Supplier Info Card – Viant Costa Rica, S.A.

Facility Address:
Viant Costa Rica, S.A.
Parque Zona Franca, Metropolitana
Edificio 2C
Barreal De Heredia, Costa Rica

Website: www.viantmedical.com

Facility Information Building 1:
Total square feet: 31,453

Facility Information Building 2:
Total square feet: 29,419

Facility Information Building 3:
Total square feet: 35,837

Facility Information Building 4:
Total square feet: 81,439

ISO Class 8 Controlled Environments:
- B1: 13,886 square feet
- B2: 7,500 square feet
- B4: 16,884 square feet

ISO Class 7 Controlled Environment:
- B3: 7,537 square feet

Company Profile:
This company was previously known as "MedPlast Medical Costa Rica S.A." The Costa Rica facility is a medical device contract manufacturing location with a Quality System certified to ISO 13485:2016, with facilities registered with the FDA.

Total Employees: ≈ 635
Manufacturing: ≈ 553
Quality Department: ≈ 82
Sterilization is not performed on site.

Primary Customer Contact:
General Manager: Alberto Meseguer
Email: alberto.meseguer@viantmedical.com
Phone: 506-2239-9298, ext. 6469
Fax: n/a

A/R Contact: Walter Picado: Controller
Email: walter.picado@viantmedical.com
Phone: 506-2239-9298, ext. 6413
Fax: n/a

Additional contacts:

Director of Quality	Juan Jose Fuentes	juan.fuentes@viantmedical.com
Director of Operations	Kevin Quiros	kevin.quiros@viantmedical.com
Director of Engineering	German Wauters	german.wauters@viantmedical.com

Juridica number: 3-101-365187

Quality Management System:
FDA Registration Number: 3005173255

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 the medical device market.
We do capacity planning for product development services.
We have an ERP 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.
Our Business Excellence group leads Lean initiatives.
We monitor our facility.
Subcontractors provide services such as sterilization, component fabrication, testing labs, etc.