

Am2P

AMERICAN PROGRAM IN
PHARMACOVIGILANCE

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



How does it work?

Course format and duration

- ⊗ Short courses are divided in several learning activities such as recorded lecture, reading, quiz, and forum that altogether **run over 2 to 3 hours**. 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.**

Attendance and progress report

- ⊗ **The Central Office can quantify and analyse** your own training attendance, workload and quiz progress per learning activity and per short course.

Evaluation and award

- ⊗ 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 Am2P certificate** for your achieved competencies.

Program accreditation

Short course Certificate of achievement is based on training course provided by experts from the NASoP chapter.

This is a qualification called "Certificate of Achievement in North America Pharmacovigilance" jointly awarded by the Academic Partners having signed the Eu2P Short Courses agreement and recognized by all Eu2P Course Providers and Associated Partners.



It is printed and issued by the Université de Bordeaux, includes the logos of Am2P Academic Partners and is signed by the chair of the Eu2P Examination Board.

The Am2P program is accredited by six Eu2P academic partners:

- ⊗ Université de Bordeaux (France)
- ⊗ Universitat Autònoma de Barcelona (Spain)
- ⊗ University of Hertfordshire (United-Kingdom)
- ⊗ Erasmus Universitair Medisch Centrum (The Netherlands)
- ⊗ Universiteit Utrecht (The Netherlands)
- ⊗ Università degli Studi della Campania Luigi Vanvitelli (Italia)

How does it cost?

Affordable prices

 Students & Professionals	 Companies
Annual registration for one short course per person from \$230 to \$350	Custom package Quote on request

Am2P 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: am2p.office@am2p-courses.com

Powered by



Am2P SHORT COURSES CATALOGUE



North America Chapter
NASoP

Basic Pharmacovigilance and Pharmacovigilance Regulations



Pharmacovigilance Regulations

- | | | |
|---|----|-------|
| 1. US/EU regulations: principles and comparison | 3h | \$350 |
| 2. Overview of the Legal Basis of PV Regulations and the Role of the QPPV | 3h | \$350 |

Labeling and Combination Products

- | | | |
|---|----|-------|
| 3. Principles of Labeling and Description of United States Prescribing Information (USPI) | 3h | \$350 |
| 4. Pharmacovigilance in Combination Products and Regulations | 3h | \$350 |

Adverse Drug Reporting

- | | | |
|--|------|-------|
| 5. From individual cases to the community impact of adverse drug reactions | 2h | \$230 |
| 6. Aggregate reporting in the US | 2.5h | \$290 |

Pharmacovigilance for Biologics



Vaccine Pharmacovigilance

- | | | |
|-----------------------------------|----|-------|
| 7. Vaccines Biologics Regulations | 3h | \$350 |
|-----------------------------------|----|-------|

Gene Therapy

- | | | |
|-----------------|----|-------|
| 8. Gene therapy | 3h | \$350 |
|-----------------|----|-------|

Pharmacovigilance for Human Cells, Tissues and Cellular and Tissue-Based Products

- | | | |
|---------------------------------|----|-------|
| 9. Pharmacovigilance for HCT/Ps | 3h | \$350 |
|---------------------------------|----|-------|

Targeted therapy

- | | | |
|---------------------------|----|-------|
| 10. Targeted Therapeutics | 3h | \$350 |
|---------------------------|----|-------|

External databases/RWD/RWE



FDA System

- | | | |
|--|------|-------|
| 11. FDA Adverse Event Reporting System (FAERS) | 2,5h | \$290 |
| 12. FDA Sentinel System | 2,5h | \$290 |

Databases

- | | | |
|--|----|-------|
| 13. Health care records from large databases as a tool to study the use of medicines | 2h | \$230 |
| 14. Integrating Pharmacovigilance and consumption data analysis - uses, limitations and potentiality | 2h | \$230 |

Benefit Risk Assessment



Benefit-risk assessment of medicines

- | | | |
|--|------|-------|
| 15. Principles and methods of benefit-risk assessment in decision-making of medicines | 2h | \$230 |
| 16. Role of benefit-risk assessment and pharmaco-economics in decision-making of medicines | 2,5h | \$290 |

Risk Management

- | | | |
|--|------|-------|
| 17. Concepts in Risk Management | 3h | \$350 |
| 18. Organization for risk management in the industry | 3h | \$350 |
| 19. Risk Evaluation and Mitigation Strategy (REMS) | 2,2h | \$260 |

APPLY ON-LINE
START YOUR COURSE NOW!

www.am2p-courses.com

