

A Deep Dive: FDA Draft Guidance on Statistical Approaches to Establishing Bioequivalence

March 14th, 2023 | 10:00 AM – 12:00 PM (ET)

AGENDA

SBIA Introduction	Forest “Ray” Ford, PharmD, BCPS <i>Captain, USPHS</i> Small Business and Industry Assistance (SBIA) Division of Drug Information (DDI) Office of Communications (OCOMM)
Introduction	Liang Zhao, PhD <i>Director</i> Division of Quantitative Methods and Modeling (DQMM) Office of Research and Standards (ORS) Office of Generic Drugs (OGD)
Overview (contents of the guidance)	Stella C. Grosser, PhD <i>Director</i> Division of Biometrics VIII (DB VIII) Office of Biostatistics (OB) Office of Translational Sciences (OTS)
Statistical Test for Population Bioequivalence	Sungwoo Choi, PhD <i>Mathematical Statistician</i> DB VIII OB OTS
Statistical Approaches to Establishing Bioequivalence – Specific Situations: An Overview of In Vitro Release Test (IVRT), In Vitro Permeation Test (IVPT), and Earth Mover’s Distance (EMD) comparative studies	Kimberly Raines, PhD <i>Branch Chief</i> Division of Biopharmaceutics (DB) Office of New Drug Products (ONDP) Office of Pharmaceutical Quality (OPQ)
Statistical Methods for Narrow Therapeutic Index and Highly Variable Drug Products	Donald Schuirmann, MS <i>Expert Mathematical Statistician</i> DB VIII OB OTS
Comparative Clinical Endpoint Bioequivalence Studies	Fairouz Makhoul, PhD <i>Deputy Director</i> DB VIII OB OTS

Bioequivalence Studies in Multiple Groups	Wanjie Sun, PhD <i>Lead Mathematical Statistician</i> DB VIII OB OTS
Adapted Design for Bioequivalence Studies	Wanjie Sun, PhD
Bioequivalence Statistics for Adhesion and Irritation Studies	Somesh Chattopadhyay, PhD <i>Lead Mathematical Statistician</i> DB VIII OB OTS
Dose Scale Analysis to Support Bioequivalence Assessment	Meng Hu, PhD <i>Team Lead</i> DQMM ORS OGD
Bioequivalence Studies Using Multiple References	Liang Zhao, PhD
Recommendations in the 2022 Revised Bioequivalence Statistical Guidance and Bioequivalence Assessments	Zhen Zhang, PhD <i>Senior Pharmacologist</i> Office of Bioequivalence (OBI) Division of Bioequivalence I (DB I) OGD
Panel Discussions	Liang Zhao, PhD, Stella C. Grosser, PhD, and Lanyan (Lucy) Fang, PhD <i>Deputy Director</i> DQMM ORS OGD Ying Fan, PhD <i>Lead Pharmacologist</i> Division of Clinical Review (DCR) Office of Safety and Clinical Evaluation (OSCE) OGD
Closing	
