METHOD DEVELPOMENT AND VALIDATION OF STABILIZERS BY USING UV-VISIBLE SPECTROPHOTOMETER AND QUANTIFICATION OF THEM IN FOOD PRODUCTS

IN PARTIAL FULFILLMENT OF THE REQUIREMENTS FOR THE AWARD
OF THE DEGREE OF



MASTER OF PHARMACY

IN

PHARMACEUTICAL ANALYISIS

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This is to certify that the dissertation entitled "Method Development and Validation Of Stabilizers by Using UV-Visible Spectrophotometer And Quantification Of Them In Food Products" is submitted to Osmania University for the partial fulfilment of requirements for the award of the degree of Master Of Pharmacy in Pharmaceutical Analysis under Faculty of Pharmacy embodies the results and studies of a bonafide research work of <u>B.Anila</u> under my supervision at RBVRR Women's College of Pharmacy and the contents of the thesis do not form the basis for the award of any other degree or diploma to the candidate from this or any other university elsewhere.

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Date: 22/12/2023

Place: Hyderabad

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UV-VISIBLE AND RP-HPLC METHOD DEVELOPMENT AND VALIDATION OF MOLNUPIRAVIR A COVID DRUG IN BULK AND PHARMACEUTICAL DOSAGE FORM

THE DISSERTATION SUBMITTED TO OSMANIA UNIVERSITY



IN PARTIAL FULFILMENT OF THE REQUIREMENTS FOR THE AWARD OF THE DEGREE OF

MASTER OF PHARMACY

IN

PHARMACEUTICAL ANALYSIS

BY

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ANALYTICAL METHOD DEVELOPMENT AND VALIDATION OF SUGAMMADEX IN BULK AND PHARMACEUTICAL DOSAGE FORM USING UV-VISIBLE SPECTROPHOTOMETRY AND RP-HPLC

THE DISSERTATION SUBMITTED TO OSMANIA UNIVERSITY



IN PARTIAL FULFILLMENT OF THE REQUIREMENTS FOR THE AWARD OF THE DEGREE OF

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IN

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DEVELOPMENT AND VALIDATION OF UV AND

RP-HPLC METHOD FOR DETERMINATION OF DAPAGLIFLOZIN AND VILDAGLIPTIN IN BULK AND COMBINED TABLET DOSAGE FORM INCLUDING STRESS STUDIES

IN PARTIAL FULFILLMENT OF THE REQUIREMENTS FOR
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IN

PHARMACEUTICAL ANALYSIS

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This is to certify that the dissertation entitled "Development and validation UV and RP-HPLC method for quantitative estimation of Dapagliflozin and Vildagliptin in Bulk and combined tablet dosage form with application to dissolution and stress studies" is submitted to Osmania University for the partial fulfilment of requirements for the award of the degree of Master Of Pharmacy in Pharmaceutical Analysis under Faculty of Pharmacy embodies the results and studies of a bonafide research work of M. Nandhini under my supervision at RBVRR Women's College of Pharmacy and the contents of the thesis do not form the basis for the award of any other degree or diploma to the candidate from this or any other university elsewhere.

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STABILITY INDICATING METHOD DEVELOPMENT AND VALIDATION FOR THE ESTIMATION OF BARICITINIB IN BULK AND IN FORMULATION USING UV-VISIBLE SPECTROSCOPY, AND RP-HPLC.

THE DISSERTATION SUBMITTED TO OSMANIA UNIVERSITY



IN PARTIAL FULFILLMENT OF THE REQUIREMENTS FOR THE AWARD OF THE DEGREE OF

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IN

PHARMACEUTICAL ANALYSIS

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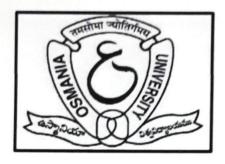
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Place: Hyderabad

by 29/12/23. CD8-y. Padmarathi)

METHOD DEVELOPMENT AND VALIDATION FOR THE QUANTIFICATION OF ARTIFICIAL SWEETENERS IN THE FOOD PRODUCTS BY USING UV-VISIBLE SPECTROPHOTOMETER

IN PARTIAL FULFILLMENT OF THE REQUIREMENTS FOR THEAWARD OF THE DEGREE OF



MASTER OF PHARMACY

IN

PHARMACEUTICAL ANALYSIS

Submitted by

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This is to certify that the dissertation entitled "Method Development and Validation for the Quantification of Artificial Sweeteners in the Food products by using UV-Visible Spectrophotometer" is submitted to Osmania University for the partial fulfilment of requirements for the award of the degree of Master Of Pharmacy in Pharmaceutical Analysis under Faculty of Pharmacy embodies the results and studies of a bonafide research work of Samudrala Saileela under my supervision at RBVRR Women's College of Pharmacy and the contents of the thesis do not form the basis forthe award of any other degree or diploma to the candidate from this or any other university elsewhere.

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NEW METHOD DEVELOPMENT AND VALIDATION FOR OXACEPROL DRUG IN BULK AND DOSAGE FORM USING UV-VISIBLE SPECTROSCOPY & RP-HPLC INCLUDING STRESS STUDIES

IN PARTIAL FULFILLMENT OF THE REQUIREMENTS FOR THE AWARD OF THE DEGREE OF



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