



IVDR Conformity Assessment Routes Guide

IVDR Conformity Assessment Routes

Notified Body Assessments

bsi.

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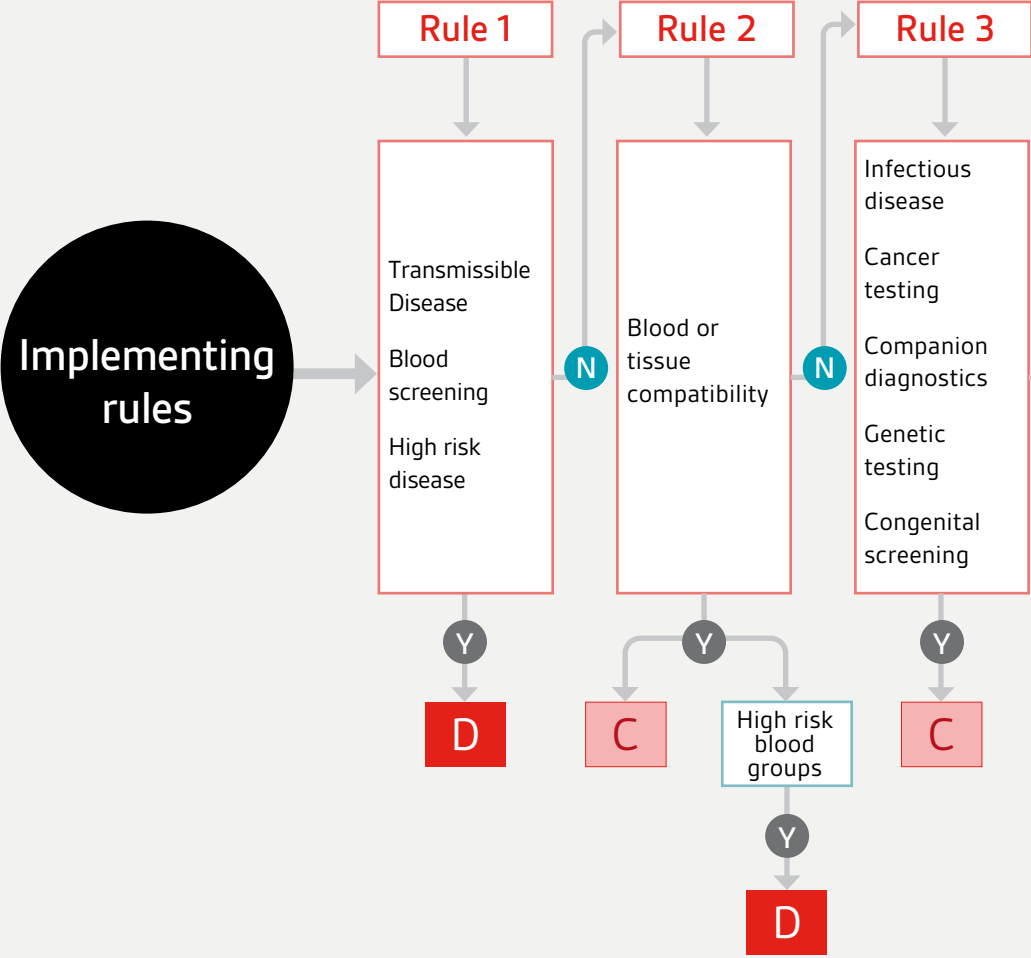
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Disclaimers:

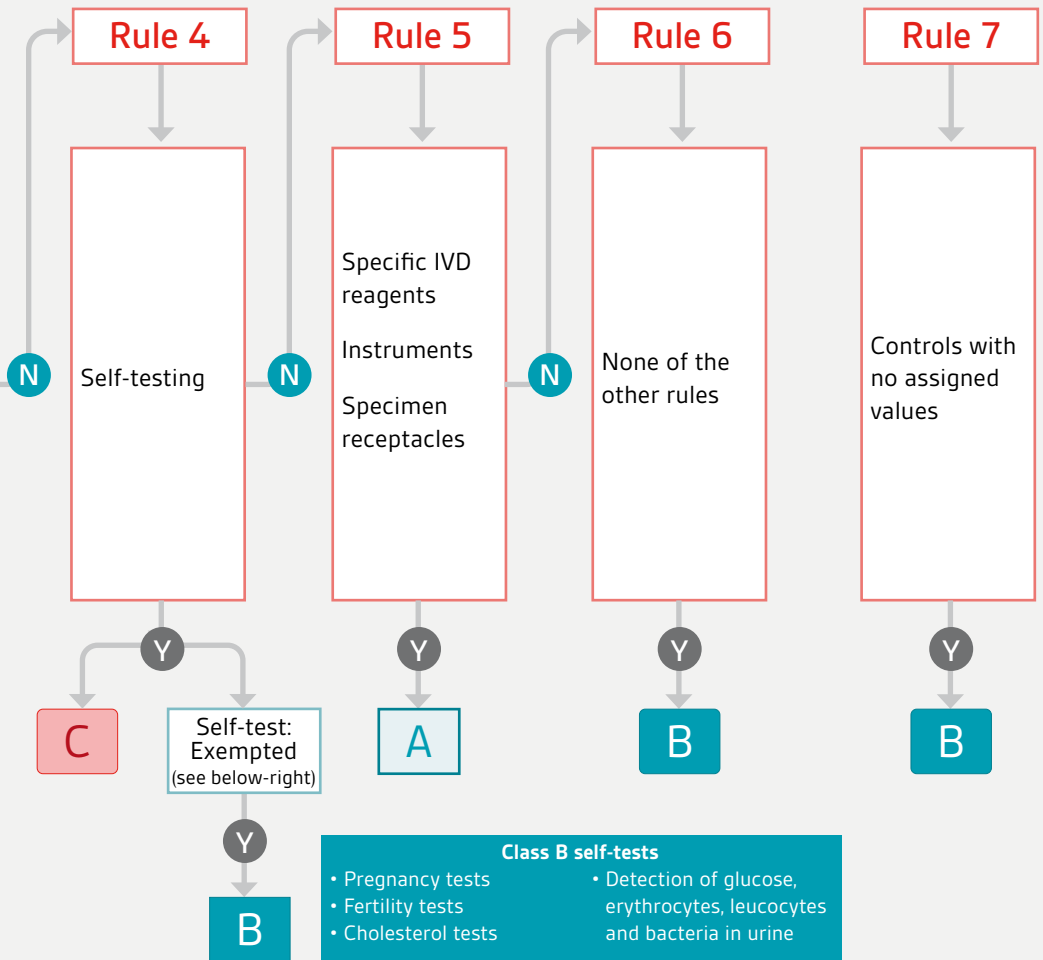
- The information presented in this brochure is based on our current understanding of the IVDR requirements at the time of publishing and is subject to change
- The tables do not cover assessments under the conformity route of Annex XI (Production Quality Assurance). BSI is not designated to Annex X (Type Examination)
- The Competent Authority may ask for verification testing by the EU Reference Laboratory for devices other than Class D

Illustration of the Classification rules as per



Note: When classifying your device, always consult the IVDR and, in particular, Annex VIII.

