

PHYSICAL CHEMISTRY 2016

13th International Conference on Fundamental and Applied Aspects of Physical Chemistry

> Proceedings Volume I

B E L G R A D E September 26 - 30, 2016



PHYSICAL CHEMISTRY 2016

13th International Conference on Fundamental and Applied Aspects of Physical Chemistry

> Proceedings Volume I

B E L G R A D E September 26-30, 2016

ISBN 978-86-82475-34-7

Title: Physical Chemistry 2016 (Proceedings)
Editors: Željko Čupić and Slobodan Anić
Published by: Society of Physical Chemists of Serbia, Studentski trg 12-16, 11158, Belgrade, Serbia.
Publisher: Society of Physical Chemists if Serbia
For Publisher: S. Anić, President of Society of Physical Chemists of Serbia
Printed by: "Jovan", Printing and Publishing Company; 200 Copies.
Number of pages: 6+502; Format B5; printing finished in September 2016

Text and Layout: "Jovan"

Neither this book nor any part may be reproduced or transmitted in any form or by any means, including photocopying, or by any information storage and retrieval system, without permission in writing from the publisher.

200 - Copy printing

CONTENT

Volume I	
Organizer	IV
Comittes	V
Sponsors	VI
Plenary Lecture	1
Spectroscopy, Molecular Structure, Physical Chemistry of	47
Plasma	
Kinetics, Catalysis	183
Nonlinear Dynamics, Oscillatory Reactions, Chaos	267
Electrochemistry	355
Biophysical Chemistry, EPR Investigations of bio-systems,	409
Photochemistry, Radiation Chemistry	



PHYSICAL CHEMISTRY 2016

13th International Conference on Fundamental and Applied Aspects of Physical Chemistry

Organized by

The Society of Physical Chemists of Serbia

in co-operation with

Institute of Catalysis Bulgarian Academy of Sciences

and

Boreskov Institute of Catalysis Siberian Branch of Russian Academy of Sciences

and

University of Belgrade, Serbia:

Faculty of Physical Chemistry Institute of Chemistry, Technology and Metallurgy Vinča Institute of Nuclear Sciences Faculty of Pharmacy Institute of General and Physical Chemistry, Belgrade, Serbia

International Organizing Committee

Chairman:	S. Anić (Serbia)			
Vice-chairman:	M. Gabrovska (Bulgaria)			
	A. A. Vedyagin (Russia)			
	S. N. Blagojević (Serbia)			
Members:	N. Cvjetićanin (Serbia), S. M. Blagojević (Serbia), M. Daković			
	(Serbia), J. Dimitrić-Marković (Serbia), T. Grozdić (Serbia), Lj.			
	Ignjatović (Serbia), D. Jovanović (Serbia), J. Jovanović (Serbia),			
	M. Kuzmanović (Serbia), D. Marković (Serbia), B.			
	Milosavljević (USA), M. Mojović (Serbia), N. Ostrovski			
	(Serbia), N. Pejić (Serbia), M. Petković (Serbia), A. Popović-			
	Bjelić (Serbia), B. Simonović (Serbia), D. Stanisavljev (Serbia),			
	M. Stanković (Serbia), Z. Šaponjić (Serbia), B. Šljukić (Serbia),			
	G. Tasić (Serbia), N. Vukelić (Serbia), V. Vukojević (Sweden)			

International Scientific Committee

Chairman:	Ž. Čupić (Serbia)
Vice-chairmans:	V. N. Parmon (Russia)
	S. Rakovsky (Bulgaria)
	B. Adnađević (Serbia)
Members:	S. Anić (Serbia), A. Antić-Jovanović (Serbia), G. Bačić (Serbia),
	R. Cervellati (Italy), G. Ćirić-Marjanović (Serbia), A. Cricenti
	(Italy), V. Dondur (Serbia), S. D. Furrow (USA), L. Gábor
	(Hungary), Vilmos Gáspár (Hungary), K. Hedrih (Serbia), M.
	Jeremić (Serbia), E. Kiš (Serbia), Lj. Kolar-Anić (Serbia), U.
	Kortz (Germany), T. Kowalska (Poland), V. Kuntić (Serbia), Z.
	Marković (Serbia), S. Mentus (Serbia), K. Novaković (UK), B.
	Novakovski (Poland), T. Parac Vogt (Belgium), M. Perić
	(Serbia), M. Plavšić (Serbia), G. Schmitz (Belgium), I. Schreiber
	(Czech Republic), P. Ševčik (Slovakia), N. Stepanov (Russia),
	M. Trtica (Serbia), V. Vasić (Serbia), D. Veselinović (Serbia),
	Á. Tóth (Hungary)

