## Biodiesel Synthesis from Coconut Oil Using Calcined Scallop Shell Waste as the Heterogeneous Catalysts

ARIF Hidayat<sup>1,a\*</sup>, NADIFAH Syachbana Sakariel<sup>2,b</sup> and LIDYA Afifa Tambunan<sup>3,c</sup>

<sup>1</sup>Chemical Engineering Department, Universitas Islam Indonesia, Indonesia
<sup>2</sup>Chemical Engineering Department, Universitas Islam Indonesia, Indonesia
<sup>3</sup>Chemical Engineering Department, Universitas Islam Indonesia, Indonesia
<sup>a</sup>arif.hidayat@uii.ac.id, <sup>b</sup>18521185@uii.ac.id, <sup>c</sup>18521188@uii.ac.id

Keywords: scallop shell, heterogeneous, catalyst, microwave, biodiesel, coconut oil.

**Abstract.** Several efforts have been performed to make the biodiesel price more competitive with fossil fuels, such as using low price raw materials, including coconut oil and the use of heterogenous catalyst. In this research, the Calcined Scallop Shell was applied as a heterogeneous catalyst for synthesizing biodiesel from coconut oil. The catalyst was obtained from calcination of scallop shell waste. The catalytic activities of Calcined Scallop Shell catalyst during transesterification reaction was influenced by several reaction conditions including methanol to oil mole ratio, reaction temperature, and concentration of catalyst. The biodiesel yield of 91.7% was obtained at a methanol to coconut oil mass ratio of 12:1, 60 °C reaction temperature, and catalyst concentration of 10% wt. of oil.

## Introduction

Efforts to find substitute renewable energy sources are continuing to repace fuels derived from fossil fuels. The limited reserves of fossil energy in the world and even Indonesia, triggering the world's attention on renewable energy needs is increasing. One of the renewable energy sources currently being studied is biodiesel. Biodiesel as a monoalkyl ester is an alternative as a substitute fuel from diesel because it has the same physical properties. Generally, monoalkyl esters from plant oils and animal fats are used to synthesize biodiesel. Vegetable oils and animal fats, as well as their derivatives, have the potential to replace diesel fuel.

Biodiesel is generally obtained from the synthesis process that mainly encountered in renewable resources which are transesterification of triglycerides or esterification of free fatty acids. To speed up the process and produce a high transformation of methyl esters, a base catalyst or an acid catalyst is added to the solution. Two types of conventional catalysts applied in biodiesel synthesis include homogeneous acids catalysts (including hydrochloric and sulfuric acids) and homogeneous alkaline catalysts (including potassium hydroxide, sodium hydroxide, and alkali metal hydroxides). The subsequent catalysts have a number of disadvantages, including being consumable, caustic, as well as being required to advanced dissociation for product separation, which causes major environmental issues. Hence, studies on heterogeneous catalysts such as low energy consumption, reusable, ease of product separation, and less separation process.

NaOH or KOH are widely used as homogeneous catalyst in the manufacture of biofuels from edible oils. However, homogeneous base catalyst have significant drawbacks, such as an advanced separation process following the reaction phase, which generates additional effluent and cannot be reused. Several investigations on the efficiency of heterogeneous catalysts in biodiesel synthesis have been investigated for many years. As a result, numerous solid catalysts such as hydrotalcite [1], metal oxides [7, 8], zeolites [9, 10], alumina [11, 12], clays [13, 14], and biomass-based catalysts [15-19] have been investigated utilizing various raw materials. The goal of the research was to develop a heterogeneous catalyst with high catalytic activities for the biodiesel synthesis. The

transesterifications of vegetable oils to produce biodiesel which involved solid catalysts had investigated by many researchers. The various type of solid catalysts such as zeolite, clay, hydrotalcite, ion exchange resin, metal oxides, biomass wastes, zirconia, heteropolyacids, activated carbon, silica, tantalum pentoxide, etc. had been studied.

