A+ Science Newsletter

“I never dreamed about success, I worked for it.”

Estée Lauder

A+ Science is proud to introduce Björn Edwall, the company’s new Quality Assurance (QA) Consultant.

Björn has more than 30 years of experience in leading positions within the pharma industry at global companies including Novartis and AstraZeneca. He has extensive Nordic hands-on experience in Quality Assurance, Patient Safety, Emergency Management, Regulatory Affairs, Clinical Trials, Medical Information and Licensing. Björn has also more than a decade’s experience of serving on the Board of Trustees for the Swedish Pharmaceutical Insurance Company.

He holds a PhD in Pharmacology and a DDS dental degree, both from Karolinska Institute in Stockholm.

A key requirement for anyone involved in the conduct of clinical research is Good Clinical Practice (GCP) training. GCP is the ethical and practical standard to which all clinical research is conducted. The principles of GCP state that: Everyone involved in conducting a trial should be qualified by education, training and experience to perform his or her respective task(s).

At A+ Science we make sure that everyone follows those principles. We have regular GCP trainings. Björn has already organised and led 5 such training sessions this year.

We are very happy with our collaboration with Oral Pharma Edwall AB, a company based in Stockholm and headed by Björn Edwall. Also, very happy to have Björn on board as QA Consultant providing independent, objective advice to A+ Science on strategies to improve the quality of its services, i.e. regulatory compliance services.

“A+ Science is a very professional CRO offering a wide range of services in the areas listed below. I see that A+ Science management is truly committed to having a quality system platform to support full regulatory compliance as well as customer needs. I look forward to my collaboration with A+ Science,” says Björn Edwall.

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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