Essentials of Pharmaceutical Regulatory Affairs

27 March 2026, London, UK
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Time	Presentation
09:30	Introduction to TOPRA
09:40	Introductions – presenters and delegates
09:55	Setting the Scene
10:20	Break (15 minutes)
10:35	Overview of Drug Development
11:05	Regulatory Control of Clinical Trials
11:25	Break (10 minutes)
11:35	Marketing Authorisation Applications
11:55	Product Information (Labelling)
12:10	Homework (SmPC)
12:40	Break (1 hour)
13:40	European Marketing Authorisation Procedures
14:30	Marketing Authorisation Strategy
14:40	Break (20 minutes)
15:00	Post- Authorisation Activities
15:35	Review of learning objectives
16:30	Q&A on the day's topics
16:45	Wrap up and feedback