# Medical devices Proficiency Testing 2024

### Accreditation ISO/IEC 17043 (A2LA)

The DRRR is an accredited proficiency testing provider by A2LA according to ISO/IEC 17043:2010. The accreditation is only valid for the matrices/parameters listed on the A2LA scope of accreditation [#5494.01]. Whether a proficiency test is covered or not covered by the scope of accreditation by A2LA can be viewed in our online portal (ODIN).

In very rare individual cases an accredited proficiency testing round will not be carried out within the scope of accreditation due to technical or organizational reasons. In these rare cases the DRRR will inform the participants before the start of the proficiency testing round, thus before the sample shipment. An immediately free cancellation for the participants is possible until the date of the sample shipment.

#### Accreditation DIN EN ISO/IEC 17043 (DAkkS)

The DRRR is an accredited proficiency testing provider by DAkkS according to DIN EN ISO/IEC 17043:2010. The accreditation is valid only for the scope listed in the annex of the accreditation certificate [D-EP-17063-01-00]. Whether a proficiency test is covered or not covered by the scope of accreditation by DAkkS can be viewed in our online portal (ODIN).

In very rare individual cases an accredited proficiency testing round will not be carried out within the scope of accreditation due to technical or organizational reasons. In these rare cases the DRRR will inform the participants before the start of the proficiency testing round, thus before the sample shipment. An immediately free cancellation for the participants is possible until the date of the sample shipment.

#### Your benefit - DRRR Medical devices program

In 2024 more than 20 medical devices proficiency testing schemes are offered in the areas sterility, microbiology and chemical characterization.

By participating in our proficiency testing schemes you have an objective and independent impression of your quality and performance of your routine testing method. Your benefits taking part in DRRR proficiency testing schemes:

- Participation in proficiency testing is required by several facilities
- Participants can compare, assure and improve their own performance and quality
- Recognition of the applied method compared to the other laboratories
- Unquestionable lab performance towards customers and certification authorities
- Saving costs of lab development and maintenance
- Saving on the working time in the lab and many other benefits

### **Registration/information**

#### Simply brilliant, your proficiency testing with ODIN.

- Convenient proficiency testing participation with ODIN easy, safe and clearly
- · Direct booking of proficiency testing schemes in our online catalogue
- · Overview about the registered proficiency testing schemes
- · Fast and secure submission of your results via ODIN
- · Online access to individual customers reports and certificates

For questions and suggestions do not hesitate to contact us!

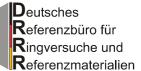
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## Medical devices proficiency tests: Registration for 2024



Art. no.	Matrix group	Proficiency testing type <sup>[A]</sup>	Period	To view pricing information please visit our
Chen	nical-physical P	Ts:		online Portal:
2010375	Sterility	Medical devices - Ethylene oxide residues (ISO 10993-7) 1	Jun-24	Login or register
2010377		Medical devices - Ethylene oxide residues (ISO 10993-7) 2	Jun-24	<u></u>
2010961		Medical devices - Protein residues	Sep-24	
2010379	Chemical	Medical devices - Extraction of metals (ISO 10993-12)	Jun-24	
2010381	characterization	Medical devices - Qual. characterization (ISO 10993-18)	Jul-24	
2010383		Medical devices - Quan. characterization (ISO 10993-18) - XRF	Jul-24	
		Medical devices - Quan. characterization (ISO 10993-18) -		
2010385		Formaldehyde	Jul-24	
		Medical devices - Degradation products from ceramics (ISO 10993-		
2010387		14) 1	May-24	
		Medical devices - Degradation products from ceramics (ISO 10993-		
2010389	-	14) 2	May-24	
2010391	_	Medical devices - Loss of polymer mass (ISO 10993-13) 1	Aug-24	
2010393	_	Medical devices - Loss of polymer mass (ISO 10993-13) 2	Aug-24	
2011168		Medical devices - Gravim. determination extractables	Jul-24	
2011169		Medical devices - Extractables & leachables	Aug-24	
2011159		Medical devices - Extraction of elements (ISO 10993-12)	Sep-24	
Micro	biological PTs:			
ivitor				
2010696		testing of sterilization of medical devices 1 (ISO 11737-1)	Jun-24	
2010964		testing of sterilization of medical devices 2 (ISO 11737-1)	Jun-24	
2010966		testing of sterilization of medical devices 3 (ISO 11737-1)	lun-24	

2010964	testing of sterilization of medical devices 2 (ISO 11737-1)	Jun-24	
2010966	testing of sterilization of medical devices 3 (ISO 11737-1)	Jun-24	
2010968	testing of sterilization of medical devices 4 (ISO 11737-1)	Jun-24	
2011171	testing of sterilization of medical devices 5 (ISO 11737-1)	Jun-24	
2010281	Tests for in vitro cytotoxicity (ISO 10993-5)	Nov-24	
2010283	Testing the germ tightness of packaging materials	Nov-24	
2010657	Identification of microorganisms using MALDI-ToF	Nov-24	
2010321	Test method Medical face masks (EN 14683)	May-24	
2010567	Microbiological analysis of endoscopes	Oct-24	

[A] = For accredited and non-accredited status please see Online portal (ODIN)

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Additional samples	s are required for the following tests:
Quantity	Art. No. / Proficiency testing type

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If you register for a large number of Proficency tests please contact us for special prices!

An offer with the total costs is needed A Purchase order from the purchasing department will follow

Registration by e-mail:

#### info@DRRR.de

Hereby we confirm obligatorily the participation in the above mentioned test(s) and the order for the additional sample sets.

	company
	additional line
	contact person
	street
	post code / city
	country
	email
	VAT-ID (EU)
Date:	

#### Deutsches Referenzbüro

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