STATUTORY INSTRUMENTS

# 2006 No. 1928

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

**Amendment of regulation 42 of the principal Regulations**

**22.** For regulation 42 of the principal Regulations (obligations of manufacturing authorisation holder), substitute—

"**42.** The holder of a manufacturing authorisation shall—

(a) comply with the principles and guidelines of good manufacturing practice;

(b) comply with the provisions referred to in regulation 40(3);

(c) allow the licensing authority access to his premises at any reasonable time; and

(d) put and keep in place arrangements which enable the qualified person to carry out his duties, including placing at his disposal all the necessary facilities.".