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Conference Series

**The 7th International Conference on
Mathematics, Science, and Education 2020
(ICMSE 2020)**

1918

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6 October 2020
Semarang, Indonesia

Editors:
Dyah Rini Indriyanti et al.

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Preface

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PREFACE

The 7th International Conference on Mathematics, Science and Education 2020 (ICMSE2020) is the international conference organized by Faculty of Mathematics and Natural Sciences Universitas Negeri Semarang (Indonesia) under the theme of *Accelerating Innovative Research from Laboratory to Industry*. This conference aims to discuss the innovation of the laboratory research as well as to downstream those innovative products.

Since the world has faced the global Covid-19 pandemic, it would be too risky to hold international conference through direct plenary meeting. Therefore, the conference that was originally conducted in Semarang, Indonesia, on 6 October 2020 but then it was converted into virtual conference to protect participants from Corona Virus transmission. In addition, travel restrictions also prevented participants from attending the conference onsite, then the virtual conference setting was decided as the best option to keep the conference ran on schedule.

The keynote and invited speakers from five countries consisting of Asst. Prof. Dr. Chalongrat Daengngam (Thailand), Assoc. Prof. Stephane Bressan (Singapore), Asst. Prof. Aditya P. Adiredja, Ph.D (USA), Assoc. Prof. Ch.M Dr. Roswanira Abdul Wahab (Malaysia), and Prof. Dr. Ir. Amin Retnoningsih, M.Si (Indonesia) delivered their talk through video conference, and the technology support to deliver the conference was "Zoom Video Conferencing" software. All keynote speakers shared the material from their countries as well as the participants.

There were three sessions which involved opening, invited speaker presentation and authors' parallel presentation. Opening session consisting of keynote speech and opening remark lasted 20 minutes, followed by each invited speaker presentation for 20 minutes and each author presentation took 10 minutes. The breakout rooms were provided to facilitate authors in presenting their research result. Every parallel session was guided by one moderator to lead the discussion. Participants still could involve themselves in the intellectual discussion in invited speakers and parallel sessions although the conference was conducted virtually as they could write the question through chat box or directly deliver their question.

Finally, the committees via review stage selected the papers-presented in this forum to be published in Journal of Physics: Conference Series (Institute of Physics Publisher) indexed in some databases, including the Conference citation index, Scopus, Chemical Abstracts Service, and Astrophysics Data System. We hope that this program will expand the mutual understanding and respect in stimulating research in Mathematics, Science, and Education; share research interest and information, and create a form of collaboration and build a trust relationship. We are delighted to be able to show the world what recent developments in the field of Mathematics, Natural Science, and Science Education through this fruitful program.

As conclusion, we would like to acknowledge to all of supporting teams, especially the organizing committee members and reviewers, for their great contribution in conference organization. We hope that this program will further encourage research publication in Mathematics, Natural Science, and Science Education and build trust relationship. We feel honoured and privileged to serve the best recent developments in the field of Mathematics, Natural Science, and Science Education to you through this exciting program.

Chairperson,

Prof. Dr. Dyah Rini Indriyanti, M.P.



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Dr. Aditya P. Adiredja






An anti-deficit perspective on the mathematical thinking of marginalized students

From counter-narratives to creative thinking

Dr. Aditya P. Adiredja
The University of Arizona, USA
ICMSE 2020, October 6th

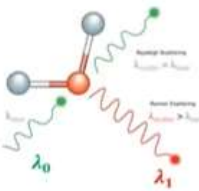


Asst. Prof. Dr. Chalongrat Daengnam

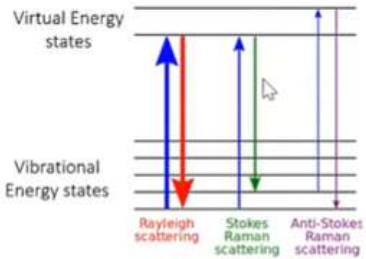


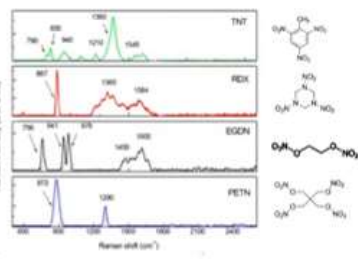
Conventional Raman Process

Target molecules



Virtual Energy states






Raman shift

$$\Delta\omega(\text{cm}^{-1}) = \left(\frac{1}{\lambda_0(\text{nm})} - \frac{1}{\lambda_1(\text{nm})} \right) \times \frac{(10^7 \text{ nm})}{(\text{cm})}$$

Raman shift is a molecular fingerprint

Raman efficiency is around 10^{-10} to 10^{-8}



Assoc. Prof. *Stephane Bressan*

Scalability Transfer Learning

Can we devise strategies to reuse a trained system of a smaller size to initialize the training of a system of a larger size? Can we transfer the weights?