Annex VIII of the IVDR



All near patient tests are classified in their own right, they can be D, C, or B, depending on intended purpose.

Useful definitions

CE 2797

Throughout this guide, our Notified Body is referenced using its assigned Notified Body number: BSI The Netherlands (2797).

Common Specifications

The European Commission provides Common Specifications to the IVDR as a means of complying with the legal obligations applicable to a device, process or system, such as the General Safety and Performance Requirements (GSPRs), the requirements for performance studies and performance evaluation, and/or post-market surveillance.

CA and EMA

In the case of companion diagnostics, the Competent Authority (CA) or the European Medicines Agency (EMA) will be consulted regarding the associated medicinal product.

EU Reference Laboratory

These have been introduced under the IVDR and are laboratories designated by the European Commission to support with the assessment of Class D IVD devices. An EU Reference Laboratory is responsible for verifying the performance of Class D IVD devices and the ongoing verification of manufactured devices.

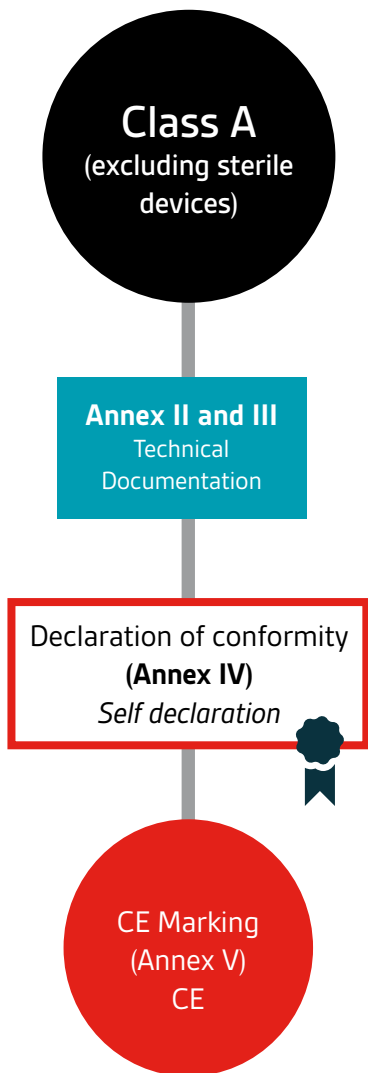
The Competent Authority may ask for verification testing by the EU Reference Laboratory for devices other than Class D.

Notified Body (NB)

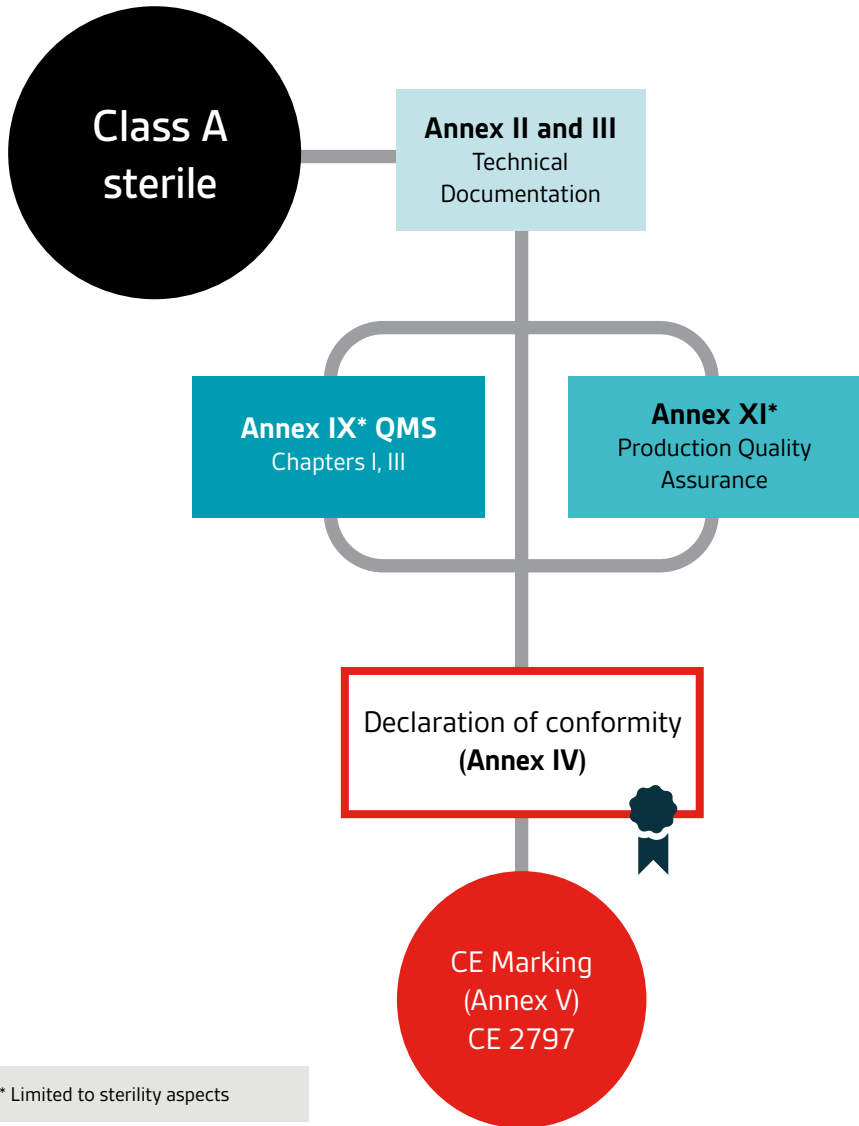
The role of BSI as a Notified Body is to conduct a conformity assessment under the IVDR. This usually requires an audit of the manufacturer's quality management system and, depending on the particular classification of the device, a review of the relevant Technical Documentation in support of the safety and performance claims for the device. The Technical Documentation is assessed against the General Safety and Performance Requirements (GSPR) within the IVDR.

