Beatrice Foti

Clinical Research Associate

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LinkedIn

Dynamic and performance-driven professional with substantial experience in overseeing entire clinical trial process, from site initiation to close-out visits. Adept at providing comprehensive guidance to clinical investigators and staff on different processes, procedures, and regulatory requirements. Demonstrated strong ability to manage site activities, oversee data integrity, and facilitate effective communication between clinical sites and research teams. Instrumental in monitoring site compliance with good clinical practice (GCP), FDA, and protocol guidelines. Proficient in latest clinical research software and tools, committed to maintaining highest levels of ethical standards and operational excellence. Excel in fast-paced environments, leveraging exceptional organisational skills and a detail-oriented approach to support ground-breaking clinical research.

Areas of Expertise

- Clinical Trial Management
- Quality Assurance & Control
- Site Staff Training & Development
- Data Collection & Analysis
- Continuous Process Improvement
- Protocol Adherence & Compliance
- Site Monitoring & Auditing
- Clinical Trial Documentation
- Cross-functional Team Collaboration

Professional Experience

Clinical Research Technology (CRT) — 2022 – 2023
Clinical Research Associate Trainee – Internship

Conduct seven remote monitoring visits across prestigious institutions, including the Agostino Gemelli University Hospital and IRCCS Istituto Nazionale Tumori "Fondazione G. Pascale", enhancing remote evaluation skills. Contribute to progress of significant clinical trials, including COMBI-TED, CAPLAND, and BIO-TAILOR, by ensuring stringent quality control and regulatory compliance. Gain hands-on experience in clinical monitoring through three days of on-site visits alongside experienced monitors at IFO - Regina Elena National Cancer Institute, Rome.

- Achieved a 2nd-level Master's degree in "Monitoring, Assurance, and Quality Control in Clinical Trials" from University of Rome "La Sapienza", aligning with industry standards and regulations.
- Demonstrated adaptability and expertise in clinical trial oversight during COVID-19 pandemic through innovative remote monitoring approaches.
- Executed on-site and remote clinical monitoring as mandated by AIFA's "Derogation from Modalities for Carrying Out Side-By-Side Visits During Pandemic", covering a total of 10 visits.
- Completed comprehensive 40-hour annual theoretical training course at Assomonitor covering critical areas including organ physiopathology, clinical trial regulations, methodology, biostatistics, and ethical aspects, fulfilling requirements of Ministerial Decree Art. 4(1)(b).

Clinical Research Technologic, Roma, Italy — 2022 – 2023
CRA Trainee

Oversaw ongoing monitoring visits whilst ensuring compliance with study protocols and regulatory requirements. Conducted comprehensive site initiation visits to assess and prepare facilities for upcoming trials. Executed close-out visits to verify proper study completion and data integrity. Assessed and validated study participant eligibility through rigorous evaluation of inclusion and exclusion criteria. Verified receipt and distribution of study materials to facilitate research operations. Monitored and documented adverse events, including serious and unexpected cases.

- Supervised proper maintenance and organisation of investigator study files.
- Managed return process for unused or expired study materials at study conclusion.
- Delivered instruction on good clinical practice standards to enhance study reliability.
- Led training and supervision of staff in maintaining integrity and confidentiality of original documents.

Farmacia Camerucci, Roma, Italy **2022 – 2023**
Pharmacist Manager

Managed pharmacy inventory with precision, optimising order and delivery workflows to align with consumer needs. Utilised ProScript Connect for streamlined management and oversight of pharmacy operations and prescription services. Conducted assessments of payment claims for NHS funding accuracy and compliance. Exhibited exceptional communication and interpersonal aptitudes, fostering meaningful interactions with diverse pharmacy clientele and team members. Provided comprehensive patient consultations, delivering insights on potential diagnoses and recommending appropriate medication options.

- Elevated pharmacy profitability by enhancing add-on sales and incentivizing customer loyalty programmes.
- Developed daily shift schedules aligned with peak hours to optimise staff efficiency and minimise customer wait times.
- Led adoption of innovative healthcare initiatives, providing comprehensive support for smoking cessation, weight management, and overall wellness services.

Education & Credentials

Master of Science: Monitoring, Assurance, and Quality Control in Clinical Trials | La Sapienza Di Roma, Roma, Italy, 2023
With 110 cum laude | 40 hours of theoretical training | Certified by assomonitor | Exploring good clinical practices

Certificate: Clinical Research Associate | Clinical research technologic, Roma, Italy, 2023
Certification as CRA, to the company clinical research technologic.

Bachelor of Science: Chimica e tecnologie farmaceutiche | Università La Sapienza di Roma, Italy, 2021

- Conducted thorough research and developed experimental thesis focusing on building predictive QSAR models utilising Machine Learning methods, specifically applied to HDAC6 inhibitors.
- Acquired proficiency in various programming languages including 'programming-foundations with-python,' 'intro-to-computer-science,' and 'intro-to-statistics-machine-learning.' Demonstrated advanced understanding and adeptness in computer programming.

CRA Certified

Languages

English – B2 (Upper Intermediate)