



Modelling of drying processes of pharmaceutical granules. Pharmaceutical Sciences for the Future of Medicines

Mortier, S.T.F.C.; Vedantam, S.; De Beer, T.; Gernaey, Krist V.; Remon, J.P.; Vervaet, C.; Nopens, I.

Publication date:
2011

Document Version
Publisher's PDF, also known as Version of record

[Link back to DTU Orbit](#)

Citation (APA):

Mortier, S. T. F. C., Vedantam, S., De Beer, T., Gernaey, K., Remon, J. P., Vervaet, C., & Nopens, I. (2011). Modelling of drying processes of pharmaceutical granules. Pharmaceutical Sciences for the Future of Medicines. Abstract from PharmSciFair Conference, Prague, .

General rights

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain
- You may freely distribute the URL identifying the publication in the public portal

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Abstract PharmSciFair Conference in Prague

Modelling of drying processes of pharmaceutical granules

Mortier S.T.F.C.^{1,2,*}, Vedantam S.¹, De Beer T.², Gernaey K.V.³, Remon J.P.⁴, Vervaet C.⁴ and Nopens I.¹

¹ BIOMATH, Department of Applied Mathematics, Biometrics and Process Control, Faculty of Bioscience Engineering, Ghent University, Coupure Links 653, 9000 Ghent, Belgium

² Laboratory of Pharmaceutical Process Analytical Technology, Department of Pharmaceutical Analysis, Faculty of Pharmaceutical Sciences, Ghent University, Harelbekestraat 72, 9000 Ghent, Belgium

³ Center for Process Engineering and Technology, Department of Chemical and Biochemical Engineering, Technical University of Denmark, Building 229, 2800 Kgs. Lyngby, Denmark

⁴ Laboratory of Pharmaceutical Technology, Department of Pharmaceutics, Faculty of Pharmaceutical Sciences, Ghent University, Harelbekestraat 72, 9000 Ghent, Belgium

* Corresponding author (severine.mortier@ugent.be)

Tablets are conventionally produced via consecutive batch process steps. Recent introduction of continuous process equipment is gaining industrial importance in pharmaceuticals. Transition to continuous production requires improved understanding of all operations, necessitating the development of mechanistic models of multi-phase systems which in the end allow process control. This contribution focuses on continuous fluidized bed drying of pharmaceutical wet granules. A step-wise approach is used in model development, starting with the drying behaviour of single granules. Experiments to determine the drying characteristics were conducted in a continuous fluidized bed dryer at several gas temperatures and velocities. The drying process was found to consist of two subsequent phases: a fast drop in moisture content followed by a slower evaporation. A mechanistic model for single granules [1] yielded good description of the data, and was calibrated and validated. The conceptual model consists of a liquid evaporation model (fast part) and a wet core - dry crust model describing the decrease of the wet core diameter and thus the moisture content (slower part). Resulting distinction between weakly and strongly bonded water in the granules yields deeper knowledge of the process. The validated model can now be used for optimization of the continuous drying process.

- [1] M. Mezhericher, A. Levy, and I. Borde, "Theoretical Drying Model of Single Droplets Containing Insoluble or Dissolved Solids," *Dry. Technol.*, vol. 25, Jun. 2007, pp. 1025-1032.