

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

Viant Medical Inc.
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:

Gary E Slack, Senior Vice President - Medical Devices

Original Registration Date: 2012-07-25

Latest Revision Date: 2020-03-25

Effective Date: 2020-05-01

Expiry Date: 2023-04-30

Page: 1 of 2



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Certificate No: **FM 590425**

| Location | Registered Activities |
|--|--|
| Viant Medical Inc. 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 Inc. 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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Page: 2 of 2

This certificate remains the property of BSI and shall be returned immediately upon request.
An electronic certificate can be authenticated [online](#). Printed copies can be validated at www.bsigroup.com/ClientDirectory
To be read in conjunction with the scope above or the attached appendix.

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