Class A devices

Note: No Notified Body involvement



Class A sterile devices



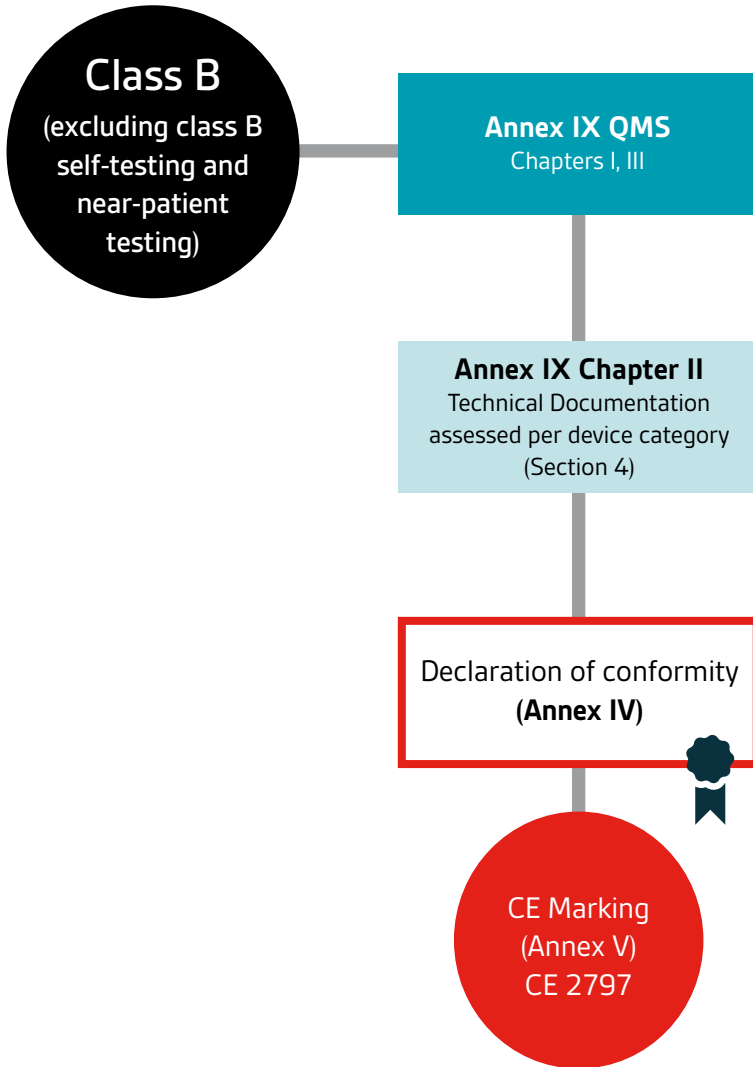
Class A sterile devices

Class A sterile devices	Initial Conformity Assessment	Surveillance				
		Y1	Y2	Y3	Y4	Y5
QMS Audits	Yes	Yes	Yes	Recertification*	Yes	Yes
Microbiology Audits	Yes	N/A	Yes	N/A	Yes	N/A
Technical Documentation Assessment	N/A	N/A	N/A	N/A	N/A	N/A
Competent Authority or EMA consultation (Annex IX, Section 5.2)	N/A	N/A	N/A	N/A	N/A	N/A
Experts consultations (article 48(6))	N/A	N/A	N/A	N/A	N/A	N/A
Verification by EU reference laboratory (Annex IX, section 4.9)	N/A	N/A	N/A	N/A	N/A	N/A
Summary of Safety and Performance (Article 29)	N/A	N/A	N/A	N/A	N/A	N/A
Performance Evaluation Report updates (Annex XIII - Part A, Section 1.3.2 and Article 56)	Not required for NB assessment					
Post Market Performance Follow-up (PMPF) updates Evaluation Report (Article 56 and Annex XIII, Part B)	Not required for NB assessment					
Post Market Surveillance (PMS) Report (Article 80)	Updated when necessary and made available to the NB upon request.					
Periodic Safety Update Report (PSUR) (Article 81)	N/A	N/A	N/A	N/A	N/A	N/A
Unannounced Audits	At least once every 5 years					

* QMS certificates are valid for three years, whilst CE certificates remain valid for a maximum of five years. The **Y3 Recertification** indicated in the table relates to the EN ISO 13485:2016 certificate cycle. Certification cycles vary and re-certification may not always occur at Y3.

Class B devices

(excluding self-testing and near-patient testing (NPT) devices)

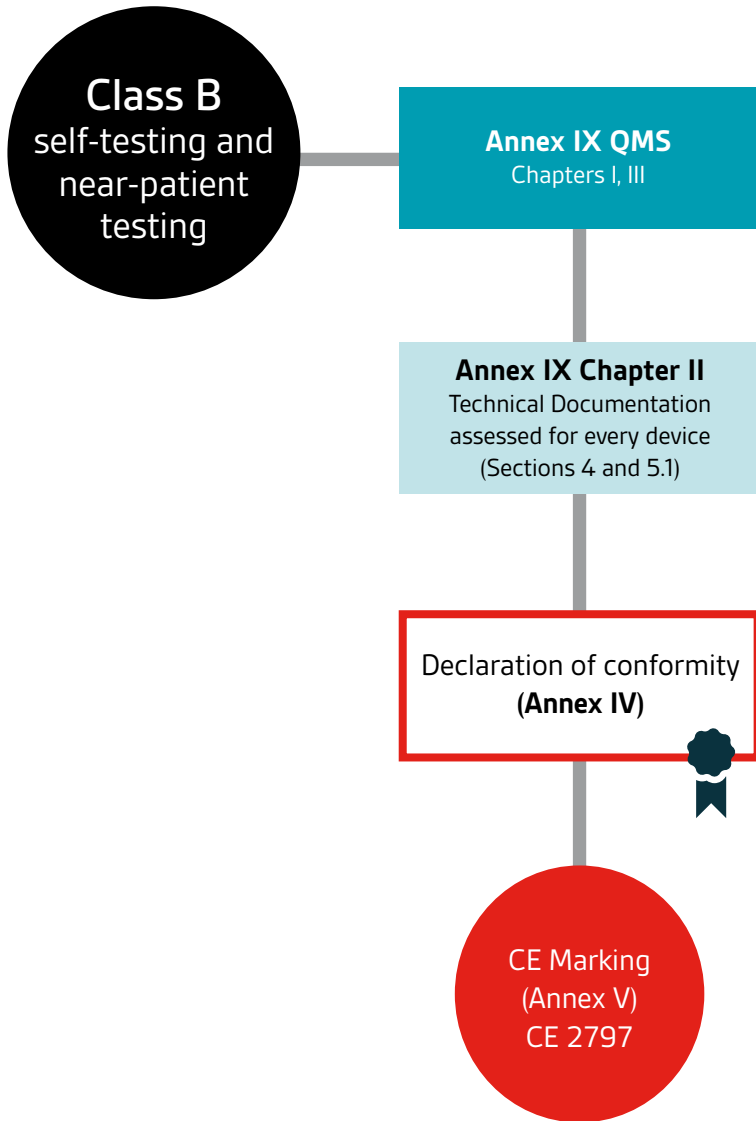


Class B devices (excluding self-testing and NPT devices)

Class B devices (excluding self-testing and NPT devices)	Initial Conformity Assessment	Surveillance				
		Y1	Y2	Y3	Y4	Y5
QMS Audits	Yes	Yes	Yes	Recertification*	Yes	Yes
Microbiology Audits	Yes (if sterile)	N/A	Yes (if sterile)	N/A	Yes (if sterile)	N/A
Technical Documentation Assessment	Sample per device category	As per the Technical Documentation Sampling Plan. A Technical Documentation surveillance audit is required every year whilst there are still devices left to be reviewed under the certificate scope.				
Competent Authority or EMA consultation (Annex IX, Section 5.2)	N/A	N/A	N/A	N/A	N/A	N/A
Experts consultations (article 48(6))	N/A	N/A	N/A	N/A	N/A	N/A
Verification by EU reference laboratory (Annex IX, section 4.9)	N/A	N/A	N/A	N/A	N/A	N/A
Summary of Safety and Performance (Article 29)	N/A	N/A	N/A	N/A	N/A	N/A
Performance Evaluation Report updates (Annex XIII - Part A, Section 1.3.2 and Article 56)		Updated as per Manufacturer's Performance Evaluation Plan; NB to review as per Technical Documentation Sampling Plan				
Post Market Performance Follow-up (PMPF) updates Evaluation Report (Article 56 and Annex XIII, Part B)		Updated as per Manufacturer's PMS, PMPF plans; NB to review as per Technical Documentation Sampling Plan. Implementation of the PMPF plan will be verified during annual surveillance visits				
Post Market Surveillance (PMS) Report (Article 80)		Updated when necessary and provided to the CA and/or NB upon request				
Periodic Safety Update Report (PSUR) (Article 81)		N/A	N/A	N/A	N/A	N/A
Unannounced Audits		At least once every 5 years				

* QMS certificates are valid for three years, whilst CE certificates remain valid for a maximum of five years. The **Y3 Recertification** indicated in the table relates to the EN ISO 13485:2016 certificate cycle. Certification cycles vary and re-certification may not always occur at Y3.

