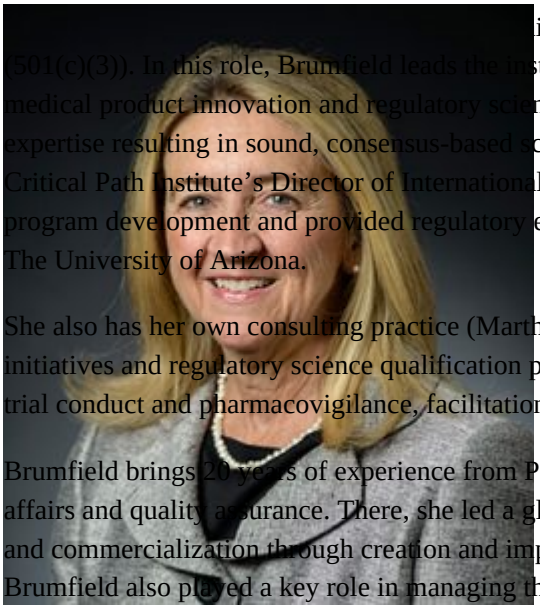
Martha A. Brumfield, PhD



ief Executive Officer of Critical Path Institute, an Arizona based non-profit 501(c)(3)). In this role, Brumfield leads the institute in its mission to catalyze the development of new tools to advance medical product innovation and regulatory science which is accomplished by leading teams that share data, knowledge and expertise resulting in sound, consensus-based science. Brumfield assumes the role of CEO after most recently serving as Critical Path Institute's Director of International & Regulatory Programs. In that position, she helped guide international program development and provided regulatory expertise to consortia. She is also Associate Professor, College of Pharmacy, The University of Arizona.

She also has her own consulting practice (Martha A. Brumfield LLC) focusing on concordance in global regulatory initiatives and regulatory science qualification programs. Other areas of focus in her practice include excellence in clinical trial conduct and pharmacovigilance, facilitation of scientific consortia and programs supporting patient access to medicines.

Brumfield brings 20 years of experience from Pfizer Inc., most recently, as senior vice president of worldwide regulatory affairs and quality assurance. There, she led a global team that supported lifecycle pharmaceutical research, development and commercialization through creation and implementation of regulatory strategies and quality assurance oversight. Brumfield also played a key role in managing the broader company relationships with global regulators, trade associations, academics and others on regulatory policy issues. She served on corporate governance initiatives including the planning and implementation of mergers and acquisitions and led her departments through these periods of significant change.

She is Chair of the Board of Directors for the Regulatory Affairs Professional Society and chairs the Global Curriculum Coordinating Committee with FDA's Office of International Policy, which is developing a curriculum for regulators in developing countries. She is also active with global nonprofits, including the Regulatory Harmonization Institute and GlobalMD, where she delivers educational workshops on regulatory and clinical trial topics in Asia. She has served on and contributed to the Institute of Medicine consensus committees, which were commissioned by U.S. FDA focusing on global regulatory systems and on falsified and substandard drugs. She also serves on the Steering Committee of the Multi-Regional Clinical Trials Center of Brigham and Women's Hospital and Harvard.

Brumfield earned a B.S. and an M.S. in chemistry from Virginia Commonwealth University, a Ph.D. in organic chemistry from the University of Maryland, and served as a post-doctoral fellow at The Rockefeller University.