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IN VITRO DIAGNOSTIC MEDICAL DEVICE REGULATION (IVDR)

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REGULATION (EU) 2017/746 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL

of April 5 2017 on

In Vitro Diagnostic Medical Devices

and repealing

Directive 98/79/EC [in vitro diagnostic devices] and Commission Decision 2010/227/EU [European databank for medical devices]

For recitals (i.e. the "preface" or "reasoning" for the regulation) see the original IVDR (pdf): https://eur-lex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:32017R0746&from=EN

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