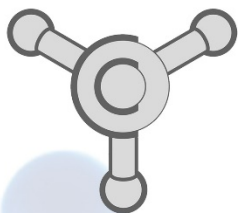


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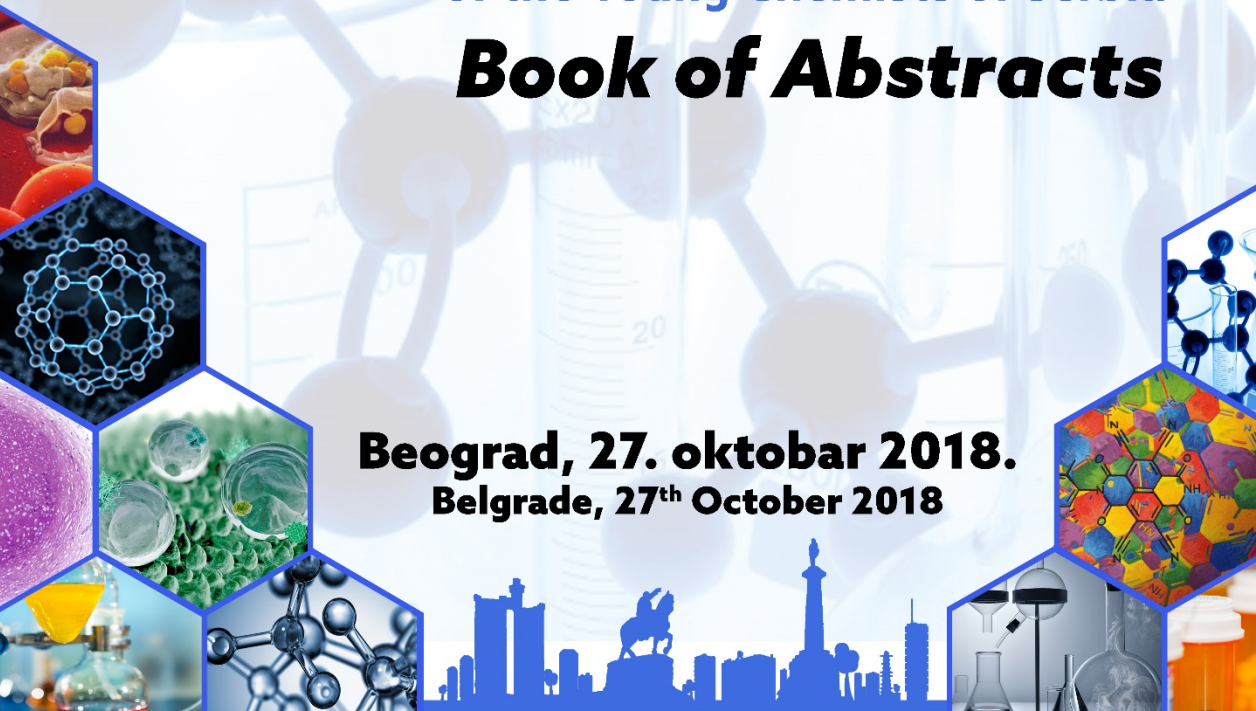


Šesta konferencija
mladih hemičara Srbije

Kratki izvodi radova

Sixth Conference
of the Young Chemists of Serbia
Book of Abstracts

Beograd, 27. oktobar 2018.
Belgrade, 27th October 2018



CIP - Каталогizacija u publikaciji
Narodna biblioteka Srbije, Beograd

54(048)(0.034.2)
577.1(048)(0.034.2)
60(048)(0.034.2)
66.017/.018(048)(0.034.2)

KONFERENCIJA mladih hemičara Srbije (6 ; 2018 ; Beograd)

Kratki izvodi radova [Elektronski izvor] = Book of Abstracts / Šestakonferencija mladih hemičara Srbije, Beograd, 27. oktobar 2018. = Sixth Conference of Young Chemists of Serbia, Belgrade, 27th October 2018 ; [organizatori Klub mladih hemičara Srbije [i] Srpsko hemijsko društvo =organizers Serbian Young Chemists' Club [and] Serbian Chemical Society] ; [urednici, editors Tamara Todorović ... et al.]. - Beograd : Srpskohemijsko društvo = Serbian Chemical Society, 2018 (Beograd :Razvojno-istraživački centar grafičkog inženjerstva TMF = Belgrade :Development and Research Centre of Graphic Engineering Faculty of Technology and Metallurgy). - 1 elektronski optički disk (CD-ROM) : tekst ; 12 cm

Sistemski zahtevi: Nisu navedeni. - Apstrakti na engl. jeziku. - Nasl. sa naslovne strane dokumenta. - Tiraž 120. - Bibliografija uz većinu apstrakata. - Registar.