Recent years, the use of catalysts from biomass waste has intensively studied for the biodiesel production. An alternative material that has not been fully explored is the solid waste of shellfish. The solid waste of shellfish is one of the potential candidates to be used as a base catalyst because it has a high calcium oxide (CaO) content. From economic viability, the use of shellfish shells has many advantages, such as widely available, the price is low, the high content of CaCO<sub>3</sub>, and stable under acidic and basic conditions. Several researchers revealed that solid-based catalysts showed comparable performance to some homogeneous catalysts when applied to biodiesel production. In this paper, scallop shell waste (*Placuna placenta*) is utilized as a raw material that contains CaO for producing biodiesel from coconut oil. Indonesia with a large sea area has an abundant amount of shellfish production. Processing this seafood will produce a huge amount of shellfish waste. So far, scallop shells have only been used to make handicrafts. The content of CaO in scallop shells can be used in the biodiesel synthesis as a base catalyst. As a catalyst, the utilization of scallop shell waste material is effective to minimize the cost of biodiesel products and providing an alternative for recycling natural mineral resources. The use of scallop shell as a catalyst has many advantages i.e., stable under acidic and basic conditions, available abundantly, cheap, and environmentally friendly.

From the literature search results, there are not many reports on the use of scallop shell waste as a new source of solid catalysts for the biodiesel synthesis. In this study the transesterifications of coconut oil using calcined scallop shell as solid catalyst was performed. The catalyst activity was assessed focusing on methanol to oil mole ratio, reaction temperature, and concentration of catalyst. The reusability of catalyst was also studied in this work.

#### Methodology

**Materials.** Coconut oil was obtained from a local market in Sleman Province, Yogyakarta, Indonesia, the scallop shell waste was collected from a seafood restaurant in Bantul Province, Yogyakarta, Indonesia. Methanol p.a. was purchased from Merck.

**Catalyst Preparation.** To ensure consistent size, the scallop shell was screened into 200 mesh. The shells were then calcined for 4 hours at 900°C in a tube furnace.

**Catalyst Performance.** The transesterification reaction of coconut oil was conducted in a batch reactor comprising a condenser, thermometer, magnetic stirrer, and a boiling flask in this study. The reactions processes were carried out in the following order: (i) coconut oil was combined with methanol before adding scallop shell that has been calcined; (ii) when the temperature was arise, the solution was mixed vigorously; (iii), the reaction product was separated from the catalyst by centrifugation after the process was finished; (iv) perform distillation to remove remaining methanol; and (v) collect biodiesel product using a separatory funnel. The the catalyst weight was varied from 1 to 5 wt.%. while the mass ratio of oil to methanol was varied from 6 to 2.

#### **Results and Discussion**

**Temperature Effect on Biodiesel Yield.** The activity test of calcined scallop shell catalyst was examined in the transesterification reactions of coconut oil with methanol. The influence of reaction temperature on biodiesel yield is depicted in Figure 1. When utilizing the calcined scallop shell catalyst, the biodiesel yield was 70.4 % and increased to 86.3% using reaction temperature of 35 - 55°C in 2 hours and a 12:1 methanol to oil mole ratio, respectively. The highest biodiesel yield of 91.7% was obtained at a 65 °C of reaction temperature.

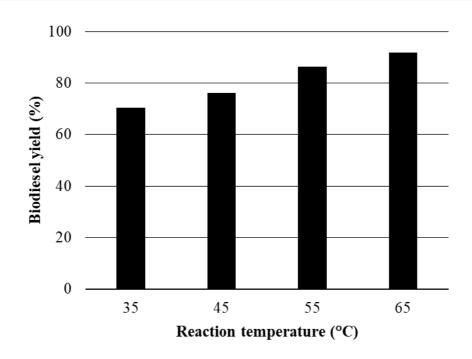


Fig. 1 Temperature Effect on Biodiesel Yield.

