The 16th International EuBIS Seminar and Training Course
Rome, Italy, 5th of November – 7th of November 2024

Quality management and inspection of blood establishments are a key stone in achieving best practice and guaranteeing the safety of blood and blood components for transfusion of patients. The European Blood Inspection System (EuBIS), initiated under the Public Health Programme of the European Commission, Directorate General SANTE, has developed a manual and training guide containing standards and guidelines for quality management and the inspection of blood establishments.

Based on the EuBIS manual and guide the EuBIS Academy has organized training seminars and courses in order to promote knowledge in the area of quality and safety of blood and blood components throughout Europe and worldwide. Following the 1st International seminar during the ISBT congress in Berlin, Germany, the seminar and training has received extensive international reputation with participants from EU member states (Austria, Belgium, Bulgaria, Croatia, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Luxembourg, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, The Netherlands, Poland, Portugal, Romania, Spain, Sweden, United Kingdom) and world-wide (Columbia, Norway, Switzerland, Kosovo, Kuwait, Turkey, Brazil, Curacao, Algeria, Lebanon, Morocco, Macedonia, Nigeria, Singapore, Saudi Arabia, Yemen, Qatar, South Africa, South Korea, Vietnam, United State of America, etc.).

*The **16th International EuBIS Seminar and Training Course** taking place from the **05th of November – 07th of November 2024 in Rome (Italy),** has been designed with the requirement of "Risk-based-Approach" to be implemented in quality systems. With the revised exercises and methodology, the course will be a further milestone in the objectives of EuBIS to give assistance in promoting its mission 'from good to best practice in transfusion medicine'.*

*This **new version of EUBIS training** aims to further expand on previous EuBIS courses and provides insights in more recent challenges and developments.*

*The training material has been revised to take into account in more detail and complexity the **Risk-based Approach as required by the Good Practice (GPG) Guideline for blood and blood components** in the European Union.*

*In addition, the training will address the new **"Regulation on standards of quality and safety for substances of human origin intended for human application (SoHO)" COM(2022)0338 – C9-0226/2022 – 2022/0216(COD),** that has been approved on the 24th of April by the European Parliament and adopted on the 27th of May 2024 by the Council, reinforcing the Risk based Approach.*

*The seminars will comprise lectures and group work in a face-to-face fashion based on cases covering several aspects of **GPG** and **GMP** such as:*

- *Risk analysis and risk management,*
- *Inspection/Audit practice,*
- *Process design and risk assessment,*
- *Validation and Qualification,*
- *Change Control and change management,*

***For registration, please use the following link**:* https://www.eubis-europe.eu/eubis_italy_nov_2024.php