European Union

a. Pharmaceutical Legislation Medicinal Products for Human Use (Directives, Regulations, Non-Legislative Acts, Miscellaneous
b. Legal framework governing medicinal products for human use in the EU
d. EU Directive 2011_62 (Amending Directive 2001/83/EC on the Community code relating to medicinal products for human use, as regards the prevention of the entry into the legal supply chain of falsified medicinal products)