PRATIK GUNWANT BOBADE

Clinical Trial Assistant | Clinical Research Associate | Clinical Research Coordinator

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PROFESSIONAL SUMMARY

Detail-oriented Clinical Data Associate and B.Pharmacy graduate with an Advanced Diploma in Clinical Research and Data Management. Proficient in EDC tools (Medidata Rave, Oracle InForm, Octalsoft), ICH-GCP E6(R3) compliance, CRF design and review, query management, and clinical documentation. Hands-on experience as a Clinical Research Coordinator Intern with patient recruitment, TMF/ISF handling, AE/SAE reporting, and site coordination. Seeking an entry-level CDA or CDM role within a CRO or pharmaceutical company to contribute to high-quality clinical trial data management.

CORE COMPETENCIES

- Subject Screening, Enrollment & Visit Coordination
- Informed Consent Process (ICF) & Re-consent Management
- CRF/EDC Data Entry, Data Cleaning & Query Resolution
- Regulatory Documentation (Essential Documents, TMF/ISF)
- IEC Submissions & Amendments
- Monitoring Visit & Audit Readiness Support
- Protocol Deviation Tracking & CAPA Support
- Source Documentation & Site File Management
- Compliance with ICH-GCP, SOPs & Protocols
- EDC TOOLS : (Octalsoft Inform)

PROFESSIONAL EXPERIENCE

Clinical Research Coordinator — Intern Dec 2025 – May 2026

SMO: Accede Clinicals | Site: Arneja Heart and Multispeciality Hospital, Nagpur, Maharashtra

- Coordinated patient recruitment and managed clinical and regulatory documentation for ongoing Phase II–IV trials, ensuring protocol compliance throughout.
- Operated EDC tools — Oracle InForm, Medidata Rave, and Octalsoft — for data entry, data cleaning, and query resolution in alignment with ICH-GCP E6(R3).
- Maintained Trial Master File (TMF) and Investigator Site File (ISF), ensuring audit-readiness and adherence to regulatory standards at all times.
- Reviewed and processed Informed Consent Forms (ICF), safeguarding patient rights and protocol adherence.
- Documented and reported Adverse Events (AE) and Serious Adverse Events (SAE) in coordination with site investigators and sponsors.
- Managed sample shipment logistics and upheld Good Documentation Practices (GDP) across the full trial lifecycle.

Pharmacist Aug 2025 – Dec 2025

Apollo Pharmacy | Nagpur, Maharashtra

- Reviewed and accurately processed prescriptions, ensuring full compliance with pharmacy regulations and dispensing standards.
- Managed billing, stock control, and drug file records, demonstrating strong data accuracy and organizational skills.

Assistant Pharmacist Oct 2019 – Oct 2022

Gajanan Medical M+M | Chimur, Chandrapur, Maharashtra

- Dispensed medications with strict prescription compliance; managed inventory, storage, and billing using Marg Software.
- Maintained accurate stock records and proper drug storage conditions, reflecting a strong attention to detail and regulatory awareness.

EDUCATION

Advanced Diploma in Clinical Research & Data Management 2025–2026

IICRM, Nagpur

Bachelor of Pharmacy (B.Pharm) 2022–2025

DBATU University | CGPA: 7.73

Diploma in Pharmacy (D.Pharm) 2021–2022

MSBTE | 65.70%

Higher Secondary Certificate (HSC) 2019

Maharashtra State Board | 45.85% 2017

Secondary School Certificate (SSC)

Maharashtra State Board | 64.20%

CERTIFICATIONS

- **GCP Certification — NIDA Clinical Trial Network** — Valid: Feb 2026 – Feb 2029
- **ICH Good Clinical Practice E6(R3)** — The Global Health Network
- **Introduction to Pharmacovigilance** — PharmUNI Certificate | Valid: Feb 2026 – Feb 2028

PROJECT & PUBLICATION

Formulation and Evaluation of Emollient Cream | Published in IJARSTC, Dec 2024

- Conducted formulation trials, stability testing, and physical/chemical evaluation of emollient cream for dermatological application.
- Authored and published a peer-reviewed research paper, demonstrating scientific writing proficiency and research methodology skills.

LANGUAGES

English (Proficient) | **Hindi** (Fluent) | **Marathi** (Native)

Preferred Location : Nagpur, Hyderabad and Overall India
Avability : Immediately Joiner