

Table 3.2: Dissolution Study of Pure Drug

Time (min.)	Cumulative % drug release
0	0.00
5	10.25±0.64
10	11.12±0.44
15	13.07±0.91
20	15.20±0.75
25	16.59±0.14
30	18.37±0.42
40	21.35±0.34

Results are the mean of three determinations

The % drug release of pure drug after 40 min was 21.35% each reading is taken was triplicate and then mean values were calculated.

2. Dissolution Profile of Solid Dispersions Prepared by Kneading Method

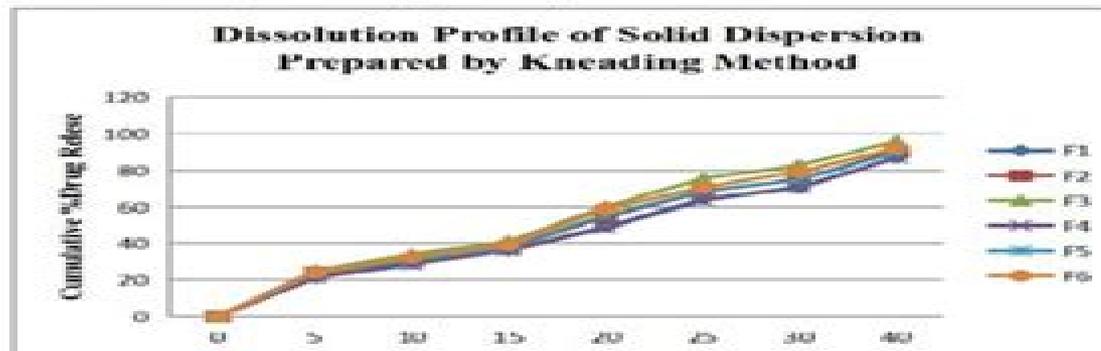
The formulations of solid dispersions prepared by Kneading method(F1-F6) were subjected to dissolution study. The percentage drug release of formulations is showed in Table 10.12 and accordingly the graph was plotted to calculate the percentage drug release of formulations in 0.1N HCl and it is shown in Figure 2.8.

Table 3.3: Dissolution Profile of Solid Dispersions Prepared by Kneading Method

Time(min)	Cumulative % Drug Release					
	F1	F2	F3	F4	F5	F6
0	00	00	00	00	00	00
5	31.54±0.21	33.27±0.25	36.43±0.22	30.31±0.57	32.04±0.35	33.45±0.20
10	44.28±0.25	47.13±0.14	48.53±0.32	43.00±0.42	45.13±0.54	48.32±0.23
15	55.02±0.13	59.04±0.46	61.08±0.71	68.06±0.35	59.10±0.26	59.25±0.45
20	68.18±0.18	70.76±0.51	72.17±0.64	75.80±0.18	69.76±0.46	71.86±0.64
25	77.96±0.94	80.17±0.15	82.50±0.24	82.50±0.78	76.03±0.37	80.39±0.32
30	85.70±0.34	87.93±0.42	90.36±0.82	85.05±0.46	84.63±0.48	88.24±0.24
40	90.93±0.21	92.92±0.81	96.61±0.37	89.02±0.63	91.85±0.78	94.61±0.37

Results are the mean of three determinations

Out of Six formulations F3 showed maximum drug release i.e. 96.61 %. Solid dispersion (F3) of Carvedilol with PEG 6000 prepared by Kneading method showed significant improvement in solubility and dissolution rate. Increased wetting and solubilizing effect of PEG 6000 as well as the molecular dispersion of drug in solid dispersion and alteration of surface properties of drug particle may be responsible for the enhanced dissolution rate of Carvedilol from solid dispersion compared to pure Carvedilol.

**Figure 2.8: Dissolution Profile of Solid Dispersions Prepared by Kneading Method**

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S Marginson



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Handbook of Modern Pharmaceutical Analysis Satinder Ahuja, Stephen Scypinski, 2001 This book describes the role modern pharmaceutical analysis plays in the development of new drugs Detailed information is provided as to how the quality of drug products is assured from the point of discovery until the patient uses the drug Coverage includes state of the art topics such as analytics for combinatorial chemistry and high throughput screening formulation development stability studies international regulatory aspects and documentation and future technologies that are likely to impact the field Emphasis is placed on current easy to follow methods that readers can apply in their laboratories No book has effectively replaced the very popular text Pharmaceutical Analysis that was edited in the 1960s by Tak Higuchi This book will fill that gap with an up to date treatment that is both handy and authoritative

A Textbook of Cosmetic Science Dr. (Prof.) Sailesh Kumar Ghatuary , Dr. Gaurav Dubey , Dr. Satkar Prasad, 2024-07-13 Introducing the book Cosmetic Science is something that fills me with an incredible amount of joy The content of this book has been meticulously crafted to adhere to the curriculum for Bachelor of Pharmacy students that has been outlined by the Pharmacy Council of India An effort has been made to investigate the topic using terminology that is as straightforward as possible in order to make it more simply digestible for pupils The book has a number of illustrations such as flowcharts and diagrams that make it simple for students to comprehend complex ideas It is the author s honest desire that both students and academicians would take something helpful away from reading this book

Pharmaceutical Drug Product Development and Process Optimization Sarwar Beg, Majed Al Robaian, Mahfoozur Rahman, Syed Sarim Imam, Nabil Alruwaili, Sunil Kumar Panda, 2020-05-01 Pharmaceutical manufacturers are constantly facing quality crises of drug products leading to an escalating number of product recalls and rejects Due to the involvement of multiple factors the goal of achieving consistent product quality is always a great challenge for pharmaceutical scientists This volume addresses this challenge by using the Quality by Design QbD concept which was instituted to focus on the systematic development of drug products with predefined objectives to provide enhanced product and process understanding This volume presents and discusses the vital precepts underlying the efficient effective and cost effective development of pharmaceutical drug products It focuses on the adoption of systematic quality principles of pharmaceutical development which is imperative in achieving continuous improvement in end product quality and also leads to reducing cost time and effort while meeting regulatory requirements The volume covers the important new advances in the development of solid oral dosage forms modified release oral dosage forms parenteral dosage forms semisolid dosage forms transdermal drug delivery systems inhalational dosage forms ocular drug delivery systems nanopharmaceutical products and nanoparticles for oral delivery

Drug Development Chris Rundfeldt, 2011-12-07 This book represents a case study based overview of many different aspects of drug development ranging from target identification and characterization to chemical optimization for efficacy and safety as well as bioproduction of natural products utilizing for example lichen In

the last section special aspects of the formal drug development process are discussed Since drug development is a highly complex multidisciplinary process case studies are an excellent tool to obtain insight in this field While each chapter gives specific insight and may be read as an independent source of information the whole book represents a unique collection of different facets giving insight in the complexity of drug development *PHARMACOGNOSY- I: A TEXTBOOK* ,2025-06-17

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