Roberto Ricciardiello

Education

CLINICAL RESEARCH TRAINING COURSE “MISSIONE CRA” | 2024 | CLINOPSHU, YGHEA CRO

50 hours Clinical Research training course as for the Ministerial Decree 15.11.2011.

Topics: Methodology and Regulation of clinical trials, Good Clinical Practice (ICH-GCP), Standards for Good Manufacturing Practice (GMP), Pharmacovigilance, Quality Systems and Quality Assurance (QA), Focus on Phase 1 quality, Tasks of Monitors referred to paragraph 5.18 of Annex 1 to the Ministerial Decree July 15, 1997 (including Pre-Study Visit, Site Initiation Visit, SiteMonitoring Visit and Close Out Visit), Tasks of Clinical Research Coordinators, Focus on Regulation (EU) N° 536/2014, Workshop on Site Selection, Pre-Study Visit, Essential Documents, Drug Accountability, Site visits simulation (PSV, SIV, SMV and COV)

PHD STUDENT | ONGOING | UNIVERSITÀ DEGLI STUDI DI PAVIA, IGM-CNR

- Genetics, Molecular and Cellular Biology
- Xeroderma Pigmentosum Variant disease (XPV), Skin cancer

MASTER’S DEGREE (LM-6)| 2018/2021| UNIVERSITÀ DEGLI STUDI DI ROMA, “LA SAPIENZA”

- Genetics and Molecular Biology
- Type 1 congenital muscular dystrophy (MDC1A), Duchenne dystrophy

BACHELOR’S DEGREE (L-13)| 2015/20218| UNIVERSITÀ DEGLI STUDI DI SALERNO

- Biological science
- Neurodegenerative diseases

LICENSES

- Licensed biologist
- GCP certified

LANGUAGES

- Italian: Native speaker
- English: B2 certificate (ESB)

WORK EXPERIENCES

STUDY COORDINATOR | ISTITUTO NAZIONALE DEI TUMORI | NOVEMBER 2024 – ONGOING

Support for the team in coordinating clinical research (medical staff, nurses, pharmacists, laboratory, ethics committee, other facilities involved in the study); Management of relationships for conducting clinical research (CRO, Monitors, Data Management, Auditors, etc.); Management of clinical study documentation (Investigator Folder); Management of data collection systems (CRF - Case Report Form); Management of IVRS systems (Interactive Voice Response System); Planning and management of study visits (SIV, COV) and monitoring activities; Support during internal and external audits and inspections by regulatory authorities; Transfer of any SAE to the Pharmacovigilance system in accordance with required timelines; Completion of the annual study report requested by the Ethics Committee; Participation in meetings; Management of updates/amendments and communications to staff involved in the study and the Ethics Committee; Management of administrative relationships

with participating centers; Support in budget preparation; Support in the financial management of clinical research

RESEARCHER | IGM-CNR | MAY 2021 - SEPTEMBER 2024

- learned GLP guidelines and gained other valuable skills. I became proficient in Microsoft Office, data processing, analysis, and creating presentations, while also developing a strong attention to detail. Managing two different projects, I learned how to troubleshoot complex technical issues and think creatively in designing experiments. I also improved my ability to work well in multidisciplinary teams, maintain professional relationships, and meet deadlines. Additionally, I had the chance to teach my work to interns at the University of Pavia. Above all, I enhanced my ability to interact with people from different countries and cultures. As of today, thanks to my work experiences, I feel confident in my ability to manage stress situations

INTERNSHIP IN RESEARCH LABORATORY | IBPM-CNR | JANUARY 2019 - JANUARY 2021

- I actively collaborated as a team member, participating in lab meetings and contributing collectively to research activities but also I obtain ability to quickly learn and apply new techniques and technologies

TRAINING INTERNSHIP AT A HOSPITAL FACILITY| OSPEDALE SAN GIOVANNI DI DIO E RUGGI D'ARAGONA, SALERNO (SA) | MARCH - JULY 2018

- I learn to operate machinery for biological samples, as well as accepting, logging, and reporting on samples