EuBIS
European Blood Inspection System
Initiated under the Public Health Programme of the
European Commission, DG Sanco, EC – GA No. 2006202

supported by the
CENTRO NAZIONALE SANGUE

8th EuBIS Seminar and Training

'Good practice in blood components and medicinal products referring to GP and GMP'

Quality management and inspection criteria for blood establishments and pharmaceutical products

5th -7th of October 2016, Rome, Italy

organised by the EuBIS Academy
in cooperation with Centro Nazionale Sangue (CNS)

supported by the
EBA European Blood Alliance

EuBIS

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8th International Seminar and Training Course - Rome, 5th - 7th of October 2016

Seminar and training - From Good Practice (GP) to Good manufacturing Practice (GMP) – blood components and medicinal products

The seminar will build on previous EuBIS training courses with the scope to train on the impact of the new regulation of Good Practice for blood and blood components in the European Union.

The seminar comprises lectures and group work in a face-to-face fashion based on cases covering several aspects of GP and GMP such as:

- GP guidelines
- Inspection/Audit
- Validation
- Change Control
- Corrective actions
- Risk assessment

Previous participation in a EuBIS/Catie course is beneficial but not a prerequisite.

Training course language: **English**

Course fees European/International participants: € 470,00
Course fee for Italian participants: *Sponsorship is given by CNS

Registration: www.eubis-europe.eu

Participants from Italy are offered an examination (in Italian language) with a certificate of successful participation by CNS.

Training course participants:

- Qualified as inspectors by a Competent Authority
- Individuals working in a blood establishment in the area of quality management.

Training places are limited: Selection of training participants and confirmation of participation will be on the basis on time of registration.

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