

ADVANCED PG DIPLOMA IN PHARMACOVIGILANCE & CLINICAL DATA MANAGEMENT

Duration: 6 Months

Prepare for the industry with specialized training in pharmacovigilance and clinical data management, led by experts. Additionally, benefit from placement assistance with leading pharmaceutical and CRO companies.

Accreditation & Recognition



#startupindia



SOCR
FOR CLINICAL RESEARCH BRILLIANCE



- 1 **Pharmacovigilance Essentials:** Learn how to identify, report, and analyze adverse drug reactions and safety signals.
- 2 **Clinical Data Review & Validation:** Gain skills in reviewing, managing, and validating clinical trial data using global standards.
- 3 **Regulatory Safety Submissions:** Understand MedDRA coding, ICSR reporting, and preparation of PSURs for regulatory compliance.
- 4 **Career-Focused Curriculum:** Get trained with industry-relevant content designed for long-term success in pharma, CROs, and healthcare analytics.

Highlights

- Industry-Relevant Global Curriculum
- Globally Recognized Certification
- Dedicated Placement Support
- Resume-Boosting Certification
- Career Oriented Training
- Free Placement Assistance
- Real time Interview Preparation

REGISTER NOW

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SCAN QR CODE
to register

IACCRE - ACCREDITED
PROGRAM



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support@pharma-research.in

https://pharma-research.in

COURSE SYLLABUS



For Registration Call
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Pharmacovigilance – Module 1

- Introduction to Clinical Research & Good Clinical Practice
- Overview ICH GCP
- Good Clinical practice
- Clinical Trails
- Institutional Reviews board
- Investigation Responsibilities in clinical trails
- Roles and Responsibilities of Sponsor & Clinical Trial Monitor
- Informed Consent Process
- Serious Adverse Event (SAE), Adverse event reporting
- Site Initiation visit
- Site close visit
- Test Your Self

The Advanced PG Diploma in Pharmacovigilance & Clinical Data Management is a 6-month online program tailored for graduates in pharmacy, life sciences, medicine, nursing, and related fields. The course is divided into two industry-aligned modules that provide a comprehensive understanding of clinical research operations and drug safety protocols. The Pharmacovigilance module covers ICH-GCP guidelines, clinical trial phases, informed consent, SAE/AE reporting, and the roles of sponsors and monitors. The Clinical Data Management module focuses on data collection, validation, medical coding, lab data handling, eCRFs, query management, and regulatory compliance systems — equipping learners with practical skills required to manage data throughout the clinical trial lifecycle.

Offered by Pharma Research, an IACCRC-accredited institution, this program is globally recognized and backed by expert-designed curriculum and real-world applications. Students receive an internationally valid certification with lifetime verification, plus a detailed performance card upon completion. The program also includes placement assistance through Pharma Research's dedicated career support and job partner [JobsAvenue.net](#), giving learners access to opportunities in top pharmaceutical companies, CROs, and healthcare organizations. This course not only prepares students for roles such as Clinical Data Analyst or Drug Safety Executive, but also builds a strong foundation for long-term career growth in the clinical research industry.



SELF-PACED VIDEO LECTURES

Learn anytime, anywhere with lifetime access to expert-curated video modules.



GLOBALLY ACCEPTED CERTIFICATION

Earn international certification accredited by IACCRC and recognized by pharma companies, CROs, hospitals, and regulatory agencies worldwide.



CAREER-ORIENTED TRAINING

Receive help with resume building, interview preparation, and job referrals through our dedicated placement cell and partners.



REAL TIME TECHNICAL INTERVIEW PREPARATION

Get industry-based real-time interview questions and answers to boost your job readiness and confidence.

Clinical Data Management Module 2

- Introduction To Clinical Data Management
- Clinical Data Management
- Medical Coding in Clinical Data
- Medical Coding And Medical Dictionaries
- Clinical Data Process
- Data Capture
- Data Collection
- Data Validation
- Lab Data Management
- Query Management
- Case Report Form
- Electronic Case Report Form
- Quality & Regulatory System
- Regulatory Systems

HOW WE

We follow a structured and student-centric approach to deliver high-quality education and career outcomes in the field of clinical research.

Personalized Learning Approach

Every learner receives step-by-step guidance through Online sessions, recorded content, and mentorship.

Industry-Aligned Curriculum

Our content is curated in collaboration with industry professionals to meet global standards.

Career-Oriented Training

From resume preparation to job interview support, we help you confidently enter the workforce.



ELIGIBILITY CRITERIA

Minimum Qualification: Candidates must have completed or be in the final year of any of the following degrees: B.Pharm, M.Pharm, D.Pharm, MBBS, BDS, BAMS, BHMS, B.Sc/M.Sc in Life Sciences, Biotechnology, Microbiology, Biochemistry, Chemistry, Nursing, or any Allied Health or Medical Science discipline.

Final-Year Students: Those currently pursuing their final year in the above-mentioned fields are also eligible to apply.

Working Professionals: Individuals working in pharma, clinical research, healthcare, or life sciences sectors seeking skill advancement are encouraged to enroll.

Basic Computer & English Proficiency: Recommended for effective learning and communication.



WHAT'S NEXT AFTER COMPLETING THE DIPLOMA?

