STATUTORY INSTRUMENTS

2014 No. 1878

The Human Medicines (Amendment) (No. 2) Regulations 2014

Amendment of regulation 48

3. In regulation 48(2) (definitions in relation to the Part on marketing authorisations), after the definition of "generic medicinal product" insert—

""parallel import licence" means a licence that—

(a) is granted by the licensing authority in compliance with the rules of European Union Law relating to parallel imports; and

(b) authorises the holder to place on the market a medicinal product imported into the United Kingdom from another EEA State;".