The Eu2P ready-to-use short courses

"What is the format and the duration of short courses?"

Short courses are divided in several learning activities such as recorded lecture, reading, quiz, and forum that altogether **run over 30 minutes to 8 h**.
The learning activities **can be followed according to your wish**.
You can start a learning activity then stop and start another one, then come back later to the one you have firstly opened.

Once open, your training remains accessible until completion, and within one year.

"How are the short courses attendance and progress reported?"

The Eu2P central office can quantify and analyse your own training attendance, workload and quiz progress per learning activity and per short course.

"How am I evaluated at the end of my training?"

When **your training progress reaches 100%**, you **perform an online graded evaluation** on your whole training content.

This evaluation consists in a **quiz covering all your learning activities**.

The training **evaluation is successful if you reach at least a 80% good score**.

You **then receive an official Eu2P certificate** for your achieved competencies.

Enjoy affordable prices

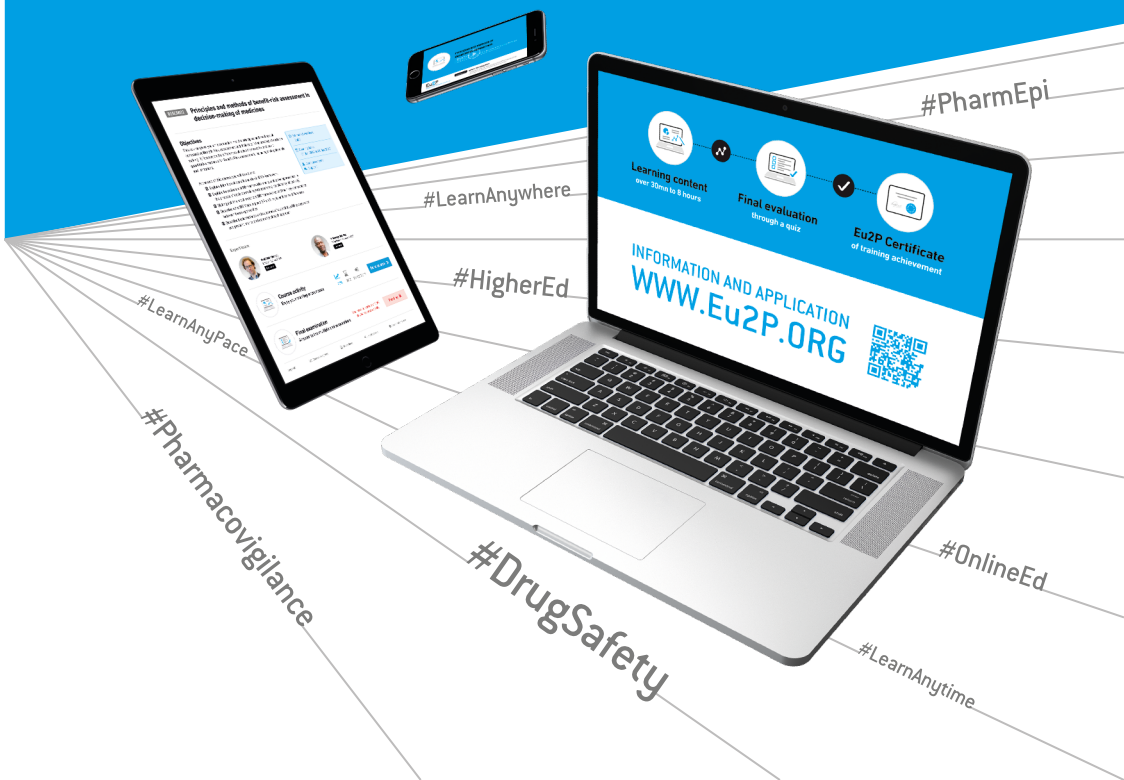
Students & Professionals	Companies
Annual registration for one short course per person from 100 € to 500 €	**Custom package** Quote on request

Eu2P supports worldwide pharmaceutical companies for their in-house training plans and provides quotes tailored to their needs in terms of content, duration and number of collaborators.

For more information, contact **Dr Karine Palin**: eu2p.office@eu2p.org

Eu2P
EUROPEAN PROGRAMME IN PHARMACOVIGILANCE AND PHARMACOEPIDEMIOLOGY

GAIN FAST AND SOLID EXPERTISE IN DRUG SAFETY WITH ONLINE SHORT COURSES



Eu2P short courses catalogue

Medicines benefit assessment

1.	Heterogeneity of treatment effect in clinical trials	2h	200€
2.	Selecting outcomes: Surrogate endpoints	2h	200€

Risk identification and quantification

3.	Concepts in risk management	3h	300€
4.	Pharmacovigilance plan versus risk minimisation plans	3h	300€
5.	Organisation for risk management in the industry	3h	300€
6.	Risk management plans and European Union regulatory perspective	2h	200€
7.	Methods and new initiatives for signals detection	6h	500€

Benefit-risk assessment

1.	Role of benefit-risk assessment and pharmacoeconomics in decision-making of medicines	2h	200€
2.	Principles and methods of benefit-risk assessment in decision-making of medicines	2h	200€
3.	Drug life cycle as a tool in drug policy analysis	1h	Free
4.	Main challenges of health economics and outcomes research	2h	200€

Medicines and public health

1.	Integrating Pharmacovigilance and consumption data analysis - uses, limitations and potentiality	2 h	200€

Risk communication

1.	How to measure the effectiveness of risk communication?	2 h	200€

Pharmacovigilance and regulatory aspects

1.	Pre-marketing pharmacovigilance legislation	2 h	200€
2.	Post-marketing legislation	2 h	200€
3.	Eudravigilance and EVPM	1 h	Free
4.	Development Safety Update Report (DSUR)	0,5 h	Free
5.	International Harmonisation Initiatives	0,5 h	Free
6.	Vaccines biologics regulations	2 h	200€
7.	Materiovigilance regulations	1 h	Free
8.	US Regulations	2 h	200€
9.	Japanese Regulations	2 h	200€
10.	French Regulations	2 h	200€
11.	Contractual agreements in PV	1,5 h	150€
12.	GVP Module I - PV system and their quality system	2 h	200€
13.	GVP Module II - Pharmacovigilance System Master File (PSMF)	2 h	200€
14.	GVP Module III - PV Inspections	1 h	Free
15.	GVP Module IV - PV audits	1 h	Free
16.	GVP Module V - Risk Management System	2 h	200€
17.	GVP Module VI - Management and Reporting of Adverse Reactions to Medicinal Products	4 h	400€
18.	GVP Module VII - Periodic Safety Update Report (PSUR)	3 h	300€
19.	GVP Module VIII - Post-Autorisation Safety Studies (PASS)	2 h	200€
20.	GVP Module IX - Signal Management	1 h	Free
21.	GVP Module XVI - Risk Minimisation Measures	2 h	200€
22.	Signal detection	1,5 h	150€
23.	Validation of computerised systems	1 h	Free
24.	Management and reporting of ADR	4 h	400€