

https://jobfever.govhelp.in/job/sun-pharma-recruitment-2024-freshers-jobs-medical-researcher-post/

Sun Pharma Recruitment 2024 - Freshers Jobs - Medical Researcher Post

Job Location

India

Remote work from: IND

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Base Salary

USD 16,500 - USD 21,000

Qualifications

12th Pass/Graduate

Employment Type

Full-time

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Description

Sun Pharma Recruitment 2024

As a Medical Researcher, you will be responsible for conducting comprehensive research on new and existing pharmaceutical products. Your work will contribute to the development and evaluation of clinical trials, ensuring the safety and efficacy of our drugs. You will be an integral part of our research team, collaborating with scientists, clinicians, and regulatory authorities to bring life-saving treatments to market.

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Sun Pharma Jobs Near Me

Responsibilities:

- Conduct literature reviews and research to stay current on the latest scientific advancements in your assigned therapeutic area.
- Design and analyze clinical trial protocols, ensuring adherence to Good Clinical Practice (GCP) guidelines.
- Manage and analyze clinical trial data, preparing reports and presentations for internal and external stakeholders.
- Collaborate with cross-functional teams, including scientists, statisticians, and regulatory affairs specialists, to ensure smooth project execution.
- Contribute to the development and implementation of research strategies to optimize clinical trial outcomes.
- · Stay abreast of regulatory requirements and ensure compliance with

Hiring organization

Sun Pharma

Date posted

January 10, 2024

Valid through

31.08.2024

APPLY NOW

relevant guidelines and regulations.

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Sun Pharma Careers

Skills:

- Master's degree in Life Sciences (e.g., Biology, Pharmacology, Biochemistry, etc.) or a related field.
- Strong understanding of medical research principles and methodologies.
- Experience with clinical trial design and data analysis.
- Excellent analytical and problem-solving skills.
- Strong communication and interpersonal skills.
- Proficiency in data analysis software (e.g., SAS, R).
- Ability to work independently and as part of a team.
- Excellent written and verbal communication skills.
- Experience with Good Clinical Practice (GCP) guidelines.
- Working knowledge of regulatory affairs processes.

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