



# EC-CERTIFICATE

(Full quality assurance system)

## DQS Medizinprodukte GmbH

hereby certifies that the company

### Christoph Miethke GmbH & Co. KG

Ulanenweg 2  
14469 Potsdam  
Germany

has implemented and maintains a full quality assurance system which applies to the products at every stage from design to final controls.

An audit, documented in a report, performed by DQS, has verified that this quality assurance system fulfils the requirements of

### **Annex II – excluding Section 4 of Council Directive 93/42/EEC concerning medical devices**

with respect to the following medical devices:

Implants and accessories for the treatment of hydrocephalus according to annex

The manufacturer is subject to surveillance according to Annex II, Section 5. The CE marking with the Notified Body Number (0297) may be affixed on the devices listed in the certificate. An EC Design Examination Certificate according to Annex II, Section 4 is required for class III devices covered by this certificate.

Certificate registration no.	009066 MR2
Certification ID	170502072
Date of certification	2010-06-23
Valid until	2013-06-22

Frank Graichen  
Managing Director

Stefan Hofmann  
Head of Certification Body

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**DQS Medizinprodukte GmbH is a Notified Body according to Council Directive 93/42/EEC concerning medical devices with the Notified Body Number 0297.**



**Annex to Certificate**  
**Registration No. 009066 MR2**  
**Certificate ID 170502072**  
**(Issued: 2010-06-23)**

## **Christoph Miethke GmbH & Co. KG**

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14469 Potsdam

**Product groups:**

**Products:**

Hydrocephalus valves:

DSV (IIb)  
MSV (IIb)  
Shunt Assistant (IIb)  
PAEDI-GAV (IIb)  
GAV (IIb)  
proGAV (IIb)  
uniGAV (IIb)  
miniNAV (IIb)  
proSA (IIb)

Accessories:

Ventricular catheters (III)  
Peritoneal catheters (IIb)  
Burr hole reservoirs (IIb)  
Flushing reservoirs (IIb)  
Deflectors (IIb)  
Connectors (IIb)

Instruments:

Tunneler for single use (IIa)  
Check Mate (Is)