
STATUTORY INSTRUMENTS

2002 No. 618

The Medical Devices Regulations 2002

PART VII

General, Enforcement and Miscellaneous

Interpretation of Part VII

59. In this Part, unless the context otherwise requires—

“registrable device” means a device in respect of which, in accordance with the Medical Devices Directives, registration is required with the competent authorities of a Member State or (where appropriate) a State which is a Party to an Association Agreement;

“relevant device” means a device that is a “relevant device” for the purposes of Part II, III or IV.