



Regulation (EU) 2017/745, Annex IX Chapter I and III

#### MDR 730376 R000

**Manufacturer:** EUROS SAS

Address: Z.E Athélia III LA CIOTAT 13600 France

Single Registration Number: FR-MF-000000605

#### Scope: See attached Device Schedule

On the basis of our examination of the quality system in accordance with Regulation (EU) 2017/745, Annex IX Chapter I and III, the quality system meets the requirements of the Regulation. For the placing on the market of Class III and Class IIb implantable devices an Annex IX Chapter II certificate is required.

For and on behalf of BSI, a Notified Body for the above Regulation (Notified Body Number 2797):

Gary E Slack, Senior Vice President Medical Devices

Gary C Stade

First Issued: **2021-12-01** Date: **2021-12-01** Expiry Date: **2026-11-30** 

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Validity of this certificate is conditional on the Manufacturer's quality system being maintained to the requirements of the Regulation as demonstrated through the required surveillance activities of the Notified Body.

This certificate was issued electronically and is bound by the conditions of the contract.

NB Contact: BSI Group The Netherlands B.V., Say Building, John M. Keynesplein 9, 1066 EP, Amsterdam, Netherlands. Tel: + 31 (0) 20 346 07 80 Corporate Contact: BSI Group Assurance Limited, registered in England under number 05435540 at 389 Chiswick High Road, London, W4 4AL, UK. A Member of the BSI Group of Companies.





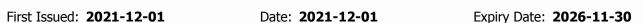
Regulation (EU) 2017/745, Annex IX Chapter I and III

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#### **Device Schedule: Class IIa, Custom-made and other devices**

Device(s)	Risk Classification	
Reusable instruments 'Orthopaedic Instruments'	Class Ir	
Custom made 'Cranial Plate'	Custom-made Class III implantable	
For Class Ir devices (Class I re-usable surgical instrument	ts) the Notified Body conformity assessment is limited to the aspects	

For Class Ir devices (Class I re-usable surgical instruments), the Notified Body conformity assessment is limited to the aspects relating to the reuse of the device.



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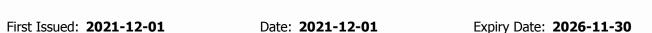
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#### **Certificate History**

(References to applicable Common Specifications, Harmonized Standards complied with, and the relevant test and audit reports that support any of the below certificate changes may be requested from Certificate. Verification@bsigroup.com)

Date	Reference Number	Action	TI Sea
Current	3221218	Issued	



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### List of Critical Subcontractors and Crucial Suppliers

Recognised as being involved in services related to the products covered by:

### MDR 730376 R000

Date: 2021-12-01

**Critical Subcontractor/Crucial Supplier** 

Service(s) supplied

Synergy Health Marseille - SAS Site de Marseille MIN 712 - LES Arnavaux MARSEILLE CEDEX 14 13323 France **Radiation (Gamma Sterilization)** 

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