Federico Goodsaid, PhD Biosketch

Federico Goodsaid is SVP for regulatory Affairs at Ariana Pharma and Principal Consultant at Regulatory Pathfinders LLC, a regulatory consulting practice focused on regulatory strategy and pathways required for precision medicine therapeutic and diagnostic products. Prior to this appointment, he was the Senior Vice President of Product Development and Regulatory Affairs at TOMA Biosciences, where we worked on the preclinical and clinical development and regulatory submission for a 130-gene targeted NGS onco panel. His previous work at Vertex focused on therapeutic product submissions in cystic fibrosis and diagnostic and biomarker development and regulatory submissions in rare diseases. He was previously Associate Director for Operations in Genomics and Biomarker Qualification Coordinator at the Office of Clinical Pharmacology/Office of Translational Sciences/ Center for Drug Evaluation and Research/ U.S. FDA, working on the regulatory application and development of genomics and biomarkers. Before joining the FDA, he served as a Senior Staff Scientist at Applied Biosystems and Lead for the Molecular Toxicology Group at the Schering-Plough Research Institute. Following his Ph.D. from Yale University in Molecular Biophysics and Biochemistry he was trained as a Postdoctoral Fellow at Cornell University and at Washington University in St. Louis.