 

Changes are coming to the way in which MHRA ensures the safety and quality of medical devices. A series of improvements are being made to modernise

the current system that will ensure better protection of public health and patient safety. The new Regulation for Medical Devices (MDR), which entered into

force on 25th May 2017, are a balancing act of proportionate responsibility and an increasingly technological approach to healthcare

 (<https://www.gov.uk/guidance/medical-devices-eu-regulations-for-mdr-and-ivdr>).

PMG members maybe interested in MHRA’s introductory interactive guide to the [new EU Regulations for medical devices (MDR)](https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/640404/MDR_IVDR_guidance_Print_13.pdf) .