# BSI update on the new UKCA and future UK regulation for Medical Devices and IVDs

Tue, Sep 29, 2020 3:00 PM - 4:00 PM CEST

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*Please note spaces on this webinar are limited. It will be recorded and made available to view on our website following the live broadcast.*

Join BSI's Dr Jayanth Katta, Senior Regulatory Lead, to hear the Notified Body perspective on the announcement regarding the new UK Conformity Assessed (UKCA) mark. The Medicines and Healthcare products Regulatory Agency (MHRA) has published new guidance on GOV.UK, which sets out how medical devices will be regulated after the transition period with the EU has ended (from 1 January 2021).

The webinar will cover:

1. The introduction of the new UK route to market and the mark of conformity for devices.
2. The transition period for products already CE marked, following confirmation of recognition of the CE mark until 30 June 2023.
3. The implications of Great Britain not implementing the EU Medical Devices Regulation (MDR) and In Vitro Diagnostics Regulation (IVDR).
4. Outline of how the requirements will impact manufacturers in the UK and EU.

Dr Katta will be joined by Gary Slack, Senior Vice President, to discuss BSI's strategic approach to the new regulation and how we will implement this alongside our EU Notified Body role.

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