

Professional Development Resources Calendar 2024

TOPRA's guide to help plan your training and budget allocation for the year ahead.



Masterclass courses can be taken as part of the MSc Regulatory Affairs (Medicines/Medical Devices). Get the recognition you deserve for the training you undertake. topra.org/qualification



Do you have teams who need training on particular topics? Do you want training tailored to the specific needs of your organisation? TOPRA's bespoke in-house training provides the solution. Find out more by emailing training@topra.org



If you would like to learn more about the opportunity to become a Chartered Scientist or Registered Scientist in Regulatory Affairs, please visit: topra.org/charteredscientist

Further information

Read more about the TOPRA Competency Framework here topra.org/competencies

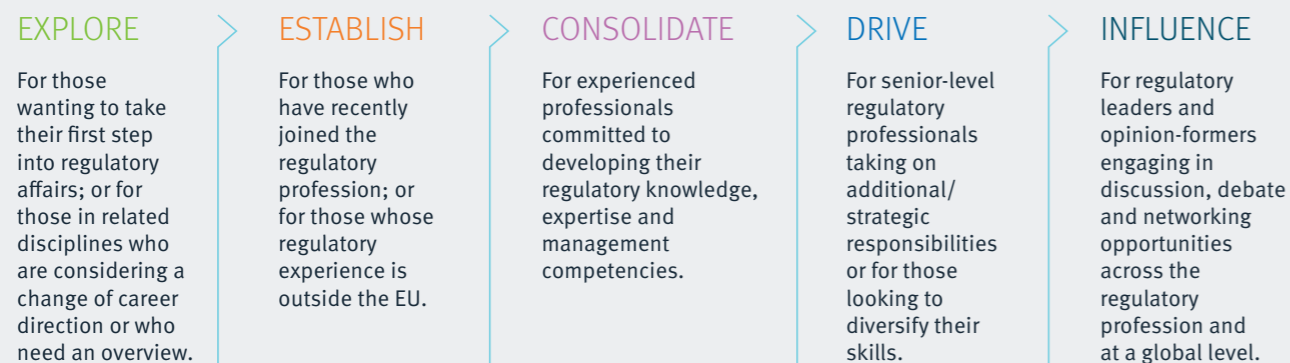


Benefits of TOPRA's training opportunities

- Top quality training designed for the needs of regulatory affairs professionals
- Expert speakers from industry-leading companies, consultancies and regulatory authorities
- Interactive opportunities to apply your new knowledge and develop key skills
- Excellent networking opportunities to help you grow your peer group
- CPD hours automatically assigned for members, which will help to maintain members' professional registration (CSci & RSci) if applicable.

We are here to support you and your team through every career stage

Although every individual career pathway is different, five key 'stages' apply to any regulatory career:



Get ready for TOPRA's must-attend event of the year



 ROTTERDAM | 

Symposium 2024

topra.org/symposium

"An amazing few days of talks, networking and exhibiting. I left feeling smarter and extremely excited about what's to come next."
 — Simon Brough, Head of Business Development, PharmiWeb

Committed to quality – TOPRA is proud to have provided gold standard training for regulatory affairs professionals for more than 40 years. Find out more at topra.org/pd

TOPRA's 2024 Courses and Conferences Calendar

KEY Basics Foundation CRED Masterclass Symposium Awards

Month	Course/Event	Course/Event	Course/Event	Course/Event	Course/Event	Course/Event	Course/Event
JANUARY/ FEBRUARY/ MARCH	Successful and Skilful Communication 23 January London, UK and online	Data Management and Digitalisation in Regulatory Affairs 31 January–2 February London, UK and online	Compiling Successful Clinical Trial Applications 7–8 February Prague, Czechia and online	Essentials of European Veterinary Regulatory Affairs 21 February London, UK and online	CRED Regulation of Drug-device Combination Products 6 March London, UK and online	Regulatory Document Writing and Management 19–20 March London, UK and online	
APRIL/MAY	Essentials of European Pharmaceutical Regulatory Affairs 10 April London, UK and online	Overview of EU Regulatory Affairs (Spring Introductory Course) 16–19 April London, UK and online	Clinical Evaluation of Medical Devices 24–26 April London, UK and online	Project Management for Regulatory Affairs Professionals 7–8 May London, UK and online	Regulatory Requirements for a New Active Substance: Quality 15–17 May London, UK and online	Managing Lifecycle and Variations Effectively 29–30 May Rotterdam, the Netherlands and online	
JUNE	Essentials of European Pharmaceutical Regulatory Affairs 3 June Brussels, Belgium and online	Strategic Planning in Regulatory Affairs 5–7 June London, UK and online	Understanding Digital Health and Electronic Products 11 June Online	Essentials of European Medical Device Regulatory Affairs 12 June London, UK and online	Principles of Medical Device Regulatory Affairs 24–26 June London, UK and online		
JULY/AUGUST	Essentials of In-Vitro Diagnostics Regulatory Affairs 3 July London, UK and online	Successfully Navigating European Regulatory Procedures 9–10 July London, UK and online	Regulatory Control of Clinical Operations 17–19 July London, UK	Essentials of European Pharmaceutical Regulatory Affairs 7 August London, UK and online			
SEPTEMBER	Getting the CMC Dossier Right 4–5 September London, UK and online	Registration of Biological, Biotechnology and Advanced Therapy Products 9–11 September London, UK and online	Regulatory Strategy: From Development to the Market Place 18–20 September London, UK and online	Successful and Skilful Communication 24 September London, UK and online	An Overview of Regulatory Product Information 25 September London, UK and online		
OCTOBER	Essentials of European Pharmaceutical Regulatory Affairs 9 October London, UK and online	Understanding Clinical Development 15–16 October London, UK and online	Post-Market Surveillance and Vigilance for Medical Devices 23–25 October London, UK and online	TOPRA Symposium 30 October–1 November Rotterdam, the Netherlands			
NOVEMBER/ DECEMBER	European IVD Regulatory Affairs 5–7 November London, UK and online	TOPRA Awards for Regulatory Excellence TBC	Overview of EU Regulatory Affairs (Autumn Introductory Course) 19–22 November TBC	Regulatory Strategy for a New Active Substance: Global Clinical Development TBC Krakow, Poland and online	Optimising Regulatory Strategies for Orphan Drugs 26 November Online	Essentials of European Medical Device Regulatory Affairs 27 November Dublin, Ireland and online	Regulatory Strategy in the Post-Market Phase 4–6 December London, UK and online



Online courses available year-round

TOPRA offers more than its face-to-face courses and conferences. We have a wide range of webinars and eLearning courses to always keep you updated. Have a look at our online training opportunities:

www.topra.org/elearning

On-demand eLearning

- Essentials of European Medical Device Regulatory Affairs
- Essentials of European Pharmaceutical Regulatory Affairs
- Governance for Non-Executive Directors
- ...and more

On-demand Webinars

- Introduction to Data Visualisation
- EU Labelling Requirements
- Medical Devices in Asia
- The Regulation of Biotechnology Products
- ...and more



Erik Smit
Client
Development
Manager

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