

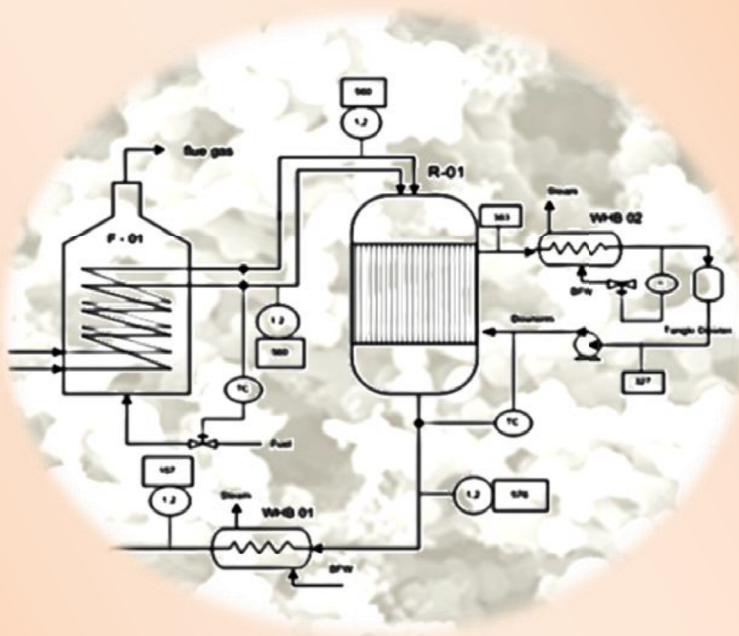


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PREFACE

This abstract book contains abstracts of research papers presented on International Conference on Chemical and Material Engineering (ICCME) 2012. The ICCME 2012 is organized by Department of Chemical Engineering Diponegoro University along with the opening of the doctoral program in the Department of Chemical Engineering. The conference took place at Grand Candi Hotel, Semarang Indonesia in September 12-13, 2012. It is designed as an international forum on fundamental and application of Chemical and Material Engineering among the researchers, students, industries and government. The ICCME offers a platform for extensive sharing and exchange of ideas, thoughts and discussions on all aspects of Chemical and Material Engineering.

The ICCME 2012 invited 65 keynote speakers from 84 and accepted 70 research papers. There are 4 (four) parallel sessions comprising 4 categories: Bioprocess and Renewable Energy (BRE), Material and Science Development (MSD), Separation and Process Engineering (SPE) and Process System Engineering (PSE). The full research papers can be obtained on the enclosed CD Proceeding with ISBN of 978-602-097-281-7

The organizing committee express our gratitude to the distinguished keynote speakers: Prof. M Ulbricht, Prof Hadi Nur, Prof Purwanto, Ir. Gunung Sardjono, Ir. Hardiono and Ichsan, MSc, PDEng for their participation to this conference. We thank to all participants for their contributions to the Conference Programme, abstract book and conference proceedings. We also express our sincere thanks to the scientific committee for reviewing and evaluating the abstracts.

It is very pleased to acknowledge to the conference sponsors: IKPT, Pertamina Gas, BP Migas, Pura Grup, C-BIORE, MER-C, D-WaRE, BCREC for their financial support to enable this conference to be accomplished.

Cover	1
Scientific and Editorial Board Members	2
Editor	3
Organizing Committee	4
Preface	6
Contents	7
Table of Abstracts	8
Programme Schedule	17
Abstract	
Invited Speakers	21
Oral Presentation	
Bioprocess and Renewable Energy (BRE)	27
Material and Science Development (MSD)	52
Process System Engineering (PSE)	80
Separation and Process Engineering (PSE)	92
Acknowledgement	

Table of Papers

Invited Speakers

- [IS – 01](#) **PROF. DR. MATHIAS ULBRICHT**
UDuE, Germany
- [IS – 02](#) **PROF. DR. PURWANTO**
Diponegoro University, Indonesia
- [IS – 03](#) **IR. R. GUNUNG SARDJONO HADI, MT**
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- [IS – 04](#) **DRS. HARDIONO, MComm**
BP Migas, Indonesia
- [IS – 05](#) **ICHSAN, MSC, PDENG**
PT Maris Sustainable Indonesia
- [IS – 06](#) **PROF. DR. HADI NUR**
UTM, Malaysia

Oral Presentation

BRE – BIOPROCESS AND RENEWABLE ENERGY

- [BRE – 01](#) **ENZYMATIC HYDROLYSIS OF ALKALINE PRETREATED COCONUT COIR**
Akbarningrum Fatmawati^a, Rudy Agustriyanto^a, Carolina Adhelia^a, Jovita Paulina^a, Yusnita Liasari^b
^a *Chemical Engineering Department, Faculty of Engineering, Surabaya University, INDONESIA*
^b *Faculty of Biotechnology, Surabaya University, INDONESIA*
- [BRE – 02](#) **OPTIMIZATION OF STREPTOMYCES SP.A11 MEDIUM CULTIVATION ON CYCLO(TYROSYL-PROLYL) PRODUCTION USING THE RESPONSE SURFACE METHODOLOGY**
Rofiq Sunaryanto
Center of Biotechnology, Badan Pengkajian dan Penerapan Teknologi (BPPT), INDONESIA
- [BRE – 03](#) **ENHANCEMENT OF BIOMASS PRODUCTION FROM SPIRULINA SP CULTIVATED IN POME MEDIUM**
Hadiyanto, Muhamad Maulana Azimatun Nur, Ganang Dwi H
Center of Biomass and Renewable Energy, Department of Chemical Engineering Diponegoro University, INDONESIA
- [BRE – 04](#) **ETHANOL PRODUCTION FROM NON FOOD TUBERS OF ILES-ILES (AMORPHOPHALLUS CAMPANULATUS) USING HYDROLYZES BY COMMERCIAL ENZYMES (A AND B AMYLASE) AND FERMENTATION BY SACCHAROMICES CEREVISEAE**
Kusmiyati^{a,b}, Asha Tridayana^a, Nurul Widya FP^a, Tri Utami^a
^a *Chemical Engineering Department, Faculty of Engineering, Muhammadiyah Surakarta University INDONESIA*
^b *Renewable Energy Research Centre, Muhammadiyah Surakarta University - Surakarta INDONESIA*
- [BRE – 05](#) **SYNTHESIS OF ZEOLITE PELLETS FROM NATURAL ZEOLITE AND STARCH AS ADSORBENT FOR FUEL GRADE BIOETHANOL PRODUCTION**
Anwar Ma'rif and Neni Damajanti
Chemical Engineering Department, Faculty of Engineering, Muhammadiyah University of Purwokerto, INDONESIA

