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Anson Group adds Gina Labella-Tulin as a Regulatory Associate

(Indianapolis, Indiana) July 13th, 2011 The Anson Group announces today the addition of Gina Labella-Tulin as a Regulatory Associate. Ms. Labella-Tulin has served in a number of project management and compliance roles in the biotechnology and diagnostics industries. She comes to Anson from Covance Central Labs, where she served as a project team manager for development and launch of projects, which included defining and implementing activities such as risk assessment, contingency planning, and resource planning.

Prior to her tenure at Covance, Ms. Labella-Tulin served in the regulatory compliance organization at Roche Diagnostics. She was responsible for ensuring that new product launches and existing product changes adhered to applicable regulatory and quality system requirements. In addition, she supported product safety compliance in Medical Device Reporting (MDR), Medwatch reporting, and escalating potential correction and removal issues.

"We're delighted to have Gina join Anson at a pivotal time in the industry's growing development of biotechnology and diagnostic products, an area in which Gina has particularly relevant experience" states Colleen Hittle, Anson's CEO. "Her enthusiasm and passion for the issues in these critical areas will have immediate and long term impact for our clients who need to maintain compliance with important FDA regulatory requirements."

About Anson Group

The Anson Group is a woman-owned independent regulatory consulting organization headquartered in Indianapolis, IN. Since 1996, the Anson Group has provided tailored, client-focused regulatory compliance solutions for medical device, pharmaceutical, biologics, combination product and most recently, connected health clients. Looking at issues in the context of our clients' resources and culture, our team of industry experts identifies available regulatory options and helps select the best pathway for overall client success.