



ADVANCING YOUR INNOVATION

Quality Assurance Reviewer

Toxikon Corporation is a preclinical contract research organization. We contract and partner with biotech, pharmaceutical and medical device industries to deliver exceptional product development services from concept to final product.

We are seeking a quality assurance reviewer who will be responsible for the implementation, functioning, and monitoring of quality assurance activities of Toxikon's In Vivo Biocompatibility services. This includes monitoring biological and chemical studies, and creating reports in conformance with GLP practices. Candidates will review both GLP and non-GLP reports for accuracy with reference to protocols, quality assurance standards, standard operational procedures, laboratory precision and accuracy limits, as well as reviewing all protocols for compliance and accuracy.

Other quality assurance responsibilities include maintaining equipment performance, calibration records and preventative maintenance logs. Included duties are performing system and phase audits according to GLP and ISO guidelines, as well as all other activities required from quality assurance.

A bachelor's degree in life sciences is required, preferably in biology. Candidates should also have at least one year experience in an FDA-regulated environment, with basic knowledge of study design and review practices. Must have strong documentation and coordination practices, be detail oriented and enjoy high work volume via computer.

Toxikon offers a comprehensive benefits program including: paid vacation, sick and personal time; 401(k) plan with company match; medical, dental, vision care, life, and short and long-term disability insurances; flexible spending accounts; employee assistance program; employee referral bonus; discount programs; and more. Toxikon is an equal opportunity employer.