EU Quality Management System Certificate

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

MDR 724378 R000

Manufacturer: Enztec Limited

Address:
3/17 Print Place
Middleton
Christchurch
8024
New Zealand

Single Registration Number: Not Available

EU Authorised Representative: Advena Limited

Address:
Tower Business Centre, 2nd Flr.,
Tower Street
Swatar, BKR 4013
Malta

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

First Issued: **2020-08-06** Date: **2020-10-19** Expiry Date: **2025-08-05**

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

Device(s)	Risk Classification
Orthopaedic instruments connected to an active device - Reusable Surgical Instruments	Class IIa
Reusable surgical instruments "Orthopaedic instruments"	Class Ir
Reusable surgical instruments "Orthopaedic instruments"	Class Ir & Im
For Class Im devices, the Notified Body conformity assessment is limited to the aspects relating to the conformity of the devices with the metrological requirements. 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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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
06 August 2020	3147250	First Issue
Current	3312997	Add Class IIa Instrument attached to an active device, to schedule

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