Local Executive Committee

Chairman:	S. N. Blagojević
Vice-chairmans:	A. Ivanović-Šašić
	 A. Stoiljković

Members:

M. Ajduković, P. Banković, N. Bošnjaković, I. N. Bubanja, D. Dimić, A. Dobrota, J. Dostanić, A. Ignjatović, S. Jovanović, Z. Jovanović, A. Jović, N. Jović-Jovičić, D. Lončarević, M. Kragović, J. Krstić, S. Maćešić, J. Maksimović, V. Marković, D. Milenković, M. Milovanović, B. Nedić-Vasiljević, M. Pagnacco, A. Pavićević, N. Potkonjak, D. Ranković, M. Ristić, B. Stanković, A. Stanojević

SPONSORS

Ministry of Education, Science and Technological Development University of Belgrade, Belgrade Institute of General and Physical Chemistry, Belgrade PRIMALAB d.o.o., Serbia

EFFECT OF BASICITY ON THE CATALYTIC PERFORMANCE OF HETEROGENEOUS CATALYSTS FOR BIODIESEL PRODUCTION

D. Marinković, M. Stanković and D. Jovanović

University of Belgrade, Institute of Chemistry, Technology and Metallurgy, Njegoševa 12, 11000 Belgrade, Serbia. (dalibor@ihtm.bg.ac.rs)

ABSTRACT

In a heterogeneous base methanolysis of vegetable oils, among of the most important characteristics of the catalysts are the basicity and the availability of basic centers. The impact of these factors on the catalytic activity was investigated in the reaction of sunflower oil methanolysis by CaO/ γ -Al₂O₃ catalyst. The particle morphology of CaO-dispersed onto γ -Al₂O₃ surface was analyzed by field emission scanning electron microscopy (FESEM), while basicity and basic strength of prepared catalysts were determined using Hammett indicators method. The results showed a strong dependance of catalytic activity on its basicity. The catalyst with the highest basicity was the most active, and decreasing basicity reduces activity of the catalyst. In addition, it was shown that the catalyst activity depends on the favorable spatial distribution of basic sites, i.e. of their availability.

INTRODUCTION

Fatty acid methyl esters (FAMEs) - biodiesel, produced from renewable resources such as vegetable oils or animal fats is expected to be one of the biomass-based alternative for fossil diesel fuel, due to its numerous advantages.

The usual method of biodiesel production is based on triacylglycerols methanolysis to FAMEs using homogeneous base catalysts. Heterogeneous solid base catalyzed processes are nowadays very promising alternatives for biodiesel production from vegetable oils [1]. Using the Ca-based catalysts for the biodiesel production is well known [2], particularly good results were obtained by loading of CaO on alumina carrier [3]. As this is a heterogeneous solid base catalyzed reaction, one of the most important factors affecting the reaction yield is catalyst basicity and the availability of basic sites on its surface.

The aim of this study was to examine the effect of basicity and basic sites distribution on the catalyst activity in the methanolysis of sunflower oil over the CaO/ γ -Al₂O₃ catalyst.

EXPERIMENTAL

Catalyst preparation

CaO/ γ -Al₂O₃ catalysts were prepared according to the modified wet impregnation (MWI) method [4], using Ca(CH₃COO)₂ as precursor salt, γ -Al₂O₃ (spherical shape) as support, and deionized water as medium. The sample designation, CaO content and preparation procedure are shown in Table 1.

