EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

5 March 2012
EMA/721630/2011
Patient Health Protection

Detailed guidance on the electronic submission of information on medicinal products for human use by marketing authorisation holders to the European Medicines Agency in accordance with Article 57(2), second subparagraph of Regulation (EC) No. 726/2004

Chapter 4: Structured Substance Information (currently not applicable)

NOTE:

The Structured Substance Information (SSI) is subject to discussion for future use. A revised chapter will be published post-July 2012.

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