Transfer learning for scalability of neural-network quantum states. Remy Zen, L. My, R. Tan, F. Hebert, M. Gattobigio, C. Miniatura, D. Poletti, S. Bressan, Phys. Rev. E, vol. 101, p. 053 301, 5 May 2020

NUS National University of Singapore

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The slide features a diagram of a neural network with two layers of nodes. The first layer has nodes labeled $h_1, h_2, h_3, \dots, h_m$ and $v_1, v_2, v_3, \dots, v_n$. The second layer has nodes labeled $h_{m'}$ and $v_{n'}$. Connections between the layers are shown with lines, and a weight $W^{(m',n')}$ is indicated between the two layers.

Assoc. Prof. *Ch.M Dr. Roswanira Abdul Wahab*

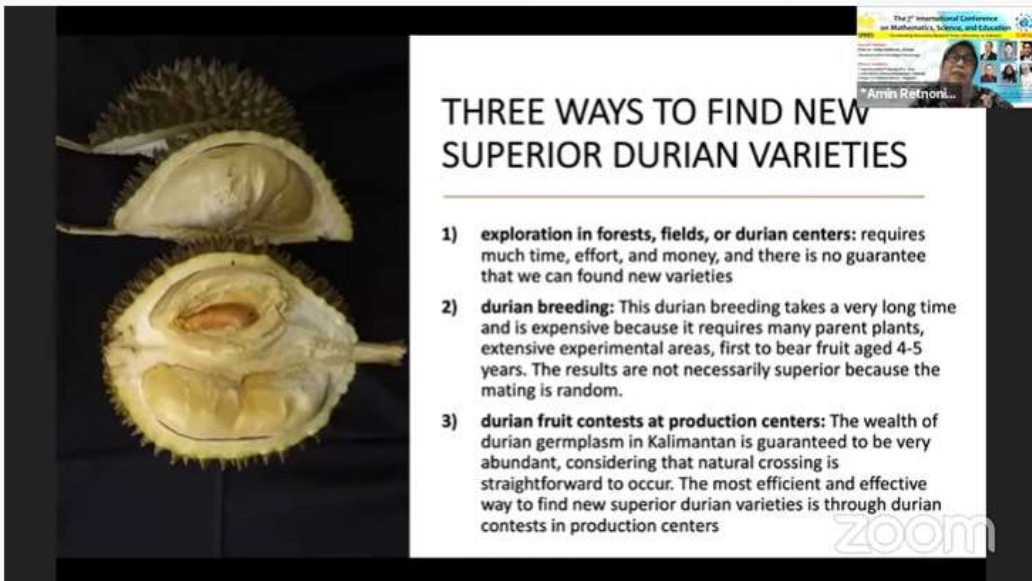
Natural sugar-based polymers

Cellulose
Neutral polysaccharides
Chitin (Chitosan)
Cellulose polysaccharides
Polysaccharides
Gellan
Lignin

Cellulose monomer
Starch monomer
Chitin monomer
Chitosan monomer
Hyaluronan monomer
Pektin
Alginate

The slide displays a flowchart on the left showing the classification of natural sugar-based polymers. It branches into Neutral polysaccharides (Cellulose, Chitin/Chitosan) and Amino polysaccharides (Chitin/Chitosan). Below these are Polysaccharides (Gellan, Lignin) and Polyphenols. On the right, several chemical structures are shown for Cellulose monomer, Starch monomer, Chitin monomer, Chitosan monomer, Hyaluronan monomer, Pektin, and Alginate.

Prof. Dr. Ir. Amin Retnoningsih, MSi



The 2nd International Conference on Mathematics, Science, and Education

Amin Retnon...

THREE WAYS TO FIND NEW SUPERIOR DURIAN VARIETIES

- 1) **exploration in forests, fields, or durian centers:** requires much time, effort, and money, and there is no guarantee that we can found new varieties
- 2) **durian breeding:** This durian breeding takes a very long time and is expensive because it requires many parent plants, extensive experimental areas, first to bear fruit aged 4-5 years. The results are not necessarily superior because the mating is random.
- 3) **durian fruit contests at production centers:** The wealth of durian germplasm in Kalimantan is guaranteed to be very abundant, considering that natural crossing is straightforward to occur. The most efficient and effective way to find new superior durian varieties is through durian contests in production centers

