

Commission Directive 2005/28/EC of 8 April 2005 laying down principles and detailed guidelines for good clinical practice as regards investigational medicinal products for human use, as well as the requirements for authorisation of the manufacturing or importation of such products (Text with EEA relevance)

CHAPTER 7

FINAL PROVISIONS

Article 32

This Directive shall enter into force on the twentieth day following that of its publication in the *Official Journal of the European Union*.