## **Accepted Manuscript**

The clinical relevance of dry powder inhaler performance for drug delivery

Pascal Demoly, Paul Hagedoorn, Anne H. de Boer, Henderik W. Frijlink

PII: S0954-6111(14)00178-4

DOI: 10.1016/j.rmed.2014.05.009

Reference: YRMED 4514

To appear in: Respiratory Medicine

Received Date: 29 January 2014

Revised Date: 8 May 2014 Accepted Date: 13 May 2014

j.rmed.2014.05.009.

Please cite this article as: Demoly P, Hagedoorn P, de Boer AH, Frijlink HW, The clinical relevance of dry powder inhaler performance for drug delivery, *Respiratory Medicine* (2014), doi: 10.1016/

This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting proof before it is published in its final form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.



#### ACCEPTED MANUSCRIPT

### **Highlights**

- $\bullet$  Optimal size of inhaled drug particles for deposition in the airways is 1.5–5  $\mu m$
- Particles >5 μm are mainly deposited in the oropharynx by inertial impaction
- Particles <1 μm have a >40% chance of being exhaled, rather than deposited
- Oropharyngeal impaction increases at higher inspiratory flow rates (IFR)
- An optimal inhaler increases its fine particle (<5 μm) output as IFR increases

#### Download English Version:

# https://daneshyari.com/en/article/6241970

Download Persian Version:

https://daneshyari.com/article/6241970

<u>Daneshyari.com</u>