- BRE – 06** **QUALITY IMPROVEMENT (FIBER CONTENT AND PROTEIN DIGESTIBILITY VALUE) OF CASSAVA PEEL BY FERMENTATION USING TAPE YEAST WITH VITAMIN B SUPPLEMENTATION**
 Wikanastri Hersoelistyorini^a, Cahya S. Utama^b
^a *Food Technology Study Program, University of Muhammadiyah Semarang, INDONESIA*
^b *Department of Animal Feed and Nutrition, Diponegoro University, INDONESIA*
- BRE – 07** **THE DETERMINATION OF SALINITY PROFILE AND NUTRITION (NAH₂PO₄) PROFILE IN UTILIZING NANNOCHLOROPSIS OCLATA TO GAIN MAXIMUM LIPID**
 Elida Purba, Kenjiro Parsaulian Siburian
Chemical Engineering Department, Faculty of Engineering, University of Lampung, INDONESIA
- BRE – 08** **DESIGN PROCESS OF ETHANOL PRODUCTION BY EXTRACTIVE -FERMENTATION TO INCREASE THE YIELD AND PRODUCTIVITY OF ETHANOL**
 Ayu Ratna Permanasari, Ririn Indriani AR, Tri Widjaja, Ali Altway
Chemical Engineering Department, Faculty of Industrial Technology, Sepuluh Nopember Institut of Technology, INDONESIA
- BRE – 09** **TRANSESTERIFICATION OF VEGETABLES OIL USING SUB-AND SUPERCRITICAL METHANOL**
 Nyoman Puspa Asri^{a,d}, Siti Machmudah^{a,b}, Wahyudiono^c, Suprpto^a, Kusno Budikarjono^a, Achmad Roesyadi^a, Mitsuru Sasaki^c, Motonobu Goto^b
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^c *Graduate School of Science and Technology, Kumamoto University, JAPAN*
^d *Chemical Engineering Department, Faculty of engineering, WR Supratman University, Surabaya, INDONESIA*
- BRE – 10** **THE INFLUENCE OF CATALYSTS TO SELECTIVITY OF PRODUCT OF PALM OIL CRACKING**
 Achmad Roesyadi, Danawati Hariprajitno, Nurjannah, Santi Dyah Savitri
Chemical Reaction Engineering Laboratory, Department Of Chemical Engineering Department Of Chemical Engineering, INDONESIA
- BRE – 11** **BIO-LUBRICANTS DEVELOPMENT: POTENTIAL USE OF BORON-CONTAINING ADDITIVES**
 Dicky Dermawan, Dyah Setyo Pertiwi
Chemical Engineering Department, Faculty of Industrial Technology, Itenas, INDONESIA
- BRE – 12** **UTILIZATION POTENCY OF EXTRACELLULAR POLYMERIC SUBSTANCE AS INDUSTRIALS BIOSORBENT AND ION EXCHANGE RESIN**
 Zainus Salimin^a, Pungky Ayu Artiani^a, Junaidi^b, and Wawan^b
^a *Radioactive Waste Technology Center, National Nuclear Energy Agency – BATAN, INDONESIA*
^b *Study Program of Environmental Engineering, Faculty of Engineering, Diponegoro University, INDONESIA*
- BRE – 13** **IMMOBILIZATION OF COW RUMEN FLUID CELLULASE BY ENTRAPMENT IN CALCIUM ALGINATE BEADS**
 Indah Hartati^a, Laeli Kurniasari^a, and Agnes Budiarti^b
^a *Department of Chemical Engineering, Faculty of Engineering, Wahid, INDONESIA*
^b *Department of Pharmacy, Faculty of Pharmacy, Wahid Hasyim University, INDONESIA*
- BRE – 14** **OPTIMIZATION PROCESS OF BIODIESEL PRODUCTION FROM NYAMPLUNG SEED (CALOPHYLLUM INOPHYLLUM L) USING IN SITU PROCESS AND ULTRASONIC ASSISTED**
 Widayat, Abdullah, Kanevi Octova Paradita and Elsanta Monaliza Tungga D
Department of Chemical Engineering Faculty of Engineering Diponegoro University, INDONESIA
- BRE – 15** **EFFECT OF PHENYLACETIC ACID ADDITION ON PRODUCTIVITY OF PENICILLIUM CHRYSOGENUM IN PENICILLIN G PRODUCTION USING PILOT SCALE REACTOR**
 Amila Pramiasandi, Rofiq Sunaryanto, Suyanto
Center for the Application of Biotechnology, BPPT, INDONESIA
- BRE – 16** **STUDY OF ENZYMATIC HYDROLYSIS OF DILUTE ACID PRETREATED COCONUT HUSK**
 Rudy Agustriyanto, Akbarningrum Fatmawati, Maria Angelina, Raissa Monica
Surabaya Unversity, INDONESIA

- BRE – 17** **KINETICS OF ETHANOL PRODUCTION FROM WHEY BY FEMENTATION USING KLUYVEROMYCES MARXIANUS**
 Dessy Ariyanti^{ab}, Desiyantri Siti Pinundi^a, Apsari Puspita Aini^a, Hadiyanto^{ab}, Djoko Murwono^a
^a *Chemical Engineering Department, Faculty of Engineering, Diponegoro University, INDONESIA*
^b *Center of Biomass and Renewable Energy (C-BIORE), Faculty of Engineering, Diponegoro University, INDONESIA*
- BRE – 18** **POTENTIAL OF SO₄²⁻ / ZnO ACID CATALYST FOR HETEROGENEOUS TRANSESTERIFICATION OF VEGETABLE OIL TO BIODIESEL**
 I. Istadi, Didi D Anggoro, Luqman Buchori, Inshani Utami, and Roikhatus Solikhah
Laboratory of Energy and Process Engineering, , Chemical Reaction Engineering an Catalysis Group, Department of Chemical Engineering, Diponegoro University, INDONESIA
- BRE – 19** **A SIMPLE METHOD FOR EFFICIENT EXTRACTION AND SEPARATION OF C-PHYCOCYANIN FROM SPIRULINA PLATENSIS**
 Noer Abyor Handayani^{ab}, Hadiyanto^{ab}, Melinda^a, Inggar^a, Amin Nugroho^a
^a *Chemical Engineering Department, Faculty of Engineering, Diponegoro University, INDONESIA*
^b *Center of Biomass and Renewable Energy (C-BIORE), Faculty of Engineering, Diponegoro University, INDONESIA*
- BRE – 20** **STUDY ON SLAUGHTERHOUSE WASTES POTENCY AND CHARACTERISTIC FOR BIOGAS PRODUCTION**
 Budiyo^a, I Nyoman Widiasta^a, Seno Johari^b, Sunarso^c
^a *Department of Chemical Engineering, Diponegoro University, INDONESIA*
^b *Faculty of Animal Science and Agriculture, Diponegoro University, INDONESIA*
- BRE – 21** **STUDY ON PRODUCTION PROCESS OF BIODIESEL FROM RUBBER SEED (HEVEA BRASILIENSIS) BY IN SITU TRANSESTERIFICATION METHOD WITH ALKALINE CATALYZED**
 Widayat^{a,b}, Agam Duma Kalista Wibowo^a, Hadiyanto^{a,b}
^a *Magister of Chemical Engineering, Diponegoro University, INDONESIA*
^b *Central of Biomass and Renewable Energy, INDONESIA*
- BRE – 22** **THERMOGRAVIMETRY CHARACTERISTICS OF MSW CHAR BRIQUETTE COMBUSTION**
 Dwi Aries Himawanto^a, Indarto^b, Harwin Saptoadi^b, Tri Agung Rohmat^b
^a *Mechanical Engineering Department Faculty of Engineering, Sebelas Maret University, INDONESIA*
^b *Mechanical and Industrial Engineering Department Faculty of Engineering, Gadjah Mada University, INDONESIA*
- BRE – 23** **PERFORMANCE OF SULFONATED POLY ETHER-ETHER KETONE (SPEEK) AND NAFION MEMBRANE IN PALM OIL MILL EFFLUENT MICROBIAL FUEL CELL**
 Nur Dianaty Nordina Abdul Halim Mohd Yusof^{a,b,c}, Ahmad Fauzi Ismail^{a,d}, Mohd Noorul Anam Mohd Norddin^{a,d}
^a *Faculty of Chemical Engineering, Universiti Teknologi Malaysia, MALAYSIA*
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^c *Department of Biology, Massachusetts Institute of Technology, UNITED STATES OF AMERICA*
^d *Faculty of Petroleum and Renewable Engineering, Universiti Teknologi Malaysia, MALAYSIA.*
- BRE – 24** **SURFACE MODIFICATION OF POLYETHERSULFONE WITH POLYVINYLPIRROLIDONE-IODINE VIA PHASE INVERSION AND UV PHOTOGRATING FOR ANTIBACTERIAL APPLICATIONS**
 Abdul Halim Mohd Yusof^{a,b}, Devanai Kannan^a
^a *Department of Bioprocess Engineering, Faculty of Chemical Engineering;Universiti Teknologi Malaysia, MALAYSIA*
^b *Department of Biology, Massachusetts Institute of Technology, UNITED STATES OF AMERICA*
- BRE – 25** **ESTERIFICATION OF JATROPHA OIL TO BIODIESEL OVER SiO₂-PHOSPHOTUNGSTIC ACID HETEROGENEOUS CATALYST**
 Nur Hidayati, Aning Tri Aisyah, Ike Sambung Sari and Prinda Widayari
Chemical Engineering Department, Universitas Muhammadiyah Surakarta, INDONESIA