Table 1. CaO content and preparation procedure for CaO/γ -Al₂O₃ samples

Sample		Preparation procedure			
		MWI		Calcination conditions	
Designation	CaO*	$C_{CaAc}{}^a$	t_{WI}^{b}	$t_{\rm C}^{\rm c}$	T_C^d
	(wt%)	(wt%)	(h)	(h)	(°C)
Ca-500/ γ -Al ₂ O ₃	4.2	25	2	4	500
Ca-550/ γ -Al ₂ O ₃	4.4	25	2	4	550
Ca-600/ γ -Al ₂ O ₃	4.1	25	2	4	600
Ca-650/ γ -Al ₂ O ₃	4.2	25	2	4	650
Ca-700/y-Al ₂ O ₃	5.5	25	2	4	700
$Ca-750/\gamma-Al_2O_3$	5.3	25	2	4	750

*Determined gravimetrically as CaC₂O₄·H₂O;^aCaAc - Ca(CH₃COO)₂; ^bImpregnation time; ^c Calcination time; ^dCalcination temperature.

Catalyst characterization - FESEM was carried out on a Tescan MIRA3 XMU with accelerating voltage of 10 kV. Samples were placed over an aluminum drum and covered with an Au and Pt film. Basic strength and basicity (H_{-}) were determined using volumetric titration with the Hammett indicators. The following Hammett indicators were used: neutral red (H_{-} =6.8), phenolphthalein (H_{-} =9.9) andthymol violet(H_{-} =11.0).

Methanolysis procedure - The activity of the prepared catalysts was examined in the methanolysis of sunflower oil. All methanolysis reactions were performed in a 250 ml three-neck glass flask equipped with a reflux condenser and a magnetic stirrer. Experiments were conducted under the following conditions: catalyst loading of 0.5 wt. % CaO (relative to the amount of oil), methanol/oil molar ratio of 12/1, at temperature of 60 °C and reaction time of 5 h.

RESULTS AND DISCUSSION

Loading CaO to γ -Al₂O₃carrier and subsequent calcination leads to the formation of cluster structure on the catalyst surface(Fig. 1). Scattered large rod-shaped crystals are present mainly on the surface of the catalyst calcined at the lowest temperature. By increasing the calcination temperature, the granular particles on the surface of catalyst become bigger. Also, it leads to the occurrence of agglomeration and clustering of particles. Evidently, on the Ca-500/ γ -Al₂O₃only some cluster of irregular shape can be seen, on the

Ca-650/ γ -Al₂O₃ catalyst the entire surface is covered with clusters of spherical shapes, while on the catalyst calcined at the highest temperature (Ca-750/ γ -Al₂O₃) spherical clusters are not visible due to massive agglomeration and sintering.

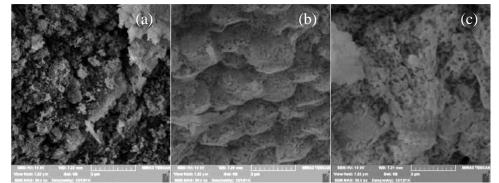


Figure 1. FESEM micrograph of the catalysts surface: (a) Ca-500/γ-Al₂O₃, (b) Ca-650/γ-Al₂O₃ and (c) Ca-750/γ-Al₂O₃

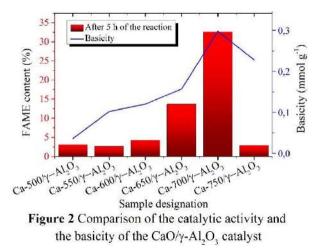
The base strength of CaO loaded onto γ -Al₂O₃ and thermally activated at different temperatures was measured by using Hammett indicators. As evident in Table 2, all catalysts are within the same range of the base strength, 9.3<*H*_<9.9.Therefore, these catalysts can be classified as materials with a medium base strength. Although belonging to the same base strength group, their basicity is not equal. Basicity of the samples increases with calcination temperature up to 700 °C at which reaches the maximum value (Table 2, Ca-700/ γ -Al₂O₃). The basicity of the catalyst obtained by calcination at the highest temperature (Ca-750/ γ -Al₂O₃) is lower, which is in accordance with the results of FESEM (Fig. 1). On the surface of Ca-750/ γ -Al₂O₃catalystit was observed the

