In addition to his role in FDA, Lawrence is an adjunct Professor at the University of Michigan and the rapporteur of ICH M4Q(R1) revision. An Associate Editor of the *AAPS Journal*, he has authored/co-authored over 150 papers and given over 400 invited presentations. He is a co-editor of the books entitled "*Biopharmaceutics Applications in Drug Development"*, "*FDA Bioequivalence Standards*", and "*Developing Solid Oral Dosage Forms: Pharmaceutical Theory and Practice, 2nd Ed.*"