**Methanol to Oil Mole Ratio Effect**. The methanol to oil mole ratio is one of the most important elements that determines biodiesel yield. The mole ratio is the proportion of methanol mol numbers to methanol oil mol numbers. The mole ratio number was used extensively to direct the process to the biodiesel product synthesis. The catalytic actions of the calcined scallop shell catalyst utilized in the transesterification process, as shown in Figure 2, show biodiesel yields of 70.1%, 83.5%, 87.6%, and 91.7%, respectively, at a reaction time of 2 hours with methanol to oil mole ratio 3:1, 6:1, 9:1, and 12:1 at 60°C reaction temperature.

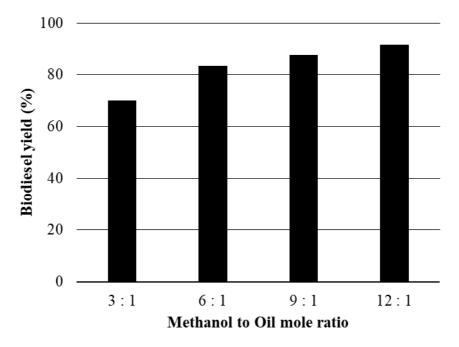


Fig. 2 Methanol to Oil Mole Ratio Effect on Biodiesel Yield.

**Catalyst concentration effect**. During the transesterification process, the catalyst concentration has an impact on the biodiesel synthesis. The amount of calcined scallop shell catalyst is changed between 1 and 10% by weight of oil. As seen in Figure 3, the increase in catalyst concentration increases the biodiesel yield. At 10% wt. of oil, the biodiesel yield achieves maximum. When the

calcined scallop shell catalyst was used in the transesterification reaction, comparable results were achieved. When the amount of calcined scallop shell catalyst is loaded from 1 to 10% wt. of oil, the yield of biodiesel increase.

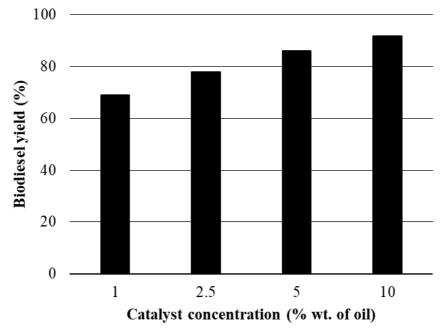


Fig. 3 Catalyst Concentration Effect on Biodiesel Yield.

**Reusability**. The Calcined Scallop Shell catalyst was employed for the transesterification process several times to assess its recyclability. The catalyst was then filtered before being utilized in a fresh reaction cycle with the required processing stages, including washing and drying. The findings for the catalyst utilized in three cycles are shown in Figure 4. The transesterification reaction was conducted under the following conditions: catalyst concentration of 10% wt. of oil, 60 °C reaction temperatures for 2 hours period, and methanol to oil mole ratio of 12:1. The activity was smaller in the 2nd and 3rd cycles compared to the 1st cycle, as seen in Figure 4. After the 3rd cycle, the catalytic activity dropped by up to 40%.

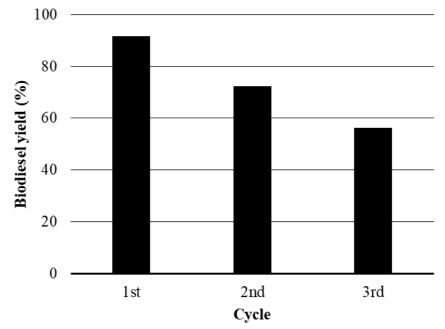


Fig. 4 The Reusability of Catalyst.