- [MSD – 01](#) **CHARACTERIZATION OF SAGO STARCH AND STUDY OF LIQUEFICATION PROCESS ON HIGH FRUCTOSE SYRUP PRODUCTION**
Anastasia Prima Kristijarti, Tony Handoko, Cindy Adelia, Lucy Andrea
Chemical Engineering Department, Faculty of Industrial Engineering, Parahyangan Catholic University, INDONESIA
- [MSD – 02](#) **POLYMER SOLAR CELLS: EFFECTS OF ANNEALING TREATMENT AND POLYMER BLENDS ON I-V CHARACTERISTICS**
Erlyta Septa Rosa and Shobih
Research Center for Electronics and Telecommunication, Indonesian Institute of Sciences (PPET-LIP), INDONESIA
- [MSD – 03](#) **THE OPTIMIZATION OF DIPHENYL METHANE DIISOCYANATE POLYMERIZATION PROCESS WITH THE USED FRYING OIL POLYALCOHOL TO FOAM POLYURETHANE USING RSM**
Faleh Setia Budi^a and Didi Dwi Anggoro^b
^a *Food Science and Technology Department, Faculty of Agricultural Engineering and Technology, IPB, INDONESIA*
^b *Chemical Engineering Department, Engineering Faculty UNDIP, INDONESIA*
- [MSD – 04](#) **NUMERICAL ANALYSIS OF A JOURNAL BEARING WITH CHEMICAL ROUGHNESS**
Mohammad Tauviqirrahman^{a,b}, Muchammad^a, Jamari^b, and Dik J. Schipper^a
^a *Laboratory for Surface Technology and Tribology, Faculty of Engineering Technology, University of Twente, THE NETHERLANDS*
^b *Department of Mechanical Engineering, University of Diponegoro, INDONESIA*
- [MSD – 05](#) **INVESTIGATION ON CENTRIFUGAL PUMP SHAFT: A COMPARISON STUDY OF THE SME AND THE IMPORTED PRODUCT**
Rifky Ismail, Sugiyanto, Mohammad Tauviqirrahman and Jamari
Mechanical Engineering Department, Faculty of Engineering, Diponegoro University, Jl. Prof Sudharto, SH-Tembalang, Semarang INDONESIA
- [MSD – 06](#) **PREPARATION OF NANOPARTICLE SILICA FROM SILICA SAND AND QUARTZITE BY ULTRAFINE GRINDING**
Agus Wahyudi^a, Teguh Nurasi^b And Siti Rochani^a
^a *R&D Centre for Mineral and Coal Technology, INDONESIA*
^b *Department of Chemistry, University of Jenderal Achmad Yani, INDONESIA*
- [MSD – 07](#) **APPLICATION OF TiO₂ FOR SELF CLEANING IN WATER BASED PAINT WITH POLYETHYLENE GLYCOL (PEG) AS DISPERSANT**
Nining Kusmahetiningsih, Dyah Sawitri
Departement of Engineering Physics, Faculty of Industrial Engineering, Sepuluh Nopember Institute of Technology, INDONESIA
- [MSD – 08](#) **USING SELF CLEANING TiO₂ PHOTOCATALYST IN THE MAKING OF CERAMIC TILES TO DECREASE AMMONIUM CONCENTRATION AND BACTERIUM GROWTH**
Ana Hidayati M^a, Sri Darmawati^a, Muh. Amin^b
^a *Health Analyst Department, Nursing and Health Faculty, University of Muhammadiyah Semarang, INDONESIA*
^b *Mechanical Engineering Department, Engineering Faculty, University of Muhammadiyah Semarang, INDONESIA*
- [MSD – 09](#) **THE EFFECT OF POLYANILINE ADDITION ON THE PROPERTIES OF CARBON-BASED POLYPROPYLENE COMPOSITE**
Lies A. Wisojodharmo, Dewi Kusuma Arti, Niya Listiani Dewi
Center for Materials Technology, Agency for the Assessment and Application Technology (BPPT), INDONESIA

- MSD – 10** **COMPUTATIONAL ANALYSIS OF WALL SLIP AND CAVITATION IN LUBRICATED SLIDING SYSTEMS**
 Muchammad^a, M. Tauviquirrahman^a, J. Jamari^b, and D.J. Schipper^a
^a *Laboratory for Surface Technology and Tribology, Faculty of Engineering Technology, University of Twente, THE NETHERLANDS*
^b *Department of Mechanical Engineering, University of Diponegoro, INDONESIA*
- MSD – 11** **FRICTION ANALYSIS ON SCRATCH DEFORMATION MODES OF VISCO-ELASTIC-PLASTIC MATERIALS**
 Budi Setiyana^a, I. Syafaat^a, J. Jamari^b, and D.J. Schipper^a
^a *Laboratory for Surface Technology and Tribology, Faculty of Engineering Technology, University of Twente, THE NETHERLANDS*
^b *Department of Mechanical Engineering, University of Diponegoro, INDONESIA*
- MSD – 12** **PREDICTION OF SLIDING WEAR OF ARTIFICIAL ROUGH SURFACES**
 Imam Syafa'at^a, Budi Setiyana^a, Rifky Ismail^a, Eko Saputra^a, J. Jamari^b, D.J. Schipper^a
^a *Laboratory for Surface Technology and Tribology, Faculty of Engineering Technology, University of Twente, THE NETHERLANDS*
^b *Department of Mechanical Engineering, University of Diponegoro, INDONESIA*
- MSD – 13** **THE APPLICATION OF B-CYCLODEXTRIN AND POLYETHYLENE GLYCOL 6000 IN THE MICRONISATION OF DRUG – POLYMER COMPOSITE WITH PARTICLE FROM GAS SATURATED SOLUTIONS (PGSS) METHOD**
 Putu Riani Pradnyandari, Rizky Tetrisyanda, Prida Novarita Trisanti, Firman Kurniawansyah, and Sumarno
Chemical Engineering Department, Faculty of Industrial Technology, Institute of Technology Sepuluh Nopember (ITS), INDONESIA
- MSD – 14** **THE EFFECT OF SONICATION ON THE CHARACTERISTIC OF CHITOSAN**
 Azra Yuliana, Linggar S. Pradeckta, Emma Savitri, Anita R. Handaratri, Sumarno
Chemical Engineering Department, Faculty of Industry Technology, Sepuluh Nopember Institut of Technology Surabaya, INDONESIA
- MSD – 15** **EFFECTS OF CHAIN EXTENDER TO THE STRUCTURE OF CASTOR OIL-BASED POLYURETHANE FOAM**
 Edhi Pratondo, Adityo Wahyu Hanggoro, Eva Oktavia Ningrum, Sumarno
Chemical Engineering Department, Faculty of Industrial Technology, Sepuluh Nopember Institute of Technology, INDONESIA
- MSD – 16** **THE SYNTHESIS OF CARBOXYMETHYL AMYLOSE GRAFTED POLYACRYLAMIDE AND ITS APPLICATION IN DRUG RELEASE ASPIRIN**
 Noor Hidayati, Naila Amanda, Karsono Samuel Padmawijaya, Firman Kurniawansyah, and Sumarno
Chemical Engineering Department, Faculty of Industry Technology, Sepuluh Nopember Institut of Technology Surabaya, INDONESIA
- MSD – 17** **EFFECT OF TWEEN 80'S EMULSIFIER CONCENTRATION WITH SPONTANEOUS DIFFUSION METHOD ON STABILITY SOLUTION TEGERAN'S NANOEMULSION NATURAL DYES**
 Heny Herawati^a, Sri Yuliani^a, Meika Syahbana Rusli^b, Ratih Purnamasari^b
^a *Indonesia Center For Agricultural Postharvest Research and Development, INDONESIA*
^b *Bogor Agricultural University, Agricultural Technology Faculty, IPB, INDONESIA*
- MSD – 18** **NANO CRYSTALLINE STARCH AND ITS ALTERNATIF IMPLEMENTATION**
 Heny Herawati
Indonesia Center For Agricultural Postharvest Research and Development, INDONESIA
- MSD – 19** **COMPOSITE SPEEK WITH NANOPARTICLES FOR FUEL CELL'S APPLICATIONS: REVIEW**
 Arief Rahman Hakim
Department of Chemical Engineering, Diponegoro University, INDONESIA
- MSD – 20** **COMPARISON ON MODELLING OF DRYING KINETICS OF GRANULAR POLYMERS PA6 BY DIFFUSION MODELS AND NORMALIZATION METHOD**
 Suherman^a, Mirko Peglow^b, and Evangelos Tsotsas^b
^a *Chemical Engineering Department, Faculty of Engineering, Diponegoro University, Jl. Prof Sudharto, SH-Tembalang, Semarang, INDONESIA*
^b *Thermal Process Engineering, Otto-von-Guericke-University, Universitätsplatz 2, 39106 Magdeburg, GERMANY*