Class B self-testing and NPT devices



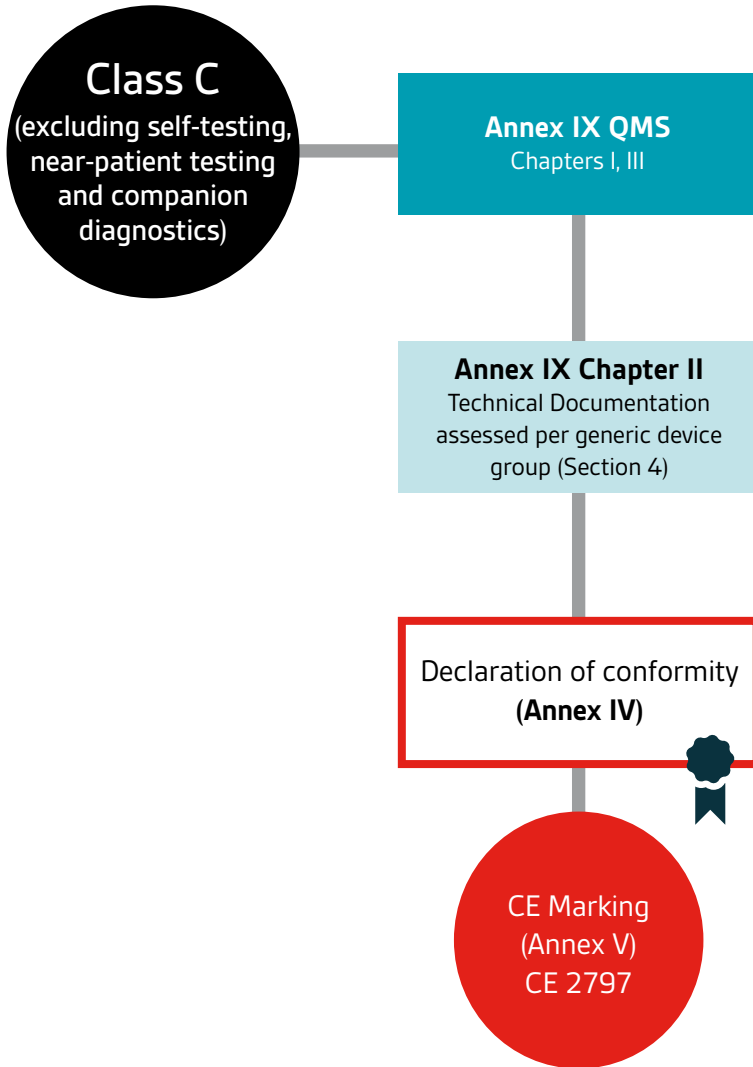
Class B devices self-testing and NPT devices

Class B devices self-testing and NPT devices	Initial Conformity Assessment	Surveillance				
		Y1	Y2	Y3	Y4	Y5
QMS Audits	Yes	Yes	Yes	Recertification*	Yes	Yes
Microbiology Audits	Yes (if sterile)	N/A	Yes (if sterile)	N/A	Yes (if sterile)	N/A
Technical Documentation Assessment	Review for every device	N/A	N/A	N/A	N/A	Recertification
Competent Authority or EMA consultation (Annex IX, Section 5.2)	N/A	N/A	N/A	N/A	N/A	N/A
Experts consultations (article 48(6))	N/A	N/A	N/A	N/A	N/A	N/A
Verification by EU reference laboratory (Annex IX, section 4.9)	N/A	N/A	N/A	N/A	N/A	N/A
Summary of Safety and Performance (Article 29)	N/A	N/A	N/A	N/A	N/A	N/A
Performance Evaluation Report updates (Annex XIII - Part A, Section 1.3.2 and Article 56)		Updated as per Manufacturer's Performance Evaluation Plan; NB to review at the time of substantial change reviews				
Post Market Performance Follow-up (PMPF) updates Evaluation Report (Article 56 and Annex XIII, Part B)		Updated as per Manufacturer's PMS, PMPF plans; NB to review at the time of substantial change reviews				
Post Market Surveillance (PMS) Report (Article 80)		Updated when necessary and provided to the CA upon request. NB to review at time of substantial change reviews				
Periodic Safety Update Report (PSUR) (Article 81)		N/A	N/A	N/A	N/A	N/A
Unannounced Audits		At least once every 5 years				

* QMS certificates are valid for three years, whilst CE certificates remain valid for a maximum of five years. The **Y3 Recertification** indicated in the table relates to the EN ISO 13485:2016 certificate cycle. Certification cycles vary and re-certification may not always occur at Y3.

Class C devices

(excluding self-testing, NPT and companion diagnostics (CDx))