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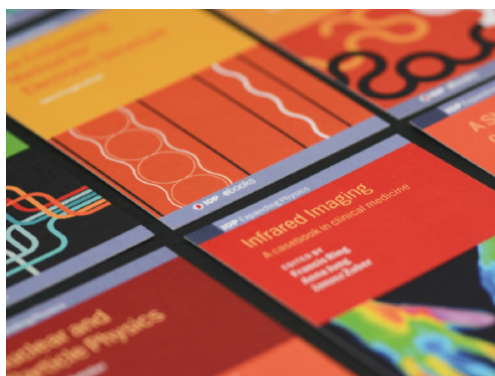
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- **Type of peer review: Single-blind / ~~Double-blind~~ / ~~Triple-blind~~ / Open / Other (please describe)**
- **Conference submission management system:** www.icmseunnes.com
- **Number of submissions received:** 513
- **Number of submissions sent for review:** 393
- **Number of submissions accepted:** 327
- **Acceptance Rate (Number of Submissions Accepted / Number of Submissions Received X 100):** $327/513 \times 100 = 63.74$
- **Average number of reviews per paper:** 3-4 reviews per paper
- **Total number of reviewers involved:** 43
- **Any additional info on review process:**

The review process of abstract was done before conference day to inform to the authors to present their works in the conference and submit their full manuscript. Having been full manuscripts submitted, they were reviewed by our reviewers. The review process of full manuscript has been done for six months, since October 10, 2020 until March 31, 2021. The manuscript must comply these requirements to be proceeded to publication stage,

 - 1) The topic of manuscript should be relevant to conference scopes,
 - 2) The manuscript must be written in appropriate English,
 - 3) The manuscript must be written using IOP template,
 - 4) The manuscript should cite totally at least 15 references with at least 80% are international journal articles. It is suggested the references are published at the last seven years,
 - 5) Citation style is numbering with the first cited the first appears,
 - 6) The section sequence generally is: Introduction, Methods, Result and Discussion, Conclusion, Acknowledgement (optional), References. There is no section number for the two last sections,
 - 7) The references must be cited predominantly in Introduction dan Results and Discussion sections. There is no citation in Abstract nor in Conclusion.
- **Contact person for queries: Dr. Masturi (email: tourfis@gmail.com)**



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Accepted papers received: 07 May 2021

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Optical properties of zinc oxide thin film fabricated by spraying deposition method

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The Atwood machine experiment assisted by smartphone acceleration sensor for enhancing classical mechanics experiments

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022010

Characterization of Mach Zehnder interferometer plastic optical fiber for intensity-based temperature sensor

Ian Yulianti, N M Dharma Putra, Fianti, H Rumiana, Z A F Latif, K E Kurniansyah and S Maimanah

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Critical thinking skills assessment instrument in physics subjects: how to develop a four tier diagnostic test?

A A Waluyo, Hartono and Sulhadi

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A Yulianto and A Rohman

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Re-use of waste components of accumulators (accu) as x-ray radiation protection

Susilo, W H Nugrahaningsih, L Prasetya, L Ni'mah, R Setiawan and Mosik

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B Astuti, I Suryaningsih, A Rusilowati, Khumaedi and H H Kusuma

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Review: nanocomposite of bioactive glass/forsterite from raw material sand and egg shell for bone and dental implants

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S Rufaida and N Nurfadilah

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The SCAPS-1D modeling of ZnO/CdS/CdTe thin film: analysis of thickness and stoichiometric fraction of absorber layer on solar cell performance

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An analysis of thinking patterns of natural sciences teacher candidate students in understanding physics phenomena using P-Prims perspective

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Experimental and kinetic study of free fatty acid esterification derived from *Ceiba pentandra* seed oil with ethanol

R D Kusumaningtyas^{1,*}, Haifah¹, D Widjanarko², H Prasetiawan¹, Y W P Budiono¹, A D H Kusuma¹, N D Anggraeni¹ and S C F Kurnita¹

¹ Chemical Engineering Department, Faculty of Engineering, Universitas Negeri Semarang, Indonesia

² Mechanical Engineering Department, Faculty of Engineering, Universitas Negeri Semarang, Indonesia

*Corresponding author: ratnadewi.kusumaningtyas@mail.unnes.ac.id

Abstract. Indonesia has an abundant feedstock of *Ceiba Pentandra* which is very potential as a source for biodiesel production. However, it contains very high free fatty acid and can not be directly converted into biodiesel through transesterification process. One of the solution is by reacting *Ceiba pentandra* oil with ethanol under acid catalyst called esterification process. In this study, the operating conditions used for esterification reaction was *Ceiba pentandra* oil to methanol molar ratio of 1:12; with reaction time of 120 minutes. The reaction temperature was varied into 40°C, 50°C, 60°C and 70°C. The results show that, highest conversion achieved was 93.5% at reaction temperature of 70°C. Based on the experimental data, pseudo-Homogeneous model was used to model the kinetic of esterification reaction. Based on this model, the activation energy was 21.319kJ/mol and the kinetic factor was 14,264.08/minutes with R² of 0.9675 and SSE of 0.0004.

