

## EU Quality Management System Certificate

The Notified Body

**MEDCERT Zertifizierungs- und Prüfungsgesellschaft für die Medizin GmbH  
Pilatuspool 2 – 20355 Hamburg – Germany**

herewith certifies that the company

**OHST Medizintechnik AG  
Grünauer Fenn 3  
14712 Rathenow  
Germany  
SRN DE-MF-000008887**

has introduced, applies and maintains a quality management system for the medical devices/groups of medical devices listed in the appendix.

The compliance of this quality management system to the requirements of the **Regulation (EU) 2017/745 on medical devices** was verified by assessment according to:

### Annex IX Chapter I

Any applicable limitations of this certification for certain medical devices are included in the appendix. This certification is subject to surveillance by MEDCERT.

**Effective date:** 2022-08-25  
**Expiry date:** 2025-12-03

Report No.: 0529IA30F  
Procedure No.: QS – 0529  
Certificate No.: 0529GB448220825

Preceding certificate No.: 0529GB448210813  
Preceding certificate date: 2021-08-13  
Identification of changes: WO-010185

Hamburg, 2022-08-25

  
\_\_\_\_\_  
MEDCERT Certification Body  
Lorenz Runge

The certificate is only valid when provided entirely with all of its pages.  
To verify the validity of this certificate, contact [info@medcert.de](mailto:info@medcert.de).

MEDCERT Notified Body Identification Number: 0482



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bei Arzneimitteln und  
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BS-MDR-096

**Appendix of EU Quality Management System Certificate**

Procedure No.: QS – 0529  
Certificate No.: 0529GB448220825

**Class IIa medical devices**

Category	EMDN code	Medical devices/groups of medical devices
MDN 1208	L091199	Orthopaedic prosthetics instruments, reusable - other

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## Appendix of EU Quality Management System Certificate

Procedure No.: QS – 0529  
Certificate No.: 0529GB448220825

### Class III custom-made implantable medical devices

<b>Category</b>	<b>Medical devices/groups of medical devices</b>
MDN 1102	Non-active osteo- and orthopaedic implants

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## Appendix of EU Quality Management System Certificate

Procedure No.: QS – 0529  
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### Class III medical devices

For placing on the market of class III medical devices covered by this certificate, an additional **EU Technical Documentation Assessment Certificate according to Annex IX Chapter II** of Regulation (EU) 2017/745 is required, which also contains the exact determination of medical devices covered by certification.

<b>Category</b>	<b>Medical devices/groups of medical devices</b>
MDN 1102	Non-active osteo- and orthopaedic implants

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