U.S. FOOD & DRUG ADMINISTRATION

CENTER FOR DEVICES AND RADIOLOGICAL HEALTH

DEVICES GOOD MANUFACTURING PRACTICE (DGMP) ADVISORY COMMITTEE MEETING

MARCH 2, 2022

Name	Affiliation	Role
Yadin David, EdD, PE	Principal Biomedical Engineering Consultants, LLC Houston, TX	Member/Chair
Jeri Culbertson, DNP	Infection Prevention Consultant Focus on Zero, LLC Rapid City, SD	Member
Lisa Dimmick, MS	Team Leader U.S. Nuclear Regulatory Commission Rockville, MD	Member
Gordon Gillerman	Director Standards Coordination Office National Institute of Standards and Technology Gaithersburg, MD	Member
Chiaoyun (Benson) Kuo, PhD	Director Regulatory Consulting Center Asst. Professor, Dept. of Regulatory & Quality Sciences School of Pharmacy, University of Southern California Los Angeles, CA	Member
Alisha Loy, LSSBB, CRCST	Quality and Operations Manager University of Iowa Hospitals and Clinics Iowa City, IA	Member
Elise Owen, MBA, PMP	Standards Executive U.S. Environmental Protection Agency Washington, DC	Member

Robert Phillips, MBA, RAC	Vice President Quality & Regulatory Siemens Healthineers Malvern, PA	Industry Representative
Scott Sardeson, RAC	International Regulatory Affairs & Quality Compliance Director 3M Company, Health Care Business St. Paul, MN	Industry Representative
Keisha Thomas, MS, MHS, CQIA, RAC	Acting Associate Director for Compliance and Quality, CDRH	Food and Drug Administration
Melissa Torres, ME, MS, CQA	Associate Director for International Affairs, CDRH	Food and Drug Administration
Kimberly Lewandowski-Walker	Senior Regulatory Officer Medical Device Single Audit Program, CDRH	Food and Drug Administration
Karen Masley-Joseph, MBA, CMQ-OE, CSSGB	Senior Advisor Office of Medical Device and Radiological Health Operations, ORA	Food and Drug Administration
Anne Reid, MS	Deputy Program Director Office of Medical Device and Radiological Health Operations, ORA	Food and Drug Administration
Jarrod Collier, MS	Office of Management Designated Federal Officer, CDRH	Food and Drug Administration