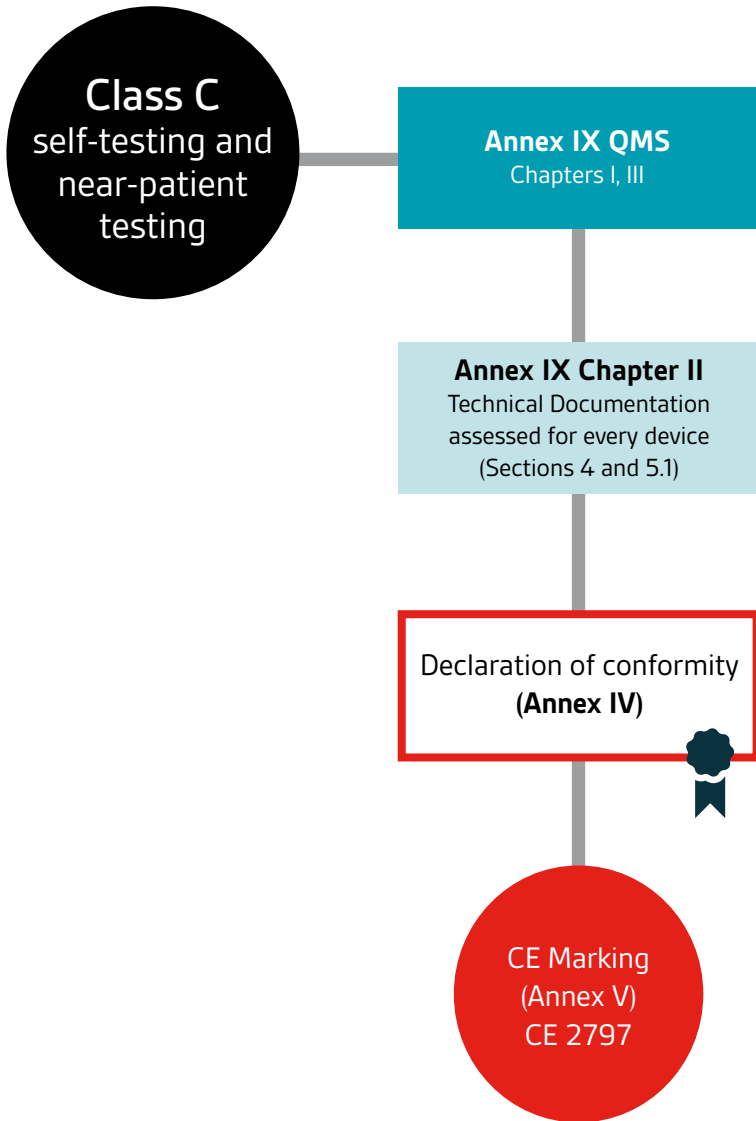


Class C devices (excluding self-testing, NPT and companion diagnostics (CDx))

Class C devices (excluding self-testing, NPT and CDx devices)	Initial Conformity Assessment	Surveillance				
		Y1	Y2	Y3	Y4	Y5
QMS Audits	Yes	Yes	Yes	Recertification*	Yes	Yes
Microbiology Audits	Yes (if sterile)	N/A	Yes (if sterile)	N/A	Yes (if sterile)	N/A
Technical Documentation Assessment	Sample per generic device group	As per the Technical Documentation Sampling Plan. A Technical Documentation surveillance audit is required every year whilst there are still devices left to be reviewed under the certificate scope				
Competent Authority or EMA consultation (Annex IX, Section 5.2)	N/A	N/A	N/A	N/A	N/A	N/A
Experts consultations (article 48(6))	N/A	N/A	N/A	N/A	N/A	N/A
Verification by EU reference laboratory (Annex IX, section 4.9)	N/A	N/A	N/A	N/A	N/A	N/A
Summary of Safety and Performance (Article 29)	Yes	Updated as soon as possible, where necessary				
Performance Evaluation Report updates (Annex XIII - Part A, Section 1.3.2 and Article 56)		Updated at least annually; NB to review as per Technical Documentation Sampling Plan				
Post Market Performance Follow-up (PMPF) updates Evaluation Report (Article 56 and Annex XIII, Part B)		Updated as per Manufacturer's PMS, PMPF plans; NB to review as per Technical Documentation Sampling Plan. NB QMS audits to verify implementation of the plan by sampling complaints, vigilance information etc.				
Post Market Surveillance (PMS) Report (Article 80)		Post-market surveillance will be captured in the Periodic Safety Update Report				
Periodic Safety Update Report (PSUR) (Article 81)		PSUR update required at least annually. The PSUR should be available to the NB upon request				
Unannounced Audits		At least once every 5 years				

* QMS certificates are valid for three years, whilst CE certificates remain valid for a maximum of five years. The **Y3 Recertification** indicated in the table relates to the EN ISO 13485:2016 certificate cycle. Certification cycles vary and re-certification may not always occur at Y3.

Class C self-testing and NPT devices

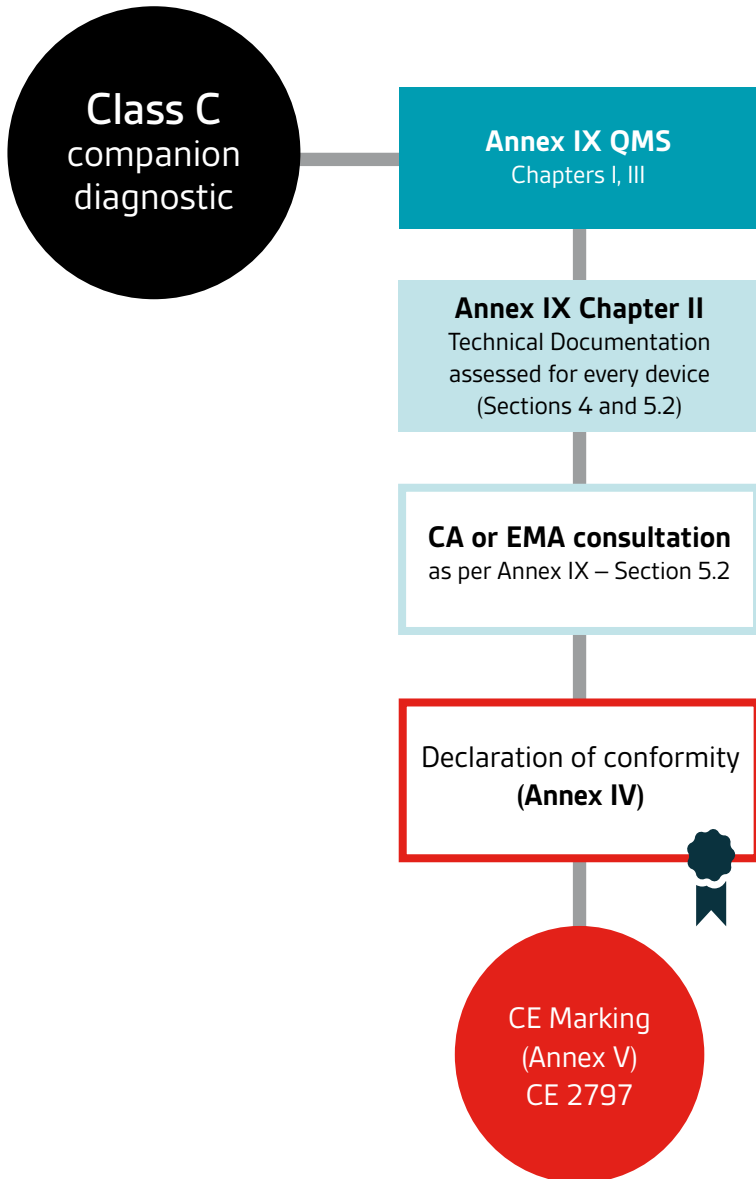


Class C devices self-testing and NPT devices

Class C self-testing and NPT devices	Initial Conformity Assessment	Surveillance				
		Y1	Y2	Y3	Y4	Y5
QMS Audits	Yes	Yes	Yes	Recertification*	Yes	Yes
Microbiology Audits	Yes (if sterile)	N/A	Yes (if sterile)	N/A	Yes (if sterile)	N/A
Technical Documentation Assessment	Review for every device	N/A	N/A	N/A	N/A	Recertification
Competent Authority or EMA consultation (Annex IX, Section 5.2)	N/A	N/A	N/A	N/A	N/A	N/A
Experts consultations (article 48(6))	N/A	N/A	N/A	N/A	N/A	N/A
Verification by EU reference laboratory (Annex IX, section 4.9)	N/A	N/A	N/A	N/A	N/A	N/A
Summary of Safety and Performance (Article 29)	Yes	Updated as soon as possible, where necessary				
Performance Evaluation Report updates (Annex XIII - Part A, Section 1.3.2 and Article 56)		Updated at least annually; NB to review at the time of PSUR reviews or substantial change reviews				
Post Market Performance Follow-up (PMPF) updates Evaluation Report (Article 56 and Annex XIII, Part B)		Updated as per Manufacturer's PMS, PMPF plans; NB to review at the time of PSUR reviews or substantial change reviews				
Post Market Surveillance (PMS) Report (Article 80)		Post-market surveillance will be captured in the Periodic Safety Update Report				
Periodic Safety Update Report (PSUR) (Article 81)		PSUR update required at least annually. The PSUR should be available to the NB upon request				
Unannounced Audits		At least once every 5 years				

