STATUTORY INSTRUMENTS

2012 No. 1916

The Human Medicines Regulations 2012

PART 5

Marketing authorisations

General provisions relating to offences

[F1Offences in connection with parallel import licence application

95A. A person is guilty of an offence if, in the course of an application for the grant, renewal or variation of a parallel import licence for a relevant medicinal product, the person—

(a) fails to provide the licensing authority with any information that is relevant to the evaluation of the safety, quality or efficacy of the product; or

(b) provides to the licensing authority any information that is relevant to the evaluation of the safety, quality or efficacy of the product but that is false or misleading in a material particular.**]**

Textual Amendments

F1 Reg. 95A inserted (E.W.S.) (1.10.2014) by The Human Medicines (Amendment) (No. 2) Regulations 2014 (S.I. 2014/1878), regs. 1, **16** and reg. 95A inserted (N.I.) (1.10.2014) by The Human Medicines (Amendment) (No. 2) Regulations 2014 (S.R. 2014/324), regs. 1(1), **16**

Changes to legislation:
There are currently no known outstanding effects for the The Human Medicines Regulations 2012, Section 95A.