1. Introduction

Biodiesel is a promising alternative fuel which can be produced from various renewable feed stocks such as oil from plants and animal fats[1]. Several studies have been utilized castor oil [2], used cooking oil[3], *Ceiba Pentandra* oil [4–6] and nyamplung oil [7] through trans-esterification process. However, these natural resources contain high free fatty acid (FFA) which leads to side reactions like saponification that have a negative effect on biodiesel production. Prior to this condition, reduction of FFA on non-edible oils should be performed through esterification process by using low molecular weight alcohol such as ethanol and methanol [8].

Several researchers have studied the conversion of *Ceiba Pentandra* oil into biodiesel [6,9–12]. Previous studies on the esterification of *Ceiba pentandra* oil also had been performed by Kusumaningtyas [13] by using methanol, the highest conversion obtained was 95.14%. However, kinetic study on the esterification of *Ceiba pentandra* oil with ethanol on the presence of H₂SO₄ has not been reported yet.

This work is going to provide experimental data and the mathematical modeling of FFA from *Ceiba Pentandra* oil esterification with ethanol took place. Effect of the esterification parameters such as reaction time and temperature were studied towards the FFA conversion.



2. Materials and Methods

2.1. Materials

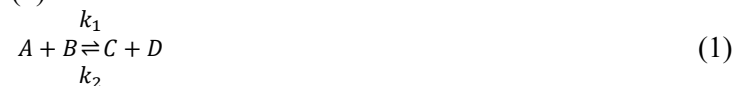
Materials used in this research were crude *Ceiba Pentandra* oil from Pati, Jawa Tengah, Indonesia, Ethanol (99.9% p.a. from Merck), H₃PO₄ (Merck), KOH (Merck) and Oxalic acid (Merck).

2.2. Methods and Analysis

In this research, the degumming process, *Ceiba pentandra* oil and esterification process were following the same steps such as the previous studies[13]. However, in the esterification process the methanol was replaced by using an ethanol as a reactant.

2.3. Kinetic Model

In this research, three different pseudo-homogeneous kinetic models were introduced i.e. irreversible 1st order, irreversible second order and reversible second order kinetic model. The esterification of FFA can be written as shown in equation (1).



where, A is the FFA (RCOOH), B is the ethanol (CH₃OH), C is the methyl ester (R'COOR), D is the water (H₂O), k₁ is the reaction rate constant of forward reaction and k₂ is the reaction rate constant of backward reaction. According to equation 1, the reaction rate of esterification reaction is shown in equation 2.

$$(-r_A) = -\frac{dC_A}{dt} = k_1[A][B] - k_2[C][D] \quad (2)$$

where, $-r_A$ is the reaction rate of A (FFA), [A] is the concentration of FFA, [B] is the concentration of the ethanol, [C] the methyl ester concentration, [D] is the concentration of water and t is the reaction time. Based on equation 2, general equation for irreversible reaction kinetic model is shown in equation 3.

$$\frac{dX_A}{dt} = kC_{A_0}^{\alpha-1}(1 - X_A)^\alpha \quad (3)$$

where, k is the reaction rate constant (1/minute), C_{A0} is the initial concentration of FFA (mol/L), X_A is the conversion of FFA and α is the order of reaction.

2.3.1. Model 1: Irreversible 1st Order Kinetic Model. In this model, the value of $\alpha = 1$. By substituting the α into equation 3 and integrating the equation, the kinetic model for irreversible 1st order reaction is shown in equation 4.

$$-\ln(1 - X_A) = kt \quad (4)$$

2.3.2. Model 2: Irreversible 2nd Order Kinetic Model. In this model, the value of $\alpha = 2$. By substituting the α into equation 3, the kinetic model for irreversible 2nd order reaction is shown in equation 5.

$$\frac{dX_A}{dt} = kC_{A_0}(1 - X_A)^2 \quad (5)$$

The differential equation of FFA conversion can be solved by using polymath 6.1 software.

2.3.3. Model 3: Reversible 2nd Order Kinetic Model. If the backward reaction is taken into account, then a second order reverse reaction can be applied[2]. The kinetics of the second order reverse esterification reaction can be seen in equation 6 [14].

$$\frac{dX_A}{dt} = k_1 \left(\frac{M}{l} - X_A \right) (1 - X_A) - k_2 (X_A)(D_o + X_A) \quad (6)$$

where, M/l is the ethanol to FFA molar ratio and D_o is the molar ratio of water to FFA in the reactor feed.

