Contact us
+44 345 080 9000
medicaldevices@bsigroup.com

Regulatory review

These five insightful webinars focussed on various aspects of the MDR, from looking at postmarket clinical follow-up, to helping you with your medical device software and when a clinical evaluation is required.

Our fifth and final webinar Post

The deadline for UKCA marking is fast approaching, with many medical device manufacturers still not applying for UKCA certification.

This webinar will offer approved body insights for all people involved in working towards a UKCA application, whether you are a novice or have more significant experience of working with an approved body.

Register from two sessions on:

09:00-10:00 Wednesday 27 April

16:00-17:00 Wednesday 27 April -

We also have lots of resources for you to use to ensure you maintain market access in the UK. Please refer to our detailed FAQ document online, which covers many questions you may have around UKCA and our website.