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MPharm Project work 2023-2024

SL NO	Programme name	Program Code	Link the documents
1	MPharm	QA	Click Here

**SEQUENTIAL DEVELOPMENT OF LURASIDONE SELF EMULSIFYING DRUG
DELIVERY SYSTEM IN A QbD PARADIGM**

M. PHARM DISSERTATION

SUBMITTED TO THE



**RAJIV GANDHI UNIVERSITY OF HEALTH SCIENCES, BENGALURU,
KARNATAKA**

BY

ANUSHA.S

22PQ053

(M. PHARM-QUALITY ASSURANCE)

UNDER THE GUIDANCE OF

Dr. KAVITHA A N

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This is to certify that the dissertation entitled is "SEQUENTIAL DEVELOPMENT OF LURASIDONE SELF EMULSIFYING DRUG DELIVERY SYSTEM IN A QbD PARADIGM" a bonafide research work done by Ms. ANUSHA S under the guidance of Dr. KAVITHA A N, Professor and HOD, Department of Quality Assurance, Krupanidhi College of Pharmacy-560035


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Place: Bengaluru

Date: 24/10/2024

Place: Bengaluru

**QbD APPROACH ON CONCEPTUALIZATION AND CHARACTERIZATION OF
ORALLY DISINTEGRATING TABLET OF ELETRIPTAN**

by

KAVYA S M

22PQ054

**Dissertation Submitted to the Rajiv Gandhi University of Health Sciences, Karnataka,
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In partial fulfillment of the requirements for the degree of

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**RP HPLC BASED METHOD DEVELOPMENT, VALIDATION AND STABILITY
INDICATING ASSAY OF OTESECONAZOLE IN BULK DRUG USING ANALYTICAL
QUALITY BY DESIGN APPROACH**

by

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**DESIGN AND DEVELOPMENT OF TRANSDERMAL PATCH OF
LUMATEPERONE IN A QUALITY BY DESIGN APPROACH**

by

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**DEVELOPMENT AND CHARACTERIZATION OF MUCOADHESIVE FAST
DISSOLVING BUCCAL FILM OF ASENAPINE MALEATE IN A QUALITY BY
DESIGN PARADIGM**

by

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**THE NOVEL APPROACH TO DEVELOP RP-HPLC METHOD VALIDATION AND STABILITY
INDICATING ASSAY OF VERICIGUAT DRUG BY ANALYTICAL QUALITY BY
DESIGN(AQbD)**

by

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**DEVELOPMENT AND VALIDATION OF STABILITY INDICATING
RP-HPLC METHOD FOR DETERMINATION OF DARIDOREXANT
DRUG USING AQbD APPROACH**

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**QbD APPROACH IN FORMULATION AND EVALUATION OF ALCAFTADINE
OPHTHALMIC *IN-SITU* GEL**

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**ANALYTICAL METHOD DEVELOPMENT, VALIDATION AND STABILITY-
INDICATING ASSAY OF FAVIPRAVIR A SAR COV-2 MOLECULE BY RP-
HPLC USING ANALYTICAL QUALITY BY DESIGN APPROACH**

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**ANALYTICAL QUALITY BY DESIGN APPROACHES TO RP-HPLC
METHOD DEVELOPMENT VALIDATION AND STABILITY
INDICATING ASSAY FOR THE DETERMINATION OF FINERENONE
IN BULK DRUG**

by

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