* QMS certificates are valid for three years, whilst CE certificates remain valid for a maximum of five years. The **Y3 Recertification** indicated in the table relates to the EN ISO 13485:2016 certificate cycle. Certification cycles vary and re-certification may not always occur at Y3.

Class C companion diagnostic (CDx) devices



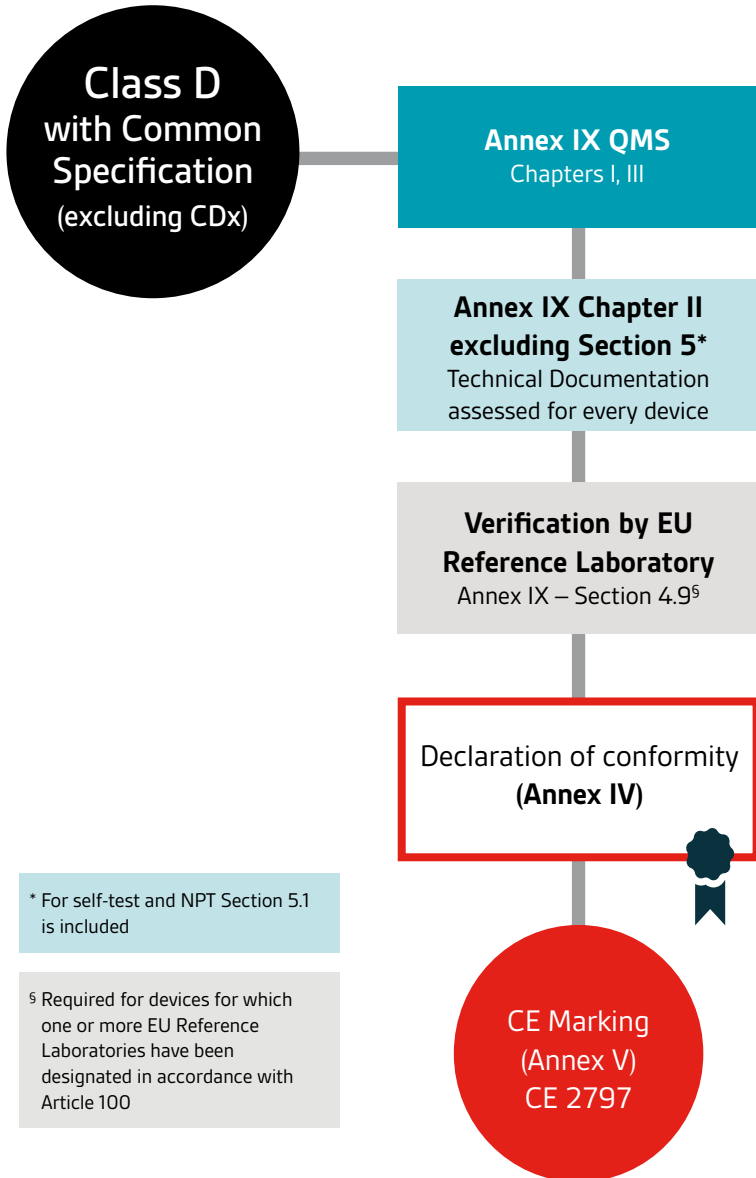
Class C companion diagnostic (CDx) devices

Class C companion diagnostic (CDx) devices	Initial Conformity Assessment	Surveillance				
		Y1	Y2	Y3	Y4	Y5
QMS Audits	Yes	Yes	Yes	Recertification*	Yes	Yes
Microbiology Audits	Yes (if sterile)	N/A	Yes (if sterile)	N/A	Yes (if sterile)	N/A
Technical Documentation Assessment	Review for every device	N/A	N/A	N/A	N/A	Recertification
Competent Authority or EMA consultation (Annex IX, Section 5.2)	Yes	Modifications to the devices may need supplementary consultations (determined on a case-by-case basis taking into account the nature of the changes proposed)				
Experts consultations (article 48(6))	N/A	N/A	N/A	N/A	N/A	N/A
Verification by EU reference laboratory (Annex IX, section 4.9)	N/A	N/A	N/A	N/A	N/A	N/A
Summary of Safety and Performance (Article 29)	Yes	Updated as soon as possible, where necessary				
Performance Evaluation Report updates (Annex XIII - Part A, Section 1.3.2 and Article 56)		Updated at least annually; NB to review at the time of PSUR reviews or substantial change reviews				
Post Market Performance Follow-up (PMPF) updates Evaluation Report (Article 56 and Annex XIII, Part B)		Updated as per Manufacturer's PMS, PMPF plans; NB to review at the time of PSUR reviews or substantial change reviews				
Post Market Surveillance (PMS) Report (Article 80)		Post-market surveillance will be captured in the Periodic Safety Update Report				
Periodic Safety Update Report (PSUR) (Article 81)		PSUR update required at least annually. The PSUR should be available to the NB upon request				
Unannounced Audits		At least once every 5 years				

* QMS certificates are valid for three years, whilst CE certificates remain valid for a maximum of five years. The **Y3 Recertification** indicated in the table relates to the EN ISO 13485:2016 certificate cycle. Certification cycles vary and re-certification may not always occur at Y3.

Class D with Common Specification

(excluding CDx)