Table 2. Basicity and basic strength (H_{-}) of the CaO/y-Al₂O₃ catalysts

the CaO/ y-Al ₂ O ₃ catalysts			
Sample	Basic strength	Basicity	
designation	(H_)	(mmol g ⁻¹)	
$Ca-500/\gamma-Al_2O_3$	9.3< <i>H</i> _<9.9	0.036	
Ca-550/y-Al ₂ O ₃	9.3< <i>H</i> _<9.9	0.102	
Ca-600/y-Al ₂ O ₃	9.3< <i>H</i> _<9.9	0.120	
Ca-650/y-Al ₂ O ₃	9.3< <i>H</i> _<9.9	0.157	
Ca-700/y-Al ₂ O ₃	9.3< <i>H</i> _<9.9	0.298	
Ca-750/y-Al ₂ O ₃	9.3< <i>H</i> _<9.9	0.228	

massive sintering of With particles. the appearance of larger particles on the catalyst surface, the number and the availability of base catalytically active sites are reduced, despite the same concentration level of catalytically active compound - CaO (Table 1). Activity of the catalyst

samples was evaluated by analysing the FAMEsyield after 5 h. As can be seen, the Fig. 2 reveals a clear correlation betweenFAMEs content after 5 h of the reaction and the basicity of the catalysts. The catalyst sample with the highest



basicity(Ca-700/ γ -Al₂O₃) showed the highest activity, and the samples with the lowest basicity (Ca-500/y- Al_2O_3 Ca-550/yand Al₂O₃)showed the lowest activity. Despite the relatively high basicityCa- $750/\gamma$ -Al₂O₃ catalyst did not show adequate activity. probably due to reduction of base sites space availability.

CONCLUSION

Heterogeneous CaO/ γ -Al₂O₃ catalyst samples were synthesized by modified wet impregnation method and thermally activated under inert atmosphere. The analysis of the catalyst activity in the reaction of sunflower oil methanolysis and basicity of synthesized catalysts a correlation between basicity and catalyst activity was observed. The catalyst with higher basicity had higher activity. The exception was the catalyst thermally activated at the highest temperature in which, probably due to sintering effect that leads to agglomeration of active sites on surface causes a reduction in its activity.

Acknowledgement

This work was supported by the Ministry of Education, Science and Technological Development of the R. of Serbia (Project III 45001).

REFERENCES

- [1] D.M. Marinković, M.V. Stanković, A.V. Veličković, J.M. Avramović, M.R. Miladinović, O.O. Stamenković, V.B. Veljković, D.M. Jovanović, Renewable and Sustainable Energy Reviews., 2016, 56, 1387-1408.
- [2] M. Kouzu, T. Kasuno, M. Tajika, Y. Sugimoto, S. Yamanaka, J. Hidaka, Fuel, 2008, 87, 2798-2806.
- [3] N. Pasupulety, K. Gunda, Y. Liu, G.L. Rempel, F.T.T. Ng, Applied Catalysis A: General, 2013, 452, 189-202.
- [4] D.M. Marinković, M.V. Stanković, A.V. Veličković, J.M. Avramović, M.D. Cakić, V.B. Veljković, Advanced Technologies, 2015, 4(1), 26-32.

CIP - Каталогизација у публикацији - Народна библиотека Србије, Београд

544(082) 66.017/.018(082) 502/504(082) 663/664:658.56(082) 615.31:547(082)

INTERNATIONAL Conference on Fundamental and Applied Aspects of Physical Chemistry (13 ; 2016 ; Beograd)

Physical Chemistry 2016 : proceedings. Vol. 2 / 13th International Conference on Fundamental and Applied Aspects of Physical Chemistry, Belgrade, 26-30 September 2016 ; [editors Željko Čupić and Slobodan Anić]. - Belgrade : Society of Physical Chemists of Serbia, 2016 (Belgrade : Jovan). - IV, 507-930 str. : ilustr. ; 24 cm

Tiraž 200. - Bibliografija uz svaki rad.

ISBN 978-86-82475-33-0

1. Society of Physical Chemists of Serbia (Beograd) а) Физичка хемија - Зборници b) Наука о материјалима - Зборници c) Животна средина - Заштита - Зборници d) Животне намирнице - Контрола квалитета - Зборници e) Фармацеутска хемија - Зборници COBISS.SR-ID 225802508

0 8, 09. 2016