2.3.4. Kinetic Parameter. The best Model was then further calculated for the kinetic parameter by using equation 7.

$$\ln k = \ln A e^{-\frac{Ea}{RT}} \tag{7}$$

Where, Ae is the kinetic factor, Ea is the energy activity, R is gas constant and T is the reaction temperature

3. Results and Discussion

3.1. Properties of *Ceiba pentandra* Oil

The raw material analysis was carried out on *Ceiba pentandra* oil before and after the degumming process. The raw material was analysed for its fatty acid content by using GC-MS. The density test is carried out to determine the density of the oil. The viscosity of the oil can be determined through the viscosity test, while the acid number test is used to determine the acid number in the oil prior to the esterification process. Density of *Ceiba pentandra* oil before degumming process is 0.942 g/ml, and after degumming process it decreases to 0.941 g/ml. The viscosity of *Ceiba pentandra* oil after degumming process decreases from 35.70 cP to 35.36 cP. While, the acid number drop significantly from 19.6 to 17.82 mg KOH/g oil which also affect the acidity of the oil from 9.78 to 8.89

3.2. Effect of Reaction Time and Temperature on the FFA Content

Figure 1 shows that there is a continuous decline in FFA as a function of reaction time. At the temperature of 70°C, it can be seen that the FFA decreased to 0.782%. These results have met the FFA standard in oil which is 2%. It can be concluded that the esterification of *Ceiba pentandra* oil can be carried out in the presence of 0.5% sulfuric acid catalyst at the temperature of 70°C.

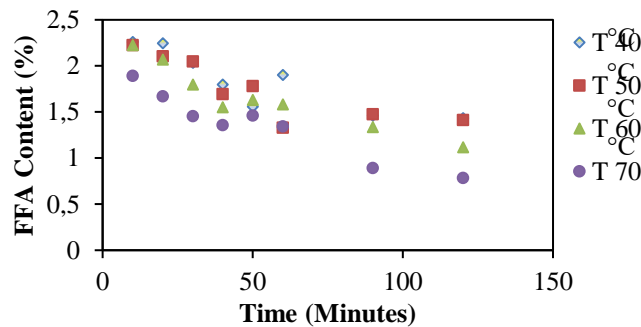


Figure 1. Effect of reaction time and temperature to the FFA content.

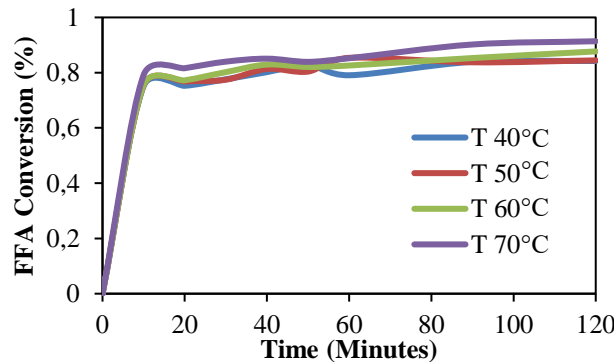


Figure 2. Effect of reaction time and temperature to the FFA conversion.

3.3. Effect of Reaction Time and Temperature on the FFA Conversion

Figure 2 shows that the FFA conversion is increasing exponentially at the first 10 minutes of reaction time, and slightly increasing for the rest of reaction time. The highest conversion of FFA was found at the reaction temperature of 70°C and 120 minutes of reaction time with an FFA conversion of 91.37%. Based on these results, the optimum time for the esterification reaction is at a temperature of 70°C for 120 minutes. Higher Temperature and longer time reaction will not affect significantly on the FFA conversion.

At temperature of 40°C the conversion of FFA is lower than the conversion at 70°C. With a higher temperature, the conversion will have a faster rate. In this reaction, the optimum temperature was found to be at 70 °C. A positive effect on FFA conversion with temperature is due to the increase in kinetic energy of the reactant molecule[8].

3.4. Kinetic Parameter of Model 1

Figure 3 shows the experimental data and the predicted result of FFA Conversion by using Model 1.

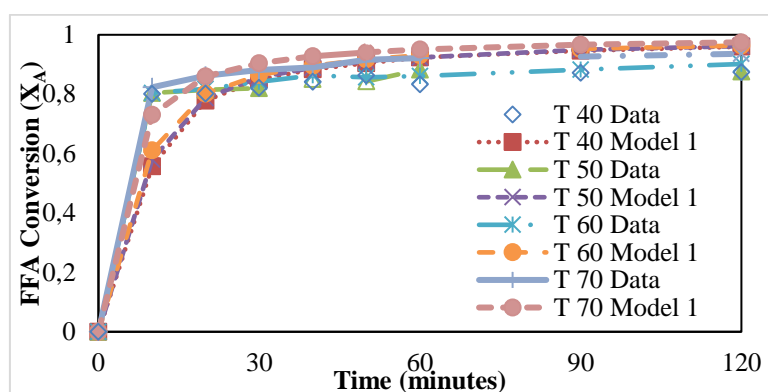


Figure 3. Experimental and calculated conversion of FFA by using Model 1.

