

Research Article of Formulation and Evaluation of Fast Dissolving Tablet of Nitrendipine

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ABSTRACT

Objective: The aim of present study is to formulate fast dissolving tablet of Nitrendipine, the drug will be directly absorbed into systemic circulation through buccal mucosa and lead to produce immediate action.

Methods: Fast dissolving tablets of Nitrendipine were prepared by wet granulation method. Required quantity of drug and other excipients were weighed and sieved from sieve no.60 for finding homogenous mixer, then a damp mass of mixer was prepared by using distilled water as a solvent. Damp mass was passed through sieve no. 10 and dried the granules at 50 °C till moisture remaining less than 2%.

Results: All the formulated tablets met the pharmacopoeias standard of uniformity of weight, percentage friability, thickness, and drug content. The in vitro disintegration and dispersion studies were also performed, which shows very good bioavailability and drug release profile. Accelerated stability studies were done for four weeks and found that no significant change in drug content and other parameters like hardness and in vitro dispersion time after four weeks even at 50 °C. It may be predicted that formulation will be stable for more than one year.

Conclusion: The present investigation successfully formulated mouth dissolving tablets of Nitrendipine with improved drug release profile. The formulation was chosen because it showed good results in terms of cumulative drug release, disintegration time, hardness and friability. The dissolution study of this formulation showed an increase in the cumulative % drug release.

KEYWORDS: Fast dissolving tablets, Nitrendipine, Bioavailability, Wet granulation method, Carr's index

INTRODUCTION

Fast dissolving tablets is a solid dosage form that dissolves or disintegrates within a minute in the oral cavity without the need of water and has a pleasant taste. FDT is also known as an orally disintegrating tablet, fast-dissolving tablet, fast-melting tablet, mouth melting tablet or fast-disintegrating tablet. Fast disintegrating dosage form has been successfully commercialized, and the growing importance was highlighted recently when the European Pharmacopoeia adapted the term 'or dispersible tablets' as a tablet to be placed in the mouth where it disperses rapidly before swallowing [1, 2]. FDTs are designed to disintegrate or dissolve rapidly on contact

with saliva, thus eliminating the need for chewing the tablet, swallowing an intact tablet, or taking the tablet with water. Although no water is needed to allow the drug to disperse quickly and efficiently, most technologies utilize the body's own salivation. This mode of administration was initially expected to be beneficial to pediatric and geriatric patients, to people with conditions related to impaired swallowing, and for the treatment of patients when compliance may be difficult (e. g. psychiatric disorders). FDT has previously been distinguished as a separate dosage form because of the specific, intended performance characteristics of such products, which are rapid oral

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Table of Contents Research Article Formulation Development And Evaluation Of

1. Understanding the eBook Research Article Formulation Development And Evaluation Of
 - The Rise of Digital Reading Research Article Formulation Development And Evaluation Of
 - Advantages of eBooks Over Traditional Books
2. Identifying Research Article Formulation Development And Evaluation Of
 - 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 Research Article Formulation Development And Evaluation Of
 - User-Friendly Interface
4. Exploring eBook Recommendations from Research Article Formulation Development And Evaluation Of

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

- Fact-Checking eBook Content of Research Article Formulation Development And Evaluation Of
- 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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