

## NINETEENTH YOUNG RESEARCHERS' CONFERENCE MATERIALS SCIENCE AND ENGINEERING

December 1-3, 2021, Belgrade, Serbia

## Program and the Book of Abstracts

Materials Research Society of Serbia &

Institute of Technical Sciences of SASA

Book title:

Nineteenth Young Researchers' Conference - Materials Science and Engineering: Program and the Book of Abstracts

Publisher:

Institute of Technical Sciences of SASA Knez Mihailova 35/IV, 11000 Belgrade, Serbia Tel: +381-11-2636994, 2185263, http://www.itn.sanu.ac.rs

Conference organizers:

Materials Research Society of Serbia, Belgrade, Serbia Institute of Technical Sciences of SASA, Belgrade, Serbia

Editor:

Dr. Smilja Marković

Technical Editor: Aleksandra Stojičić

Cover page: Aleksandra Stojičić and Milica Ševkušić

Cover: Milica Ševkušić

Printing:

Gama digital centar Autoput No. 6, 11070 Belgrade, Serbia Tel: +381-11-6306992, 6306962 http://www.gdc.rs

Publication year: 2021

Print-run: 120 copies

CIP - Каталогизација у публикацији

Народна библиотека Србије, Београд

66.017/.018(048)

## YOUNG Researchers Conference Materials Sciences and Engineering (19; 2021; Beograd)

Program; and the Book of abstracts / Nineteenth Young Researchers' Conference Materials Science and Engineering, December 1-3, 2021, Belgrade, Serbia; [organized by] Materials Research Society of Serbia & Institute of Technical Sciences of SASA; [editor Smilja Marković]. - Belgrade: Institute of Technical Sciences of SASA, 2021 (Belgrade: Gama digital centar). - XVIII, 86 str.: ilustr.; 23 cm

Tiraž 120. - Registar.

ISBN 978-86-80321-36-3

а) Наука о материјалима -- Апстракти б) Технички материјали – Апстракти

COBISS.SR-ID 51231241

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## Antiradical activity of graphene quantum dots enriched albumin hydrogel: An EPR study

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Graphene quantum dots (GQD) are single to multilayer carbon-based nanoparticles. They possess high chemical and physical stability, high biocompatibility, low toxicity, and are easily dispersible in water. These properties make them a novel material for the therapeutical application. Due to their antiradical scavenging activity, GOD are considered to be a promising material in terms of inhibiting oxidative damage caused by the elevated level of free radicals. Serum albumin is the primary carrier of various solutes in blood plasma and is commonly used in solubilizing therapeutics. It is suitable for the production of biological hydrogels which are biocompatible and have medical significance as efficient site-located drug delivery systems. This research aims to examine the antiradical activity of GQD incorporated into the albumin hydrogel towards 'OH radicals and their ability to induce prolonged scavenging effects. Since 'OH radicals are short-lived species, the electron paramagnetic resonance (EPR) spin-trapping technique (and spin-trap DEPMPO) has been used for their detection. Gellation of bovine serum albumin (BSA) solution containing GQD has been thermally induced. The obtained gel has been introduced into the Fenton reaction ('OH radicals generator) system containing the spin-trap and gently stirred. To estimate the antiradical effect arising from GQD, control experiments were performed using BSA hydrogel without GOD as well as by replacing BSA hydrogel with water. The obtained results indicate that BSA itself possesses certain antiradical activity, since it eliminated 19.84 % of 'OH radicals from the system. However, the most significant elimination of 'OH radicals has been observed in the system containing GOD (99.86 %), where 80.02 % of 'OH radicals have been eliminated entirely by GQD. Our results show the significant potential of albumin hydrogels as GQD carriers, as well as the compelling anti-hydroxyl radical scavenging efficiency of the GQD. These results give a promising perspective in the light of the synergical effect of GQD albumin hydrogels as site-located radical scavengers.

This research was supported by Ministry of Education, Science and Technological Development of Republic of Serbia Contract number: 451-03-9/2021-14/200146 and the Science Fund of the Republic of Serbia, PROMIS, #6062285, PHYCAT.