A+ Science Newsletter

"When you reach the end of your rope, tie a knot and hang on".

Abraham Lincoln

We hope all our readers have had a wonderful relaxing summer. We are back with loaded batteries and loads of energy and are looking forward to start working with several new projects.

In this month's newsletter we are proud to announce that Karuna Vuppalapati has just joined our team to strengthen our Pharmacovigilance Department. We are all delighted to have her at A+ Science.

Karuna Vuppalapati is a clinician and researcher with medical degree (M.B.B.S) from Dr. NTR University of Health Sciences, in India. To pursue a career in research she moved to Sweden and obtained master's in Molecular Medical Biology from Örebro University and PhD in Medicine from Karolinska Institute. She then got post graduate diploma in Diabetes from University of South Wales and worked as a clinician with focus on diabetes prevention and management in India. Karuna moved to Sweden in 2020 and pursued courses such as clinical trials in Medicine, pre-clinical safety assessment and pharmacovigilance from Uppsala University. Karuna joined the A+ Science team just before summer.

Karuna Vuppalapati

“I found that the coordination and collaboration among the team members, work culture, office atmosphere and timely response by everyone to be very balanced, organised and structured’’. I am very happy to be a part of the great team at A+ Science and I am delighted with the warm gestures from everyone. The current opportunity as a drug safety associate at A+ Science provided a chance to explore a different career path. The opportunity gave me a detailed insight about pharmacovigilance, and I am excited for the new challenges”, says Karuna Vuppalapati.

Are you aware of 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 and MHRA, Regulatory intelligence, literature search and review (global and local), ICSR collection from EVWEB, monitoring of MHRA ICSR portal, XEVMPD and 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 reach your goals and objectives.

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