Study Coordinator General Toxicology

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 General Toxicology Department performs a full range of studies to support the pharmaceutical, chemical, agrochemical and veterinary industries.

Our team consists of senior scientists with an average of 20 years of experience in toxicology, pharmacokinetics and safety pharmacology, and wide knowledge of the current regulatory guidelines.

We offer scientific advice on study types, study design and choice of dose levels before starting with first-in-human trials.

As a GLP-certified and AAALAC-accredited center, we fully embrace the 3Rs of animal welfare. We are strongly convinced that our animal welfare policies have a positive impact on our studies

Position

We are looking for a Study Coordinator for general toxicology, safety assessment, reprotoxicology, pharmacokinetics, and veterinary regulatory studies. AnaPath is accredited to work with pharmacological, chemical, food additives and phytosanitary products.

The team of the Toxicology Department, 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 Study Coordinator at the General Toxicology Department. Functions will include assisting the Study Director in the coordination of all study-related work (study plans and planning of activities among the various departments involved); planning of study-related activities in LIMS; preparation of tables for reports and Common Technical Document (CTD); monitoring of activities performed to ensure compliance with Standard Operating Procedures (SOPs) and Study Plan and/or Amendment(s); preparation of documentation for archiving; collaboration in generation of SEND tables; cooperation in the monitoring of the Calibration/Maintenance Plan for laboratory equipment. All the above must meet our Quality standards and the requirements of any audits received

Requirements

Life sciences degree or equivalent and experience in working effectively as Study Coordinator

Qualification for staff responsible for directing animal experiments (C) (according to EU Directive 63/2010, RD 53/2013 (Royal Decree) and ECC/566/2015 (Official State Gazette) (Frist provisional regulation, Categories recognized according to Royal Decree 1201/2005) to perform: a) procedures on animals b) designing of procedures and projects)

Good knowledge of good laboratory practice (GLP) standards

Good knowledge of Animal Welfare regulations

Knowledge of regulatory toxicology will be an asset

Good communication skills

Advanced English level

Work permit for Spain

Own transport

Terms of Employment

Indefinite contract

Full-time

Work location: Santa Perpètua de Mogoda and Castellar del Vallès (Barcelona).

Salary conditions: depending on the applicant's qualifications

Contact

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