
Status: Point in time view as at 31/01/2020.

Changes to legislation: *There are currently no known outstanding effects for the Commission Decision of 27 November 2009 amending Decision 2002/364/EC on common technical specifications for in vitro diagnostic medical devices (notified under document C(2009) 9464) (Text with EEA relevance) (2009/886/EC), Article 3. (See end of Document for details)*

Commission Decision of 27 November 2009 amending Decision 2002/364/
EC on common technical specifications for in vitro diagnostic medical devices
(notified under document C(2009) 9464) (Text with EEA relevance) (2009/886/EC)

Article 3

This Decision shall apply from 1 December 2010 for those devices first placed on the market prior to 1 December 2009.

It shall apply from 1 December 2009 for all other devices.

However, Member States shall allow manufacturers to apply the requirements set out in the Annex before the dates set out in the first and second paragraphs.

Status:

Point in time view as at 31/01/2020.

Changes to legislation:

There are currently no known outstanding effects for the Commission Decision of 27 November 2009 amending Decision 2002/364/EC on common technical specifications for in vitro diagnostic medical devices (notified under document C(2009) 9464) (Text with EEA relevance) (2009/886/EC), Article 3.