# Supplier Info Card – Vention Medical, Inc. (Michigan)

**Facility Address:**
Building 1
Vention Medical, Inc.
620 Watson Street SW
Grand Rapids, MI 49504

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

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

**Phone:**616-325-2514
**Fax:** 616-325-2507

**Website:** www.ventionmedical.com

**Facility Information Building 1 (Watson)**:
Total Square ft: 98,000

**Facility Information Building 2 (33rd)**:
Total Square ft: 81,650

ISO Class 8 Controlled Environment
Watson: 1846 Square ft
33rd: 1150 Square ft
Total: 2996 Square ft

ISO Class 7 Controlled Environment
Watson: 8846 Square ft
Total: 8846 Square ft

**Company Profile:**

This Company was previously known as "ATEK Medical, LLC, a Vention Medical Company", and has merged with "The MedTech Group, Inc., a Vention Medical Company." The new company is *Vention Medical, Inc.* The Grand Rapids, MI facilities are Medical Device Contract Manufacturing locations with a Quality System certified to ISO 13485:2003, with facilities registered with the FDA.

**Total Employees:** ≈ 300
**Manufacturing:** ≈ 175

**Quality Department:** ≈ 30
Sterilization is not performed on site

**Primary Customer Contact:**

**Operations Manager:** Brian Kimble
**Email:** bkimble@ventionmedical.com

**Phone:616-643-5541**
**Fax: 616-643-1094**

**A/R Contact**: Diana Lanning
**Email:** dlanning@ventionmedical.com

**Phone:616.643.5220**
**Fax: 616-643-1094**

**Additional Contacts**:

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|---|---|---|
| Director of Quality/Regulatory | Carla Krause | 616-643-7326 |
| Value Stream Manager | Don Goetzinger | 616-643-5261 |
| Value Stream Manager | Todd Heald | 616-643-5230 |
| Value Stream Manager | Mike Kitchen | 616-325-1494 |
| Supply Chain Manager | Tom Annis | 616-643-5557 |

**Quality Management System**:
**FDA Registration Number**: Current FDA Facility Registrations and device listings will remain intact. There are two Establishment registration numbers: 1419629 for Building 1 (Watson) and 3009493875 for building 2 (33rd St). Each will be updated in accordance with 21 CFR 807 – *Establishment Registration and Device Listing*, §30 *Updating device listing information*.

**ISO Registration**: 13485:2003 Certified – Quality System Registrations will remain intact. They will be updated with name changes in accordance with our registrar's standard procedures. A copy of the updated cert will be available on our Vention Medical Website in accordance with our registrar's standard procedures.

**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 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 require our suppliers to sign an NDA
We manage Quality Agreements with all customers
Laboratory Services such as BI testing and LAL testing
Facility Monitoring
Lean Initiatives led by the Business Excellence group

Subcontractors provide services such as; sterilization, laboratory services, component fabrication, passivation, laser welding, and injection molding.