

# Clinical Trials in the European Union: Selected Legislation, Guidelines, and the Declaration of Helsinki

## Legislation

- Directive 2001/20/EC ... relating to the implementation of good clinical practice in the conduct of clinical trials (Repealed 27 May 2014)
- Commission Directive 2005/28/EC laying down principles and detailed guidelines for good clinical practice
- Regulation (EU) No 536/2014 on clinical trials on medicinal products for human use, and repealing Directive 2001/20/EC

## Implementing Guidances for the EU Directives

- Detailed guidance on the collection, verification and presentation of adverse event/reaction reports arising from clinical trials on medicinal products for human use (CT-3) June 2011
- Detailed guidance on the application format and documentation to be submitted in an application for an ethics committee opinion on a clinical trial on medicinal products for human use February 2006, Revision 1
- Questions and Answers on Clinical Trials in the EU August, 2011

## ICH Guidelines (Step 5, EU)

- E6 (R2): Good Clinical Practice (Integrated Addendum)
- E2A: Clinical Safety Data Management: Definitions and Standards for Expedited Reporting
- E8: General Considerations for Clinical Trials
- E2F: Development Safety Update Reports

## Declaration of Helsinki

World Medical Association, 2013