

# CERTIFICATE

## TUV USA, Inc.

hereby certifies that

**Fi Company Seattle**  
**3300 Airport Way South**  
**Seattle, WA 98134 USA**

has established and applies a quality system for

**Manufacturer and assembler of custom  
industrial plastic products**

Proof has been furnished that the requirements according to

**ISO 9001:2015**

are fulfilled.

Further clarifications regarding the scope of this certificate and the applicability of  
ISO 9001:2015 requirements may be obtained by consulting the organization.

**Certificate Registration No.**

**56 100 18560082**

**Initial Certification Date: October 21, 2015**

**Issue Date: October 22, 2021**

**Expiry Date: October 21, 2024**



Quality Systems Division  
215 Main Street  
Salem, NH 03079 USA



# Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 9001:2015

This is to certify that:

Southwest Fabricators S. de R.L. de C.V.  
Calle 11 Norte No. 650  
Tijuana  
Baja California  
22444  
Mexico


Holds Certificate No:

**FS 701029**

and operates a Quality Management System which complies with the requirements of ISO 9001:2015 for the following scope:

Fabrication and electronic integration of precision sheet metal enclosures (kiosks, frames, cabinets and chassis) for indoor and outdoor applications.

For and on behalf of BSI:

  
\_\_\_\_\_  
Carlos Pitanga, Chief Operating Officer Assurance – Americas

Original Registration Date: 2019-03-21

Effective Date: 2022-03-21

Latest Revision Date: 2022-02-23

Expiry Date: 2025-03-20

Page: 1 of 1



...making excellence a habit.™

# Certificate

acc. to **ISO 13485:2016**

Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes

Certificate Registration No.: **23-1602-Q**

TUV USA, Inc. hereby certifies that the quality management system of the company mentioned below is in conformance with **ISO 13485:2016** for Medical Devices-Quality Management Systems-Requirements for Regulatory Purposes.

**The Fi Company**  
**3300 Airport Way South**  
**Seattle, Washington 98134, USA**

Additional sites covered by QM System: [See Annex 1](#)

Scope:

**Manufacture of Medical Device Components Through Reaction Injection Molding, Thermoforming, and Custom Fabrication Processes**

The validity of this certification document can be obtained by contacting the TUV USA, Inc. Office.

TUV USA, Inc. (a Member of the TÜV NORD Group)

215 Main Street, Suite 1, Salem, NH 03079, USA

Tel: 001-603-870-8023, Fax: 001-603-870-8026, Email: [medical-usa@tuv-nord.com](mailto:medical-usa@tuv-nord.com)



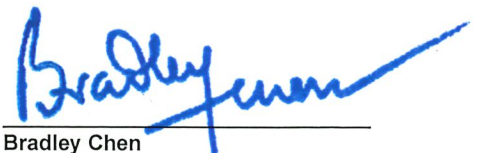
Audit Report Reference No.: **22-4002 CA**

Initial Certification Date: **2023-05-19**

Current Cycle Start Date: **2023-05-19**

Effective Date:  
**2023-05-19 / ed. 1**

Valid Until:  
**2024-05-18**



Bradley Chen  
Vice President – Medical, Americas  
Medical Products Division  
TUV USA, Inc.

**Annex 1, page 1 of 1**

*(Annex 1 MUST be displayed with the main certificate)*

**Certificate Registration No. :** 23-1602-Q / ed. 1  
**Company Name:** The Fi Company  
**Central Office Address:** 3300 Airport Way South, Seattle, Washington 98134, USA



**Additional Site(s) covered by the QM System:**

**Location**

**Scope of Certification**

**The Fi Company**  
3300 Airport Way South  
Seattle, Washington 98134, USA

Molding, Assembly, Administrative,  
Management, HR, Purchasing, Audits,  
Storage, Monitoring

**The Fi Company**  
17 South Nevada Street  
Seattle, Washington 98134, USA

Thermoforming, CNC, Assembly

---End of list---

The validity of this certification document can be obtained by contacting the TUV USA, Inc. Office.

**TUV USA, Inc. (a Member of the TÜV NORD Group)**  
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Bradley Chen  
Vice President – Medical, Americas  
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