



Memo No RIMS, Ranchi, Dated

Website Advertisement for Short term fellowship program in Good Clinical Practice

1. Applications are invited from Faculty members/Residents/PhD students for **Short term fellowship program in Good Clinical Practice** organized by RIMS, Ranchi tentatively in 3 modules of 2 weeks each from **11th November 2024 to 25th January 2025**. This Fellowship program is funded by Department of Health Research, Indian Council of Medical Research (DHR, ICMR) New Delhi.
2. For expression of interest in this program, please submit the filled application form, concept note for clinical trial and bio-data to **email id- rims.dhrproject@gmail.com** by 27.10.2024. Format of application form and concept note is provided in the program brochure as Appendix 1 and 2.
3. Faculty members/Residents/PhD/ PG students from RIMS, Ranchi may also apply for the training program.

Ali
17/10/24

Dr. Arpita Rai,
Principal Investigator,

S.B. Singh
17/10/2024

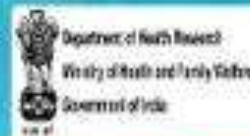
Dr. S.B. Singh
Co-Investigator

DHR Project -Short term fellowship program in Good Clinical Practice.

* Attachments: - Program brochure

Copy forwarded to:

1. Dr. S.B. Singh, Incharge, Website, RIMS, Ranchi for Information & necessary action.
2. RIMS, Ranchi Notice board for General information.



*Rajendra Institute of Medical
Sciences, Ranchi*
organizes

**Short Term Fellowship Program
in Good Clinical Practice**

supported by Department of Health Research,
New Delhi

MODULE 1: 11th NOVEMBER 2024 – 23rd NOVEMBER 2024

MODULE 2: 09th DECEMBER 2024 – 21st DECEMBER 2024

MODULE 3: 13th JANUARY 2024 – 25th JANUARY 2024

LIMITED SEATS AVAILABLE

Venue: 3rd Floor Research & Administration Hall,
Administrative Block, RIMS, Ranchi



Invitation

Dear Friends,

Greetings from Rajendra Institute of Medical Sciences, Ranchi

It's our great pleasure and honor to invite you to Short Term Fellowship Program in Good Clinical Practice, to be held in RIMS, Ranchi from 11/11/2024 to 25/01/2025. Eminent faculty will provide candidates a platform to interact and exchange of views on good clinical practice and evidence-based practice. The fellowship program will strengthen your understanding of Clinical research process. Looking forward to welcoming you to Ranchi.

Best Regards,



Prof. (Dr.) Raj Kumar
Director & CEO, RIMS, Ranchi
Rajendra Institute of Medical
Sciences, Ranchi

List of Mentors: -

Dr. Arpita Rai

Principal investigator, DHR PROJECT, RIMS
Assoc. Professor, Dental Institute, RIMS

Dr. Kameshwar Prasad

Co-Investigator, DHR Project, RIMS
HOD Neurology, Fortis Hospital, Vasant Kunj, Delhi

Dr. S. B. Singh

Co-Investigator, DHR PROJECT, RIMS
Addl. Professor, PSM, RIMS

Prof. Santanu K Tripathi

Dean (Academic), Netaji Subhas Medical College and Hospital, Patna.

Prof. Bimal Kumar Mishra

Principal Adarsh College, Giridih, Jharkhand

Dr. P. K. Bhattacharya

Dean (Research), RIMS

Dr. Sandhya Kumari

Professor, Anatomy, RIMS

Dr. Bhoopendra Singh

Professor, FMT, RIMS

Dr. Archana Kumari

Professor, Obs & Gynae, RIMS

Dr. Lakhan Manjhi

Addl. Professor, Pharmacology, RIMS

Dr. Anupa Prasad

Addl. Prof., Genetics & Genomics, RIMS

Dr. Amit Kumar

Assoc. Professor, Lab Medicine, RIMS

Dr. Dewesh Kumar

Assoc. Prof. PSM, RIMS

Dr. Amit Vasant Mahuli,

Assoc. Professor, Dental Institute, RIMS

Dr. Ganesh Chouhan

Assoc. Prof. Genetics & Genomics, RIMS

Introduction to Department of Health Research (DHR):

Department of Health Research (DHR) was created as a separate Department within the Ministry of Health & Family Welfare by an amendment to the Government of India (Allocation of Business) Rules, 1961 on 17th Sept, 2007. The Department of Health Research was formally launched on 5th October 2007 by the Minister for Science & Technology and Earth Sciences in a function presided by the Minister for Health & Family Welfare, in the presence, inter-alia, of the Minister of State for Health & Family Welfare. The Department became functional from November 2008 with the appointment of first Secretary of the Department. The aim of the DHR is to bring modern health technologies to the people through research and innovations related to diagnosis, treatment methods and vaccines for prevention; to translate them into products and processes and, in synergy with concerned organizations, introduce these innovations into public health system.