- MSD – 21 **CURRENT DENSITY PERFORMANCES IN POLY ETHER ETHER KETON MEMBRANE FOR DIRECT METHANOL FUEL CELLS**
Tutuk Djoko Kusworo^a, E. L. Dewi^b, D. K. Arti^a, A. Dhuhita^a, A. Fauzi Ismail^c and M.N.A. Mohd Norddin^c
^a *Chemical Engineering Department, Engineering Faculty, Diponegoro University, INDONESIA*
^b *Agency of Assessment and Application of Technology*
^c *Advanced Membrane Technology Research Centre, Faculty of Chemical and Natural Resources Engineering, Universiti Teknologi Malaysia, MALAYSIA*
- [MSD – 22](#) **PREPARATION AND CHARACTERIZATION OF ZEOLITE MEMBRANE**
Aprilina Purbasari^a, Titik Istirokhatun^b, Heny Kusumayanti^a, Ariestya M. Devi^a, Lulluil Mahsunah^a, Heru Susanto^a
^a *Chemical Engineering Department, Faculty of Engineering, Diponegoro University, INDONESIA*
^b *Environmental Engineering Department, Faculty of Engineering, Diponegoro University, INDONESIA*
- MSD – 23 **EFFECT OF SOIL BURIAL ON THE MECHANICAL PROPERTIES OF HEAT TREATED AND UNTREATED RED BALAU SAW DUST FILLED LDPE COMPOSITES.**
Ruth Anayimi Lafia-Araga, Aziz Hassan, Rosiyah Yahya, Normasmira Abd. Rahma
Chemistry Department, Faculty of Science, University of Malaya, Malaysia
- MSD – 24 **IMPACT AND DSC PROPERTIES OF HEAT TREATED AND UNTREATED RED BALAU (SHOREA DIPTEROCARPACEAE)/LDPE COMPOSITES**
Aziz Hassan, R.A. Lafia-Araga, R. Yahaya and N.A. Rahman
Department of Chemistry, University of Malaya, Malaysia
- MSD – 25 **COMPUTATION OF INTERFACIAL SHEAR STRENGTH AND OTHER TENSILE RELATED PROPERTIES OF INJECTION MOULDED CARBON FIBRE/POLYAMIDE 6,6**
Rosiyah Yahya, Aziz Hassan, Zulkifli Abu Hasan
Department of Chemistry, University of Malaya, 50603 Kuala Lumpur, MALAYSIA.
- MSD – 26 **IMPACT PROPERTIES OF INJECTION MOULDED ARAMID/CARBON HYBRID FIBRE REINFORCED POLYPROPYLENE COMPOSITES**
MIM Rafiq and Aziz Hassan
Department of Chemistry, University of Malaya, MALAYSIA
- MSD – 27 **HARDNESS OF CONTINUOUS HARD ANODIZING OF ALUMINIUM 6061 AFFECTED BY VOLTAGE AND TIME PROCESS USING TITANIUM CATHODE**
Endi Sutikno, Ellen Desta Purwanendra, Putu Hadi Setyarini
Mechanical Engineering Department, Brawijaya University, INDONESIA
- MSD – 28 **SURFACE CHARACTERISTICS OF ALUMINIUM HARD ANODIZING USING TITANIUM CATHODE**
Putu Hadi Setyarini, Riviero Givenchy, Endi Sutikno
Mechanical Engineering Department, Brawijaya University, INDONESIA
-

PSE – PROCESS AND SYSTEM ENGINEERING

- [PSE – 01](#) **THE PROCESS TRANSPORT OF DRYING CORN WITH MIXED-ADSORPTION DRYING**
Mohamad Djaeni, Luqman Buchori
Chemical Engineering Department, Faculty of Engineering, Diponegoro University, INDONESIA
- [PSE – 02](#) **SIMULATION OF DUPLEX HEAT TREATMENT NB3SN COMPOUNDS IN CU-NB-SN SUPERCONDUCTING MULTIFILAMENTARY WIRE**
Andini Nur Vania Swari, Arimaz Hangga, Doty Dewi Risanti
Department of Engineering Physics, Faculty of Industrial Technology, ITS, INDONESIA

- PSE – 03** **INVENTORY ANALYSIS OF RADIOLOGICAL FOR GRAPHITE THERMAL COLUMN FROM TRIGA MARK II REACTOR, BANDUNG**
Mulyono Daryoko
Radioactive Waste Technology Center, BATAN, INDONESIA
- PSE – 04** **DYNAMIC SIMULATION AND COMPOSITION CONTROL IN A 10 L MIXING TANK**
Yulius Dedy Hermawan
Chemical Engineering Department, Faculty of Industrial Technology, UPN “Veteran” Yogyakarta, INDONESIA
- PSE – 05** **THE EFFECT OF LOADING ON THE CONTACT STRESS OF UHMWPE MATERIAL FOR ARTIFICIAL HIP JOINT BEARING**
Eko Saputra^a, Rifky Ismail^a, Jamari^a and Iwan Budiwan Anwar^b
^a *Laboratory for Engineering Design and Tribology, Department of Mechanical Engineering, University of Diponegoro, INDONESIA*
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- PSE – 06** **SIMULATION OF COUNTER BLOW PROCESS OF PBL QUARTZ BOTTLE FABRICATION**
Arimaz Hangga, Lizda Johar Mawarani, Doty Dewi Risanti
Department of Engineering Physics, Faculty of Industrial Technology, Institut Teknologi Sepuluh Nopember, INDONESIA
- PSE – 07** **CADMIUM METALS PARTICLES-COVERED POLYSTYRENE NANOSPHERES THIN FILM MATERIAL:FABRICATION, ANALYSIS AND MODEL**
Pratama Jujur Wibawa^{a,b,c}, Hashim Saim^d, Moh^d. Arif Agam^d and Hadi Nur^e
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- PSE - 08** **DISTRIBUTION TEMPERATURE OF ANALYSIS ON CH4-CO2 GAS MIXED IN DOUBLE PIPE HEAT EXCHANGER BY CONTROLLED FEEZE OUT AREA METHODE**
Fatma Y. Hasyim, Novi E. Mayangsari, and Sumarno
Material of Laboratory
Chemical Engineering Department, Faculty of Industry Technology, Sepuluh Nopember Institut of Technology Surabaya, INDONESIA
- PSE – 09** **SIMULATION ON THE ACTIVITY OF BIOMASS IN ACTIVATED SLUDGE IN THE PERFORMANCE OF NON IDEAL FLOW MEMBRANE BIOREACTOR SUBMERGED**
Aisyah Endah Palupi
Mechanical Engineering Department, Engineering Faculty, State University of Surabaya 60231 – INDONESIA
- PSE – 11** **EFFICACY AND HEAT TRANSFER EFFECTS OF ARTOCARPUS ALTILIS MALE INFLORESCENSE AS MOSQUITO VAPORIZING MATS**
Muhammad Khamim Asy’Ari^a, Nadhifa Maulida^a, Wilujeng Fitri Alfiah^a, Aisyiah Nur Isaneni^a Titin Yulia Riska^a, and Doty Risanti^b
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- [PSE – 12](#) **SELECTION OF NATURAL DYE PHOTOSENSITIZER FOR QUASI-SOLID STATE DYE-SENSITIZED MESOPOROUS TiO₂ SOLAR CELL (DSC) FABRICATION**
Ruri Agung Wahyuono, Doty Dewi Risanti
Department of Engineering Physics, Institut Teknologi Sepuluh Nopember Surabaya, INDONESIA
- [PSE – 13](#) **AN INVESTIGATION OF CHAR FORMATION AND SHRINKING VOLUME BY VISUALIZATION TECHNIQUE INDUCED BY PYROLYSIS**
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SPE – SEPARATION PROCESS ENGINEERING