**Biodiesel characterization**. Figure 5 displays the FTIR spectrum for the biodiesel from coconut oil. The FTIR spectra was examined in the 600 cm<sup>-1</sup> to 3500 cm<sup>-1</sup> region. FTIR is an instrument that can supply information of compound that has functional group. As demonstrated in Figure 5, the existence of biodiesel is represented by CH<sub>3</sub> asymmetric bending about 1425–1500 cm<sup>-1</sup> and O-CH<sub>3</sub> extending around 1100–1200 cm<sup>-1</sup>. The C=O bending of the ester group in biodiesel is represented by a wavenumber approximately 1700 to 1800 cm<sup>-1</sup>. The occurrence of an ester group about 2800–3000 cm<sup>-1</sup> can be seen by looking at the CH peak. The =C–H stretching, C–H bending, and the C–H bond stretching are indicated by wave numbers of 675–1000 cm<sup>-1</sup>, 1350–1480 cm<sup>-1</sup>, 2800–3000 cm<sup>-1</sup> to 1651 cm<sup>-1</sup>. The existence of an alcohol group and an aromatic group is shown by the peaks at 3000–3100 cm<sup>-1</sup> and 3470 cm<sup>-1</sup>.

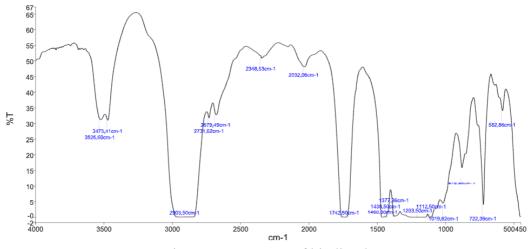


Fig. 5 FTIR spectra of biodiesel.

#### Conclusions

In this study, biodiesel synthesis from coconut oil was investigated using a Calcined Scallop Shell Waste as a solid catalyst. The biodiesel yield of 91.7% was obtained at a 65 °C reaction temperature, methanol to coconut oil mass ratio of 12:1, and catalyst concentration of 10% wt. of oil.

#### References

[1] J. Nowicki, J. Lach, M. Organek, E. Sabura, Transesterification of rapeseed oil to biodiesel over Zr-dopped MgAl hydrotalcites, Appl. Catal. A: Gen. 524 (2016) 17-24.

[2] M. Sirajuddin, M. Tariq, S. Ali, Organotin (IV) carboxylates as an effective catalyst for the conversion of corn oil into biodiesel, J. Organomet. Chem. 779 (2015) 30-38.

[3] E. Dahdah, J. Estephane, R. Haydar, Y. Youssef, B. El Khoury, C. Gennequin, A. Aboukais, E. Abi-Aad, S. Aouada, Biodiesel production from refined sunflower oil over Ca–Mg–Al catalysts: Effect of the composition and the thermal treatment, Renew. Energy 146 (2020) 1242-1248.

[4] L. Wu, T.Y. Wei, Z.F. Tong, Y. Zou, Z.J. Lin, J.H. Sun, Bentonite-enhanced biodiesel production by NaOH-catalyzed transesterification of soybean oil with methanol, Fuel Process. Technol. 144 (2016) 334-340.

[5] Y.S. Pradana, A. Hidayat, A. Prasetya, A. Budiman, Biodiesel production in a reactive distillation column catalyzed by heterogeneous potassium catalyst, Energy Procedia 143 (2017) 742-747.

[6] T. Qiu, X. Guo, J. Yang, L. Zhou, L. Li, H. Wang, Y. Niu, The synthesis of biodiesel from coconut oil using novel Brønsted acidic ionic liquid as green catalyst, Chem. Eng. J. 296 (2016) 71-78.

[7] A. Hidayat, N.I.F. Mukti, B. Handoko, B. Sutrisno, Biodiesel production from rice bran oil over modified natural zeolite catalyst, Int. J. Technol. 9 (2018) 400-411.

[8] A. Hidayat, B. Sutrisno, Free fatty acids esterification on palm oil sludge using zirconia-supported Indonesian natural zeolite as heterogeneous catalyst, Orient. J. Chem. 34 (2018) 2464-2470.