It can be seen that, with the increase in reaction temperature it also increases the reaction rate and reduces the reaction time[15]. At the temperature of 40°C, it requires 90 minutes to obtain $\pm 90\%$ conversion while it only needs 45 minutes to get the same conversion at 70°C. The kinetic reaction rate constants, sum square of errors (SSE) and the coefficients of determination (R^2) are shown in Table 1.

Table 1. Kinetic parameter of Model 1.

Temperature (°C)	k (L/mol.min)	SSE	R^2
40	1.36×10^{-2}	0.1318	0.296
50	1.44×10^{-2}	0.1135	0.325
60	1.56×10^{-2}	0.085	0.398
70	2.00×10^{-2}	0.053	0.666

3.5. Kinetic Parameter of Model 2

Figure 4 shows the experimental data and the predicted result of FFA Conversion by using Model 2.

The kinetic reaction rate constants, sum square of errors (SSE) and the coefficients of determination (R^2) are shown in Table 2.

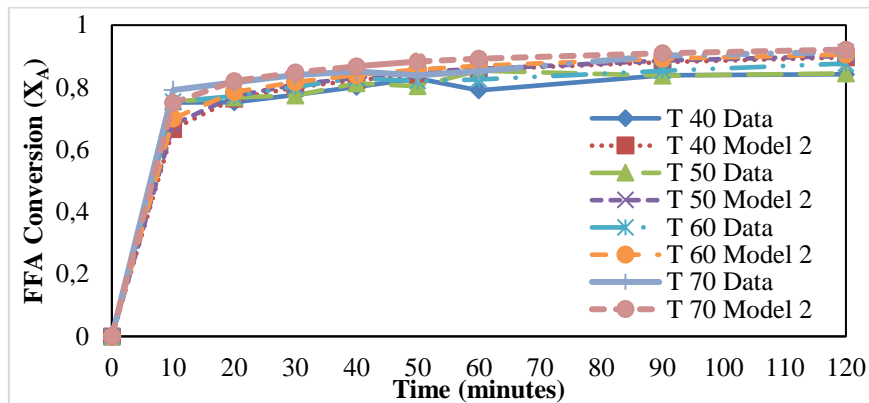


Figure 4. Experimental and calculated conversion of FFA by using Model 2.

Table 2. Kinetic parameter of Model 2.

Temperature (°C)	k (L/mol.min)	SSE	R ²
40	2.2398	0.017	0.5184
50	2.4655	0.014	0.5690
60	2.7877	0.009	0.6975
70	4.2198	0.005	0.7156

The SSE of Model 2 is much lower compared to Model 2. It is in line with the *Le Chatelier* principle where the excess of ethanol cause the concentration of the reactants is higher than the concentration of the products. Due to this condition, the reaction will be shifted towards the product side. [13]

3.6. Kinetic Parameter of Model 3

Figure 5 shows the experimental data and the predicted result of FFA Conversion by using Model 3.

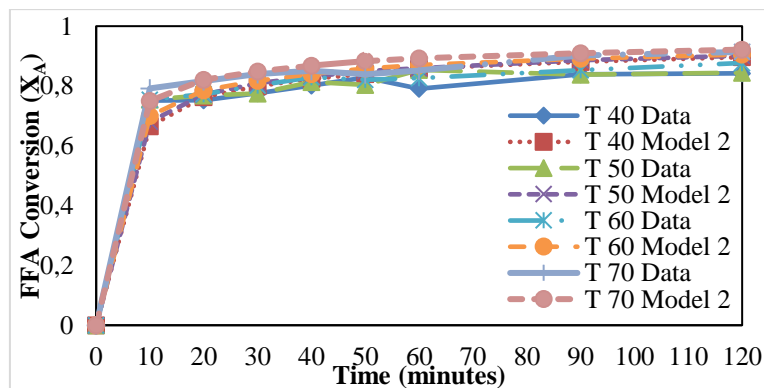


Figure 5. Experimental and calculated conversion of FFA by using Model 3.

The kinetic reaction rate constants, sum square of errors (SSE) and the coefficients of determination (R²) are shown in Table 3.

Based on Table 1, 2 and 3 the average SSE for Model 1, 2 and 3 were 0.0958, 0.01125 and 0.071 respectively. From Table 3, it can be seen that the kinetic reaction rate of forward and backward reaction have almost the same value. It probably, the backward reaction is also can be considered in

the esterification reaction[5]. However, SSE of Model 2 is much lower compared to the Model 3, it indicates that the model is particularly in a good agreement with the experimental data for FFA esterification process. By using arrhenius equation, it can be obtained that this reaction had an energy activity of 17.74 kJ/mol and kinetic factor of 2,015.85/minute.

