TO MAGNIFICO RETTORE OF UNIVERSITA' DEGLI STUDI DI MILANO **ID CODE - 6862**

I the undersigned asks to participate in the public selection, for qualifications and examinations, for the awarding of a type B fellowship at **Dipartimento di _ Scienze Farmacologiche e Biomolecolari dell'Università degli Studi di Milano**

Scientist-in-charge: **Prof. Casula Manuela**

[Name and surname]

CURRICULUM VITAE

PERSONAL INFORMATION

Surname	Gandhare
Name	Renuka

PRESENT OCCUPATION

Appointment	Structure
Student	University of Florence

EDUCATION AND TRAINING

Degree	Course of studies	University	year of achievement of the degree
Degree	Bachlers in Pharmacy	Nagpur University	2014
Specialization			
PhD			
Master	Masters in Pharmcology	Nagpur University	2016
Degree of medical specialization			
Degree of European specialization			
Other	Masters in advanced molecular science	University of Florence	2024

REGISTRATION IN PROFESSIONAL ASSOCIATIONS

Date of registration	Association	City
31/Dec/2019	Maharashtra State of Pharmacy Council	Nagpur

FOREIGN LANGUAGES

Languages	level of knowledge
English	C2
Hindi	C2
Marathi	C2
italian	A1

TRAINING OR RESEARCH ACTIVITY

description of activity

1. Thesis Studies on the role of withania somnifera in progesterone withdrawal induced anxiety in rats
2. Establishment of a selectable prostate cancer cell line expressing EGFP for vivo cell tracking.

CONGRESSES AND SEMINARS

Date	Title	Place
04/01/2013 06/01/2013	45th Annual Conference of Indian Pharmacological Society	Nagpur
06/01/2015	Rajiv Gandhi National Institute of Intellectual Property Management	Nagpur
25/09/2015	Advocacy and Awareness of Clinical Trials in India	Nagpur
26/05/2014	Clinical Research Emerging Carrier Avenues	Nagpur
20/12/2013	Pharmaceutical Dosage Form Design and Development: Challenges and Regulatory Constraints	Nagpur

OTHER INFORMATION

DRUG SAFETY ASSOCIATE COGNIZANT - Responsible for handling the case like Spontaneous Reports: Led the processing and evaluation of spontaneous reports, facilitating the identification of potential safety signals and contributing to risk assessment activities.

Clinical Trials: Managed adverse event data from clinical trials, ensuring compliance with regulatory requirements and providing critical insights into the safety profile of investigational drugs.

Literature Cases: Conducted comprehensive literature reviews and analysis to identify relevant adverse event data published in scientific literature, supporting ongoing pharmacovigilance monitoring efforts. Social Media Monitoring: Implemented strategies for monitoring social media platforms for discussions related to adverse events and medication safety, enhancing signal detection capabilities and proactive risk management.

05/05/2018 – 13/07/2019 Mysore, India

SENIOR DRUG SAFETY ASSOCIATE BIOCLINICA - Lead a high-performing case processing team, overseeing the collection and evaluation of adverse event reports for pharmaceutical products in compliance with regulatory standards. Implemented streamlined processes and quality control measures resulting in a significant reduction in report processing time and increased efficiency. Developed and maintained comprehensive risk management plans for products, collaborating cross functionally to identify and mitigate potential safety risks. Ensured timely submission of adverse event reports to regulatory authorities, consistently meeting or exceeding specified timelines and regulatory requirements. Actively participated in industry conferences and workshops to stay updated on evolving pharmacovigilance regulations and best practices, driving continuous improvement initiatives within the organization

13/08/2019 – 13/09/2022 Mumbai, India

SENIOR PROCESS ASSOCIATE TCS - Document Management: Managing the organization, indexing, and filing of electronic documents within the eTMF, including essential trial documents such as protocols, investigator brochures, informed consent forms, and regulatory submissions. Quality Control: Implementing quality control measures to ensure the accuracy, completeness, and integrity of documents stored in the eTMF. This involves conducting regular reviews, audits, and reconciliation of documents to identify and address any discrepancies or deficiencies.

Declarations given in the present curriculum must be considered released according to art. 46 and 47 of DPR n. 445/2000.

The present curriculum does not contain confidential and legal information according to art. 4, paragraph 1, points d) and e) of D.Lgs. 30.06.2003 n. 196.

Please note that CV WILL BE PUBLISHED on the University website and It is recommended that personal and sensitive data should not be included. This template is realized to satisfy the need of publication without personal and sensitive data.

Please DO NOT SIGN this form.

Place and date: _____, _____