

ILMAT

2023

U. PORTO

7TH INTERNATIONAL Conference on Ionic Liquid Based Materials

21st to 24th November 2023

Porto, Portugal
Instituto Pernambuco

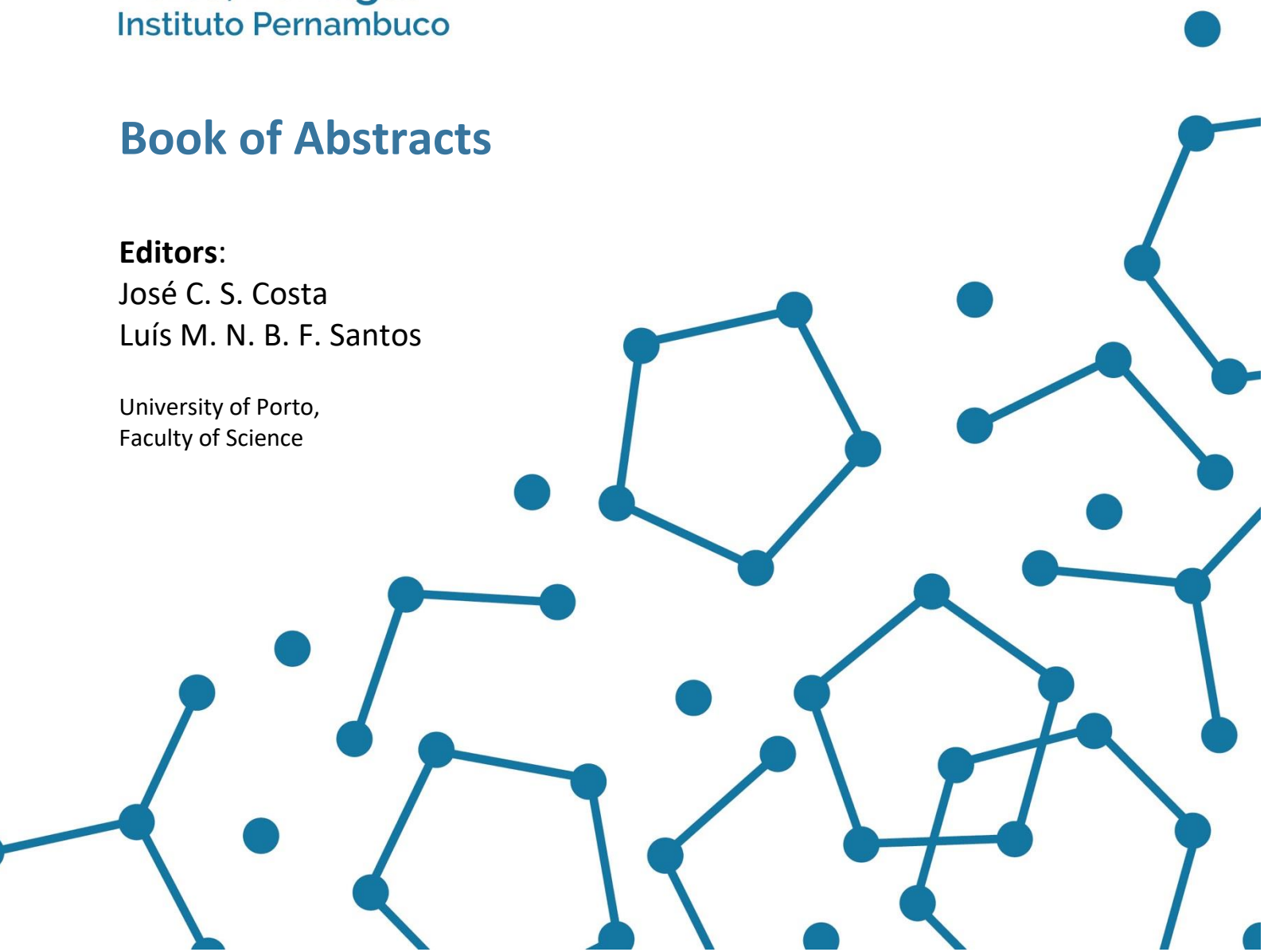
Book of Abstracts

Editors:

José C. S. Costa

Luís M. N. B. F. Santos

University of Porto,
Faculty of Science



ILMAT 2023

7TH INTERNATIONAL
Conference on Ionic Liquid
Based Materials

21st to 24th November 2023
Porto, Portugal
Instituto Pernambuco

Book of Abstracts of the 7th International Conference on Ionic Liquid-Based Materials

