

Senior/ Principal Biostatistician

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Company: Avance Clinical

Location: bangalore

Category: other-general

Role Title: Senior/ Principal Biostatistician

Location: Bangalore

Number of open positions: 2

Who are we?

We're Avance Clinical a Contract Research Organization. We offer specialized services supporting drug development for the pharmaceutical and biotechnology industry. We have a rich history in this area with our founding roots dating back to the 1980's. If you already work in this industry you'll know that we are experts in our field.

Why should you join Avance Clinical?

We appreciate that our most important asset is our people. We offer a fun, friendly work environment and a culture that values honesty and respect. We have a strong sense of team. We genuinely appreciate each individual but recognize the collective when achieving success.

We are committed to creating a flexible work environment for our people by supporting them to balance their work commitments and personal needs, both now and into the future.

Broad Purpose

The Senior/ Principal Biostatistician is responsible for performing statistical design, analysis and reporting of clinical studies, and also mentoring of Avance staff in these activities.

Core Responsibilities

Provide statistical advice on study design.

Provide statistical input in the development of study protocols.

Prepare/review randomization schedules.

Prepare/review statistical analysis plans.

Prepare/review listings, tables and figures of study data.

Perform/review statistical analyses of study data.

Prepare/review efficacy sections of clinical study reports.

Client liaison for statistics or study design related issues.

Participate in client meetings.

Mentor role for statistics team members, and also for staff in the Clinical and Data teams, for development of statistics skills.

Provide Business Development support for proposals including statistical presentation/analysis. Other duties as directed by Line Management or delegate

Qualifications, Skills and Experience

Bachelor's degree or higher with an emphasis in life sciences, medical/pharmaceutical sciences and/or statistics.

Minimum of 7- 10 years statistical analysis experience.

Minimum 7-10 years' experience in the pharmaceutical industry / Contract Research Organization (CRO) working in a GCP environment.

Minimum of 3 years' experience in tertiary education and/or equivalent mentoring

Formal GCP training and experience with regulatory filing of statistics.

Experience in clinical or pharmacological research with a focus in a technical writing environment. Proven experience with SAS together with advanced skills in MS Word and MS Excel.

Ability to communicate with clients, senior management and other stakeholders.

Commercial links with pharmaceutical and biotechnology industry (desirable)

What we offer

Stimulating work

Project diversity

An intellectual challenge

An agile & flexible workplace

Opportunity to progress

Stability

Sense of community

A leadership team that are working together on a common goal

Flexible work options

Great tools / tech to do your job

A competitive salary

What next?

As a growing business we're looking for likeminded people to join us – we hope that's you.

To join our team please submit your CV & cover letter as one Microsoft Word document

Alternatively, you could e mail your CV to ramya.yadehalli@avancecro.com

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