



Amita Yadav  
Kamal Singh Rathore  
Geeta M. Patel

# Formulation, Evaluation and Optimization of Mouth Dissolving Tablets



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# Formulation Evaluation Of Mouth Dissolving Tablets Of

**Dr.G.SANDHYARANI Guggilla**



## **Formulation Evaluation Of Mouth Dissolving Tablets Of:**

*Current Advances in Drug Delivery Through Fast Dissolving/Disintegrating Dosage Forms* Vikas Anand Saharan, 2017-05-11 Fast Dissolving Disintegrating Dosage Forms FDDFs have been commercially available since the late 1990s FDDFs were initially available as orodispersible tablets and later as orodispersible films for treating specific populations pediatrics geriatrics and psychiatric patients Granules pellets and mini tablets are among latest additions to these dosage forms which are still in the development pipeline As drug delivery systems FDDFs enable quicker onset of action immediate drug delivery and sometimes offer bioavailability benefits due to buccal sublingual absorption With time FDDF have evolved to deliver drugs in a sustained and controlled manner Their current market and application is increasing in demands with advances in age adapted dosage forms for different patients and changing regulatory requirements that warrant mandatory assessments of new drugs and drug products before commercial availability This book presents detailed information about FDDFs from their inception to recent developments Readers will learn about the technical details of various FDDF manufacturing methods formulation aspects evaluation and methods to conduct clinical studies The authors also give examples of marketed fast disintegrating dissolving drug products in US Europe Japan and India This reference is ideal for pharmacology students at all levels seeking information about this specific form of drug delivery and formulation

*Formulation and Evaluation of Mouth Dissolving Tablets* Reshu Gupta, 2013 Oral delivery is currently the gold standard in the pharmaceutical industry where it is regarded as the safest most convenient and most economical method of drug delivery having the highest patient compliance This tablet format is designed to allow administration of an oral solid dose form in the absence of water or fluid intake Such tablets readily dissolve or disintegrate in the saliva generally within **"Fast**

**dissolving tablets "** Dr.G.SANDHYARANI Guggilla, 2019-02-14 FORMULATION AND EVALUATION OF FAST DISSOLVING TABLETS ON RIZATRIPTON **Advances in Pharmaceutical Product Development** Keerti Jain, Awesh K.

Yadav, 2025-03-18 This book discusses the stages involved in pharmaceutical product development including the importance requirement and effect of each stage and process It also covers prototype development for pharmaceutical formulations scale up studies optimization testing packaging and commercialization of different dosage forms for pharmaceutical products like tablets suspensions emulsions coating inhalational products sterile products and herbal formulations The book also presents advancements in tablet production and tablet coating including materials material handling granulation and granulation technologies process automation processing problems in tablet production and troubleshooting advances in equipment for coating and coating materials Further the chapter explores the advances in the formulation and development of aerosols nebulizers inhalers metered Dose Inhalers MDI and dry powder Inhalers DPIs Towards the end the book examines the challenges formulation development testing stability and regulatory guidelines in the development of herbal formulations This book provides a valuable source of information for the researcher scientists students and people working in the area

mainly focused on the challenges in pharmaceutical product development      *Formulation, Evaluation and Optimization of Mouth Dissolving Tablets* Amita Yadav, Kamal Singh Rathore, Geeta M. Patel, 2014-08-09      **Handbook of Pharmaceutical Granulation Technology** Dilip M. Parikh, 2021-05-11 This fully revised edition of Handbook of Pharmaceutical Granulation Technology covers the rapid advances in the science of agglomeration process control process modelling scale up emerging particle engineering technologies along with current regulatory changes presented by some of the prominent scientist and subject matter experts around the globe Learn from more than 50 global subject matter experts who share their years of experience in areas ranging from drug delivery and pharmaceutical technology to advances in nanotechnology Every pharmaceutical scientist should own a copy of this fourth edition resource Key Features Theoretical discussions covering granulation and engineering perspectives Covers new advances in expert systems process modelling and bioavailability Chapters on emerging technologies in particle engineering Updated Current research and developments in granulation technologies      **Annual Report** University of Poona, 2010      Formulation and Evaluation Mouth Dissolving Tablets of Tramadol HCL Raja Sridhar Rao Ponugoti, Chandrashekara Rao Gonugunta, 2015-08-11 According to United States Pharmacopoeia the orodispersible tablets may be defined as solid dosage form containing medicinal substance or active ingredient which disintegrates rapidly within a matter of seconds when placed upon the tongue This means that the tablets dissolve or disintegrate in the oral cavity without use of water In this regard the tablets need to improve disintegration time dispersion time drug release studies bioavailability and patient compliance and also need to mask the bitter taste of the drug and to maintain the drug stable at accelerated condition i e 40 C 75% RH up to 6 months period as per ICH guidelines Tramadol HCl is centrally acting synthetic opioid analgesic for the treatment of moderate to severe pain and is readily soluble in water The half life of the drug is around 5.5 hours The MDT's place a major role for rapid onset of action for geriatrics pediatrics and the patients who have less access of water The drug itself having bitter taste so the present authors developed mouth dissolving tablets of tramadol HCl with the aim to mask the bitter taste of the drug to minimize the disintegration time and improve the drug release rate      *Indian Science Abstracts*, 2011-03      **Journal of Scientific and Industrial Research**, 2014      **Preparation and Evaluation of Mouth Dissolving Film of Metoclopramide Hcl** Dhaneshwar Kumar Vishwakarma, 2011-08 Oral drug delivery is the most widely utilized route of administration among the entire route that has been explored for the systemic delivery of drug via various pharmaceutical products of different dosage forms The conventional tablet seems to be most popular because of its ease of transportability and comparatively low manufacturing cost but poor patient compliance in case of pediatrics and geriatrics patients who experienced difficulties in swallowing in response to this mouth dissolving drugs delivery system MDDs were developed as an alternative to tablet cap syrups A variety of MDDs like mouth dissolving tablets and mouth dissolving film MDFs were commercialized MDFs evolved over the past few years from by the confection and oral care market in the form of breath strips became a novel widely

