

## EU Technical Documentation Assessment Certificate

Regulation (EU) 2017/745, Annex IX Chapter II

### MDR 758295 R000

**Manufacturer:** Argentum Medical LLC

**Address:**

2571 Kaneville Court  
Geneva  
Illinois  
60134  
USA

**Single Registration Number:** US-MF-000003391

**EU Authorised Representative:** Emergo Europe B.V.

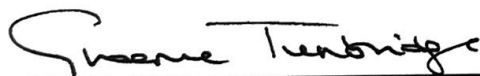
**Address:**

Westervoortsedijk 60  
6827 AT Arnhem  
The Netherlands

**Scope:** See attached **Device Schedule**

On the basis of our assessment of the technical documentation in accordance with Regulation (EU) 2017/745, Annex IX Chapter II, the technical documentation meets the requirements of the Regulation. For the placing on the market of these devices an additional Annex IX Chapter I and III certificate is required.

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



Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

First Issue Date: **2025-02-14**

Current Issue Date: **2025-02-14**

Starting Validity Date: **2025-02-14**

Expiry Date: **2030-02-13**

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