



America

CERTIFICATE

No. QS6 074222 0014 Rev. 00

Certificate Holder: **Creagh Medical Ltd**
 IDA Business Park
 Ballinasloe, Co Galway, H53 K8P4
 IRELAND

Certification Mark:



Scope of Certificate: **Design, Development and Production of Percutaneous Transluminal Angioplasty and Percutaneous Transluminal Coronary Angioplasty Catheters; Drug Coating and Production of Drug Coated Balloon Catheters**

Standard(s): **ISO 13485:2016**

Regulatory Authority(ies): **Australia TGA, Brazil ANVISA, Health Canada, USA FDA, MHLW / PMDA. See attached for listing of specific regulatory requirements.**

The Certification Body of TÜV SÜD America Inc. certifies that the quality management system of the manufacturer listed above has been audited against the stated criteria and found to conform to those criteria for the scope of certification listed. Validity of this certificate can be obtained by visiting the website www.tuvsud.com/ps-cert
 TÜV SÜD America Inc. is an MDSAP Recognized Auditing Organization.

DUNS No: **93-350-1909**

Effective Date: **2020-09-09**

Expiry Date: **2023-09-08**

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Date of Issue: 2020-10-02

(Tina Israel)
 Manager, US Certification Body,
 Medical and Health Services

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Regulatory Requirements: Audit/Certification Criteria

Australia

Therapeutic Goods (Medical Devices) Regulations 2002
 - Schedule 3, Part 1 (excluding Part 1.6) – Full Quality Assurance Procedure

Brazil

- RDC ANVISA n. 16/2013
 - RDC ANVISA n. 23/2012
 - RDC ANVISA n. 67/2009

Canada

- Medical Device Regulations – Part 1- SOR 98/282

Japan

- MHLW Ministerial Ordinance 169, Article 4 to Article 68
 - PMD Act

United States

- 21 CFR Part 803
 - 21 CFR Part 806
 - 21 CFR Part 807 – Subparts A to D
 - 21 CFR Part 820

Facility(ies):

Creagh Medical Ltd
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Facility Scopes:

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