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**FOR IMMEDIATE RELEASE**

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**Pearl Pathways Hires Michele Taylor**

*Experienced mechanical engineer and life science project manager joins Pearl Pathways*

INDIANAPOLIS, INDIANA – November 4, 2016 — [Pearl Pathways](http://www.pearlpathways.com/) announces the hiring of Michele Taylor as a regulatory compliance Analyst serving medical device and biopharmaceutical life science companies.

Taylor brings 12 years of experience in Project Management in Life Sciences, Medical Device Product Development, and Manufacturing to the Pearl Pathways team. Previously at Beckman Coulter, she managed project work on custom integrated lab automation systems for the drug discovery and research industries. Taylor managed new Product Development projects and provided engineering support for existing product lines and manufacturing of medical devices at Cook Urological and Baxter.

Taylor delivers regulatory services for Pearl Pathways’ clients including quality compliance support, regulatory pathway assessments, and eCTD publishing for global health authority submissions. Taylor conducts auditing services helping companies make improvements to establish more effective Quality Management Systems. She also provides project management support to Pearl’s service delivery.

Taylor has a Bachelor of Science in Mechanical Engineering from Indiana University – Purdue University at Indianapolis and is Six Sigma certified. She is a trained project manager and adept at eCTD publishing.

Diana Caldwell, President and CEO shares, “Michele brings strong engineering and manufacturing background to the Pearl team. Our clients will benefit from her product development experience and incredible project management skills. She has broad work experience having worked with pharmaceuticals, traditional medical device, and in-vitro diagnostics (IVD) companies. We are thrilled to have her join the team.”

## About Pearl Pathways

***Pearl Pathways*** is a comprehensive life science product development services company. Our experienced team is obsessed with expediting life science product development regulatory pathways. We have three business units to serve you:

***Pearl IRB*** is a full service commercial Independent Review Board that provides human research IRB reviews, IRB exemptions and waivers, and also offers support for research protocol/ICF medical writing, site assessments, and monitoring services.

***Pearl ReGXP*** is a regulatory and quality compliance consulting practice that provides regulatory filing guidance, conducts global health authority negotiations, develops/improves quality systems, and delivers GMP/GLP/GCP auditing services.

***Pearl IDEAS*** provides strategic product development assistance, third party vendor selection and management strategies, due diligence services, and sales and marketing services for drug, biologic and device companies.

To learn more, please visit us at [www.pearlpathways.com](http://www.pearlpathways.com), call us at (317) 899-9341, or email [contact@pearlpathways.com](mailto:contact@pearlpathways.com). Pearl Pathways is located in Indianapolis, Indiana, and is AAHRPP accredited and a WBENC certified woman owned business. For media inquiries, contact [contact@pearlpathways.com](mailto:contact@pearlpathways.com).