accepted form by consumers It consist very thin oral strips which release active ingredients immediately after uptake into oral cavity Today MDFs are a proven accepted technology for the systemic delivery of active pharmaceuticals ingredients

**Chemical Abstracts** ,2002      *Taste Masking and Formulation of Ondansetron Mouth Dissolving Tablets* Poudel Bhupendra Kumar,Subdei Shyam Raj,Thapa Panna,2015-11-23 The present work is an attempt to design mouth dissolving tablets of Ondansetron hydrochloride Drug Resin Complex was formulated by direct compression into mouth dissolving tablet using microcrystalline cellulose PH 102 as diluent and croscarmallose sodium and sodium starch glycolate as superdisintegrants while aspartame was used as sweetening agent to enhance palatability Taste masking was done by using polacrilex resin Tulsion 335 The prepared formulation can be useful for patients who have swallowing difficulty      *The Formulation and Evaluation of Orally Disintegrating Tablets* Stephanie M. Chillas,2013 Orally disintegrating tablets are a vital tool in keeping our children and elderly population healthy Their ease of use and accurate dosing allow higher patient compliance and more reliable therapeutic effects Superdisintegrants are the fundamental element contained in orally disintegrating tablets and are responsible for their unique ability to quickly disintegrate and dissolve on the surface of the tongue without the use of any additional liquid In order to determine the most effective type and optimal amount of superdisintegrants for orally disintegrating tablets manufactured by direct compression the following tablet parameters were tested based on the standard USP 30 methods hardness thickness friability disintegration time and wetting time Four superdisintegrants were tested namely Kollidon CL SF Primojel Ac Di Sol and Polyplasdone XL and the most efficient superdisintegrant was selected based on the above mentioned studies Polyplasdone XL outperformed the other superdisintegrants in nearly all the concentrations and all of the testing parameters The ideal concentration was 5% Polyplasdone XL for the orally disintegrating tablets before the API diphenhydramine HCl was added Various concentrations were analyzed using the aforementioned parameters and 10% Polyplasdone XL was selected for the final formulation Tablets were tested over eight weeks of accelerated stability testing at various temperatures to determine percent of diphenhydramine HCl released in dissolution disintegration time wetting time hardness thickness and weight variation The tablets remained stable and within the acceptable range of drug release 90% 110% at the three temperatures analyzed Additionally tablets were analyzed at different levels of relative humidity for percent weight change and change in hardness The final formulation of diphenhydramine HCl tablets was stable between 3 40 C and at 33% relative humidity or less for eight weeks      *The Lancet* ,1962      *Genetic Engineering News* ,2003      *Compendium of Pharmaceuticals and Specialties (Canada)* . ,1987 Presents information supplied by manufacturers and distributors about pharmaceuticals available for use in Canada Organized by trade name Includes a section on Canadian drug regulations      **Formulation And Evaluation Of Nebivolol Hcl Fast Dissolving Tablets** Vanitha Kondi,2015-06-16 The objective of the present study was the formulation and evaluation of Nebivolol Hcl fast dissolving tablet by solid dispersions Fast dissolving tablets are novel types of tablets

that dissolve disintegrate disperse in saliva within few seconds without water The major category of Nebivolol Hcl is in the treatment of hypertension adrenergic beta antagonist and vasodilator It is a poorly soluble and require enhancement of solubility and dissolution rate in its formulation development Drug Delivery to the Oral Cavity Tapash K. Ghosh,William R. Pfister,2005-02-28 With contributions from recognized authorities in industry academia and government this reference presents the state of the art in the testing formulation and clinical evaluation of intraoral drug delivery products summarizing intraoral dosage forms in various stages of research as well as products currently on the market **Formulation and Evaluation of Fast Dissolving Tablet of Thiabendazole** Pareek Ashish Kumar,2015-11-30 Fast Dissolving Tablets of Thiabendazole is designed for Providing the better and effective treatment against Helminthiasis Fast Dissolving Tablet of Thiabendazole is designed with the aim to enhance the bioavailability of the dosage form Helminthiasis infection is very common in urban areas and particularly in the childrens that are playing in soil so the Fast dissolving tablet of Thiabendazole provide cidal action by inhibiting the enzyme fumarate reductase so it provide a safest action and effective treatment

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