2026 EDQM VIRTUAL TRAINING PROGRAMME

Independent modules on the Ph. Eur., reference standards and the CEP Procedure
(CHEMICALLY DEFINED ACTIVE SUBSTANCES)

Week 1 - Programme

Monday, 30 November - Module 1: General methods, general chapters and general monographs (live webinar)

Duration: 1 hour 30 mins

10:00-11:00 ***General methods, general chapters and general monographs***
11:00-11:30 ***Live Q&A session***

Tuesday, 1 December - Module 2: Individual monographs - Focus on chemically defined active substances and medicinal products containing them (live webinar)

Duration: 1 hour 30 mins

10:00-10:30 ***Individual monographs for chemically defined active substances***
10:30-11:00 ***Individual monographs for medicinal products containing chemically defined active substances***
11:00-11:30 ***Live Q&A session***

Thursday, 3 December - Module 3: Impurity control in the Ph. Eur. (live webinar)

Duration: 1 hour 30 mins

10:00-11:00 ***Impurity control in the Ph. Eur.: theory and practical examples***
11:00-11:30 ***Live Q&A session***

Friday, 4 December - Module 4: Ph. Eur. reference standards (live webinar)

Duration: 2 hours 10 mins

10:00-10:40 ***Ph. Eur. reference standards: establishment and use***
10:40-10:50 ***What GMP inspectors expect concerning reference standards***
10:50-11:40 ***Handling, dispatch, where to find useful information and other practicalities***
11:40-12:10 ***Live Q&A session***

***

Week 2 - Programme

Monday, 7 December - Module 5: Fundamentals of the CEP procedure (live webinar)

Duration: 1 hour 45 mins

10:00-10:30 ***General overview of the CEP procedure including the fast track and reliance processes***

10:30-11:15 ***How to read and use a CEP***

11:15-11:45 ***Live Q&A session***

Tuesday, 8 December - Module 6: Building successful CEP dossiers: new applications (live webinar)

Duration: 1 hour 30 mins

10:00-11:00 ***How to build a robust new CEP application***

11:00-11:30 ***Live Q&A session***

Tuesday, 8 December - Module 7: Building successful CEP revision applications (live webinar)

Duration: 1 hour 30 mins

14:30-15:30 ***How to build a robust CEP revision application***

15:30-16:00 ***Live Q&A session***

Thursday, 10 December - Module 8: Control of impurities: CEP approach (live webinar)

Duration: 1 hour 30 mins

10:00-11:00 ***Control of impurities: CEP approach***

11:00-11:30 ***Live Q&A session***

Friday, 11 December - Module 9: The EDQM inspection programme (live webinar)

Duration: 2 hours

10:00-10:45 ***Introduction to the EDQM inspection programme***

10:45-11:30 ***How to prepare for an inspection – Most common GMP deficiencies***

11:30-12:00 ***Live Q&A session***