ISBN 978-86-7132-072-6

1. Klub mladih hemičara Srbije (Beograd) 2. Srpsko hemijsko društvo (Beograd)

a) Хемија - Апстракти b) Биохемија - Апстракти c) Биотехнологија - Апстракти d) Наука о материјалима - Апстракти
COBISS.SR-ID 269395724

6. KONFERENCIJA MLADIH HEMIČARA SRBIJE

6th CONFERENCE OF THE YOUNG CHEMISTS OF SERBIA

Beograd, 27. oktobar 2018. / BELGRADE, 27th OCTOBER 2018

KRATKI IZVODI RADOVA / BOOK OF ABSTRACTS

Izdaje / Published by

Srpsko hemijsko društvo / Serbian Chemical Society

Karnegijeva 4/III, 11000 Beograd, Srbija

Tel./fax: +381 11 3370 467; www.shd.org.rs; office@shd.org.rs

Za izdavača / For Publisher

Vesna MIŠKOVIĆ-STANKOVIĆ, predsednica Društva / President

Urednici / Editors

Tamara TODOROVIĆ

Ljubodrag VUJISIĆ

Jelena RADIVOJEVIĆ

Vuk FILIPOVIĆ

Dizajn, slog i kompjuterska obrada / Page Layout and Design

Vuk FILIPOVIĆ

Tiraž / Circulation

120 primeraka / 120 copies

ISBN 978-86-7132-072-6

Umnožavanje / Copying

Razvojno-istraživački centar grafičkog inženjerstva,

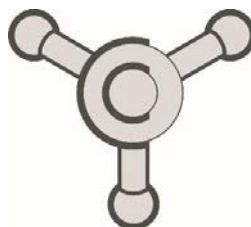
Tehnološko-metalurški fakultet, Karnegijeva 4, Beograd, Srbija

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Konferenciju su podržali / Supported by



**Ministarstvo prosvete, nauke i tehnološkog
razvoja Republike Srbije**
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Sadržaj / Table of Contents

Plenarno predavanje / Plenary Lecture

Petar Ristivojević

Planar chromatography today: application in natural product and food analysis __ 5

Predavanja po pozivu / Invited Lecture

Aleksandra Mitrović

Covalent functionalization of black phosphorus _____ 6

Nina Božinović

Interaction of human immunoglobulins with metalloporphyrins _____ 7

Saopštenja / Contributions

Biohemija i biotehnologija / Biochemistry and biotechnology _____ 7

Hemija hrane / Food chemistry _____ 14

Hemija makromolekula i nanotehnologije /
Chemistry of macromolecules and nanotechnology _____ 19

Hemija u Nastavi / Educational chemistry _____ 25

Hemijska analiza / Chemical analysis _____ 29

Hemijska sinteza / Chemical synthesis _____ 46

Industrijska i primenjena hemija / Industrial and applied chemistry _____ 67

Medicinska hemija / Medicinal chemistry _____ 74

Nauka o materijalima / Materials science _____ 91

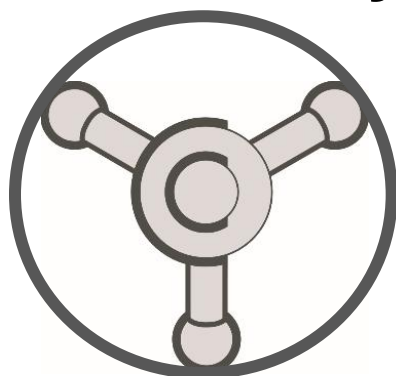
Teorijska hemija / Theoretical chemistry _____ 104

Indeks autora / Author Index _____ 113

Kratki izvodi radova ***Book of Abstracts***



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Hemijska sinteza

Chemical Synthesis

HS10 PE 8

Synthesis and characterization of 1-(4-substitutedbenzyl)-3',4'-dihydro-2'H-spiro[imidazolidine-4,1'-naphtalene]-2,5-dione derivatives as potential anticonvulsant agents

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Among the diverse pharmacological activity reported for spirohydantoin and the closely related spirosuccinimides, their anticonvulsant and antiproliferative activities are commonly encountered. Of all cyclic ureide derivatives, Phenytoin (5,5-diphenylhydantoin) is one of the well-known commercially available anticonvulsants, which is also widely used as an antiarrhythmic agent. However, the clinical use of 5,5-diphenylhydantoin is limited by its central nervous adverse effects in addition to a wide variety of drug interactions [1]. With the aim of developing new analogs with a more efficient therapeutic effect, a series of 1-(4-substitutedbenzyl)-3',4'-dihydro-2'H-spiro[imidazolidine-4,1'-naphtalene]-2,5-diones, have been synthesized, characterized by melting points, FT-IR, ¹H and ¹³C NMR spectroscopic techniques, and evaluated for anticonvulsant activity. The objective of the present investigation is to identify the pharmacological impact of the spirocyclic attachment of a semi-rigid tetralin residue at C-5 of the imidazolidine-2,4-dione ring and to explore the effect of substitution at N-3 on anticonvulsant activity.

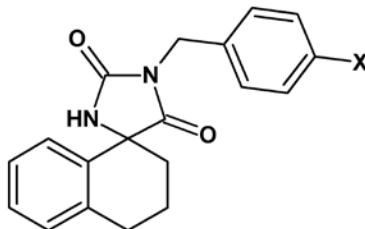


Figure 1. Structure of the investigated spirohydantoin derivatives

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Acknowledgments

This work was supported by the Ministry of Education, Science and Technological Development, of the Republic of Serbia (Project no. 172013).