EMERGO
by UL

The UK MHRA maintains a large database of guidance documents on their website. Although not legally binding, these guidance documents present the Agency's current thinking on a wide variety of topics related to medical device regulation, the post-Brexit transition, and COVID-19 in the United Kingdom and may help clarify a particular policy or regulatory issue.

To access the MHRA guidance documents, click the following links:

- **Medical Devices Regulation and Safety:** https://www.gov.uk/topic/medicines-medical-devices-blood/medical-devices-regulation-safety
- **MHRA Guidance on Post-Brexit Transition:** https://www.gov.uk/government/collections/new-guidance-and-information-for-industry-from-the-mhra
- **MHRA Guidance on COVID-19:** https://www.gov.uk/government/collections/mhra-guidance-on-coronavirus-covid-19
- **Regulating Medical Devices in the UK:** https://www.gov.uk/guidance/regulating-medical-devices-in-the-uk
- **Approved Bodies for Medical Devices:** https://www.gov.uk/government/publications/approved-bodies-for-medical-devices/approved-bodies-for-medical-devices
- **Medical Devices - Conformity Assessment and the UKCA Mark:** https://www.gov.uk/guidance/medical-devices-conformity-assessment-and-the-ukca-mark
- **Register Medical Devices to Place on the Market:** https://www.gov.uk/guidance/register-medical-devices-to-place-on-the-market
- **Medical Devices - Guidance for Manufacturers on Vigilance:** https://www.gov.uk/government/collections/medical-devices-guidance-for-manufacturers-on-vigilance