

Education and Training Services

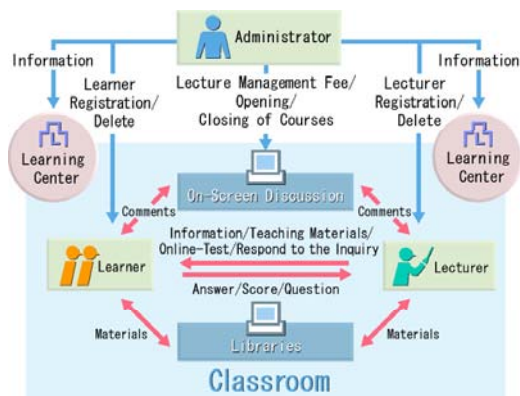
By Utilizing e-Learning and Classroom Learning with Experienced Know-How in Clinical Studies, We Perform Practical and Workable Services

Based on experienced CRO know-how, we provide effective uniform education and training services via e-learning and practical classroom lectures by clinical development specialists. We also provide a practical and workable training system which has proven to reduce cost and time.

e-Learning

【Benefits】

- Reduces cost and time of finding physical lecture locations
- Computerized training materials
- Upkeep of staff high skills
- One-to-One training
- Efficient training by one-to-one education
- Evaluation and record of course completion and comprehension



System Characteristics

- Easy to use even for internet novices
- Easy-to-manage lecture records and scores
- Real-time discussion is possible via email and bulletin board
- Mentoring support by lecturers

Contents Characteristics

- Practical content utilizing CRO experience in clinical studies
- Online test, immediate scoring and evaluation are available
- Customized lecture is provided upon your request.

●Lectures

- Introduction to Clinical Study-CRC
- Practicality in Clinical Study-CRC
- GCP Online Test
- Introductory Training of a Monitor

Classroom Learning

【Introductory Training of a Monitor (CRA)】

- Monitors are required not only knowledge of regulatory requirements of GCP and logical way of thinking but also communication skills with client and persons involved in the clinical study.
- We have a 3-week basic training in how to be a professional monitor.
- Lecturers have clinical development experience in pharmaceutical companies and leaders of monitors.
- We provide role-playing to enhance communication skills with investigators or coordinators involved in the study.

●Lectures

- Introduction to Clinical Studies
- Monitoring-Related Regulation
- Investigator's Brochure
- Briefing on Informed Consent
- Insurance and Compensation
- Clinical Study Algorithm and Organization
- GCP Guideline
- Protocol and CRF
- Clinical Study Budget
- Collection of Pharmacovigilance Data

●Role-playing

- Selection of Medical Institutions and Investigators (Surveillance Requirements)
- Study Contract Request
- Explanatory Meeting for Medical Office
- SDV (Source Document Verification) with CRF

ADVANTAGE of EPS

- We provide practical content through experienced Asian study know-how not generally available in other company.
- Full customized flexible design is feasible.
- We are able to provide any kind of e-learning and lecture topics to all professionals who work in clinical study.

