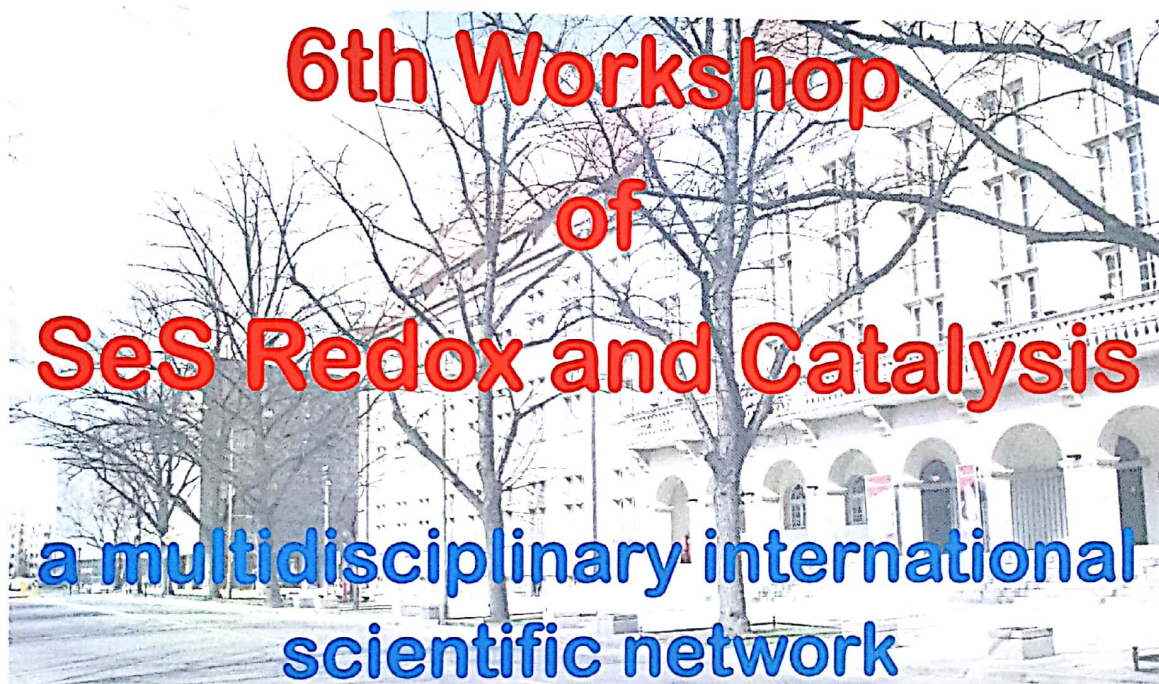


WSeS-6



**Wrocław University
of Science and Technology
Wrocław, Poland**

22-23 September 2017

**6th Workshop
of SeS Redox and Catalysis**

**a multidisciplinary international
scientific network**

**Wrocław University
of Science and Technology
Wrocław, Poland**

22-23 September 2017

Book of Abstracts of 6th Workshop
of SeS Redox and Catalysis
Editors: Elżbieta Wojaczyńska, Jacek Wojaczyński

All rights reserved. No part of this book may be reproduced, stored in a retrieval system, or transmitted in any form or by any means, without the prior permission in writing of the Publisher.

© Copyright by Oficyna Wydawnicza Politechniki Wrocławskiej, Wrocław 2017

ISBN 978-83-7493-991-1

Oficyna Wydawnicza Politechniki Wrocławskiej
Wybrzeże Wyspiańskiego 27, 50-370 Wrocław
<http://www.oficyna.pwr.edu.pl>
oficwyd@pwr.edu.pl

Print and binding: beta-druk, www.betdruk.pl

P18: Synthesis of alkyl benzisosenazol-3(2H)-ones and their sulfur analogues

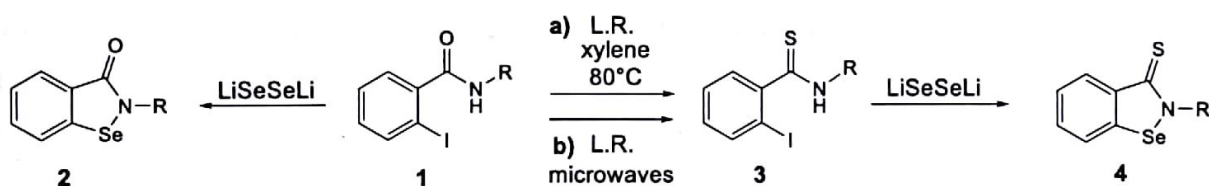
Magdalena Obieziurska, Agata J. Pacula, Jacek Ścianowski

Department of Organic Chemistry, Faculty of Chemistry, Nicolaus Copernicus University, 7 Gagarin Street, 87-100 Toruń, Poland;
m.obieziurska@wp.pl



Benzisoselenazolones are important group of organoselenium compounds particularly due to their antioxidant properties. They can control the imbalance between the production and elimination of reactive oxygen species (ROS), which can lead to several frequently occurring diseases such as cancer, diabetes, atherosclerosis, hypertension, Alzheimer's disease.¹

The aim of this study was the synthesis of alkyl benzisosenenazole-3(2H)-ones and their sulfur analogues. In the first step *N*-substituted *o*-iodobenzamides **1** by the reaction of amines with *o*-iodobenzoic acid chloride were obtained. Then they were transformed into benzisosenazol-3(2H)-ones **2** by the reaction with dilithium diselenide² or into the corresponding *N*-substituted *o*-iodobenzothioamides **3** by two alternative methods. The first one **a** was the reaction of **1** with Lawesson's reagent in xylene at 80 °C.³ The second one **b** was based on the reaction without the presence of solvent under the influence of microwave radiation.⁴ The corresponding thiocarbonyl benzisosenazole-3(2H)-ones **4** were prepared by the reaction *N*-substituted *o*-iodobenzothioamides **3** with dilithium diselenide.²



Scheme 1. Synthesis of benzisosenenazole-3(2H)-ones and thiocarbonyl derivatives

Acknowledgements

The work was supported from NCN grant nr UMO-2015/17/B/NZ7/03058.

References:

1. M. Iwaoka *Organoselenium Chemistry: Between Synthesis and Biochemistry* ed. C. Santi, Bentham Science Publishers **2014**, 18, 355-361.
2. Pacula, J. Ścianowski, K. Aleksandrak *RSC Adv.* **2014**, 4, 76, 48959-48962.
3. S. Scheibye, J. Kristensen, S.O. Lawesson *Tetrahedron* **1979**, 35, 1339-1343.
4. R.S. Varma, D.Kumar *Org. Lett.* **1999**, 1, 697-700.