



Human Fertilisation and Embryology Act 1990

1990 CHAPTER 37

Licence conditions

[^{F1}15A Duties of the Authority in relation to serious adverse events and serious adverse reactions

- (1) The Authority shall investigate serious adverse events and serious adverse reactions and take appropriate control measures.
- (2) In investigating any serious adverse event or serious adverse reaction, the Authority shall, where it is appropriate to do so, arrange for—
 - (a) any premises to which a licence relates and any relevant third party premises to be inspected on its behalf, and
 - (b) a report on the inspection to be made to it.
- (3) [^{F2}If the Authority, in relation to Northern Ireland, receives a request from a competent authority in an EEA state] to carry out an inspection in relation to a serious adverse event or serious adverse reaction, the Authority must arrange for such an inspection to be carried out, for a report to be made of the inspection and for appropriate control measures to be taken.]

Textual Amendments

- F1** S. 15A inserted (25.5.2007 for certain purposes, otherwise 5.7.2007) by [The Human Fertilisation and Embryology \(Quality and Safety\) Regulations 2007](#) (S.I. 2007/1522), regs. 1, **18**
- F2** Words in s. 15A(3) substituted (31.12.2020) by [S.I. 2019/482](#), **reg. 2(9)** (as substituted by [The Human Fertilisation and Embryology \(Amendment\) \(EU Exit\) Regulations 2020](#) (S.I. 2020/1307), regs. 1, **10**); 2020 c. 1, **Sch. 5 para. 1(1)**

Changes to legislation:

There are currently no known outstanding effects for the Human Fertilisation and Embryology Act 1990, Section 15A.