- [SPE – 01](#) **UTILIZATION OF COAL FLY ASH AS CO GAS ADSORBENT**
Ayu Lasryza, Dyah Sawitri
Department of Engineering Physics, Faculty of Industrial Technology, ITS, INDONESIA
- [SPE – 02](#) **PHENOL REMOVAL FROM AQUEOUS SOLUTIONS BY ELECTROCOAGULATION TECHNOLOGY USING IRON ELECTRODES: EFFECT OF SOME VARIABLES**
Mohammad Ali Zazouli^a, Mahmoud Taghavi^b
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^b *Department of Environmental Health Engineering, Faculty of Health and Health Sciences Research Center, IRAN*
- [SPE – 03](#) **THE OFF GAS TREATMENT IN THE PROCESS OF VITRIFICATION AND INCINERATION OF NUCLEAR WASTE**
Herlan Martono, Aisyah
Radioactive Waste Technology Centre, BATAN, INDONESIA
- [SPE – 04](#) **ESSENTIAL OIL EXTRACTION OF FENNEL SEED (FOENICULUM VULGARE) USING STEAM DISTILLATION**
Astrilia Damayanti and Eko Setyawan
Chemical Engineering Department, Faculty of Engineering, Semarang State University, INDONESIA
- [SPE – 05](#) **PREPARATION AND CHARACTERIZATION OF NANOFILTRATION MEMBRANE FOR WATER TREATMENT**
Tutuk Djoko Kusworo, Eka Cahya Muliawati and Ardian Dwi Yudhistira
Department of Chemical Engineering, Diponegoro University, INDONESIA
- [SPE – 06](#) **THE APPLICATION OF NITROGEN LASER ON EXTRACTION OF URANIUM IN THE LONG LIFE OF HIGH LEVEL RADIOACTIVE LIQUID WASTE USING TBP-KEROSENE SOLVENT**
Gunandjar
Radioactive Waste Technology Center, National Nuclear Energy Agency of Indonesia (BATAN), INDONESIA
- [SPE – 07](#) **EFFECT OF PH AND STIRRING SPEED ON THE COLLAGENOUS PROTEIN EXTRACTION FROM CHICKEN BONE WASTE IN A WELL AGITATED EXTRACTION SYSTEM**
Andri Cahyo Kumoro, Beatrice L. M. Tanjung, Fadilla H. Utami, Diah Susetyo Retnowati and Catarina Sri Budiayati
Department of Chemical Engineering, Diponegoro University, INDONESIA
- [SPE – 08](#) **DRYING OF CURCUMA (CURCUMA XANTHORRHIZA ROXB) USING DOUBLE PLATE COLLECTOR SOLAR DRYER**
Tjukup Marnoto, Mahreni, Wasir Nuri, Bayu Ardinanto, Ratna E. Puspitasari
Chemical Engineering Department, Faculty of Industrial Technology, UPN "Veteran" Yogyakarta University, INDONESIA

- SPE – 09** **SOLUBILITY EXAMINATION OF PALM KERNEL OIL IN SUPERCRITICAL CO₂ AND ITS CORRELATION WITH SOLVENT DENSITY BASED MODEL**
 Wahyu Bahari Setianto^a, Priyo Atmaji^b and Didi Dwi Anggoro^b
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- SPE – 10** **Oil Extraction Process From Solid Waste Rubber Seed By Soxhletation and Extraction Solvent by Stirring Methods**
 Achmad Wildan, Devina Ingrid A, Indah Hartati
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- SPE – 11** **FOULING AND REJECTION BEHAVIOUR OF ULTRAFILTRATION FOR OIL IN WATER EMULSION SEPARATION**
 Nita Aryanti^a, Agus Hadiyanto^a, Wiharyanto Oktiawan^b
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- SPE – 12** **THE EFFECT OF CHEMICAL ADDITIVES ON MOBILITY OF HEAVY METALS (PB, CD AND ZN) IN SOIL**
 Abdoliman Amouei^a, Amirhossein Mahvi^b, Masoumeh Tahmasbizadeh^b, Mohammad Ali Zazouli^c
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- SPE – 13** **CHARACTERISTICS OF HOSPITAL WASTEWATER IN BABOL UNIVERSITY OF MEDICAL SCIENCES AND EFFECTS ON THE ENVIRONMENT**
 Abdoliman Amouei^a, Hosseinali Asgharnia^a, Hourieh Fallah^a, Aliakbar Mohammadi^a, Mohammad Ali Zazouli^b
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- SPE – 14** **QUANTITY AND QUALITY OF SOLIDWASTES IN THE HOSPITALS OF BABOL UNIVERSITY OF MEDICAL SCIENCES, NORTH OF IRAN**
 Abdoliman Amouei^a, Masoumeh Tahmasbizadeh^b, Hosseinali Asgharnia^a, Hourieh Fallah^a, Aliakbar Mohammadi^a, Mohammad Ali Zazouli^c
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- SPE – 15** **EVALUATION OF PHENOL REMOVAL FROM AQUEOUS SOLUTIONS BY AZZOLA**
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- SPE – 16** **SILICA EXTRACTION FROM BAGGASE FLY ASH WITH ALKALI FUSION METHOD**
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Study On Production Process of Biodiesel from Rubber Seed (*Hevea Brasiliensis*) by In Situ Transesterification Method with Alkaline Catalyzed

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Abstract :

Biodiesel is methyl or ethyl fatty acids generated from vegetable oils (edible and non edible) or animal fats. Production of biodiesel from rubber seeds by in situ alkaline catalyzed method from non edible raw materials with a major goal for alternative fuels was studied. The objective of this research is to investigate the influence of reaction time, concentration of alkaline catalyst and ratio raw materials : methanol to the production of biodiesel. The first stage was carried out in order to get reaction time based on the density and viscosity of mixture. In this process, KOH 0.5% (w/v) was used as catalyst with the ratio rubber seed to methanol (1:2). Experiments followed by process with catalyst concentration variation in range 0.1-1% (w/v) and ratio rubber seed to methanol in range 1:1.5-1:3. Research method included, the preparation of samples, biodiesel production, biodiesel separation, and biodiesel characterization include density, viscosity, GC analysis, acid value and Iodine number. The results show that operation time for biodiesel production by using in situ method with alkaline catalyzed was 120 minutes and maximum yield of Fatty Acid Methyl Ester (FAME) was obtained at 52.86%.

