Mary Anne Gfell, MBA, CCRP

Mary Anne Gfell serves as a Regulatory Compliance Advisor for Pearl Pathways. She has over 15 years of managing pharmaceutical and medical device product development, non-clinical and clinical research, and leading teams. She is a certified and experienced project manager having led complex multidisciplinary programs both in her work at Covance and Indiana University.

Previously, she was a Global Project Manager for Covance, a multi-national CRO where she managed central laboratory services global trials. She led a cross functional team and managed complex global trials requiring her to integrate multiple stakeholder's goals, complex regulatory requirements, and aggressive deadlines. She is adept at regulatory requirements, clinical trials, medical and technical writing, creation and oversight of GXP quality systems, and project management for outsourced services for pharmaceutical companies.

Gfell received her Masters of Business Administration and Bachelors of Science from Indiana University. Gfell holds her project management certification and is an active certified member of the Society of Clinical Research Associates.

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