

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)

Done at Brussels, 8 April 2005.

*For the Commission*

Günter VERHEUGEN

*Vice-President*