
STATUTORY INSTRUMENTS

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.”