

Formulation Development and in Vitro Evaluation of Capecitabine Immediate Release Tablets

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ABSTRACT

The aim of this study is to formulate and significantly improve the bioavailability and reduce the side effects of immediate release tablets Capecitabine. The precompression blends of Capecitabine were characterized with respect to angle of repose, bulk density, tapped density, Carr's index and Hausner's ratio. The precompression blend of all the batches indicates good to fair flowability and compressibility. Immediate release tablets were prepared with various disintegrants like PEG 6000, Croscarmellose sodium and Sodium-starch glycolate at different concentration ratios and were compressed into tablets. The formulated tablets were evaluated for various quality control parameters. The tablets were passed all tests. Among all the formulations F7 formulation containing, drug and Croscarmellose sodium showed good result that is 98.12 % in 45 min. Hence from the dissolution data it was evident that F7 formulation is the better formulation.

KEYWORDS: Capecitabine, PEG 6000, Croscarmellose sodium and Sodium-starch glycolate, Immediate release Development

INTRODUCTION:

Oral route is the most convenient and extensively used for drug administration. Oral administration is the most popular route for systemic effects due to its ease of ingestion, pain avoidance, versatility and most importantly, patient compliance suitable for industrial production, improved stability and bioavailability. The concept of immediate release tablets emerged from the desire to provide patient with more conventional means of taking their medication when emergency treatment is required. Recently, immediate release tablets have gained prominence of being new drug delivery systems. The oral route of administration has so far received the maximum attention with respect to research on physiological and drug constraints as well as design and testing of product. Drug delivery systems (DDS) are a strategic tool for expanding markets/indications, extending product life cycles and generating opportunities. Most immediate release tablets are intended to disintegrate in the stomach, where the pH is acidic. Several orally disintegrating

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tablet (ODT) technologies based on direct compression. In pharmaceutical formulation includes any formulation in which the rate of release of drug from the formulation is at least 70% (preferably 80%) of active ingredient within 4 hours, such as within 3 hours, preferably 2 hours, more preferably within 1.5 hours, and especially within an hour (such as within 30 minutes) of administration. In Formulation of immediate release the commonly Superdisintegrants used are Croscarmellose, sodium, Sodium Starch glycolate and Crospovidone.¹

Oral route of administration is the most popular route for systemic effects due to its ease of ingestion, pain avoidance, versatility and most importantly, patient compliance. Also solid oral delivery systems does not need sterile conditions and are therefore, less expensive to manufacture. Patient compliance, high precision dosing, and manufacturing efficiency make tablets the solid dosage form of choice. There is

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Table of Contents Formulation Development And Evaluation Of Immediate

1. Understanding the eBook Formulation Development And Evaluation Of Immediate
 - The Rise of Digital Reading Formulation Development And Evaluation Of Immediate
 - Advantages of eBooks Over Traditional Books
2. Identifying Formulation Development And Evaluation Of Immediate
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Formulation Development And Evaluation Of Immediate
 - User-Friendly Interface
4. Exploring eBook Recommendations from Formulation Development And Evaluation Of Immediate

- Personalized Recommendations
 - Formulation Development And Evaluation Of Immediate User Reviews and Ratings
 - Formulation Development And Evaluation Of Immediate and Bestseller Lists
5. Accessing Formulation Development And Evaluation Of Immediate Free and Paid eBooks
 - Formulation Development And Evaluation Of Immediate Public Domain eBooks
 - Formulation Development And Evaluation Of Immediate eBook Subscription Services
 - Formulation Development And Evaluation Of Immediate Budget-Friendly Options
 6. Navigating Formulation Development And Evaluation Of Immediate eBook Formats
 - ePub, PDF, MOBI, and More
 - Formulation Development And Evaluation Of Immediate Compatibility with Devices
 - Formulation Development And Evaluation Of Immediate Enhanced eBook Features
 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Formulation Development And Evaluation Of Immediate
 - Highlighting and Note-Taking Formulation Development And Evaluation Of Immediate
 - Interactive Elements Formulation Development And Evaluation Of Immediate
 8. Staying Engaged with Formulation Development And Evaluation Of Immediate
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Formulation Development And Evaluation Of Immediate
 9. Balancing eBooks and Physical Books Formulation Development And Evaluation Of Immediate
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection Formulation Development And Evaluation Of Immediate
 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
 11. Cultivating a Reading Routine Formulation Development And Evaluation Of Immediate
 - Setting Reading Goals Formulation Development And Evaluation Of Immediate
 - Carving Out Dedicated Reading Time
 12. Sourcing Reliable Information of Formulation Development And Evaluation Of Immediate

- Fact-Checking eBook Content of Formulation Development And Evaluation Of Immediate
 - Distinguishing Credible Sources
13. Promoting Lifelong Learning
- Utilizing eBooks for Skill Development
 - Exploring Educational eBooks
14. Embracing eBook Trends
- Integration of Multimedia Elements
 - Interactive and Gamified eBooks

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