[9] M. Hajek, J. Kocik, K. Frolich, A. Vavra, Mg-Fe mixed oxides and their rehydrated mixed oxides as catalysts for transesterification, J. Clean. Prod. 161 (2017) 1423-1431.

[10] C.S. Castro, L.C.F. Garcia Jr, J.M. Assaf, The enhanced activity of Ca/MgAl mixed oxide for transesterification, Fuel Process. Technol. 125 (2014) 73-78.

[11] N.A. Negm, G.H. Sayed, F.Z. Yehia, O.I. Habib, E.A. Mohamed, Biodiesel production from one-step heterogeneous catalyzed process of Castor oil and Jatropha oil using novel sulphonated phenyl silane montmorillonite catalyst, J. Mol. Liq. 234 (2017) 157-163.

[12] L. Zatta, L.P. Ramos, F. Wypych, Acid-activated montmorillonites as heterogeneous catalysts for the esterification of lauric acid acid with methanol, Appl. Clay Sci. 80-81 (2013) 236-244.

[13] E.G. Silveira Junior, V.H. Perez, I. Reyero, A. Serrano-Lotina, O.R. Justo, Biodiesel production from heterogeneous catalysts based  $K_2CO_3$  supported on extruded  $\gamma$ -Al<sub>2</sub>O<sub>3</sub>, Fuel 241 (2019) 311-318.

[14] A. Ramesh, K. Palanichamy, P. Tamizhdurai, S. Umasankar, K. Sureshkumar, K. Shanthi, Sulphated Zr–Al<sub>2</sub>O<sub>3</sub> catalysts through jatropha oil to green-diesel production, Mater. Lett. 238 (2019) 62-65.

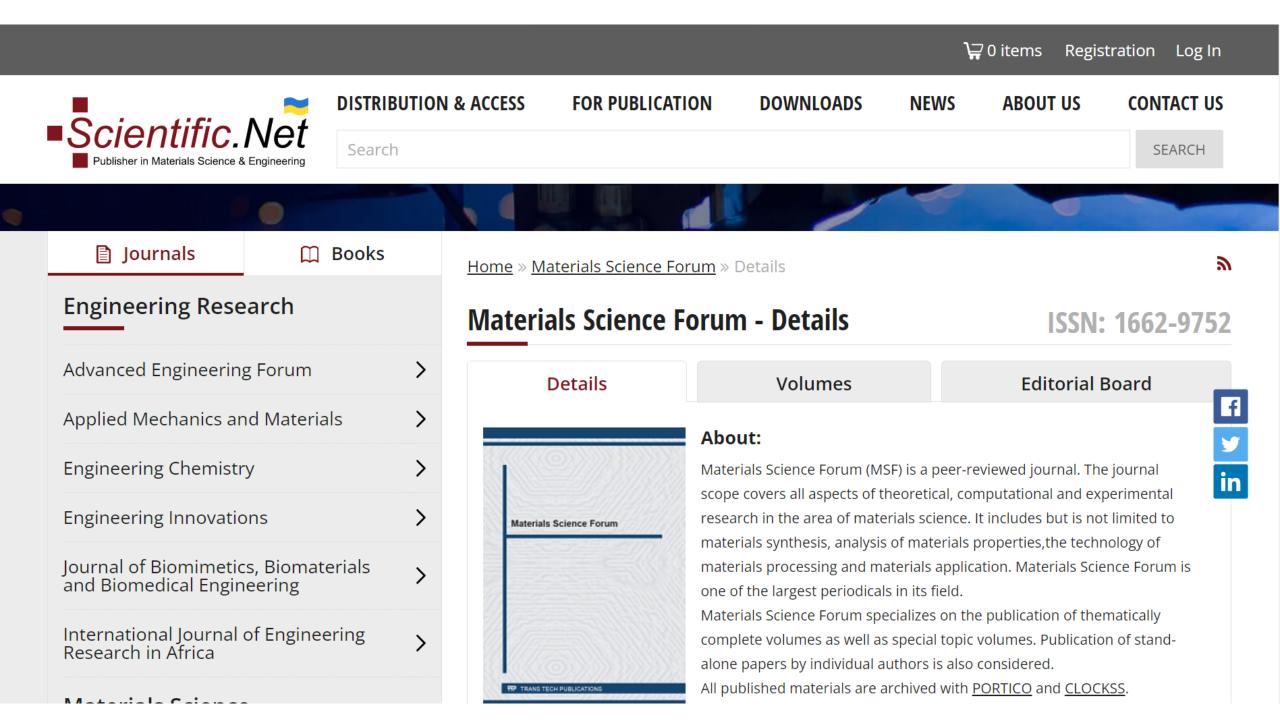
[15] A. Hidayat, Rochmadi, K. Wijaya, A. Budiman, Reaction kinetics of free fatty acids esterification in palm fatty acid distillate using coconut shell biochar sulfonated catalyst, AIP Conf. Proc., 1840 (2017), 050001.