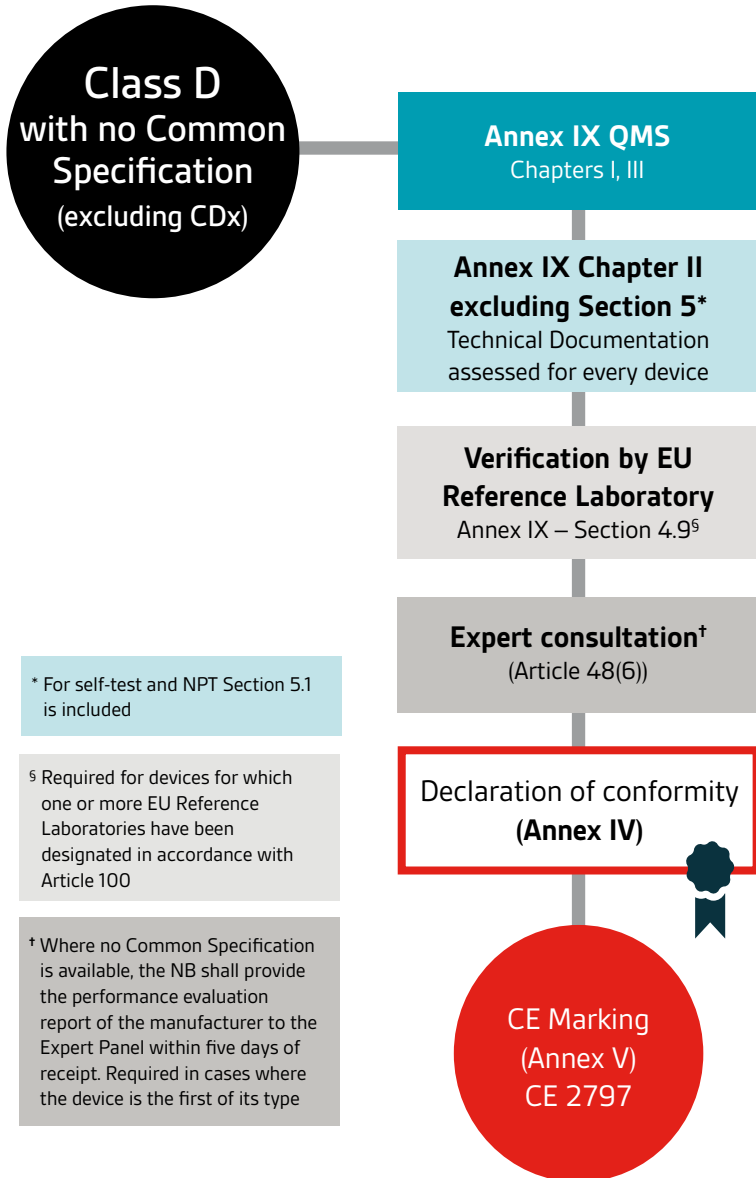
Class D with Common Specification (excluding CDx)

Class D with Common Specification (excluding CDx)	Initial Conformity Assessment	Surveillance				
		Y1	Y2	Y3	Y4	Y5
QMS Audits	Yes	Yes	Yes	Recertification*	Yes	Yes
Microbiology Audits	Yes (if sterile)	N/A	Yes (if sterile)	N/A	Yes (if sterile)	N/A
Technical Documentation Assessment	Review for every device	N/A	N/A	N/A	N/A	Recertification
Competent Authority or EMA consultation (Annex IX, Section 5.2)	N/A	N/A	N/A	N/A	N/A	N/A
Experts consultations (article 48(6))	N/A	N/A	N/A	N/A	N/A	N/A
Verification by EU reference laboratory (Annex IX, section 4.9)	Yes	Modifications to the devices may need supplementary verifications (determined on a case-by-case basis taking into account the nature of the changes proposed)				
Summary of Safety and Performance (Article 29)	Yes	Updated as soon as possible, where necessary				
Performance Evaluation Report updates (Annex XIII - Part A, Section 1.3.2 and Article 56)	Updated at least annually. NB to review at the time of PSUR reviews or substantial change reviews					
Post Market Performance Follow-up (PMPF) updates Evaluation Report (Article 56 and Annex XIII, Part B)	Updated as per Manufacturer's PMS, PMPF plans; NB to review at the time of PSUR reviews or substantial change reviews					
Post Market Surveillance (PMS) Report (Article 80)	Post-market surveillance will be captured in the Periodic Safety Update Report					
Periodic Safety Update Report (PSUR) (Article 81)	PSUR update required at least annually; submitted to the NB via EUDAMED for NB review					
Unannounced Audits	At least once every 5 years					

* QMS certificates are valid for three years, whilst CE certificates remain valid for a maximum of five years. The **Y3 Recertification** indicated in the table relates to the EN ISO 13485:2016 certificate cycle. Certification cycles vary and re-certification may not always occur at Y3.

Class D with no Common Specification

(excluding CDx)

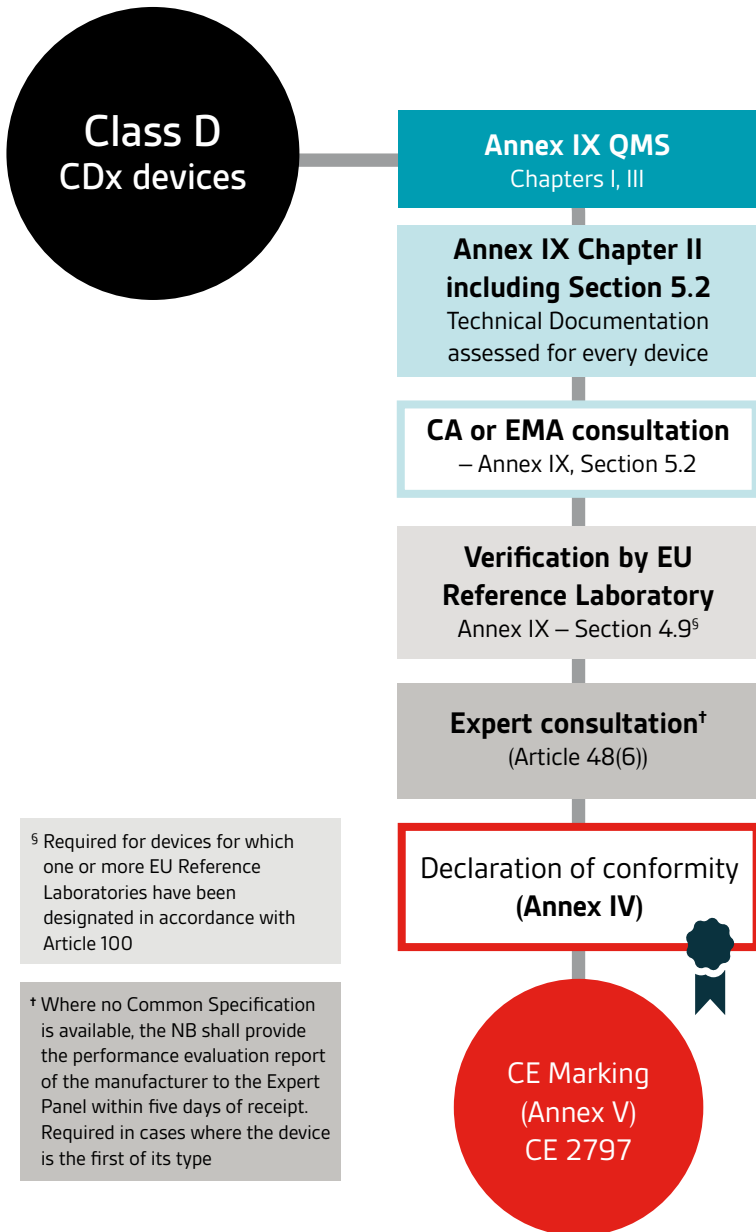


Class D with no Common Specification (excluding CDx)

