A+ Science Newsletter

“Talent wins games, but teamwork and intelligence win championships.”

Michael Jordan

We are very happy to announce the appointment of Eva Fredriksson as our new Head of Clinical Operations at A+ Science.

Eva Fredriksson has extensive experience from leading positions in life science industry and healthcare. Eva comes most recently from the Stockholm Region, where she was in charge of developing collaboration between the National Quality Register and the Life Science industry, as well as work with change management. Eva has also worked for many years with clinical trials at Roche, the pharmaceutical company. During the last years at Roche Eva was responsible for the entire clinical trial operations in Sweden. Prior to that and for several years, she worked with quality development and research projects in oncology.

“I started at A+ Science in the beginning of November and after being here for just couple of weeks it feels very good, it’s really an exciting company. With my background as a research nurse and having worked at a pharmaceutical company involved with clinical trials, it already feels like I'm at the right place.

It also feels exciting and tempting to work at a Swedish company in the CRO industry. Through our work we get the opportunity to support and ensure that patients and healthcare will benefit by new drugs and new technologies as well

as contribute to the continuous development of Swedish life science," says Eva Fredriksson.

Eva Fredriksson

Eva be responsible for the oversight, management and delivery of clinical trials. She will provide strong leadership to her team and build on the encouraging progress. We are excited to see her step up to this key role at A+ Science

Are you aware of all the services A+ Science provides?

- Our services include clinical trial management, we can conduct the entire clinical study, from start to finish or parts of it, depending on our customers' needs.
- We also provide full pharmacovigilance services, pre-and post-marketing, including 24/7 availability of QPPV, Deputy QPPV and Drug Safety Physician, safety database license including MedDRA license, ICSR processing (pre-and post-marketing) including medical review and submission to Eudravigilance, Monitoring of new legislation, literature search and review (global and local, ICSR collection from EVWEB, entry and maintenance into the XEVMPD, PSMF maintenance, signal management, writing of SOPs, PSURs, DSURs and RMPs.
- We can empower your team by outsourcing personnel (short- and long-term solutions).

- Since 2004, we offer services within Continuing Medical Education (CME) in collaboration with University Hospital of Umeå and the Sahlgrenska Academy, University of Gothenburg.

Contact us at info@a-plusscience.com to find out more about how our team at A+ Science can help you conduct your clinical trial. Our aim is to assist you reach your goals and objectives.

Earn Trust- Make Difference

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