BIDESH KARAN

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CAREER OBJECTIVE

As a highly organized, collaborative, and initiative-driven **PharmD** graduate with a strong foundation in Pharmaceutical sciences, I aim to secure a challenging role in **Regulatory Affairs** to utilize my knowledge of regulatory submission process, and compliance to contribute to the success of a dynamic organization.

EDUCATION

COURSE OF STUDY	BOARD/ UNIVERSITY	INSTITUTE	YEAR OF PASSING	PERCENTAGE
PHARM D (Doctor of pharmacy)	Jawaharlal Nehru Technological University Hyderabad	Pratishta Institute of Pharmaceutical Sciences, Suryapet	2024	70%
HSC	West Bengal Council of Higher Secondary Education	Sabang Saradamoyee H.S. High School	2017	73%
SSC	West Bengal Board of Secondary Education	Dhaneshwarpur Gopal Chandra Siksha Sadan	2015	81%

EXPERIENCE

Clinical pharmacist Intern

Vijay Krishna multi-speciality hospital , Suryapet

During my internship, I gained hands-on experience in patient care, medication management, and pharmacy operations. I assisted with medication orders, conducted medication safety checks, and provided patient counseling and education. Overall, my internship experience taught me the importance of collaboration, communication, and patient-centered care.

TRAINING AND CERTIFICATION:

- Completed Advanced Certification Course in Regulatory Affairs, Clinical Research and Clinical Data Management from Gratisol Labs. [GLCRCDM527]
- Completed Clinical Data Management skill development course from Climed Research Solutions and Curio Training & Research Institute [CTRI].[CU03CDM050]
- Advance certification course in Multi-Vigilance program from Medxury.

PROJECT WORK (DURING PHARM D)

Title: EXAMINING TRENDS OF PRESCRIPTION PATTERNS FOR RHEUMATOID ARTHRITIS MEDICATION-A CROSS-SECTIONAL STUDY.

Duration: **06 months** No. of Subject : **60**

CURRICULAR ACTIVITES

- Coordinated and organized competitions and seminars during Pharmacy Week (annual event).
- Provided volunteer support at Vijay Krishna Multi-Speciality Hospital,[General Medicine 8 months, Orthopedic 2 months, Pediatric 2 months] Suryapet, assisting with patient data management tasks.

PROFESSIONAL SKILLS

✓ Knowledge on Drug development (Phase I-IV) of clinical trial/ Drug approval process.

✓ Knowledge of regulatory guidelines and compliance standards [FDA,EMA,PMDA]

✓ Familiarity with submission process and document requirements.

✓ Knowledge of CTD and eCTD.

✓ Knowledge of DMF and its purpose.

✓ Knowledge of 180 days exclusivity.

✓ Understanding of Regulatory requirements for clinical research and drug safety.

✓ Knowledge of 505(b)(2) NDA, 505(b)(1) NDA, 505(j) Application and its purposes.

SOFT SKILLS

✓ Excellent verbal and written Communication skills

✓ Strong analytical and research skills

✓ Strong time management skills

✓ Team leadership & MS Word, Excel, Power point (Windows-xp,Windows7).

✓ Skilled problem solver

LANGUAGES

English (fluent), Hindi (fluent), Bengali (native), Telugu (communicable).

DECLARATION

I do hereby declare that the particulars of information and facts stated herein above are true, correct and complete to the best of my knowledge and belief.

BIDESH KARAN