

# Nikita Gavnekar

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#### **CAREER OBJECTIVE**

Highly motivated and detail-oriented Microbiology graduate seeking a challenging position in the field of Microbiology. Eager to apply my knowledge and laboratory skills to contribute to research and development in the areas of healthcare, pharmaceuticals, and biotechnology.

#### **EDUCATION**

**Advanced Post Graduate Program**: Clinical Trial Management, Clinical Data Management, Pharmacovigilance, and Medical Writing, Clini-India Pune - April 2020

**M.Sc Microbiology:** Shivaji Science College, Nagpur: with a percentage of **67** 

**B.Sc Microbiology**: Satpuda College of IT and BT, Sausar with a percentage of **77.2** 

**Higher Secondary Certificate:** School for Excellence, Sausar with a Percentage of **62** 

## **ACADEMIC PROJECT**

Screening and characterization of phenol degrading of bacteria

#### PROFESSIONAL SKILL SUMMARY:

- **laboratory Techniques**: Aseptic Culturing, PCR, Microscopy
- Microbial Analysis: API Strip, Biochemical Tests, Microbial Identification
- Pharmacovigilance
- Clinical Trial and Clinical Data Management
- Developing and Implementing Quality Management Systems

### **WORK EXPERIENCE**

Biology Teacher at Career Academy, Sausar, from 2016 to 2019, teaching high school students.

#### **PERSONAL DETAILS:**

• Father's Name: Mr. Suresh Gavnekar

Date of Birth: 01 Jan 1991

• Gender: Female

• Language Known: English, Hindi, Marathi

 Address: Kadu Colony, Ward no. 11, Sausar, Chhindwara, Madhya Pradesh

#### **SKILLS**

#### **Technical Skills**

- Microbiological Techniques: Aseptic Culturing, Streak Plating, Microscopy, Gram Staining, etc.
- Molecular Biology: PCR, DNA Extraction, Gel Electrophoresis
- Health and Safety Compliance: Biosafety Protocols, Chemical Handling
- Laboratory Safety and Quality Control
- Clinical Research: ICH-GCP guidelines, CRF Designing, Data Collection, Query Management
- Clinical Data Management: Electronic Data Capture, CDM Process Flow, Data Validation, Data Archiving
- Clinical Trial Management: Site Selection, Investigational Brochure, CRF Completion, Data Query Resolution
- **Pharmacovigilance:** ADR, SAE, MedDRA, Case Assessment, Narrative Writing, Drug Safety Principles
- Scientific Writing and Presentation
- Proficient in data entry and data validation activities
- Proficiency in Microsoft Excel, Word, and PowerPoint

#### **Soft Skills**

- Strong attention to detail and accuracy
- Effective written and verbal communication
- Ability to work in a team environment and collaborate with crossfunctional teams
- · Good time management and organizational skills
- Strong analytical and problem-solving skills

#### **CERTIFICATIONS**

- Completed Advanced Post Graduate Program in Clinical Research & Management. (Acquired Knowledge Proficiency in Pharmacovigilance, Clinical Research, Clinical Data Management, and Medical Writing)
- Certificate in 'Introduction to Pharmacovigilance' from Uppsala Monitoring Centre.
- Completed rigorous and intensive 6-month **Medical Scribe** training at Medvoice Transcript Solutions, Bangalore.
- Certificate on **Diploma in computer application**