Senior or Principal Data Manager

Could you be our new Senior or Principal Data Manager?

Well, if you:

- Are considering a change of scenery.
- Want a job with a significant degree of variation and flexibility.
- Value the freedom it gives to have the opportunity of working from home and perhaps even on part time basis - if preferred
- And can see yourself thriving in a small “Professional Family” of 12 other - really friendly, forthcoming, highly skilled and helpful - colleagues.

… the answer could actually be a “yes” – and you should definitely consider reading on - and getting in touch with us.

Who we are – professionally:

We are a Contract Research Organization (CRO) who have specialized in Data Management, Statistical Programming and Statistics – for more than 10 years.

We offer all Biometrics services required in relation to clinical and device trials.

We are data managers, statistical/SAS programmers and statisticians, and our office is located in Birkerød.

Who we are – personally:

When we founded BioStata in 2010 it was with the clear vision of establishing a company build on a deliberately small team of around 15.
This was chosen in order to combine the strength of deep expertise normally found in large organizations – with the flexibility and freedom only found in small companies.

Having lived this vision for more than a decade now, we can safely say that some of the keywords describing our working atmosphere are: Trust, helpfulness, open-mindedness, kindness, responsibility, opportunity for influence – and a high degree of professional dedication to what we do.

A job in the Life Science industry is often associated with a lot of pressure, hectic deadlines, and stress.

At BioStata we deliberately want to take this in a better direction - creating a more meaningful and human-friendly work life.
We are therefore very mindful of having personal flexibility, setting realistic timelines – and aiming for a workload that is as manageable and smooth as possible, given that we are a consultancy company.

What your important contribution to the team will be:

You should – preferably - have one or more of the following areas of competencies:

- Handling day-to-day tasks on an ongoing clinical trial such as:
  - eCRF setup. (We use Zelta or Viedoc)
  - Authoring and reviewing DM documentation.
  - Handling of external data and Query management.
  - Data cleaning in cooperation with our programmers and statisticians.
  - Project management and communication with sponsors and other stake holders.
  - Database lock.
- Provide input on timelines, resources, and process improvements.
- Perform DM oversight on trials carried out by other CROs on behalf of sponsor.
- Experience with CDISC.

In addition to the above-mentioned competencies, we need you to be:

- Fluent in English.
- Able to work structured, independently and with a flexible mindset.

You will be working together with our other programmers, data managers and statisticians in professional trial teams, where we all benefit from combining decades of knowledge and experience with the bright ideas from the younger members of our professional family.

You will be involved – as a part of the trial team – in a great variety of projects, tasks, and customers, where the work is mainly carried out at our office, from home or sometimes at the customer site, depending on the nature of the task.

So… are you tempted to join us?

Well – at this point you probably have a much better idea of this than you did just a few minutes ago...

If you tend towards “yes” – or if you are curious about getting to know more about us and the position – please contact Tu Duyen le Thi, Director – Biometrics, tl@biostata.com, tel.: +45 2774 8202.

We are very much looking forward to hearing from you – and perhaps welcoming you as an important part of our professional family.