RAPS

2021 Conferences and Events

Online Workshop | European Online Workshop | Regulatory Conference | RAC Exam

JANUARY

19–20
Biologics CMC: Future Trends, Current Trends and Regulatory Challenges

20–21
Learn the Ins and Outs of E-Filing With the China NMPA

FEBRUARY

9–10
Biologics CMC: Phase-Appropriate Product Development

11–12
IVDR: Practical Implementation of the New Regulation for IVDs

16–17
Root Cause Investigation for CAPA

18–19
Pediatric Drug Development: Developing a Successful Strategy Through Case Studies

MARCH

9
Postmarket Surveillance in Europe

10–11
The Impact of the EU MDR on Legacy Devices

15–16
China NMPA Regulations and Processes for Medical Devices

17–18
Cybersecurity Unauthorized

18–19
Design Controls and Human Factors

22 Mar–30 Apr
Spring RAC Exams

22–23
Introduction to Electronic Regulatory Submissions in the eCTD Format

24
Data Management and Digitalization in Regulatory Affairs

25–26
Intermediate Course for Regulatory Submissions in eCTD Format

APRIL

13–14
Biologics CMC: Future Trends, Current Trends and Regulatory Challenges

14–15
Software as a Medical Device

MAY

5–6
US Regulatory Essentials for Medical Devices

10–14
2021 EuroConvergence

12–13
Clinical Pathways for China NMPA Registration

25–26
The Impact of the EU MDR on Legacy Devices

For our most up-to-date calendar, visit **RAPS.org/events.**

Online Workshop | Virtual Event | Regulatory Conference | RAC Exam

JUNE

8–9
Expedited Pathways (US/Global) — Medical Devices

9
Regulatory Essentials, Pharmaceuticals

16–17
In Vitro Diagnostic Regulation (IVDR)

JULY

12 Jul–20 Aug
Summer RAC Exams

13–14
Biologics CMC: Phase-Appropriate Product Development

27–28
Software as a Medical Device

AUGUST

16–17
China NMPA — In Vitro Diagnostics (IVD)

31 Aug–Sep 1
The Impact of the EU MDR on Legacy Devices

For even more live virtual learning from RAPS, check out our webcasts and other special events at **RAPS.org/events.**

SEPTEMBER

11–14
2021 Convergence

OCTOBER

5–6
Cybersecurity Unauthorized

7–8
Regulatory Intelligence

11–12
Introduction to Electronic Regulatory Submissions in the eCTD Format

14–15
Intermediate Course for Regulatory Submissions in eCTD Format

19–20
Design Controls and Human Factors

20–21
Developing a Global Unique Device Identification (UDI) Program

NOVEMBER

1 Nov–10 Dec
Autumn RAC Exams

2–3
Biologics CMC: Future Trends, Current Trends, and Regulatory Challenges

10–11
Rollercoaster Ride Ahead: Time Is Short for EU IVDR Transition

15–16
Expedited Pathways (US/Global) — Medical Devices

DECEMBER

1–2
Planned Changes to Regulations Impacting Medical Devices

14–15
Biologics CMC: Phase Appropriate Product Development

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