SCHEDULE

AMENDMENTS TO OTHER ENACTMENTS

10. In the Medical Devices Regulations 2002(**1**), in regulation 2 (interpretation), in paragraph (1), in the definition of "Directive 2001/83", after "human use" insert "as amended by Directive 2002/98/EC of the European Parliament and of the Council setting standards of quality and safety for the collection, testing, processing, storage and distribution of human blood and blood components, Commission Directive 2003/63/EC amending Directive 2001/83/EC on the Community code relating to medicinal products for human use, Directive 2004/24/EC of the European Parliament and of the Council amending, as regards traditional herbal medicinal products, Directive 2001/83/EC on the Community code relating to medicinal products for human use and Directive 2004/27/EC of the European Parliament and of the Council amending Directive 2001/83/EC on the Community code relating to medicinal products for human use".

(**1**) S.I. 2002/618.