<https://dhr.gov.in/about-us/about-department>

Introduction to Rajendra Institute of Medical Sciences (RIMS), Ranchi:

The Rajendra Institute of Medical Sciences (RIMS) is a premier medical institute located at Ranchi, the capital of Jharkhand. The Institute is an autonomous body established in 2002 under an Act of Jharkhand state Assembly. The Institute started as Rajendra Medical College in the year 1960 and was named after Dr. Rajendra Prasad, the first President of India. The Medical College Hospital came into existence in February 1965 and has excelled ever since. The Motto of the Institute is **“SARVE SANTU NIRAMAYA”**. RIMS Ranchi has a very large and spacious campus. It has a 1500 bedded multi- specialty hospital with State of the Art equipment. The institute is a tertiary care centre that has the excellent infrastructure for patient care as well opportunity to acquire knowledge and skills from experienced faculties. The Institute also has a 100 bedded Trauma Centre, the first of its kind in Eastern India with most modern equipment. Under PMSSY scheme, a 240 bedded Super Specialty block was established to provide care for the patients in the field of Oncology, Cardiology, Urology, Pediatric Surgery and Cardio-thoracic Surgery. The Dental College, RIMS, Ranchi was founded in the year 2017 and is the first autonomous Dental College under Government of Jharkhand. RIMS conduct teaching programs in medical, dental and para-medical courses. Besides undergraduate courses, RIMS also offers post - graduate courses, PhD and super-specialty courses offered in various disciplines. RIMS also run a College of Nursing and are affiliated to Ranchi University.

<https://www.rimsranchi.ac.in/>



About The Project: -Short Term Fellowship Program in Good Clinical Practice (GCP)

RIMS, Ranchi has received funding under the Support to Institute for imparting training (Human Resource Development for Health Research) scheme of DHR to conduct short term fellowship in Good Clinical Practice for 3 years (2022-24). The first fellowship was conducted between April to June 2022 and 32 successful candidates were awarded the fellowship. The second fellowship was conducted between April to June 2023 and 38 successful candidates were awarded the fellowship.

Good Clinical Practice is a set of guidelines for biomedical studies which encompasses the design, conduct, termination, audit analysis, reporting and documentation of the studies involving human subjects. The guideline seeks to establish two cardinal principles: protection of the rights of human subjects and authenticity of biomedical data generated. These guidelines have been evolved with consideration of WHO, ICH, USFDA and European GCP guidelines as well as the Ethical Guidelines for Biomedical research on Human Subjects issued by the Indian Council of Medical Research. They should be followed for carrying out all biomedical research in India at all stages of drug development, whether prior or subsequent to product registration in India.

This Good Clinical Practice (GCP) training is designed to prepare medical/dental college faculty, PhD scholars and post-graduate students in the conduct of clinical trials with human participants. This 6-week training will be based on International Council on Harmonization Good Clinical Practice (ICH-GCP) Principles.

Objectives: -

1. To provide an overview and practical advice on the application and implementation of internationally accepted principles for GCP and clinical research in human subjects.
2. To provide a reference tool for participants by providing the necessary background and insight into the reasons for the requirements of GCP and their efficient application.
3. To get a comprehensive understanding of the clinical research process

Criteria of Selection:

We seek **interested post graduates, PhD students and faculties working in medical/dental colleges from across India**. The training will be free of cost for the participants. However, it will follow a strict selection criterion. An elementary knowledge of research methodology would be preferred; however, the workshop does not entail any theoretical background of textual knowledge. Applicants will be required to provide the following by email rims.dhrproject@gmail.com by **27th Oct 2024**:

- Application form for expression of interest (Appendix 1)
- A concept proposal related to the clinical trial as per the attached format. The ideas/interventions can be drug molecules, therapeutics, vaccine candidates, biological, nutritional products, diagnostics, or community based interventions and others. The concept notes will be peer-reviewed by the Advisory Committee for Clinical Research, RIMS, and Ranchi. The following criteria will be used to evaluate proposals - Novelty, applicability and justification for research (Appendix 2)
- Brief Curriculum Vitae / resume highlighting relevant skills, experience, and training.