Keywords: Biodiesel, rubber seed, in situ, (trans) esterification, Fatty Acid Methyl Ester (FAME).

1. Introduction

Recently, fossil fuel is the source of energy in Indonesia with the largest consumption among other energy sources. Fossil fuel consumption in Indonesia now has reached 363.52 million BOE (Barrels of Oil Equivalent) or approximately 36.41% of the total consumption of energy amounted to 998.53 million BOE. The magnitude of the oil consumption is reverse to the petroleum reserves in Indonesia which were reduced annually. In 2011 total remaining of fossil oil reserves is 7,73 billion barrels, decrease 0.3% from 2010 [1]. To reduce dependence on fossil oil and support the global environment safety program, the only way is to develop eco-friendly alternative fuels. One of the alternative energy is biodiesel.

Biodiesel is a methyl or ethyl esters of fatty acids generated from vegetable oils (edible and non edible) or animal fats. Biodiesel can be produced from various raw materials such as vegetable oils (e.g., soybean, cottonseed, coconut, nuts, canola/rapeseed, sunflower, safflower, coconut, castor), animal and used frying oil [2]. Meanwhile, Indonesia is a country with a largest rubber land area of the world's with vast acreage totalled 3.4 million hectares, with rubber production reached 2.6 million tons in 2009 [3]. On the other side rubber seeds are not much utilized, whereas oil levels contained are quite high 40-50% [4]. Therefore, utilization of rubber seeds as raw material for biodiesel production is very promising.

Currently, biodiesel is produced from Crude Oil Palm Oil (CPO) using methanol and base catalyst. CPO is edible so it can interfering the human needs. Alternative raw materials which not colliding is non edible oil such as canola/rapeseed, sunflower, rubber seed etc. In biodiesel production was occurs esterification or/and transesterification reaction. Esterification is reaction between alcohol with carboxylic acid (free fatty acid). Transesterification is reaction between triglyceride and alcohol with eliminate glycerol which presented in Figure 1. Biodiesel production process can be done by using a homogenous acid catalyst process [5], supercritical process [6], enzymatic process [7], heterogeneous acid catalyst [8] and sonochemical [9].

The biodiesel production from rubber seed oil was using two stage method of esterification-transesterification [4,10]. Biodiesel production process can save cost if use in situ (trans)esterification. In this process, costs for solvent extraction and purification of oil can be reduced, so biodiesel production are becoming more simple [11].

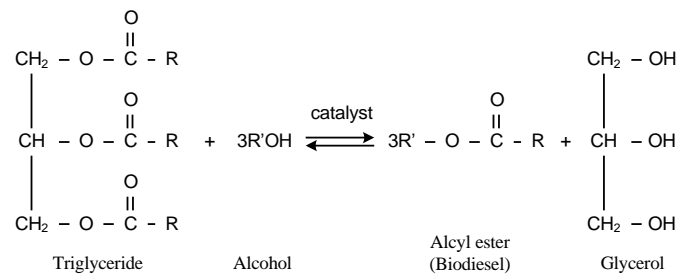


Fig. 1. Transesterification reaction of biodiesel production

In situ Esterification or transesterification is a method to produce biodiesel by contacting raw materials (seed) directly with alcohol/methanol assisted with acid/alkaline catalyst. In the process of in situ transesterification with either an acid or alkaline catalyst, molar ratio of methanol/oil is much higher than the value calculated based on the stoichiometry of the transesterification reaction, for instance, 532:1 [12], 300:1 [13] and 543:1 [14]. Excess methanol will play the role of the extraction solvent [12].

In situ transesterification was introduced by Harrington & Evans [12]. They use sunflower seeds as a raw material. Marinkovic., et.al [13] did experiments with the same process. Ozgul and Selma [15,16] used in situ esterification with rice bran as raw materials and ethanol and methanol as solvent and so Ginting., et.al [17] did in situ transesterification from castor seed (*Jatropha curcas*). The objective of this research will produce biodiesel from rubber seeds with in situ process and study influence of reaction time, ratio rubber seed to methanol and alkaline catalyst concentration.

2. Materials and methods

2.1. Materials

The rubber seed was obtained from Perkebunan Karet Kendal Indonesia. Methanol that was used in the experiment is technical grade, while KOH as catalyst is analytical grade (Merck, Germany).

2.2. Experimental procedure

Rubber seeds samples were peeled. Seed contents was macerated, blended and dried in an oven at temperature of 55°C for 2 hours. One hundred grams samples were intake to bottle equipped with mixer (see Fig. 2). Methanol solution and KOH were added and then heated and mixed. The heating of solution in 60°C temperature at atmospheric pressure. In situ processing (extraction reaction process) did until 120 minutes. The product was filtered and methanol was separated with distillation. Biodiesel product was analyzed for weight, viscosity, density and concentration. Concentration of methyl ester in biodiesel product was analyzed by GC.

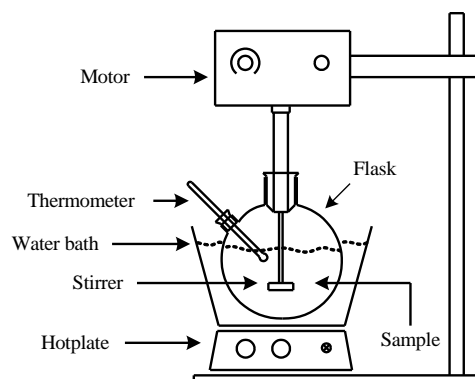


Fig. 2. Experiment tool sets

The experiment was divided into three stages. The first stage (preliminary study) was carried out to obtain reaction time for in situ processing. In this step, response variable was analyzed on density and viscosity. The experiment was done in condition of catalyst concentration of KOH 0.5% (w/v) and ratio of rubber seed to methanol (1:2) and every 15 minute was analyzed until constants condition. In second stage, experiment was investigated with catalyst concentration of KOH about 0.1; 0.25; 0.5; 0.75 and 1% (w/v). The experiments was done in variable ratio of rubber seed to methanol (1:2) and reaction time 120 minutes. In third stage, experiment conducted about effect of ratio rubber seed to methanol ie; 1:1.5; 1:1.75; 1:2; 1:2.5 and 1:3 (w:v).

Weight of product was used to calculated about biodiesel yield. Yield of FAME calculated using this equation [17,18]:

$$\text{Yield of FAME} = \frac{\text{actual weight of biodiesel (g)}}{\text{theoretical weight of biodiesel (g)}} \times 100\%$$

3. Results and discussion

3.1. Preliminary study

Figure 3 shows density and viscosity versus reaction time. Density and viscosity are physics properties that depend with composition in solution. (Trans)esterification reaction was produced methyl ester and glycerol. So, the in situ process can be obtained methyl ester, glycerol, and vegetable oil as products. Figure 3 shows that increasing time followed increasing the density and viscosity of product reaction. Product has constant density and viscosity at 120 minutes. It shows which biodiesel production and extraction of oil have been formed completely. Thus from this stage it was obtained 120 minutes as a maximum reaction time, which would be a reference time for the second and third stages.

