Supplier Info Card – Viant Medical, Inc.

Facility Address:

Building 1:
Viant Medical, Inc.
620 Watson Street SW
Grand Rapids, MI 49504

Phone: 616-643-5200
Fax: 616-643-1094

Building 2:
Viant Medical, Inc.
5079 33rd Street SE
Grand Rapids, MI 49512

Phone: 616-325-2514
Fax: 616-913-1501

Building 3:
Viant Medical, Inc. (Sterilization Services)
520 Watson Street SW
Grand Rapids, MI 49504

Phone: 616-643-5261
Fax: 616-643-5292

Website: www.viantmedical.com

Facility Information Building 1 (Watson):
Total square feet: 97,000
ISO Class 7 Controlled Environment: 1,846 square feet

Facility Information Building 2 (33rd St.):
Total square feet: 110,000
ISO Class 8 Controlled Environment: 8,022 Square Feet

Facility Information Building 3 (Sterilization Services):
Total square feet: 26,000

Company Profile:

Viant Medical, Inc. was formerly known as "MedPlast Medical, Inc." The Grand Rapids, Michigan 620 Watson and 5079 33rd street facilities are medical device contract manufacturing locations with a Quality System certified to EN ISO 13485:2016, and registered with the FDA. The Grand Rapids, Michigan 520 Watson facility is a contract sterilization location certified to EN ISO 13485:2016, EN ISO 11135-1:2007, ISO 11135:2014, and registered with the FDA.

Total employees: 485
Manufacturing: 348
Quality Department: 40

Primary Customer Contact:

Director of Operations: Ryan Mack
Email: Ryan.Mack@viantmedical.com

Phone: 616-325-2526

A/R Contact: Open
Email:

Phone:

Additional Contacts:

Sr. Director Quality Assurance & Regulatory Affairs	Bryan Curry	210-316-3950
Operations Manager, 33rd Street	Ryan Mack	616-325-2526
Director of Supply Chain	Open	

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Quality Management System:
FDA Registration Number: There are two establishment registration numbers: 1419629 for 620 Watson and 520 Watson and 3009493875 for 5079 33rd street.

ISO Registration: 13485:2016 Certified

Customer Quality System Audits - This information card is intended to be used by our customers as a "letter to file" that will allow previously conducted Quality System audits to remain valid until the next scheduled audit.

Additional Facility Information:
100% of our business is in the medical device market.
We do capacity planning for operations.
We have an MRP system in place.
Customer-supplied documents are controlled.
Customer-supplied fixtures and gauges are controlled.
Customer-supplied materials are controlled.
We manage Quality Agreements with all customers.
We monitor our facilities and conduct laboratory services such as BI testing and LAL testing.
Our Business Excellence group leads Lean initiatives.
Subcontractors provide services such as sterilization, laboratory services, component fabrication, passivation, laser welding, and injection molding.