Pamela J. Weagraff, MBA
Director, MedTech Regulatory

Pam leads the MedTech Regulatory team for IQVIA. Pam delivers strategic and tactical consulting services to optimize business strategies.

Pam's primary focus is on commercialization of innovative technologies:

- Novel drug delivery devices for combination products
- Digital Health: software platforms, mobile medical apps, wearables and implantables
- Innovative electro-mechanical software-controlled devices, e.g., new imaging modalities
- Active and passive implantables

Pam brings 25 years of medical device experience in development and deployment of strategies throughout the product life cycle in early stage marketing, product management, regulatory, clinical and quality. Her industry experience included a variety of start-ups, such as MediSpectra and T2 Biosystems, midsize and large companies such as Hewlett Packard's Medical Products Group (now Philips Health), as well as extensive international experience from working and living in the EU and travels to the Asia Pacific region.