[16] V.A.S. Augustia, R.A. Djalal, B. Sutrisno, A. Hidayat, Kinetic study of free fatty acid in Palm Fatty Acid Distillate (PFAD) over sugarcane bagasse catalyst, IOP Conf. Ser. Earth Environ. Sci., 105 (1) (2018), 012065.

[17] A. Hidayat, B. Sutrisno, Esterification free fatty acid in sludge palm oil using  $ZrO_2/SO_4^{2-}$  - Rice husk ash catalyst, AIP Conf. Proc. 1840 (2017) 050001.

[18] M.S.A. Farabi, M.L. Ibrahim, U. Rashid, Y.H. Taufiq-Yap, Esterification of palm fatty acid distillate using sulfonated carbon-based catalyst derived from palm kernel shell and bamboo, Energy Convers. Manag. 181 (2019) 562-570.

[19] G.Y. Chen, R. Shan, J.F. Shi, B.B. Yan, Transesterification of palm oil to biodiesel using rice husk ash-based catalysts, Fuel Process. Technol. 133 (2015) 8-13.



0 items

Search

# Volumes

# **Materials Science Forum Vol. 1073**

DOI:	<u> https://doi.org/10.4028/v-0788xf</u>	
ToC:	Table of Contents	
Search		
<b>1</b> 2 3 >		
Paper Title		Page
<u>Preface</u>		
Solution Authors: Kumar P. Chandra Choudhary, Kamal Prasad	7 <mark>Ca<sub>0.3</sub>TiO<sub>3</sub>-BaZr<sub>0.2</sub>Ti<sub>0.8</sub>O<sub>3</sub> Solid a, Janki N. Singh, Ajit R. Kulkarni, R.N.P. structure, Fourier transformed infrared generated energy harvesting</mark>	3
Nano Al <sub>2</sub> O <sub>3</sub> / Micro ZrO <sub>2</sub> Pa Study of Mechanical and T Authors: K. Sekar Abstract: AA7050 aluminum al	orid Composite Reinforced with Articles by Stir Casting Method and Tribological Properties loy with desirable mechanical properties resistance, and toughness widely used in	11

