

SEC is recruiting for an experienced Biostatistician to join our pharmaceutical client on permanent basis. This role is full time at our client's site in Switzerland. Please contact Louise Beka for full details of client and location.

The role responsibilities:

- You will act as the trial statistician for 1 or more clinical trials. You will be responsible for protocol development; case report form review, randomization, IVRS specifications, review of clinical data management documents, writing of the statistical analysis plan, blind review before database lock, statistical analysis and QC of statistical programming.
- You will follow up on the status of the various activities for each of the studies in a project and ensure standardization across studies within the project
- You will work on several clinical trials at the same time and/or manage safety poolings for IND updates, ISS/ISE including all trials performed historically. You will also prepare or supervise preparation of Tables, Figures and Listings required for Summary Documents
- You will understand the regulatory requirements with respect to design and analysis of trials in his/her project.

What we are looking for:

An experienced biostatistician professional with a minimum MSc Qualification in Statistics. We are really looking for professional who has CRO / Pharma experience only. Experience or specialization in Oncology, CNS or Immunology therapeutic area will be highly advantageous. To secure this position you should ideally have experience acting as a Lead Statistician on one or more studies within Phase I-IV.

My client offers highly competitive salary with relocation package for this position, according to expertise and the opportunity to work on some high profile and challenging projects.

To apply please forward your CV to Louise Beka with a short cover note. Alternatively call Louise Beka for further information and a confidential discussion of this opportunity.

Louise Beka
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