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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):
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