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# 3.3.2. Total number of books and chapters in edited volumes/books published and papers in national/ international conference proceedings year wise during last five years

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1	Pharmaceutical Dosage form Technology	Dr.T.E Gopala Krishna Murthy	Brillion Publishing	978-81- 19238- 28-6	2023
2	Molecular Docking Studies to improve Bioavailability of p-gp Substrate	Dr.T.E Gopala Krishna Murthy	LAP LAMBERT Academic Publishing GmbH & Co. KG, (Germany)	978-620- 6-78443- 2	2023

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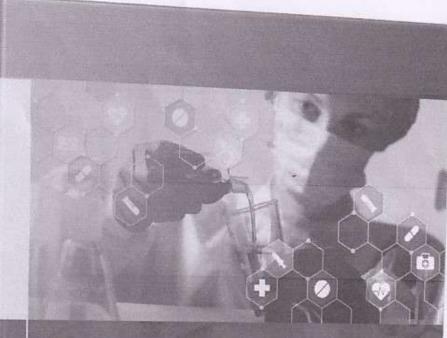
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2	Pharmaceutical Dosage form Technology	B.Sudheer Chowdary	Liquid Dosage Forms	Brillion Publishing	978- 81- 19238- 28-6	2023
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PHARMACEUTICAL DOSAGE FORM TECHNOLOGY



# PHARMACEUTICAL DOSAGE FORM TECHNOLOGY

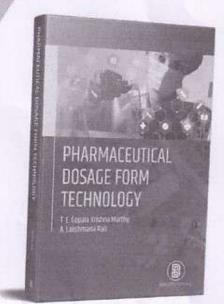
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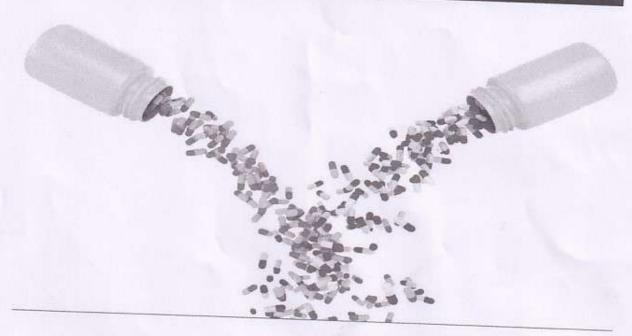
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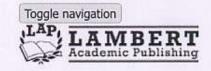


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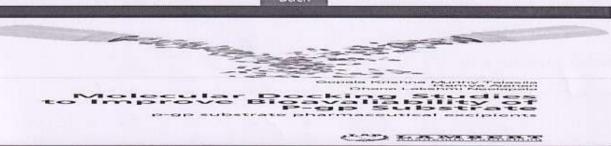
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p-gp substrate pharmaceutical excipients





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# Molecular Docking Studies to Improve Bioavaliability of P-gp Substrate

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P-glycoprotein (P-gp) known as multi drug resistance (MDR) protein was discovered by Juliano and Ling in 1976 and belong to the adenosine triphosphate (ATP)-binding cassette sub-family B member 1 (ABCB1). Active pharmaceutical ingredients, natural constituents, and pharmaceutically inert excipients have been widely studied as P-gp inhibitors. Three excipients ghee, badam oil and fenugreek oil were selected as excipients as there are enriched with p-gp substrates. Molecular docking studies were conducted to identify the binding energy involved in between p-gp and the components of selected excipients. The selected drugs affinity with p-gp was also estimated by molecular docking studies. The diffusion of selected drugs through the intestine membrane was also estimated. Good correlation was observed between the docking score and the estimated diffusion rate from the diffusion data. Thus, these studies conclude that the molecular docking studies are beneficial to select the suitable excipients for improving the bioavailability of reported p-gp substrates.