After completing the Advanced PG Diploma in Pharmacovigilance & Clinical Data Management, learners become eligible for a wide range of job opportunities within the pharmaceutical, clinical research, and healthcare regulatory sectors. Common entry-level roles include Pharmacovigilance Associate, Clinical Data Coordinator, Drug Safety Executive, Clinical Trial Assistant, and Regulatory Affairs Support Officer. These positions are available across Contract Research Organizations (CROs), pharmaceutical companies, biotech firms, hospitals, and regulatory agencies.

Fresh graduates and early-career professionals can expect starting salaries in the range of ₹3.0 to ₹5.5 LPA, depending on location, employer, and educational background. With 2–3 years of domain experience, professionals can progress into higher roles such as Senior Pharmacovigilance Executive, Clinical Data Manager, Safety Team Lead, or Regulatory Affairs Specialist, often earning ₹7 to ₹12 LPA or more. The industry-aligned training, practical exposure, and placement assistance offered by Pharma Research provide learners with a strong foundation to grow confidently in one of the fastest-evolving and globally critical fields in healthcare and pharmaceuticals.

PLACEMENT ASSISTANCE

(We don't promise job guarantees — we promise genuine job support and industry-driven assistance)

At Pharma Research, we are committed to supporting the career journey of every student through our comprehensive placement assistance program. As part of this initiative, we offer:

- **Industry-Oriented Resume Building:** We help students craft professional resumes aligned with the latest industry standards in clinical research, pharmacovigilance, and pharmaceuticals.
- **Resume Promotion to Employers:** Your resume is shared with over 150+ pharmaceutical companies, CROs, hospitals, and research organizations globally, enabling employers to shortlist candidates as per their requirements.
- **Interview Preparation Support:** Access real-time interview questions and expert-crafted answers to help you prepare confidently for technical and HR rounds.
- **Free Job Search & Alerts:** Through our placement partner JobsAvenue.net, students receive free job alerts via email and can explore verified openings across the industry.

We provide 100% free placement assistance to all enrolled students. For full details, please visit our Placement Assistance Policy: <https://pharma-research.in/placement-policy/>





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CALL FOR APPOINTMENT

TOP RECRUITERS & HIRING PARTNERS



READY TO START YOUR CAREER IN CLINICAL RESEARCH?

Join thousands of learners who've advanced their careers with Pharma Research.

With expert-led training, international certification, and strong placement support — your success begins here.

☒ **100% Online Program**
☒ **Globally Accepted Certification**
☒ **Free Placement Assistance**
☒ **Lifetime Verification Support**



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CALL FOR APPOINTMENT

WHAT OUR STUDENTS SAY

0 Omprakash Ghakade

★★★★★

I recently completed the Advanced PG Diploma in Pharmacovigilance & Clinical Data Management from Pharma Research and am thrilled to share that I've been placed as a Clinical Data Manager at PAREXEL. The course was fully online, and the video lectures were extremely clear, practical, and easy to follow. I really appreciated the flexibility to learn at my own pace, which made it easier to balance with my other commitments. The support team was highly responsive and helped me promptly with all my technical queries during the course. Pharma Research not only provided strong academic content but also guided me throughout the placement process. Highly recommended for anyone serious about building a career in clinical research or data management.



Loreta Gherman

1 review

★★★★★ a year ago

Concerning the CCRA course I very much appreciate the provision of very relevant documentary sources. The information concerning career guidance for a perspective of work was also quite welcome.

S Steav Johnson

1 review

★★★★★ a week ago NEW

Thank you, Pharma Research, for supporting me as a career advisor and in placement journey. I am post Graduate Master of science in Biotechnology and was specifically for work-from-home job opportunities. During my online search, I came across Pharma Research and enrolled in their Advanced Postgraduate Diploma in Data Management. The program was conducted online, highly flexible, and structured with detailed content.

After completing the course, the placement assistance team guided me with valuable tips and interview strategies, which helped me confidently face interviews. I'm now working as a Clinical Data Manager in a hybrid role—working two days in the office and the rest remotely.

I, 100% recommend this institute and their course to anyone looking to build in clinical research and data management.

A Ahmed Usman

1 review · 1 photo

★★★★★ 2 years ago

Following my passion for clinical research and developing my career in this field, PHARMA RESEARCH has played an incredible role, with the opportunity to be enrolled in such extraordinary, structured, well-organized, and dynamic training programs, led by Clinical Research Professionals. Recently, I have been a graduate in ADVANCED PG DIPLOMA IN CLINICAL RESEARCH Program. Definitely, from this program, I have gained the skills of learning and thinking at the same time through assignments and knowledge checks. As Confucius said: "Learning without thinking is useless, thinking without learning is dangerous."

? FREQUENTLY ASKED QUESTIONS

Q: How is the course delivered?

A: 100% online via structured video-based modules.

Q: Is the certificate valid internationally?

A: Yes. It's accredited by IACCRC and accepted by global pharma and CRO employers.

Q: Do you offer job support?

A: Yes. We provide complete placement assistance including resume support and job alerts.

SAMPLE CERTIFICATE



 **Contact Us**



Guntur | Hyderabad | Bengaluru | Dubai |
USA | Australia | Ireland

Website: <https://pharma-research.in>

Email: support@pharma-research.in

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✨ For Quick Registration: Scan the QR code below or visit our website to register now and begin your journey toward a rewarding healthcare career!



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OUR COLLABORATIONS



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