Quality Assurance Auditor

Job Offer

AnaPath Research

AnaPath Research is a CRO located in Barcelona, with extensive experience in carrying out preclinical trials for pharmaceutical laboratories, the chemical industry and other research organizations. In our more than 30 years of activity we have worked with the main pharmaceutical industries being part of different multinationals (RCC, Harlan and Envigo). In November 2019, AnaPath Services acquired the company and re-founded it as AnaPath Research, thus undertaking together a new project of scientific quality and close contact with new and old sponsors.

With a multidisciplinary team of scientific experts, AnaPath Research covers most fields of preclinical pharmaceutical development and chemical and food safety.

Department

The Quality Assurance (QA) Department works closely with Operations departments to ensure that the studies performed at the company comply with the Good Laboratory Practice (GLP) regulations.

Our team consists of senior scientists with an average of 15-20 years of experience auditing toxicology, pharmacokinetics, safety pharmacology and bioanalytics studies.

Position

We are looking for a junior Auditor for our QA department. AnaPath is GLP certified to work with pharmacological, chemical, food additives and phytosanitary products.

The QA team, like the rest of the company, is committed to animal welfare and possess an extensive knowledge in the preclinical development of drugs.

Responsibilities

Act as QA Auditor at the QA Department. Functions will include, but will not be limited to:

- Evaluation of the level of compliance of the Quality Management System implemented according to GLP
- Planning, organizing and executing of internal study audits: study plans, critical phases and study reports
- Planning, organizing and executing of facility-based audits, process-based audits and computerized systems audits
- Planning, organizing and executing of external audits to suppliers and/or subcontracted companies
- Writing of detailed and accurate audit reports that will be given to the personnel involved and Management
- CAPA follow-up and monitoring
- Management and support for qualification and validation of computerized systems and equipment
- Contribute to the Quality Management System by defining SOPs, policies, standards, etc.

Requirements

Life sciences degree, Scientific PhD or MSc

Knowledge of good laboratory practice (GLP) standards

Bioanalysis and/or Toxicology background

Knowledge of computerized system validation processes in accordance with GLP and 21 CFR Part 11 is an asset

Good communication skills (verbal and writing) across functional departments and various levels of management

Pro-active problem solver, ability to work in a team environment and to adapt to organizational changes

Detail oriented and strong ethics and values

Own transport

Advanced English level

Work permit for Spain

Terms of Employment

Indefinite contract

Full-time

Morning work shift

Salary conditions: depending on the applicant's qualifications

Contact

rrhh@anapathresearch.com