



The European Association Medical
Devices - Notified Bodies

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Statement in the framework of the implementing acts	

In the framework of the new regulations of the 5th April 2017, 2017/745 on medical devices and 2017/746 on in vitro diagnostic medical devices, where there are presently no implementing and/or delegated acts, CAMD documents or guidance documents from the Commission available, then Team NB members will continue to consider the latest existing guidance as valid. For example, the present NBOG Best Practice Guides/MEDDEV documents.

These documents will be considered to be the only available guidance to help to address any specific topics that is within their remit.