The list of selected candidates will be updated on RIMS website.

Tentative Program

Week 1: -Overview of the clinical research process and clinical trial

- Overview of the clinical research process
- Formulating research question: PICO and its variation; Criteria for good research question
- Study Design
- Sample Size
- Randomization process and blinding
- Selecting endpoint/response variables
- Baseline assessment of study participants
- Confounders in clinical trial
- Internal vs external validity in clinical trial
- Reporting and interpretation of RCT (CONSORT checklist)

Week 2: -Systematic Review & meta-analysis

- Statistical methods used in interim monitoring
- Analysis of RCT
- Issues related to data analysis
- Survival analysis
- Search strategy
- Data extraction
- GRADE:
- Fundamentals of GRADE
- GRADE: Rating the certainty of evidence, Sub Group Analysis
- Network Meta-Analysis: Overview
- GRADE: Guideline Development overview, Strong Vs Weak Recommendation.

Week 3: -Ethics in clinical research, Research misconduct

- ✚ Ethics committee/Institutional Review Board (composition, membership & responsibilities)
- ✚ Investigator's responsibilities to Ethics committee
- ✚ Criteria for ethical approval of research
- ✚ Informed consent
- ✚ Documentation requirements in informed consent
- ✚ HIPAA rights, Privacy and Enforcement
- ✚ Confidentiality of Clinical Trial Participant Records
- ✚ Exceptions to Confidentiality Requirements
- ✚ Identifying Research Misconduct & Possible Penalties for Research Misconduct

- ✚ Investigating allegations & Responding to Allegation of Research Misconduct
- ✚ Safeguard for Informants & accused Person
- ✚ Protection from conflict of interest
- ✚ Publication bias, suppression and delays in reporting
- ✚ Conflict of interest & publication

Week 4: -Regulatory issues, Research protocol & documentation. Roles and responsibilities

- ❖ Regulatory bodies – CDSCO, CTRI, DHR
- ❖ Acts and rules pertaining to clinical trials
- ❖ Content of research protocol
- ❖ Protocol Amendment & Violation
- ❖ Documentation requirement for sponsored trial

- ❖ Regulatory issues pertaining to new drugs, investigational new drugs & subsequent new drugs, vaccine, medical devices and diagnostics, biologics.
- ❖ General & Specific responsibilities of sponsor, General & Specific responsibilities of Principal investigator

Week 5: - Conduction of trial, patient safety/ adverse event, Health Related Quality of Life

- Site Selection & Site initiation
- Recruitment & Recruitment Strategies
- Retention & Retention strategies
- Assessment of Harm/adverse event
- Classification of Adverse Events
- Analysing Adverse Event, Reporting of Harm, Regulatory Considerations
- Health Related Quality of Life (HRQL): overview, Methodological and design Issues, Selection of HRQL Instruments
- Study closeout: Termination Procedures, transfer of post-trial Care and Post Study Follow up, Data & Records Dissemination of Results.
- Visit to Central Medical Record Department & Training centre

- ❖ Role and responsibilities of research site staff
- ❖ Multi-centric clinical trial, Global Clinical Trial, Clinical Trial with Commentary & alternative medicine,
- ❖ Visit to Research lab at RIMS, Ranchi

Week 6: - Quality Control

- Problems in data collection, Minimizing Poor Quality Data, Electronic Source Data
- Monitoring board/committee structure, Monitoring board/Committee functions, Monitoring Visit Procedures
- Quality Monitoring of Data, Procedures and Drug Handling.
- Various types of audit in clinical trials
 1. Clinical Trials Audit Observations and study of Critical observations
 2. Breach reports and Compliance guidelines
- Competency Assessment
- *Research Protocol Presentation by Fellows*

Last Date of Application: 27th Oct 2024

Certification:

Upon completion of the training, certificates will be provided to the fellows. The trainees should fulfill the following conditions for being eligible for certification:

- a) A minimum attendance of 85%
- b) Submission of pre and post training assessment.
- c) Scoring a minimum of 75% marks in training end competency assessment. (In case a fellow fails to achieve 75% marks in the competency assessment, he can give two further attempts for the competency exam in online or offline mode at an interval of one month)
- d) Submission of Ad-hoc research proposal (ICMR format) based on the concept note submitted at the time of selection.
- e) Presentation of the Ad-hoc research proposal.

