



Study Director

Toxikon Corporation, originally founded in 1977, is a Contract Research and testing company with laboratories in Bedford, MA and Belgium. Our mission continues to focus on providing quality pre-clinical services, competitively priced, with on-time delivery for the Pharmaceutical, Medical Device and Biotech industries.

Toxikon is seeking a Study Director. The overall responsibility of the Study Director is for the technical conduct of safety testing studies according to GLP regulations and represents the principal point of study control for activities related to FDA/ISO/Japanese compliance for in-vitro/in-vivo studies. Responsibilities include but are not limited to: protocol approval, conduct of the study experiment according to protocol, documentation of protocol deviations and amendments, reviewing data, approving data analysis, ensuring accurate incorporation of results and conclusions into the study report prior to submission to the Sponsor, working closely with the Sponsor to complete the study, and archiving of study records. The Study Director is responsible for meeting study timelines and quality standards, and eventually for training staff scientists in study-related standard operating procedures.

The successful candidate will have 2 -5 years experience, possess excellent written and oral communication skills, with the ability to be a team player. A MS or Doctorate in a Life Science discipline is required.

For additional information about Toxikon, please view our website at www.toxikon.com

Submit resumes to:

Human Resources
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Email: hr@toxikon.com

Toxikon employs 160 people and offers a generous benefit program including health, dental, fsa, std, ltd, life insurance and matching 401K plan. Toxikon is an EOE.