



# Certificate

No. Q5 104938 0001 Rev. 02

**Holder of Certificate:** **MIPM Mammendorfer Institut  
für Physik und Medizin GmbH**  
Oskar-von-Miller-Str. 6  
82291 Mammendorf  
GERMANY

**Certification Mark:**



**Scope of Certificate:** **Design and Development, Manufacturing,  
Distribution and Servicing of Patient  
Monitoring Systems (including sensors and  
accessories) for vital and non-vital  
physiological parameters.  
Service Provider for infusion systems,  
breathing systems and anesthesia systems**

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: [www.tuvsud.com/ps-cert?q=cert:Q5 104938 0001 Rev. 02](http://www.tuvsud.com/ps-cert?q=cert:Q5_104938_0001_Rev._02)

**Report No.:** 713210868

**Valid from:** 2022-05-11

**Valid until:** 2025-05-10

**Date,** 2022-05-10

Christoph Dicks  
Head of Certification/Notified Body

# Certificate

No. Q5 104938 0001 Rev. 02

**Applied Standard(s):** EN ISO 13485:2016  
Medical devices - Quality management systems -  
Requirements for regulatory purposes  
(ISO 13485:2016)  
DIN EN ISO 13485:2016

**Facility(ies):** MIPM Mammendorfer Institut für Physik und Medizin GmbH  
Oskar-von-Miller-Str. 6, 82291 Mammendorf, GERMANY

Design and Development, Manufacturing, Distribution  
and Servicing of Patient Monitoring Systems  
(including sensors and accessories) for vital and  
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systems and anesthesia systems

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# Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016 & EN ISO 13485:2016

This is to certify that:

Viamed Ltd  
15/17 Station Road  
Cross Hills  
Keighley  
BD20 7DT  
United Kingdom

Holds Certificate Number:

**MD 78787**

and operates a Quality Management System which complies with the requirements of ISO 13485:2016 & EN ISO 13485:2016 for the following scope:

Service of Supra-Maximal Peripheral Nerve Stimulator, Simulation Equipment, Infant TPiece Resuscitators, Apgar Timer and Resuscitation Cabinets.  
Distribution and Service of electromedical devices.



For and on behalf of BSI:

Gary E Slack, Senior Vice President - Medical Devices

Original Registration Date: 2004-01-27

Latest Revision Date: 2022-02-16

Effective Date: 2022-02-26

Expiry Date: 2025-02-25

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