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

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

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Am2P

AMERICAN PROGRAM IN PHARMACOVIGILANCE

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

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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

1. Vaccines Biologics Regulations	3h	$350

Gene Therapy

2. Gene therapy	3h	$350

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

3. Pharmacovigilance for HCT/Ps	3h	$350

Targeted therapy

4. Targeted Therapeutics	3h	$350

External databases/RWD/RWE

FDA System

1. FDA Adverse Event Reporting System (FAERS)	2,5h	$290
2. FDA Sentinel System	2,5h	$290

Databases

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

Benefit Risk Assessment

Benefit-risk assessment of medicines

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

Risk Management

3. Concepts in Risk Management	3h	$350
4. Organization for risk management in the industry	3h	$350
5. Risk Evaluation and Mitigation Strategy (REMS)	2,2h	$260

Medication errors

1. Scope and Background on Medication Errors	2h	$230
2. Medication Error Pharmacovigilance	2h	$230
3. Minimizing Medication Errors Related To Nomenclature	2h	$230
4. Minimizing Medication Errors Related To Product Design	2h	$230
5. Optimizing Labeling and Packaging to Minimize Medication Errors	2h	$230
6. Human Factors and its Role in Preventing Medication Errors	2h	$230

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