

# Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

Viant Medical LLC  
620 Watson SW  
Grand Rapids  
Michigan  
49504  
USA

Holds Certificate No:

**FM 590425**

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

Production, assembly and packaging of disposable and implantable non-active medical devices and active medical devices including sterile medical devices. Operation of microbiology laboratory to support sterility and environmental control testing for medical devices.



For and on behalf of BSI:

Stewart Brain, Head of Compliance & Risk - Medical Devices

Original Registration Date: 2012-07-25

Latest Revision Date: 2019-05-30

Effective Date: 2017-05-01

Expiry Date: 2020-04-30

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Certificate No: **FM 590425**

Location	Registered Activities
Viant Medical LLC 620 Watson SW Grand Rapids Michigan 49504 USA	Production, assembly and packaging of disposable and implantable non-active medical devices and active medical devices including sterile medical devices. Operation of microbiology laboratory to support sterility and environmental control testing for medical devices.
Viant Medical LLC 5079 33rd St. SE Grand Rapids Michigan 49512 USA	Production, assembly and packaging of disposable and implantable non-active medical devices and active medical devices including sterile medical devices. Operation of microbiology laboratory to support sterility and environmental control testing for medical devices.



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An electronic certificate can be authenticated [online](#). Printed copies can be validated at [www.bsigroup.com/ClientDirectory](http://www.bsigroup.com/ClientDirectory)  
To be read in conjunction with the scope above or the attached appendix.

Americas Headquarters: BSI Group America Inc., 12950 Worldgate Drive, Suite 800, Herndon, VA 20170-6007 USA  
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