Stephanie Ranck
Sr Research Scientist, Global Regulatory Affairs-CMC Biotech/Insulins

Ms. Ranck brings 25 years of experience to the pharmaceutical industry, including QA/QC for generic API manufacturing, contract manufacturing and testing, and sterile drug product manufacturing. For the past six years, she has applied her knowledge of sterile pharmaceutical manufacturing as a Chemistry, Manufacturing Controls regulatory scientist. Registration experience includes new products, line extensions, and post-approval change management for multi-product facilities (manufacturing biologics), and globally registered parenteral products.