STATUTORY INSTRUMENTS

2007 No. 1523

The Human Tissue (Quality and Safety for Human Application) Regulations 2007

PART 4

OBLIGATIONS OF THE AUTHORITY

Register of serious adverse events and serious adverse reactions

19.—(1) The Authority shall keep a register containing information provided to it under these Regulations 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.