ILMAT 2023

7th International Conference on Ionic Liquid-Based Materials - 21st to 24th November 2023

University of Porto, Faculty of Science

Porto

Portugal

Editors

José C. S. Costa

Luís M. N. B. F. Santos

©Organizing Committee of **ILMAT 2023** and Authors

ISBN

978-989-35015-8-0

Publisher

Organizing Committee of **ILMAT 2023**

Poster P_B1.5

Synthesis and physicochemical characterization of antimalarial surface-active ionic liquids

Ana Teresa Silva^{1*}, Isabel Oliveira², Ricardo Ferraz^{1,3}, Eduardo F. Marques², Paula Gomes¹

¹ LAQV-REQUIMTE, Departamento de Química e Bioquímica, Faculdade de Ciências, Universidade do Porto, Porto, Portugal

² CIQUP, IMS (Institute of Molecular Sciences), Departamento de Química e Bioquímica, Faculdade de Ciências, Universidade do Porto, Porto, Portugal

³ Center for Translational Health and Medical Biotechnology Research (TBIO), School of Health (ESS), Polytechnic of Porto, Porto, Portugal.

*up201303026@edu.fc.up.pt

Ionic liquids are a particular class of compounds that attract interest in medicinal chemistry due to the simplicity of their preparation. Novel structures with biological activity can be achieved through simple, cost-effective reactions.¹ Reusing old ionizable drugs and improving their characteristics can be achieved economically and simply by mixing them with molecules of opposite charge. This approach is attractive for reviving old antimalarials, not only because of the prevalence of malaria in low- to middle-income countries, but also because several of these drugs are associated with malaria parasite resistance. In this context, our work has been focusing on synthesizing ionic liquids with potential antimalarial activity by mixing antimalarial aminoquinolines, specifically chloroquine, and primaquine, with natural lipids.^{2, 3} More recently, using an acid-base reaction between chloroquine and bile acids (Figure 1), we synthesized surface-active ionic liquids (SAILs), which proved to possess significant antiplasmodial activity *in vitro*. The presence of an amphiphilic anion in the ionic pair confers surface-active and self-aggregation properties to the ionic liquids. The interfacial and aggregations properties of these SAILs have been characterized by surface tension, electric conductivity, dynamic light scattering, and differential scanning microcalorimetry. Moreover, the interactions of SAILs with micelles of the block copolymer F127 have been studied with the aim of designing an efficient, robust, and biocompatible nanocarrier system for the encapsulation and *in vivo* release of these antimalarial ionic liquids.

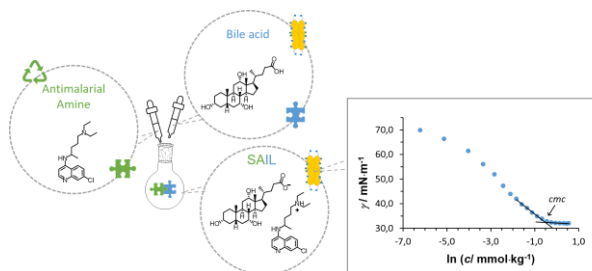


Figure 1: Synthesis and surface tension assays of a SAIL derived from the antimalarial drug chloroquine and cholic acid.

Acknowledgements

The authors acknowledge Fundação para a Ciência e Tecnologia (FCT) for financial support through projects UIDP/50006/2020, UID/QUI/0081/2020 and IMS (LA/P/0056/2020). ATS thanks FCT and Sociedade Portuguesa de Química for her doctoral grant SFRH/BD/150649/2020.

References

1. R. Ferraz, J. Noronha, F. Murtinheira, F. Nogueira, M. Machado, M. Prudencio, S. Parapini, S. D'Alessandro, C. Teixeira, A. Gomes, C. Prudencio and P. Gomes, *Rsc Adv.*, 2016, **6**, 56134.
2. A. T. Silva, L. Lobo, I. S. Oliveira, J. Gomes, C. Teixeira, F. Nogueira, E. F. Marques, R. Ferraz and P. Gomes, *Int. J. Mol. Sci.*, 2020, **21**, 5334.
3. A. T. Silva, I. S. Oliveira, J. Gomes, L. Aguiar, D. Fontinha, D. Duarte, F. Nogueira, M. Prudencio, E. F. Marques and C. Teixeira, *ChemMedChem*, 2022, **17**, e202100650.