Application of Friction Stir Spot Welding on Additive Manufactured Carbon Fiber Composite Authors: Muhammad Farhan Raza, Guney Guven Yapici Abstract: Friction stir spot welding as a solid state processing technique is used to join dissimilar and similar metals, composites and more	23
Analysis of Mass Concentration and Morphology of Fume Particles during ECDM of CFRP Composites Authors: Santosh Kumar Yadav, Abhishek Singh, Kishore Debnath Abstract: Electrochemical discharge machining (ECDM) is a hybrid method used to generate micro-features in hard and brittle materials more	29
Wear Optimization of Aluminium and Hybrid Reinforcement Metal Matrix Composites Using Response Surface Methodology Authors: S. Muzeer, S. Sivaganesan Abstract: Aluminum 6061 alloy-based alloys were used to make various motor vehicle parts such as connecting wire, O-ring, circular blocks, more	37
Antimicrobial Biodegradable Polymeric Materials for Food Packaging Applications: Current Status and Future Directions Authors: S. Achutha, Elizabath Johnson, S. Kumari Nisha, Sarika Sivakumar Abstract: The production and consumption of non-biodegradable, petroleum-based plastics had an enormous increase in the past few more	49
Investigation on Mechanical Properties of Regular and Engineered Fiber Built up Polymer Composites Authors: Makarand B. Shirke, Santosh N. Shelke Abstract: Standard fibers have attracted the attentiveness of technocrats, specialists, experts, and researchers throughout the globe more	57
<u>Design and Analysis of Electric Bike Chassis Using Glass Fibre</u> <u>Composites</u> Authors: L.S. Sudharson, Prabakaran, C.A. Arjun	67

Abstract: Electric motorcycles and scooters are alternates for the existing Internal combustion engine which eliminates emissions and ...more

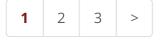
<u>Shear Thickening Fluids, Nano-Polymer Materials and their</u> <u>Application Methods for Textile Substrates</u>

## Authors: Gurumurthy Ramaiah, Dani Asfaw, Seblework Mekonnen, Welamu Tesfay, Eshetu Solomon

Abstract: Shear Thickening Fluid (STF) is a highly preferred phase change material that helps in absorbing high impact shock waves and

<u>...more</u>

Showing 1 to 10 of 22 Paper Titles



**DISTRIBUTION & ACCESS** 

FOR PUBLICATION

INSIGHTS

**DOWNLOADS** 

ABOUT US

**POLICY & ETHICS** 

**CONTACT US** 

IMPRINT

**PRIVACY POLICY** 

SITEMAP

ALL CONFERENCES

**ALL SPECIAL ISSUES** 

**ALL NEWS** 

Scientific.Net is a registered brand of Trans Tech Publications Ltd © 2023 by Trans Tech Publications Ltd. All Rights Reserved 81

0 items

Search
Journals
Engineering Research
Advanced Engineering Forum
Applied Mechanics and Materials
Engineering Chemistry
Engineering Innovations
Journal of Biomimetics, Biomaterials and Biomedical Engineering
International Journal of Engineering Research in Africa
Materials Science
Advanced Materials Research
Defect and Diffusion Forum
Diffusion Foundations and Materials Applications
Journal of Metastable and Nanocrystalline Materials
Journal of Nano Research
Key Engineering Materials
Materials Science Forum

Nano Hybrids and Composites
Solid State Phenomena
Engineering Series
Advances in Science and Technology
Construction Technologies and Architecture
Books
Special Book Collections
Foundations of Materials Science and Engineering
Scientific Books Collection
Specialized Collections
Retrospective Collection

# **Materials Science Forum - Editorial Board**

ISSN: 1662-9752

Volumes	
Editorial Board	

Founding Editor

## Fred H. Wohlbier

Honorary Editor

## Prof. Graeme E. Murch

University of Newcastle, Centre for Mass and Thermal Transport in Engineering Materials, School of Engineering; Callaghan, Australia, NSW 2308;

Editor(s) in Chief

### Prof. Iulian Antoniac

University Politehnica of Bucharest, Faculty of Materials Science and Engineering; 313 Splaiul Independentei, Bucharest, 060042, Romania;

#### Prof. Guillermo Requena

#### ORCID

German Aerospace Center (DLR), Institute of Materials Research ; Köln, DE-51170, Germany;

**Editorial Board** 

#### Prof. Dezső L. Beke

University of Debrecen, Department for Solid State Physics; Bem tér 18/b, Debrecen, 4026, Hungary;

