

**Institute of Pharmaceutical Technology and
Regulatory Affairs
Faculty of Pharmacy
University of Szeged**

I. Symposium of Young Researchers on Pharmaceutical Technology, Biotechnology and Regulatory Science

Szeged, Hungary



**31th January
2019**



**I. Symposium of Young Researchers on Pharmaceutical Technology,
Biotechnology and Regulatory Science**

Institute of Pharmaceutical Technology and Regulatory Affairs

Faculty of Pharmacy

University of Szeged

Szeged, Hungary

January 31th 2019

DOI: [10.14232/syrptbrs.2019.af](https://doi.org/10.14232/syrptbrs.2019.af)

Edited by Tivadar Bíró, Ildikó Csóka

I. Symposium of Young Researchers on Pharmaceutical Technology, Biotechnology and Regulatory Science

January 31th 2019. Szeged, Hungary

OP-13

DOI: 10.14232/syrptbrs.2019.op13

Formulation and investigation of novel, carrier-based dry powder inhalation system

Edit Benke, Rita Ambrus, Piroska Szabó-Révész

Institute of Pharmaceutical Technology and Regulatory Affairs, Interdisciplinary Excellence Centre, University of Szeged, Szeged, Hungary

Administration of the drug via the lung has several advantages in the treatment of lung diseases, such as cystic fibrosis and chronic obstructive pulmonary disease. Thus, the development of dry powder systems (DPIs) containing antibiotic [1] and non-steroidal anti-inflammatory [2] agents is on every account warranted. In the case of development, it should be borne in mind that the active ingredient particles have a good morphology and low density, between the average particle size of 1-10 μm and the cohesion between the particles is as small as possible. Furthermore, the product must have adequate stability and be compatible with the used inhaler and capsule. The aim of this work is to produce and investigate an antibiotic-containing novel, carrier-based DPI system, that combines the advantageous behaviours of traditional, carrier-based; and carrier-free DPI systems with the use of a combined formulation technique [3]. The studies show that the combined formulation has favourable physical properties, high lung deposition (*in vitro*, *in silico*), it is compatible with the used capsule (high emitted fraction) and stable.

References

1. Benke E. et al. Acta Pharm. Hung. 87, 49-58 (2017)
2. Szabó-Révész P. Drug Discov. Today Technol. 27, 87-93 (2018)
3. Ambrus R. et al. 2018. 123. 20-27 Eur. J. Pharm. Sci. 123, 20-27 (2018)

Acknowledgment

This project was supported by the UNKP-18-3 New National Excellence Program of the Ministry of Human Capacities and acknowledged by the EFOP-3.6.3-VEKOP-16-2017-00009 project.