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71)Name of Applicant:

1)Mr. Munna Singh

Address of Applicant :Assistant Professor, School of Pharmaceutical Sciences, Faculty of Pharmacy, IFTM University Lodhipur Rajput, Moradabad, Uttar Pradesh, Pin Code: 244102. -

2)Dr. Sushil Kumar

3)Mr. Raj Kumar Singh Bharti

4)Mr. Sateesh Kumar 5)Mrs. Soniya Sagar

6)Mrs. Deepa Sharma

7)Mr. Saiyad Arsh Zia 8)Dr.Prashant Kumar

9)Mr. Manzar Abbas

10)Mr. Anesh Sagar

Name of Applicant : NA

Address of Applicant : NA (72)Name of Inventor :

1)Mr. Munna Singh

Address of Applicant :Assistant Professor, School of Pharmaceutical Sciences, Faculty of Pharmacy, IFTM University Lodhipur Rajput, Moradabad, Uttar Pradesh, Pin Code: 244102. -

2)Dr. Sushil Kumar

Address of Applicant :Professor, School of Pharmaceutical Sciences, Faculty of Pharmacy, IFTM University

Lodhipur Rajput, Moradabad, Uttar Pradesh, Pin Code: 244102. 3)Mr. Raj Kumar Singh Bharti

Address of Applicant : Assistant Professor, School of Pharmaceutical Sciences, Faculty of Pharmacy, IFTM

University Lodhipur Rajput, Moradabad, Uttar Pradesh, Pin Code: 244102. -

4)Mr. Sateesh Kumar

Address of Applicant : Associate Professor, Radha Govind College of Pharmacy, Near 2km RTO Office, Moradabad, Ûttar Pradesh, Pin Code: 244102 -

5)Mrs. Soniya Sagar

Address of Applicant : Assistant Professor, Radha Govind College of Pharmacy, Near 2km RTO Office, Moradabad, Uttar Pradesh, Pin Code: 244102 -------

6)Mrs. Deepa Sharma Address of Applicant :Assistant Professor, Radha Govind College of Pharmacy, Near 2km RTO Office, Moradabad, Uttar Pradesh, Pin Code: 244102 -------

7)Mr. Saiyad Arsh Zia

Address of Applicant :Assistant Professor, Chandpur Pharmacy College, Chandpur, Bijnor, Uttar Pradesh, Pin Code:246725.

8)Dr.Prashant Kumar

Address of Applicant : Assistant Professor, Sahoo Onkar Saran School of Pharmacy, Faculty of Pharmacy,

IFTM University Lodhipur Rajput, Moradabad, Uttar Pradesh, Pin Code:244102. -

9)Mr. Manzar Abbas

Address of Applicant :Assistant Professor, Rukmani College of Pharmacy, Lodhipur Rajput, Moradabad, Uttar

Pradesh, Pin Code: 244102 -

10)Mr. Anesh Sagar

Address of Applicant :Assistant Professor, School of Pharmaceutical Sciences, Faculty of Pharmacy, IFTM University Lodhipur Rajput, Moradabad, Uttar Pradesh, Pin Code: 244102. ----

(57) Abstract:

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The present invention relates to the preparation of itraconazole-loaded microfibres using a centrifugal melt spinning process, incorporating varying drug concentrations (20%, 30%, 50%, and 100% w/w) with sucrose as a carrier. The microfibres exhibit high drug loading, efficient incorporation, and maintain the drug in an amorphous state, enhancing solubility and bioavailability. Stability tests under high-humidity conditions demonstrate the microfibres' structural integrity and resistance to recrystallization. The dissolution profiles of the microfibres show controlled and sustained drug release. The fabrication method yields high-quality microfibres with minimal material loss and is scalable for large-scale production. This invention offers a versatile and efficient platform for formulating itraconazole in microfibre form, with applications in pharmaceutical drug delivery systems.

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