STATUTORY INSTRUMENTS

2006 No. 1928

The Medicines for Human Use (Clinical Trials) Amendment Regulations 2006

Amendment of regulation 40 of the principal Regulations

20. In regulation 40 of the principal Regulations (grant or refusal of manufacturing authorisation), in paragraph (1), in sub-paragraph (a), for head (ii) substitute—

"(ii) has at his disposal—

(aa) the services of staff, and

(bb) suitable and sufficient premises, technical equipment and control facilities,

complying with the requirements of Commission Directive 2003/94/EC, as regards the manufacture or import, and control, of the products to which the authorisation relates and the storage of such products,".