As a Director of Regulatory Affairs Regulatory Compliance Associates® Inc, Linda's responsibilities include evaluating and preparing both USP global regulatory strategies and submission preparation for marketed drug products and generic drug programs, as well as client engagement. Her background includes comprehensive laboratory experience, R&D and product development; successful regulatory strategy and submission preparation; FDA inspection and 483 / Warning Letter remediation; as well as project management and supervisory experience in the pharmaceutical industry. Linda earned a Bachelor of Science in Biology from the University of Kansas., is RAC certified by the RAPS and serves as a volunteer on the RAPS Education Committee.