\_\_\_\_\_

#### **Prof. Giorgio Benedek**

University of Milano Bicocca, Department of Materials Science; U5, Universitá di Milano-Bicocca, Via R. Cozzi 55, Milano, 20125, Italy;

## Dr. Giacomo Benvenuti

<u>ORCID</u>

ABCD Technology; Switzerland;

#### Prof. Anil K. Bhatnagar

University of Hyderabad, School of Physics and School of Engineering; Hyderabad, India, 500046;

\_\_\_\_\_

### **Prof. Chi Ming Chan**

Hong Kong University of Science and Technology, Department of Chemical and Biomolecular Engineering, Clear Water Bay; Kowloon, China;

#### Dr. Abel S. Fenta

<u>ORCID</u>

CERN; Genève, CH-1211, Switzerland;

#### **Roberto B. Figueiredo**

Federal University of Minas Gerais, Department of Metallurgical and Materials Engineering; Belo Horizonte, MG, 30, Brazil, 31270-901;

### Prof. Hermann G. Grimmeiss

Lund University, Department of Solid State Physics; Box 118, Lund, 221 00, Sweden;

#### **Prof. Jerzy Jedlinski**

AGH University of Science and Technology, Faculty of Materials Science and Ceramics, Department of Physical Chemistry and Modeling of Processes and Surface Engineering; al. Mickiewicza 30, Kraków, 30-059, Poland;

## Prof. Megumi Kawasaki

#### <u>ORCID</u>

Oregon State University, School of Mechanical, Industrial and Manufacturing Engineering; 204 Rogers Hall, Corvallis, USA, 97331;

## **Prof. Terence G. Langdon**

University of Southampton, Faculty of Engineering and the Environment; Lanchester Building (Bldg. 7), Highfield Campus, Southampton, United Kingdom, SO17 1BJ;

## Prof. Jai Sung Lee

Hanyang University, Department of Metallurgy and Materials Science; 55 Daehak-no, Sangnok-gu, Ansan, Korea, South, 426-791;

-----

## Prof. Eric J. Mittemeijer

Max Planck Institute for Intelligent Systems; Heisenbergstrasse 3, Stuttgart, 70569, Germany;

## Prof. Stephen J. Pearton

University of Florida, Department of Materials Science and Engineering; Gainesville, USA, 32611-6400;

#### **Prof. Vassilis Pontikis**

Commissariat à l'Energie Atomique et les Energies Alternatives (CEA), CEA-Saclay; Bdg. 524, Gif-sur-Yvette, 91191, France;

## Prof. András Roósz

Hungarian Academy of Sciences, Miskolc University (HAS-MU); Miskolc-Egyetemváros, 3515, Hungary;

#### Prof. David N. Seidman

Northwestern University, Department Materials Science and Engineering; Cook Hall, 2220 Campus Drive, Evanston, USA, 60208;

## Dr. Ching Hua Su

NASA/Marshall Space Flight Center, EM31 NASA/Marshall Space Flight Center; Huntsville, USA, 35812;

#### **Prof. David Tomanek**

Michigan State University, Physics and Astronomy Department; 567 Wilson Road, East Lansing, USA, MI 48824-6455;

#### Prof. A.S. Wronski

University of Bradford, School of Engineering, Design and Technology; West Yorkshire, Bradford, United Kingdom, BD7 1DP;

## Emeritus Prof. David J. Young

University of New South Wales, School of Materials Science and Engineering; Sydney, Australia, NSW 2052;

DISTRIBUTION & ACCESS FOR PUBLICATION INSIGHTS DOWNLOADS ABOUT US POLICY & ETHICS CONTACT US IMPRINT PRIVACY POLICY SITEMAP ALL CONFERENCES ALL SPECIAL ISSUES

Scientific.Net is a registered brand of Trans Tech Publications Ltd © 2023 by Trans Tech Publications Ltd. All Rights Reserved