

Mastering Regulatory Sterilization & Microbiology

September 21, 2022

8:00–8:30	Check-In and Breakfast Buffet	
8:30–9:30	Cleanroom Essentials for Manufacturers	Joe Brinkman
9:30–10:15	Cleanroom Investigation Clue	Joe Brinkman
10:15–10:30	Break	
10:30–12:00	EO Validations, Standard Updates, Residuals, Tolerable Contact Limit & Risk Assessment	Joe Brinkman
12:00–1:00	Lunch	
1:00–1:45	EO Validation Shoots and Ladder	Joe Brinkman
1:45–2:45	Cleaning	Staci DeMoss
2:45–3:00	Break	
3:00–4:00	Disinfection	Staci DeMoss
4:00–5:00	Sterilization	Staci DeMoss

September 22, 2022

7:30–8:30	Breakfast Buffet	
8:30–9:15	What Are Human Factors & What Are They Doing in My Device?	Staci DeMoss
9:15–10:15	Workshop: Reusable Devices	Staci DeMoss
10:15–10:30	Break	
10:30–11:30	Understanding Product Adoption – Ethylene Oxide & Radiation	Joe Brinkman
11:30–12:00	Antimicrobial Products	Staci DeMoss
12:00–1:00	Lunch	
1:00–1:45	Packaging Validation	Joe Brinkman
1:45–2:30	Current FDA LAL Guideline – The guideline, USP, AAMI, TIR, FDA Q&A	Joe Brinkman
2:30–4:30	Break & Final Exercise	Joe Brinkman / Staci DeMoss