Other Terms and Conditions: -

1. *There is no registration fee for the candidate.*
2. *No TA DA will be provided by the host Institute.*
3. *No accommodation facility will be provided by the host Institute. Guest house on self paid basis may be provided subject to availability. Please contact Miss Anita Kumari, Project Assistant (Contact no- 8809197830)*
4. *No application will be accepted after the last date.*
5. *Decision made by the Director RIMS will be final.*



Appendix 1 - Application form for expression of interest SHORT TERM FELLOWSHIP PROGRAM IN GOOD CLINICAL PRACTICES

1. Name :

2. Contact Details (Mobile No & Email Id) :

3. Age/Gender:

4. Name of the Institution:

5. State :

6. Present Designation/Last Post Held :

7. Have you attended any workshop/training program related to GCP?

Yes

No

8. Are you involved in any clinical trial?

Yes

No

9. If you are involved what was your role?

10. Tick reasons for interests in this training program:

To Gain Knowledge about GCP

To enhance your knowledge about this concept

Developing new Prowess

Having Fun While Learning

Network with like-minded researchers

Appendix 2 – Format for concept note

1. *Title of the proposed research project:* – should be concise and yet sufficiently descriptive and informative. Title may include study design such as randomized controlled trial.
2. *Rationale up to 350 words:* –Describe the current knowledge available on the subject area, critical gaps in knowledge, its relevance and application to local, national and international context and the research questions which this project aims to address.
3. *Project Description (up to 700 words):* – Describe study setting, study design, sampling strategy, sample size, feasibility, work plan, expected outcomes, time line.

Glimpses of fellowship Program (2022)

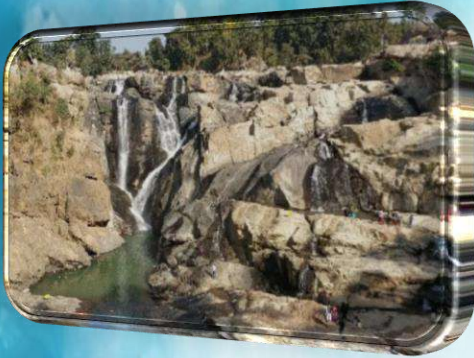


Glimpses of Fellowship Program 2024

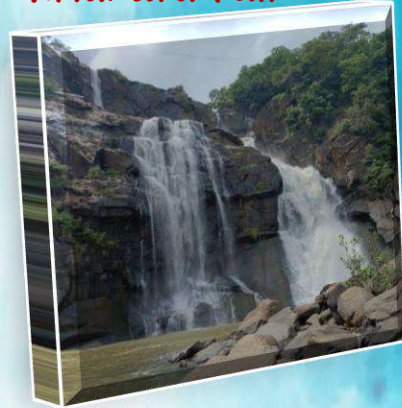


Discover Ranchi

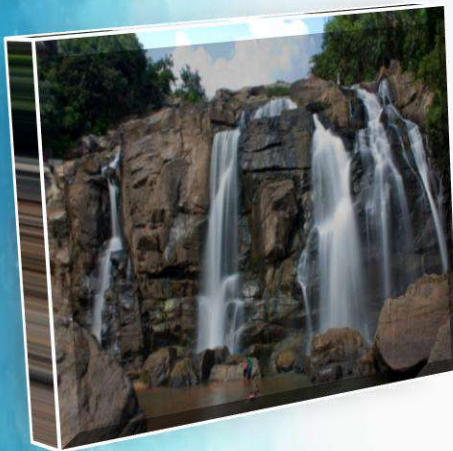
1. Dassam Fall



4. Hundru Fall



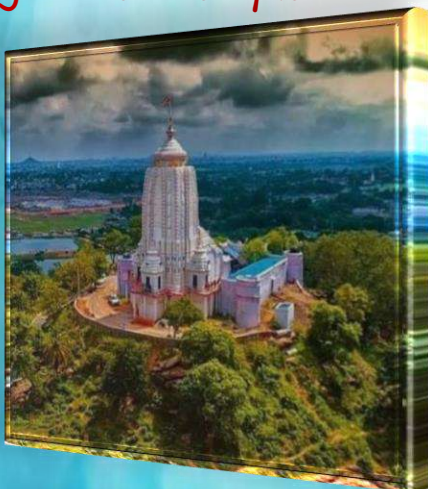
2. Jonha Fall



5. Sureshwar Mahadev Temple



4. Jagannath Temple



6. Rajrappa Temple

