



EC Design-Examination Certificate

Directive 93/42/EEC on Medical Devices, Annex II Section 4

No. **CE 720268**
Issued To: **Medos International SARL**
Chemin Blanc 38
Le Locle
CH-2400
Switzerland

In respect of:

CEREBASE DA Guide Sheath

BSI has performed a design examination on the above devices in accordance with the Council Directive 93/42/EEC, Annex II Section 4. The design conforms to the requirements of this directive. For marketing of these products an additional Annex II excluding Section 4 certificate is required.

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

Gary E Slack, Senior Vice President Medical Devices

First Issued: **2020-04-13**

Date: **2020-04-13**

Expiry Date: **2024-05-26**

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Page 1 of 3

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive 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.

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Supplementary Information to CE 720268

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Catalogue Number	Device Name	Model, Type	Intended purpose per IFU	Classification
GS9095SD	CEREBASE DA Guide Sheath	95cm	The CEREBASE DA Guide Sheath is indicated for the introduction of interventional devices into the neuro vasculature	Class III
GS9090SD		90cm		
GS9080SD		80cm		
GS9070SD		70cm		

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Certificate History

Date	Reference Number	Action
Current	3097316	First Issue.

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Page 3 of 3

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