birac
Ignite Innovate Incubate

सत्यमेव जयते
DEPARTMENT OF BIOTECHNOLOGY
Ministry of Science & Technology
Government of India

nbm
NATIONAL BIOPHARMA MISSION

National Biopharma Mission supported Hands-on Programme on Good Clinical Laboratory Practice (GCLP)

DATE	30 – 31 January, 2020	VENUE	CDSA, Faridabad

ABOUT THE PROGRAMME

Good Clinical Laboratory Practice (GCLP) is a set of standards that provide guidance on implementing Good Laboratory Practice (GLP) and Good Clinical Practice (GCP) principles to the analysis of samples from a clinical trial. By combining the GLP and GCP sets of guidelines, GCLP ensures the reliability, quality, consistency, and integrity of the clinical trial data generated by laboratories. Compliance with this standard provides assurance that the data and reported results are credible and accurate and that the rights, safety, and confidentiality of trial subjects are protected

HIGHLIGHTS

- Workshop organized by CDSA which is a pioneer in building capacity and capability in the area of clinical development and translational research in India.
- Programme shall offer comprehensive guidance for implementing GCLP in laboratories.
- Talks by reputed national and international domain experts with hands-on exercises.

Who can attend

Personnel working in the area of laboratory (clinical/medical), clinical trials, clinical research and research & development may benefit from this programme.

- Sponsorship from National Biopharma Mission includes participation, workshop material and accommodation for out-station participants.

Last date for submitting application:13th January, 2020

APPLY NOW – www.nbmworkshops.com

For any queries:
nbmworkshop@biotech.co.in or
011-23219064-67

Organising Partner
CDSA
LEARN TO LEAD

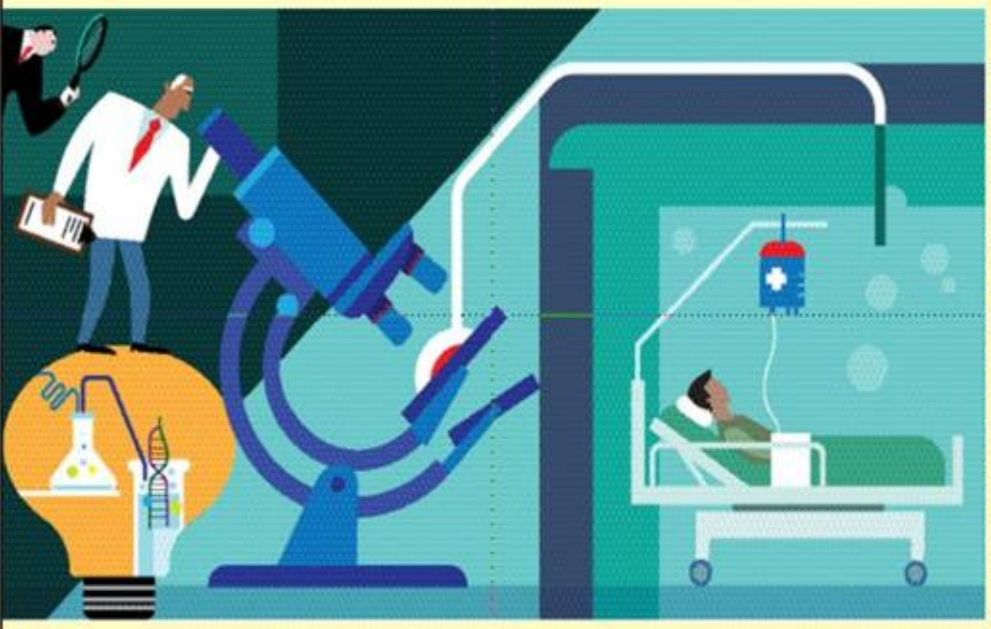
Management Partner
BCIL

birac
Ignite Innovate Incubate

DEPARTMENT OF BIOTECHNOLOGY
Ministry of Science & Technology
Government of India

nbm
NATIONAL BIOPHARMA MISSION

Training Programme on
GOOD CLINICAL PRACTICE (GCP)
supported by National Biopharma Mission



Good Clinical Practice (GCP) is an international ethical and quality standard for the design, conduct, performance, monitoring, auditing, recording, analyses, and reporting of clinical trials that provides assurance that the rights, safety, and well-being of human research participants are well protected, data and reported results are credible and accurate. Creating awareness of GCP is mandatory when it comes to innovation in medicine and healthcare. The recent launch of the New Drugs and Clinical Trials Rules, 2019 (NDCT), on March 19, 2019, opens up a new set of opportunities to accelerate innovations in India. Poor understanding of the current regulatory requirements mixed with inadequate compliance to best global practices may lead to delays and failures which can be significantly reduced through awareness programmes.

Date	28 - 29 January, 2020	11 - 12 February, 2020
Venue	Clinical Development Services Agency, Translational Health Science and Technology Institute, Faridabad, Haryana	Venture Center, NCL Innovation Park, Pune, Maharashtra
Last date for submitting applications	13 January, 2020	20 January, 2020

Highlights

- Workshop organized by CDSA which is a pioneer in building capacity and capability in the area of clinical development and translational research in India.
- The workshop shall create awareness and understanding of GCP to enable compliance with the current guidelines and regulatory requirements.
- Talks by reputed faculty, present and former regulators, national and international domain experts.
- Sponsorship from National Biopharma Mission includes participation, workshop material and accommodation for out-station participants.

Who can attend

Personnel working in the area of drug development, clinical trials, and clinical research may benefit from this programme.

APPLY NOW:
www.nbmworkshops.com
For queries, please contact:
nbmworkshop@biotech.co.in

Organizing Partner
CDSA

Management Partner
BCIL

For detailed programme and eligibility criteria, please visit: www.nbmworkshops.com

Biotechnology Industry Research Assistance Council
(A Government of India Enterprise)
1st floor, MTNL Building,9, CGO Complex, Lodhi Road, New Delhi-110003
Tel. +91 11 24389600 | Fax. +91 11 24389611
Website: www.birac.nic.in

जैव प्रौद्योगिकी उद्योग अनुसंधान सहायता परिषद
(भारत सरकार का उपक्रम)
पहली मंजिल, एमटीएनएल बिल्डिंग, 9, सीजीओ कॉम्प्लेक्स, लोधी रोड, नई दिल्ली -110003
दूरभाष। +91 11 24389600 | फैक्स। +91 11 24389611
वेबसाइट: www.birac.nic.in