

2023 RSQ Schedule at a Glance

Breakout Session Topics:

- Regulatory
- Quality
- Science
- Dietary Supplements

Monday, September 18

10:00 AM - 11:45 AM

Quality/Manufacturing
Committee Meeting
(Members Only)

10:00 AM - 11:45 AM

Regulatory and Scientific Affairs
Committee/PPS Meeting
(By Invitation Only)

12:00 PM - 3:45 PM

Regulatory and Scientific Affairs
Committee Meeting & Lunch
(Members Only)

4:30 PM - 6:00 PM

Early Arrivers Reception

Tuesday, September 19

7:00 AM - 4:30 PM

Registration

7:00 AM - 8:00 AM

Breakfast

8:00 AM - 8:15 AM

Welcome Remarks:
David Campbell & Lisa Parks
Award Presentation: Lauren Quinn

8:15 AM - 9:00 AM

General Session: FDA CDER
Leadership Update

9:00 AM - 10:00 AM

General Session: ChatGPT:
Opening Bell for a New Era of AI

10:00 AM - 10:30 AM

Break

10:30 AM - 11:30 AM

General Session: FTC Update

11:45 AM - 12:30 PM

Driving Innovation with RWE from
Consumer Generated Content ●

FSVP for DS & OTC: The Alphabet
Soup of Import Quality
Compliance ●

Principles of Science-Led Self-Care ●

Regulatory Modernization of the
Self-Care Aisle: What is Now? &
What is Next? ●

12:30 PM - 1:15 PM

Lunch

1:15 PM - 2:00 PM

General Session: Reagan-Udall
Foundation Update

2:15 PM - 3:45 PM

Consumer Medical Device
Innovation ●

Impurities: Benzene, Nitrosamine
& Other Compounds ●

Plant Tissue Culture as a Source
for Bioactive Compounds Used in
CH Products ●

Using RWE in Self-Care Product
Research & Submissions ●

2023 RSQ Schedule at a Glance

Tuesday, September 19

3:45 PM - 4:00 PM

Break

4:00 PM - 5:30 PM

An Overview of CDRH Regulatory Initiatives ●

Best Practices for Compliant Handling and AER Reporting for OTCs & DS ●

Drug Take-Back: State of Play ●

OTC Monograph under OMUFA: First 3 Years & A Look Ahead ●

5:30 PM - 6:30 PM

Reception

Wednesday, September 20

7:00 AM - 12:30 PM

Registration

7:00 AM - 8:00 AM

Breakfast

8:00 AM - 9:00 AM

General Session: Color Additives and Flavors in OTCs

9:15 AM - 10:45 AM

Developing Beneficial Dietary Supplements for Women ●

Established Ingredients Facing New Scientific, Regulatory & Legal Challenges ●

Key Strategies for Enhancing Diversity in Clinical Trials ●

9:15 AM - 10:45 AM (cont'd)

Protecting the Supply Chain and Product Integrity ●

10:45 AM - 11:15 AM

Break

11:15 AM - 12:45 PM

General Session: FDA Leadership

12:45 PM - 1:45 PM

Lunch

1:30 PM - 3:00 PM

Workshop: Applying the Principles of RWE/RWD

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