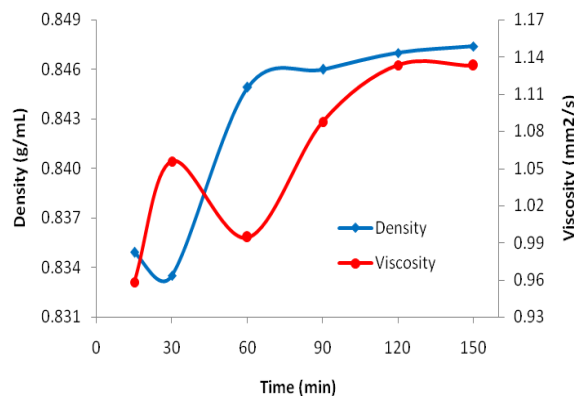


Fig. 3. Graph of time versus viscosity and density of biodiesel

3.2. Effect of catalyst concentration

In this stage, experiment was conducted to investigate the influence of catalyst concentration (KOH). Concentration of KOH was varied in range 0.1-1% (w/v). The results of these experiments are shown at Figure 4 & 5. Maximum yield of oil obtained at KOH concentration was 0.75% (w/v) with value 12.53% and the lowest on the concentration of KOH 0.5% (w/v) with 6.52%. According to Ramadhas et al., [4] and Ketaren, [19] the rubber seeds contain 40-50% of the oil. When it is compared with the results of this study, it was too small. This is because the low quality of rubber seeds and it was expected that the oil contained therein was small. Rubber seeds will quickly rots if its skin already peeled, but this can be avoid with dry it quickly and samples to be more durable and can be stored in a long time. As mentioned by Ramadhas., et al. [4] from physical appearance the higher catalyst concentration, the darker also biodiesel produced. Therefore the addition of a suitable catalyst concentration is important to the physical appearance of biodiesel.

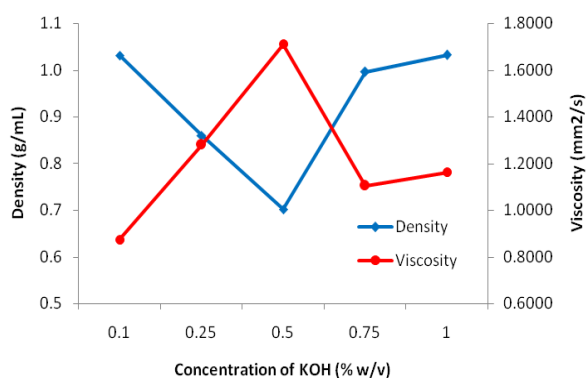


Fig. 4. Effect of catalyst concentration (%) on density & viscosity of biodiesel

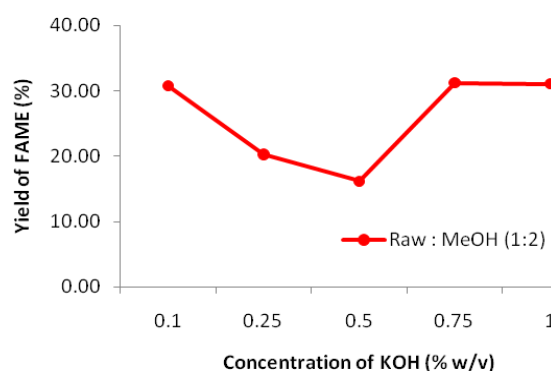


Fig. 5. Effect of catalyst concentration to yield of FAME

Figure 4 is a graph of the influence of catalyst concentration of KOH to biodiesel density and viscosity. Density of biodiesel was obtained the lowest value on KOH 0.5% (w/v) at 0.7014 g/mL, and the highest on KOH 1% (w/v) at 1.0323 g/mL. Density of biodiesel according SNI 04-7182-2006 is 0.85-0.90 g/mL. Density values obtained in this experiment is different from standard biodiesel. This is because the impurity still present in the product range. The product is the result of the extraction reaction, so in addition to biodiesel slightly even to the other products such as resin. Biodiesel product that obtained with catalyst concentration KOH 0.75% (w/v) is 0.9957 g/mL which slightly higher with biodiesel standard. The density of fuel has some effect on the break up of the fuel injected into the cylinder. In addition, more fuel is injected by mass as the fuel density increases [20]. The injected fuel amount, injection timing and injection spray pattern are directly affected by these parameters [21]. With increasing density, the diameter of the fuel droplets increases. Since the inertia of the big droplets is big, their penetrations in the combustion chamber will be higher, as well [22]. As fuel with lower density and viscosity is injected, it will improved atomization and better mixture formation can be attained [20].

The viscosity of an engine fuel is one of the most critical fuel features. It plays a dominant role in the fuel spray, mixture formation and combustion process. The high viscosity interferes with the injection process and leads to insufficient fuel atomization. Moreover, the mean diameter of the fuel droplets from the injector and their penetration increases with increasing fuel viscosity [22]. The inefficient mixing of fuel with air contributes to incomplete combustion in the engine. In addition to all these, high viscosity can cause early injection due to high line pressure, which moves the combustion of the fuel closer to top dead center, increasing the maximum pressure and temperature in the combustion chamber [21, 22].

Viscosity of any fuel is related to its chemical structure. Viscosity increases with the increase in the chain length and decreases with the increase in the number of double bonds (unsaturation level) [23, 24, 25]. The viscosity increases with increasing concentrations of KOH up to 0.5% (w/v) (as shown in Figure 4) become the highest viscosity 1.7108 mm²/s whereas the lowest viscosity 0.8729 mm²/s at KOH 0.1% (w/v). Overall value of biodiesel viscosity (40°C) is still under the standard value of the SNI 04-7182-2006 with range at 2.3-6.0 mm²/s, it means more dilute than biodiesel standard and value its worth even closer with a solar standard viscosity 1.6-5.8 mm²/s range at SNI so that this becomes no problem. In spite of, low viscosity causes rapid wear of engine parts such as injection pump and fuel injector as expressed by Chigier [26]. Viscosity of methyl ester from rubber seed oil is lower at higher temperatures and almost equal to the diesel fuel. This helps the combustion as air entrainment increases, as spray cone angle increases due to reduction in viscosity [27].

Figure 5 shows that variation of catalyst concentration (KOH) effects on yield of FAME. The highest yield of FAME was obtained when in situ process did in KOH concentration 0.75% (w/v) with value of yield of FAME is 31.18%. Yield of FAME which was produced wasn't high because in this process used ratio of raw materials to methanol (1:2). Methanol used for solvent and reactant to small while compared with experiments that did by the other researcher [12-14]. Methanol used in situ process is very small in quantity, so the yield of oil and yield of FAME obtained also slightly small. It was supposed that less of methanol extraction and transesterification the oil into biodiesel. When the experiment was used ratio of raw materials to methanol (1:3), it was obtained maximum yield of FAME 52.86% as presented in Figure 7.

3.3. Effect of ratio rubber seed to methanol

The effect of ratio rubber seed to methanol was presented in Figure 6 and 7. Increasing ratio rubber seed to methanol can affect to amount of methanol that used to in situ processing. Methanol that used increase can cause density of biodiesel increase. Density of biodiesel was obtained the highest value 1.0034 g/mL on the ratio rubber seed to methanol (1:2.5) and the lowest value 0.9129 g/mL on the ratio rubber seed to methanol (1: 1.5). When compared with standard biodiesel SNI 04-7182-2006, density of biodiesel is 0.85-0.90 g/mL, then the overall value that near to the value of standard is at ratio of raw materials to methanol (1:1.5) until (1:2) and (1:3) Methanol in layers of esters can lower the flash point of biodiesel [4].

Viscosity biodiesel has tends to constant in the ratio rubber seed to methanol (1:1,5) until (1:75) from 1.2049 mm²/s to 1.2596 mm²/s but at ratio of raw materials to methanol (1:2) decrease to 1.1048 mm²/s which still lower if compared to solar standard with viscosity range at 1.6-5.8 mm²/s on SNI 04-7182-2006. At ratio rubber seed to methanol (1:3), the viscosity was obtained at 1.1035 mm²/s which is still lower than solar standard at SNI and this is not a problem because as in the previous discussion, when a fuel with lower density and viscosity is injected, improved atomization and better mixture formation can be attained [20].

