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STATUTORY INSTRUMENTS

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**2007 No. 1522**

The Human Fertilisation and Embryology  
(Quality and Safety) Regulations 2007

**PART 2**

AMENDMENTS TO THE 1990 ACT

**Other registers to be kept by the Authority**

**24.** After section 31, insert—

**“31A. The Authority's register of licences**

- (1) The Authority shall keep a register recording the grant, suspension or revocation of—
- (a) every licence under paragraph 1 or 2 of Schedule 2 authorising activities in relation to gametes or embryos intended for use for human application, and
  - (b) every licence under paragraph 1A of Schedule 2.
- (2) The register shall specify, in relation to each such licence—
- (a) the activities authorised,
  - (b) the address of the premises to which the licence relates,
  - (c) the name of the person responsible and, if applicable, the nominal licensee, and
  - (d) any variations made.
- (3) The Authority shall make such of the information included in the register as it considers appropriate available to the public in such manner as it considers appropriate.

**31B. The Authority's register of serious adverse events and serious adverse reactions**

- (1) The Authority shall keep a register containing information provided to it under this Act about any serious adverse event or serious adverse reaction.
- (2) The Authority shall make such of the information included in the register as it considers appropriate available to the public in such manner as it considers appropriate.”

**Changes to legislation:**

There are currently no known outstanding effects for the The Human Fertilisation and Embryology (Quality and Safety) Regulations 2007, Section 24.