



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

Introduction and objectives of the workshop

Medication-errors workshop
London, 28 February – 1 March 2013

Presented by: Dr. Peter Arlett
Head, Pharmacovigilance and Risk Management
European Medicines Agency

An agency of the European Union





Sessions

Session 1 Medication errors in the product life-cycle and special populations.

Session 2 Reporting.

Session 3 Analysis of medication errors resulting in harm.

Session 4 Regulatory tools for managing the risk of medication errors.

Session 5 Implementation of preventative measures.

Session 6 The way forward.



Programme Committee

Peter Arlett

Francesca Cerreta

Emer Cooke

Brian Edwards

Henry Fitt

Mick Foy

Thomas Goedecke

Dolores Montero

Isabelle Moulon

Paolo Tomasi



Participation

142 + 27 speakers

Academic	12
Regulatory	34
Public body	8
HCP	16
Patient Con	6
Learnt Society	1
Industry	34
Service provider for industry	1
PSQCWG	10
Total	122



Objectives of the workshop

- 1. Raising awareness** amongst the stakeholders involved in the reporting, evaluation and prevention of medication errors.
- 2. Common understanding** of
 - what constitutes a medication error and,
 - the new legal requirements for reporting cases of medication error at EU level.
- 3. Better understanding of how medication errors are managed** at national level.
- 4. Sharing best practice for the prevention** of medication errors.
- 5. Proposals to facilitate stakeholder co-operation** at national and international level.



Probable outputs

- **Workshop report**
- **Key recommendations**
- **Likely ‘best-practice’ document**
- **Likely operational improvements**



“Let the public health debate commence”