Cathy Craft, RAC
Regulatory Consulting
Myraqa Senior Director, Illumina

Cathy Craft has provided regulatory consulting services for Myraqa since October 2012, bringing her 35 years of experience in the medical device manufacturing industry to IVD consulting. In this role Cathy has provided consultation on regulatory and clinical issues and strategies for a wide range of IVD products and services.

Prior to joining Myraqa, Cathy retired from her position as the Vice President of Regulatory Affairs at Siemens Healthcare Diagnostics, Inc. In that role, Cathy had responsibility for directing global regulatory affairs strategy and activities for all Siemens *in vitro* diagnostic product lines which included: General Chemistry, Immunochemistry, Hemostasis, Plasma Proteins, Molecular, Point of Care, Microbiology and Lab Automation. Her organization developed and implemented strategies to achieve global registration of new and modified products, as well as compliance with changing global regulatory requirements. Cathy led the integration of three regulatory organizations: Bayer, DPC and Dade Behring into a cohesive Siemens Diagnostics regulatory organization, implementing the registration and labeling transition to enable marketing all products under the corporate name.

In previous positions with Dade Behring as well as with DuPont Medical Products Department, Cathy has had extensive cross functional business experience including management positions in Technical Marketing, Manufacturing, and Quality Assurance.

Cathy received her Bachelor of Science in Medical Technology from the University of Texas at Austin. She began her career as a Medical Technologist at Methodist Hospital, San Antonio, Texas. Cathy has held a Regulatory Affairs Certification (R.A.C.) through the Regulatory Affairs Professional Society since 1998 and has been on the Board of Directors for AMDM since 2001.

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