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forms or medicines. An

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scientists who are

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available.

The suspension dosage  
form has long been used  
for poorly soluble  
active ingredients for  
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of stable suspensions  
over the shelf life of  
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continues to be a  
challenge on many  
fronts. A good  
understanding of the

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fundamentals of disperse  
systems is essential in  
the development of a  
suitable pharmaceutical

suspension. The  
development of a s-  
pension dosage form

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follows a very  
complicated path. The  
selection of the proper  
excipients (surfactants,  
viscosity imparting  
agents etc.) is  
important. The particle

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size distribution in the  
finished drug product  
dosage form is a

critical parameter that  
significantly impacts  
the bioavailability and  
pharmacokinetics of the

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product. Appropriate  
analytical methodologies  
and instruments

(chromatographs, visco-  
meters, particle size  
analyzers, etc.) must be  
utilized to properly

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characterize the s-  
pension formulation. The  
development process

continues with a

successful scale-up of

the manufacturing

process. Regulatory

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agencies around the world require clinical trials to establish the safety and efficacy of the drug product. All of this development work should culminate into a

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regulatory filing in  
accordance with the  
regulatory guidelines.

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organization, follows  
the development approach  
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a suitable vehicle.

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of formulations, regulatory issues,  
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is the 'art of the apothecary' or, in

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of disease.

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Textbook of Pharmaceutics follows  
the same goals as those of the  
previous edition, albeit in a new  
look. The content of the old edition  
has been updated and expanded

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and several new chapters, viz.  
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per ICH Guidelines, Parenteral  
Formulations, New Drug Delivery  
Systems and Pilot Plant  
Manufacturing, have been included,  
with an intention to make the book

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more informative for the modern  
pharmacists. The book has six  
sections: Section I deals with the  
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new chapters: Complexations and  
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have been added to make it more

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informative. Section II conveys the information regarding pharmaceutical unit operations and processes. Section III describes the area of pharmaceutical practice.

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this section. Two new chapters:  
Parenteral Formulations and New  
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Section VI contains the formulation and packaging aspects of pharmaceuticals. Pilot Plant

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the pilot plant model.

Absorption, Distribution,  
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processes and their relationship  
with the design of dosage forms  
and the success of  
pharmacotherapy form the basis of

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engineers, medicinal chemists) who  
might be working in a positions in  
pharmaceutical companies or

whose work might benefit from

basic training in the ADME

concepts and some biological

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such as objectives, keywords,  
discussion questions, summaries  
and case studies add valuable  
teaching tools. This book will  
provide not only general knowledge  
on ADME processes but also an  
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such as drug transporters, multi-  
drug resistance related to  
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generation pharmaceutical carriers  
(nanopharmaceuticals), in vitro and  
in vivo bioequivalence studies,  
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pharmacogenomics, drug-drug and food-drug interactions, and in silico and in vitro prediction of ADME

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processes. Each of these  
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burgeoning fields has a separate  
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University School of Medicine,  
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Pharmacy, University of Maryland,  
University of Bath). Additionally,

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each of the initial chapters dealing with the generalities of drug absorption, distribution, metabolism and excretion would include relevant, classic examples related to each topic with appropriate illustrations (e.g. importance of

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active absorption of levodopa,  
implications in levodopa

administration, drug drug

interactions and food drug

interactions emerging from the

active uptake; intoxication with

paracetamol as a result of

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glutathione depletion, CYP  
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induction and its relationship with  
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acute liver failure caused by  
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paracetamol, etc). ADME  
Processes and Pharmaceutical  
Sciences is written as a core  
textbook for ADME processes,

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systems could only be  
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biological systems is made available, we are realizing the non-linearity of these systems. The concepts and techniques of nonlinear analysis allow for more realistic and accurate

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Dosage Form Design  
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