al. (2021).

The potassium (K<sub>2</sub>O) content result was fewer than carbon content. It is accordance with Herfiani et al. (2017), Indigosol Blue does not contain K<sub>2</sub>O\_[3]. The potassium was found because the potatoes were contained in PBD media. Potatoes have high K<sub>2</sub>O. The value of the potassium content of potatoes according to UMHS Patient Food and Nutrition Services (2016) is 510 milligrams in a cup medium [21].

The result of assay potassium (K2O) and C-organic content in degradation indigosol indigo sol effluent on Table 3.1. showed assay the content of K<sub>2</sub>O by spectrophotometry and AAS methods was 0.01%. The Carbon (C-organic) by spectrophotometric methods obtained 0.24%. The results were not reaching the minimum standard of the previously mentioned Agriculture Minister Regulation [10], namely, the minimum quality requirement for LOF is 3-6% of K2O and a minimum of 6% C-organic. Besides the degradation of indigosol-indigo sol blue effluent, other batik dyes effluent also has the potential as LOF. Industrial waste for making batik natural dyes that are no longer used in the process, in the form of indigo paste (Strobilanthes eusiacuisine), can be reused as raw materials for other industries, such as for LOF with the composting process. It was studied by [20] Indrayani et al. (2019). Nitrogen (N) content anaerobically and aerobically has almost the same value, namely 0.11

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percent and 0.16 percent, while the value of Phosphorus ( $P_2O_2$ ) anaerobically and aerobically also has almost the same value, which is 4.87 ppm and 4.86 ppm, and Potassium ( $K_2O$ ) in liquid organic fertilizer (compost) 2137.53 mg per kg for the aerobic process and 556.30 mg per kg for the anaerobic process.

The degradation result may be used as LOF if added with have high nutritional content by others fermentation. Nutrients that can be added are obtained from fermenting banana peels or fermenting cow urine. Research to Widyabudiningsih et al. (2021), fFermented content of banana peel waste added EM4 has K<sub>2</sub>O 5.80% and Corganic 13.40% [22]Widyabudiningsih et al. (2021), The content of potassium and carbon reached the minimum standard of LOF production. While fermented cow urine of Desa Dhusun Thekelan added EM4 in 6 days-studied by Kun (2011) \_contains 7.80% organic C-content and 1.18% potassium [23]. The content of potassium was low, but the C-content was reaching the minimum standard of LOF production.

Using wastewater as fertilizer by Fraunhofer-Gesellschaft (2012) develops environmentally friendly, chemical-free processes that allow the recovered salt to be converted directly into organic nutrients for food crops [24]. Wwastewater, sewage sludge, and liquid manure are sources of fertilizer that can be used for food production. The process that occurs is electrochemical which does the precipitation of struvite (magnesium ammonium phosphate), through electrolysis of a solution containing phosphorus and nitrogen. This process does not require the addition of a synthetic base or salt. In this electrolysis event, water molecules break down into negatively charged hydroxyl ions at the cathode. oxidation then occurs at the anode to form struvite, as magnesium ions migrate through the water and react with the phosphate and ammonium molecules in solution. The struvite is then precipitated from the process water in the form of small crystals that can be directly used as fertilizer, without further processing. This process benefits because it saves energy and is free from chemical processes.

Nevertheless, it has the potential as LOF, therefore the test was carried out on plants. The plant used is spinach. Spinach was watered regularly with degradation products, fermented degradation products, and then observed the frequency of wilting and death (Table 2).

Table 2. The Results of Treatment in Plants

Treatment	Frequency of	Frequency of	L
	wilted plants (%)	dead plants (%)	
Water	40%	0%	
Degradation	0%	0%	
product			
Degradation	0%	0%	
Fermentation			
Effluent	0%	40%	

The results of fermentation of degradation products caused 0% wilting frequency and plant death as well as degradation products when compared with treatments using water (40% plant wilt frequency) and effluent (40% plant mortality frequency). This shows that the degradation product of batik liquid waste has the potential as a LOF but must be redeveloped with certain treatments to achieve the required composition as fertilizer.

The occurrence of death in the treatment of giving waste could be due to a lack of macronutrients such as N. Nitrogen (N) as the main nutrient for plants is generally indispensable for the formation or growth of vegetative parts of plants, such as leaves and roots. Deficiency can result in severely impaired growth rate, causes the leaves to turn yellow or experience chlorosis, and a strong deficiency causes browning and death [25].

#### 4. CONCLUSION

The product degradation of indigosol-indigo sol blue batik effluent had a total nitrogen content of 0.11%, K<sub>2</sub>O of 0.01%, organic carbon of 0.24%, C/N ratio of 10.13%, while for phosphorus content of 0.01%. Fermentation of degradation products causes 0% frequency of wilted and dead—plants, better—when—compared—to—water—(40%– frequency of wilted plants) and effluent (40% frequency of wilted and dead plants).

## **AUTHORS' CONTRIBUTIONS**

RSD came up with the presented idea and supervised the findings of this work. MK assists in project work and makes analytical methods. All authors contributed to processing the research data, writing the manuscript, and playing a role in working on the final manuscript.

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