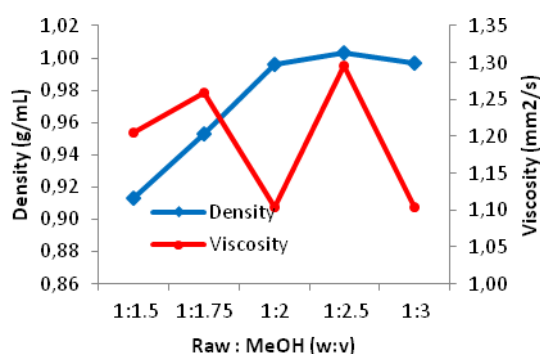


Fig. 6. Effect of raw materials to methanol on density & viscosity of biodiesel

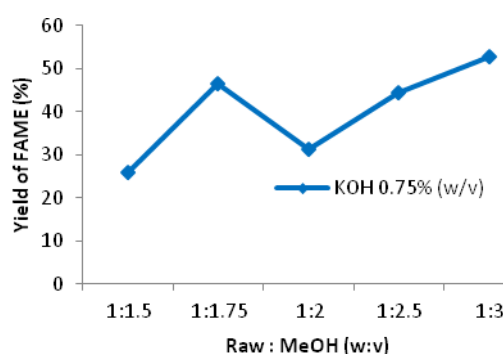


Fig. 7. Effect of raw materials to methanol on yield of FAME

As expressed by Özgül-Yücel and Türkay [16], the role of methanol in the in situ process is very important because the methanol has a dual role: as a solvent for oil extraction as well as reactant, so the addition should be an excess. Figure 7 shows that yield of FAME increase significantly at the ratio rubber seed to methanol (1:2,5) and (1:3) by 44.55% and 52.86% respectively. The amount of methanol, however, should not be too exaggerated, as expressed by Ramadhas., et al [4], methanol on ester layers can lower the flash point of biodiesel. Therefore the purification process and removal of methanol with distillation or washing should be perfect. Distillation process should be concerned (± 1 hour) because if it is too long causing a condensed biodiesel and hard to pour it at room temperature.

Acid number (acid value) is the mass of potassium hydroxide (KOH) in milligrams that is required to neutralize one gram of chemical substance. The acid number is a measure of the amount of carboxylic acid groups in a chemical compound, such as a fatty acid, or in a mixture of compounds. Acid value on biodiesel product are still quite high with the lowest value 13.73 mg KOH/g biodiesel at concentrations of KOH 0.1% (w/v) and the highest value 71.46 mg KOH/g biodiesel at concentration of KOH 0.25% (w/v) like presented in Figure 8. Figure 8 shows that acid value decrease starting from concentration of KOH 0.25 until 1% (w/v) and also from ratio of raw materials to methanol (1:1.5) until (1:3) in Figure 9. This shows that there are still many free fatty acids that have not been converted into methyl ester. Free fatty acids obtained from the extraction process. Thus the extraction process is more dominant compared to the transesterification reaction. Figure 9 shows that with increasing methanol, then KOH will more and more causing decline in acid number. As for the reaction to form methyl ester have been low. High acid value not only will make deposits in the fuel system but also decrease the quality components in the fuel system [28]. So, to increasing quality of biodiesel or decreasing of acid number, product of in situ process must be followed by esterification processing.

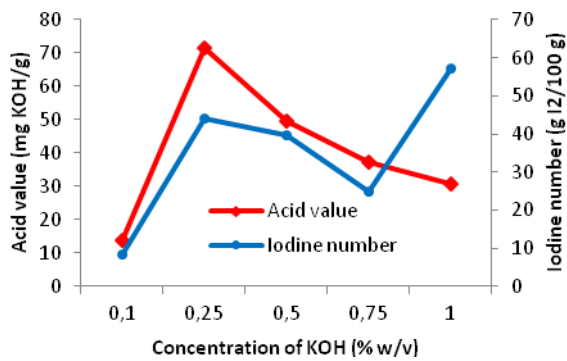


Fig. 8. Effect of catalyst concentration to acid number & Iodine number

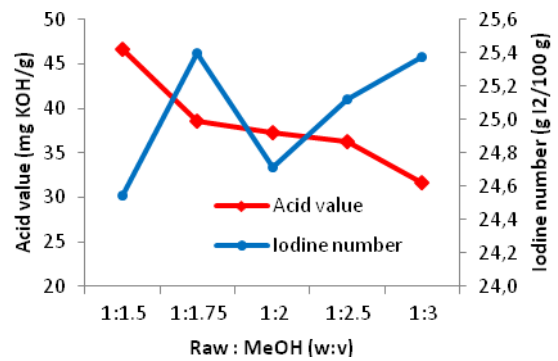


Fig. 9. Effect of ratio rubber seed to methanol on acid number & Iodine number

Iodine number is mass of iodine in grams that is consumed by 100 grams of a chemical substance. Iodine numbers are often used to determine the amount of unsaturation in fatty acids. This unsaturation is in the form of double bonds, which react with iodine compounds. The higher iodine number, the more C=C bonds are present in the fat [29]. Iodine number was relatively stable at ratio variations of raw materials to methanol as seen in Figure 9. In contrast to the Figure 8, Iodine number varies at different concentrations of catalyst. At the concentration of KOH 0.75% (w/v) had an Iodine number 24.72 g I₂/100 g biodiesel and at ratio of raw materials to methanol (1:3) had an Iodine number 25.38 g I₂/100 g biodiesel which means the unsaturation of biodiesel was relatively small and quite well and all of the Iodine numbers at all variations was still under the SNI 04-7182-2006 standard with a maximum value of 115 g I₂/100 g biodiesel.

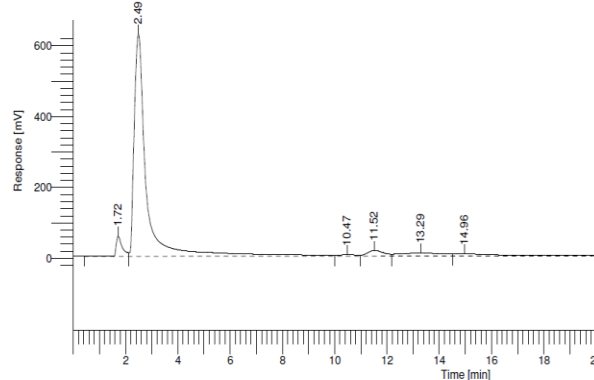


Fig. 10. GC chromatogram of biodiesel on the ratio raw materials to methanol (1:3) (w:v)

Chromatogram of Fatty Acid Methyl Ester (FAME) was detected at 10-14 retention time minutes and it was appropriate with the standard biodiesel chromatogram with FAME in minutes 6-14 retention time. When compared to standard, the expected results of biodiesel analysis with GC 10.47 minutes was methyl nonadecanoic (C_{19:0}), 11.52 minutes was methyl linolenic (C_{18:3}), 13.29 minutes was methyl eicosadienoic (C_{20:2}) and 14.96 minutes was methyl timnodonic (C_{20:5}). Although all of its peak wasn't obvious, but this shows that the FAME was indeed formed in biodiesel sample and the oil/triglyceride wasn't detected again.

4. Conclusion

In situ process that involve transesterification reaction of biodiesel production from rubber seeds with in situ method takes 120 minutes at 60°C with concentration of KOH 0.75% (w/v) was obtained maximum yield of FAME 31.18% and at ratio of raw materials to methanol (1:3) was obtained maximum yield of FAME 52.86%. Based on the results, ratio of raw materials to methanol is quite important to increase yield of FAME significantly.

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