Publication date : 01/18/2024

Course : Reporting adverse events associated with healthcare (AEAS)

Practical course - 2d - 14h - Ref. EVI
Price : 940 € E.T.

Reporting is very important for the detection and management of effects, adverse events and alerts associated with healthcare. We need to improve the relevance, completeness and responsiveness of reports. This training course will help you improve your facility's vigilance system.

Teaching objectives

At the end of the training, the participant will be able to:

- Organizing vigilance and risk management coordination within your facility
- Improving the completeness and responsiveness of declarations and reports
- Provide guidelines for in-depth cause analysis methods
- Reinforcing the quality of declaration content

Intended audience

Tout professionnel de santé, coordonnateur de la gestion des risques associés aux soins, responsable qualité.

Prerequisites

No special knowledge required.

Course schedule

PARTICIPANTS

Tout professionnel de santé, coordonnateur de la gestion des risques associés aux soins, responsable qualité.

PREREQUISITES

No special knowledge required.

TRAINER QUALIFICATIONS

The experts leading the training are specialists in the covered subjects. They have been approved by our instructional teams for both their professional knowledge and their teaching ability, for each course they teach. They have at least five to ten years of experience in their field and hold (or have held) decision-making positions in companies.

ASSESSMENT TERMS

The trainer evaluates each participant’s academic progress throughout the training using multiple choice, scenarios, hands-on work and more.
Participants also complete a placement test before and after the course to measure the skills they’ve developed.

1 Regulations and implementation of the vigilance organization

- Understand the context and challenges of risk management and vigilance.
- Understand the regulatory aspects and issues related to health vigilance.
- Understand the regulatory aspects, systems and procedures relating to healthcare-associated infections (HAIs).
- Understand the regulatory aspects, issues and coordination of risks associated with healthcare.
- Get to grips with reporting circuits, media and tools.
- Set up coordination to coordinate the organization of vigilance and risk management.

Hands-on work

Workshop: case studies and reporting exercises.

2 Care safety from acculturation to application

- Implement a global policy for managing Serious Adverse Events (SAEs).
- Apply root cause analysis methods: practical case studies.
- Develop a preventive risk culture: risk mapping, methods and tools.
- Integrate risk prevention into the overall approach.
- Generalizing the culture of care safety: from concept to practice.

Hands-on work

Reflection and discussion on the situation at the facility. Practical application of methods and tools to concrete cases. Debriefing.

TEACHING AIDS AND TECHNICAL RESOURCES

• The main teaching aids and instructional methods used in the training are audiovisual aids, documentation and course material, hands-on application exercises and corrected exercises for practical training courses, case studies and coverage of real cases for training seminars.
• At the end of each course or seminar, ORSYS provides participants with a course evaluation questionnaire that is analysed by our instructional teams.
• A check-in sheet for each half-day of attendance is provided at the end of the training, along with a course completion certificate if the trainee attended the entire session.

TERMS AND DEADLINES

Registration must be completed 24 hours before the start of the training.

ACCESSIBILITY FOR PEOPLE WITH DISABILITIES

Do you need special accessibility accommodations? Contact Mrs. Fosse, Disability Manager, at psh-accueil@orsys.fr to review your request and its feasibility.