Debra J. Rasmussen, MBA, RAC · Senior Director of Global Regulatory Affairs, Johnson & Johnson's Janssen Pharmaceuticals

Deb's regulatory and quality career has spanned both large and small companies and both US and global programs. She has been in medical devices for over 25 years and gained her experience working at Bio-Rad Laboratories, Roche Molecular Systems, Chiron Corporation, Applied Imaging Corporation, and Visible Genetics. She joined Johnson and Johnson in 2003. Johnson and Johnson has recently established a new diagnostic group in Janssen Pharmaceuticals to support all therapeutic area. This new diagnostic group, Janssen Diagnostics, is integral in the companion diagnostic efforts. For Janssen Diagnostics, she is responsible for development of regulatory strategy, policy, and registration of the *in vitro* diagnostic devices associated with Johnson and Johnson drug development and commercialization.