Table 3. Kinetic parameter of Model 2.

Temperature (°C)	k_1 (L/mol.min)	k_2 (L/mol.min)	SSE	R^2
40	3.22×10^{-2}	7.43×10^{-2}	0.099	0.638
50	3.30×10^{-2}	7.04×10^{-2}	0.0881	0.673
60	3.41×10^{-2}	6.58×10^{-2}	0.074	0.788
70	3.58×10^{-2}	4.47×10^{-2}	0.023	0.690

4. Conclusion

The conversion free fatty acid is affected by reaction time and temperature. Higher reaction temperature increases the FFA conversion. Best FFA conversion obtained was 91.37% at reaction temperature of 70°C and 120 minutes reaction time. The best kinetics model to predict FFA conversion in this esterification model was an irreversible second-order model with the reaction rate constant with SSE of 0.01125 and R^2 of 0.625. The kinetics parameter of *Ceiba pentandra* esterification reaction with a 0.5% sulfuric acid catalyst had an energy activity of 17.74 kJ / mol and kinetic factor of 2,015.85/minute.

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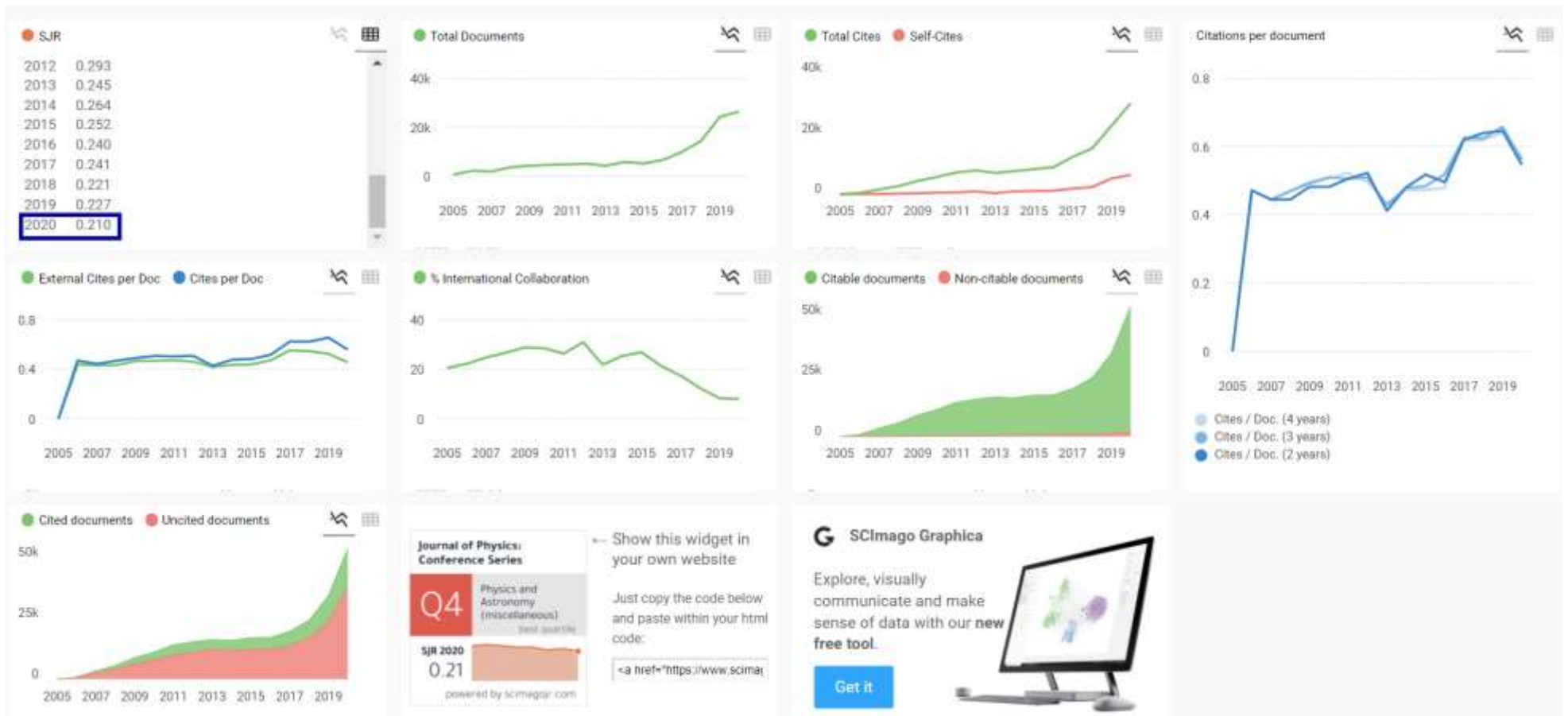
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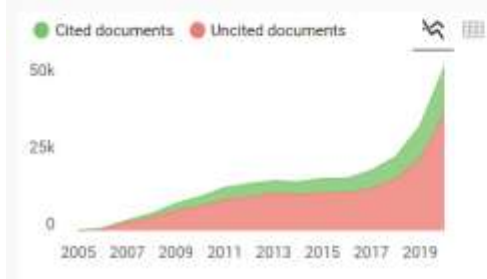
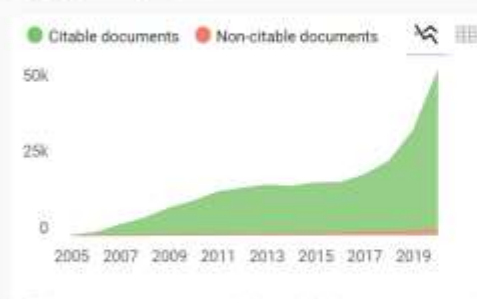
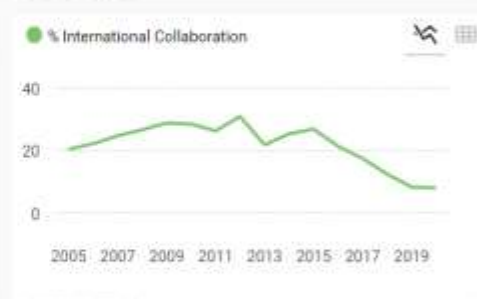
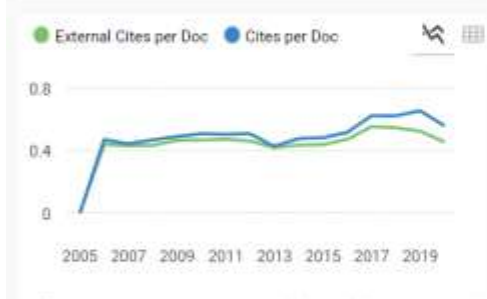
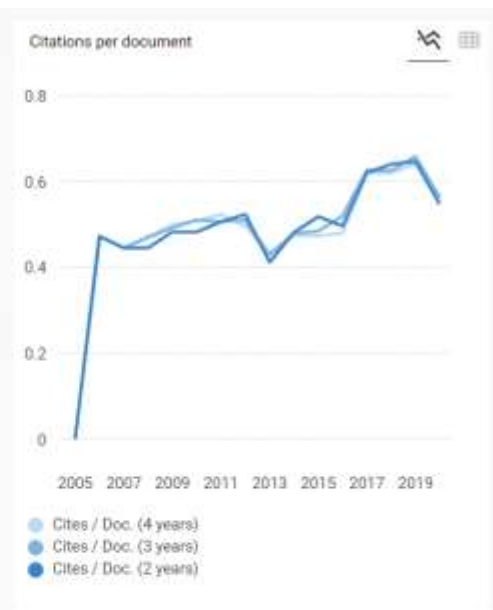
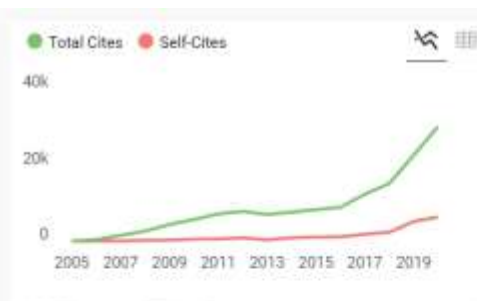
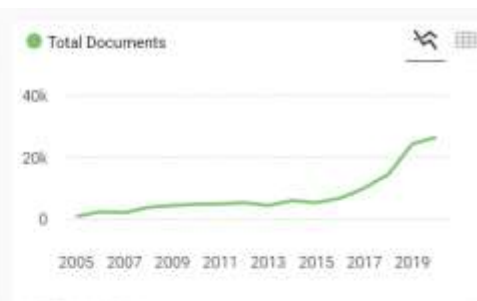
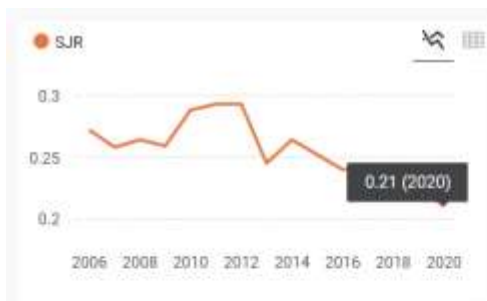
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