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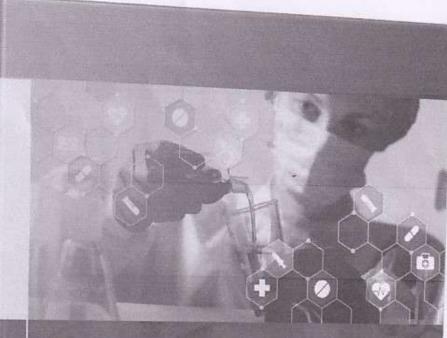








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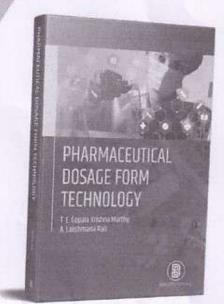
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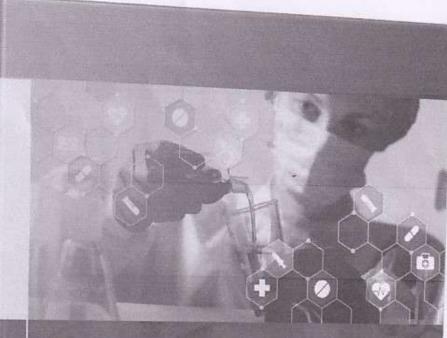
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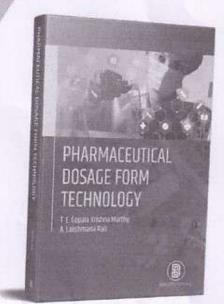
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# Current Trends in Drug Discovery, Development and Delivery (CTD4-2022)№

Edited by Manikanta Murahari; Buchi N Nalluri; G Chakravarthi

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This publication is based on peer-reviewed manuscripts from the 2022 Conference on Current Trends in Drug Discovery, Development and Delivery (CTD4-2022) held at KL University, India. Providing a wide range of up to date topics on the latest advancements in drug design and discovery technologies, this book ensures the reader receives a good understanding of the scope of the field. Aimed at scientists, students, regulators, academics and consultants throughout the world, this book is an ideal resource for anyone interested in the state of the art in drug design and discovery.

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Development and Validation of Stability Indicating Related Substances
Method for Montelukast Sodium and Levocetirizine Dihydrochloride
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p304-315

By K Ranjith; D. Rama Sekhara Reddy; TEGK Murthy

Abstract ∨

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Development of RP-HPLC-PDA Method Validation for the Simultaneous Estimation of Zidovudine, Lamivudine, and Nevirapine in Bulk and Dosage Forms and in Dissolution Samples

p316-325

By Potu. Kalyani; Buchi N. Nalluri; M. Chinna Eswaraiah; Tata Prasanna Kumari

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Part 3: Novel Drug Delivery Systems

Design, Development and Characterization of Eberconazole Proniosomes

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Design and Evaluation of Mucoadhesive in Situ Liposomal Gel for Sustained Ocular Delivery of Difluprednate Using Two Steps Factorial Design

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By K Ranjith; D. Rama Sekhara Reddy; TEGK Murthy DOI: https://doi.org/10.1039/9781837671090-00304

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In this paper authors presented the development and validation of RP-HPLC method for determination of related substances present in Montelukast sodium and Levocetirizine dihydrochloride combined tablets. Chromatographic separation was achieved on Hypersil ODS (150 x 4.6 mm),5  $\mu$  column with flow rate 1 mL/min, detection wavelength 238 nm, injection volume 10  $\mu$ L, column temperature 40°C and run time 75 mins. Gradient programming was used with mobile phase-A (MP-A)-50mM acetate buffer pH 5.5 and mobile phase-B(MP-B)-methanol: MP-A (90:10 %v/v). Developed method was validated as per ICH Q2(R1) guideline. As the method was able to analyse forced degradation samples successfully, it proved to be stability-indicating.

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# Pathophysiology of Neurodegenerative Diseases: Basics to Advanced

Sathish Kumar Manoharan & Poonguzhali Sathish Kumar

Chapter | First Online: 22 September 2020

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## Abstract

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