Supplier Info Card – Vention Medical Costa Rica, S.A.

Facility Address:
Vention Medical Costa Rica, S.A**.**
Parque Zone Franca, Metropolitana
Edificio 2C,
Barreal De Heredia, Costa Rica

Website: www.ventionmedical.com

Facility Information Building 1:
Total Square ft: 31.215

Facility Information Building 2:
Total Square ft: 26.156

Facility Information Building 3:
Total Square ft:33.658

Facility Information Building 4:
Total Square ft:84.000

ISO Class 8 Controlled Environment
B1: 13,850 sq ft
B2: 7,000 sq ft
B3: 10,000 sq ft
B4: 7,000 sq ft
ISO Class 7 Controlled Environment

Company Profile:
The MedTech Group, Inc., and ATEK Medical, LLC are wholly owned subsidiaries of Vention Medical. The former MedTech Costa Rica, S.A. and former ATEK Medical Costa Rica LIMITADA have merged. The new entity will now be known as *Vention Medical Costa Rica, S.A*. The business purpose of Medical Device Contract Manufacturing including Injection Molding and Assembly remains the same.

Total Employees: ≈ 615
Manufacturing: ≈ 500

Quality Department: ≈ 67
Sterilization is not performed on site

Primary Customer Contact:
General Manager, Yesenia Frago
Email: YFrago@ventionmedical.com
Phone: (506)-2239.9202
Fax: (506)-2338.9897

A/R Contact: Walter Picado, Controller
Email:wpicado@ventionmedical.com
Phone: (506) 2239.92.98
Fax: (506) 2239.12.31

Juridica number: 3-101365187 (this is the former MedTech Costa Rica, S.A.'s Juridica number)

Quality Management System:
FDA Registration Number: Current FDA Facility Registrations and device listings will remain intact. There are two Establishment registration numbers: 300573255 for Building 1 and 2; and 3006785981 for buildings 3 and 4. Each will be updated in accordance with 21 CFR 807 – *Establishment Registration and Device Listing*, §30 *Updating device listing information*.

ISO Registration: 13485 Certified – Quality System Registrations will remain intact. And will be updated with name changes 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 product development services
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 a confidentiality agreement
Subcontractors provide services such as; sterilization, component fabrication, testing labs, etc.