GREESHMA S PATTABHI

M.Pharm (Pharmacology)

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Professional Summary

M.Pharm (Pharmacology) graduate with professional experience at Novo Nordisk, supporting global postmarketing quality processes while ensuring compliance with GMP, ICH, FDA, and EMA quality standards requirements. Strong understanding of clinical research, data quality, regulatory documentation, and cross-functional collaboration gained through both industry and academic experience. Hands-on experience in preclinical pharmacology, including study execution, data collection, analysis, and scientific reporting. Detail-oriented professional with excellent analytical, problem-solving, and documentation skills, eager to contribute to Clinical Data Management by ensuring accurate, reliable, and high-quality clinical data.

Education

- Master of Pharmacy in Pharmacology, Sri Adichunchanagiri College of Pharmacy (2023-2025) – CGPA: 8.85
- Bachelor of Pharmacy, Sri Adichunchanagiri College of Pharmacy (2019–2023) – CGPA: 7.6

Work Experience

Novo Nordisk

Contract Employee – Quality Department *Feb 2026 – Aug 2026*

- Managed global post-marketing Product Quality Complaints (PQCs) for pharmaceutical products and medical devices, ensuring timely and compliant case handling in the NOVGLOW quality management system.
- Performed complaint evaluation, categorization, and coding in line with corporate QMS and global regulatory requirements.
- Collaborated with manufacturing sites, affiliates, and global QA teams to support investigations and root cause analysis.
- Coordinated with cross-functional stakeholders to obtain required information and ensure timely complaint closure.
- Supported complaint trend analysis, batch reviews, and CAPA implementation while maintaining accurate documentation in NOVGLOW and Vault Quality.

Karnataka Antibiotics & Pharmaceuticals Limited

Intern *Aug 2022 – Sep 2022*

- Gained knowledge about the Raw Materials Production department – Oral dosage forms, Solid dosage forms, Parenterals, etc.
- Exposure to Packaging materials and Quality Control processes.

Projects

- Evaluation of invitro antidiabetic activity using silver nano particles of Hemidesmus indicus extract (B Pharm)
- Exploring the Therapeutic Potential of Acorus Calamus Transferosomal Hydrogel as a Burn Healing Agent (M Pharm)

Certificates & Conferences

- Basic Research in Zebrafish
- National conference on 'Current Perspectives of 4Rs in Alternatives to Animal Testing' (presented)
- International conference on Principles of Research Concepts, Biostatistics and Scientific Publishing (presented)
- International e-Conference on Emerging Trends in Pharmaceutical Sciences – Research to Market (presented)

Technical Skills

- Basic knowledge of Clinical Data Management
- Good Pharmacovigilance Practices (GVP)
- Knowledge of regulatory guidelines for drug metabolism and pharmacokinetics
- Knowledge of clinical trial phases & study workflow

- Knowledge of ICH-GCP, FDA, EMA guidelines
- Understanding of ICH E2A/E2D guidelines
- SOP development and compliance management
- Knowledge about protocol preparation
- Adverse event reporting (AER)
- Proficient in MS Word, Excel, PowerPoint

Professional Skills

- Effective communication and teamwork
- Problem-solving
- Time management
- Self-motivated
- Attention to detail and data accuracy

Linguistic Proficiency

- English – Fluent
- Kannada – Fluent
- Telugu – Basic
- Hindi – Basic