Webinar on

Continuing your exports to UK: Transitioning to the UKCA marking post-Brexit and updates on UK legislation for medical devices

Date : Thursday, 8 April 2021

Time : 3.00pm to 5.00pm (SG/MY/PH) | 2.00pm to 4.00pm (VN/ID/TH)

In view of Brexit, the EU's CE marking will not be accepted in UK after 31 December 2021. The UKCA marking will be the new UK product certification marking that will be used for goods being placed on the market in UK. This webinar will provide general updates and guidance for obtaining the UKCA marking if you wish to continue to export to UK.

This webinar will also touch on Medical Device regulations coming into effect from 1 January 2021, and the new technical specifications to be met by businesses to demonstrate conformity for goods to be exported into the UK market.

Topics covered:

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

Agenda

3.00pm	Welcome address by Emmanuel Herve, ASEAN Managing Director
3.15pm	General update of the UK Conformity Assessed (UKCA) marking by Shahm Barhom, Group Product Certification Director
3.45pm	Post Brexit UK Legislation for medical devices by Gary Slack, Senior Vice President of Medical Devices
4.15pm	Q&A session
5.00pm	End of webinar

Book your place

Speakers

Emmanuel Herve
ASEAN Managing Director, BSI

Shahm Barhom
Group Product Certification Director, BSI

Gary Slack
Senior Vice President of Medical Devices, BSI