Class D with no Common Specification (excluding CDx)	Initial Conformity Assessment	Surveillance				
		Y1	Y2	Y3	Y4	Y5
QMS Audits	Yes	Yes	Yes	Recertification*	Yes	Yes
Microbiology Audits	Yes (if sterile)	N/A	Yes (if sterile)	N/A	Yes (if sterile)	N/A
Technical Documentation Assessment	Review for every device	N/A	N/A	N/A	N/A	Recertification
Competent Authority or EMA consultation (Annex IX, Section 5.2)	N/A	N/A	N/A	N/A	N/A	N/A
Experts consultations (article 48(6))	Yes, if the device is the first of its type	N/A	N/A	N/A	N/A	N/A
Verification by EU reference laboratory (Annex IX, section 4.9)	Yes	Modifications to the devices may need supplementary verifications (determined on a case-by-case basis taking into account the nature of the changes proposed)				
Summary of Safety and Performance (Article 29)	Yes	Updated as soon as possible, where necessary				
Performance Evaluation Report updates (Annex XIII - Part A, Section 1.3.2 and Article 56)		Updated at least annually. NB to review at the time of PSUR reviews or substantial change reviews				
Post Market Performance Follow-up (PMPF) updates Evaluation Report (Article 56 and Annex XIII, Part B)		Updated as per Manufacturer's PMS, PMPF plans; NB to review at the time of PSUR reviews or substantial change reviews				
Post Market Surveillance (PMS) Report (Article 80)		Post-market surveillance will be captured in the Periodic Safety Update Report				
Periodic Safety Update Report (PSUR) (Article 81)		PSUR update required at least annually; submitted to the NB via EUDAMED for NB review				
Unannounced Audits		At least once every 5 years				

* QMS certificates are valid for three years, whilst CE certificates remain valid for a maximum of five years. The **Y3 Recertification** indicated in the table relates to the EN ISO 13485:2016 certificate cycle. Certification cycles vary and re-certification may not always occur at Y3.

Class D CDx devices



Class D CDx devices

Class D CDx devices	Initial Conformity Assessment	Surveillance				
		Y1	Y2	Y3	Y4	Y5
QMS Audits	Yes	Yes	Yes	Recertification*	Yes	Yes
Microbiology Audits	Yes (if sterile)	N/A	Yes (if sterile)	N/A	Yes (if sterile)	N/A
Technical Documentation Assessment	Review for every device	N/A	N/A	N/A	N/A	Recertification
Competent Authority or EMA consultation (Annex IX, Section 5.2)	Yes	Modifications to the devices may need supplementary consultations (determined on a case-by-case basis taking into account the nature of the changes proposed)				
Experts consultations (article 48(6))	Yes, if no CS and the device is the first of its type	N/A	N/A	N/A	N/A	N/A
Verification by EU reference laboratory (Annex IX, section 4.9)	Yes	Modifications to the devices may need supplementary verifications (determined on a case-by-case basis taking into account the nature of the changes proposed)				
Summary of Safety and Performance (Article 29)	Yes	Updated as soon as possible, where necessary.				
Performance Evaluation Report updates (Annex XIII - Part A, Section 1.3.2 and Article 56)		Updated at least annually; the NB will provide it to the expert panel as needed. NB to review at the time of PSUR reviews or substantial change reviews				
Post Market Performance Follow-up (PMPF) updates Evaluation Report (Article 56 and Annex XIII, Part B)		Updated as per Manufacturer's PMS, PMPF plans; NB to review at the time of PSUR reviews or substantial change reviews				
Post Market Surveillance (PMS) Report (Article 80)		Post-market surveillance will be captured in the Periodic Safety Update Report				
Periodic Safety Update Report (PSUR) (Article 81)		PSUR update required at least annually; submitted to the NB via EUDAMED for NB review				
Unannounced Audits		At least once every 5 years				

* QMS certificates are valid for three years, whilst CE certificates remain valid for a maximum of five years. The **Y3 Recertification** indicated in the table relates to the EN ISO 13485:2016 certificate cycle. Certification cycles vary and re-certification may not always occur at Y3.

Our website provides a wealth of resources including guidance documents, training courses, webinars and whitepapers.

To find out more, visit bsigroup.com/IVD

BSI Transitions
In Vitro Diagnostic Regulation

In-Vitro Diagnostic Regulation (IVDR) Readiness Review

Company Name _____
Address _____
Contact Name _____
Job Title _____
Telephone _____
Email _____
Certification No. _____
Date: _____

How ready are you for the IVD Regulation?
The IVD industry is undergoing significant change. The IVDR, which replaces the IVD Directive (98/79/EC), entered into force on May 26, 2022. EU Directives by their own end result that must be achieved in every Member State. National authorities have to adapt their laws to the IVDR.

BSI Transitions
In Vitro Diagnostic Regulation

IVDR Documentation Submissions
Best Practices Guidelines

bsi. ...making excellence a habit™



The EU IVDR Date of Application is approaching

Are you ready for the May 2022 deadline?

The **In Vitro Diagnostic Regulation (IVDR) EU 2017/746** entered into force in May 2017 with a five-year transition period. Manufacturers have the duration of the transition period to update their Technical Documentation to meet the requirements and comply with the Regulation before the Date of Application of the IVDR in May 2022.

Conduct assessments from a full scope EU IVDR Notified Body

As a manufacturer of an IVD medical device, you must ensure that your product meets the requirements of the IVDR. BSI, as a Notified Body, can help you ensure that your product meets the requirements of the IVDR.



Unrivalled expertise from an EU Notified Body and UK Approved Body

As a manufacturer of an In Vitro Diagnostic (IVD) medical device, you must ensure that you meet the relevant regulatory requirements before placing your product onto the market. For the EU, these are defined in **EU In Vitro Diagnostic Regulation (IVDR) EU 2017/746** and, for the UK, the **UK Medical Devices Regulations (UK MDR) 2002**.

It is critical to work with a notified body that understands the industry and has the experience to review and confirm your product's readiness for market – efficiently, promptly and robustly. At BSI, our technical specialists have extensive experience and can support you through the process of certifying your IVD medical device.

BSI The Netherlands (27379) is a leading Notified Body, we review medical devices to ensure that they conform to the requirements of the European Directives and Regulations. BSI UK (01080) is a UK-Approved Body able to provide conformity assessments under the new UKCA scheme.



Inspiring trust for a more resilient world.

Notes



Your resource for excellence

Talk to BSI

- We have 4,600 colleagues globally
- Offices in 31 countries around the world
- Over 84,000 clients operating in 193 countries
- Together our clients account for 75% of the FTSE 100 49% of the Fortune 500 and 77% of the Nikkei 225 listed companies
- More than 750 colleagues in BSI Medical Devices

Additional services

Medical device newsletter service

Keep updated on what's happening in the industry and changes in regulatory and quality requirements. You can take advantage of this free service by signing up on our website.

Informative webinars

Hear regular updates from our experts on key topics; listen live or listen back.

Comprehensive whitepapers

Our technical specialists collaborate with external experts to bring you the latest views and understanding on complex regulatory issues. Download your complimentary copies now.

Medical device guidance documents

Our online guidance documents provide assistance in understanding the regulatory requirements for medical devices.

Standards

BSI British Standards delivers leading-edge best practice solutions through the development and publication of more than 59,000 Standards and related products.

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