

Certificate US18/81827348.00



The management system of

Classic Industries dba Technimark El Paso

425B Pan American Dr.
El Paso, TX 79907, United States

has been assessed and certified as meeting the requirements of

ISO 13485:2016 EN ISO 13485:2016

For the following activities:

**Manufacturer of plastic injection-molded components for active
and non-active medical devices, contract manufacture
of non-active medical device components.**

This certificate is valid from 26 March 2021 until 26 March 2024
and remains valid subject to satisfactory surveillance audits.

Re certification audit due before 15 November 2023.

Issue 4. Certified since 26 March 2018.

This is a multi-site certification.

Additional site details are listed on the subsequent page.

Authorised by

SGS United Kingdom Ltd

Rossmore Business Park, Ellesmere Port, Cheshire, CH65 3EN, UK
t +44 (0)151 350-6666 f +44 (0)151 350-6600 www.sgs.com

HC SGS 13485 2016 0118 M2

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**Classic Industries
dba Technimark El Paso**

**ISO 13485:2016
EN ISO 13485:2016**



Issue 4

Additional facilities:

420B Pan American Drive, El Paso, TX 79907, United States

**Scope: Shipping and receiving, storage and inventory control of
materials used for manufacture and packaging of medical
device products**

**Calle Cultura Jumanos # 940 Colonia Heroes de Mexico, Ciudad
Juarez, Chihuahua, 32590, Mexico**

**Scope: Shipping, receiving and warehousing of raw materials
and finished goods for the manufacturing plant, 425B Pan American
Drive site.**

420A Pan American Drive Suite 1, El Paso, TX 79907, United States

**Scope: Shipping and receiving, storage and inventory control
of materials used for manufacture and packaging of medical
device products**



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Certificate US18/81827348.01

SGS

The management system of

Technimark El Paso

420B Pan American Drive
El Paso, TX 79907, United States

has been assessed and certified as meeting the requirements of

ISO 13485:2016
EN ISO 13485:2016



For the following activities:

**Shipping and receiving, storage and inventory control of materials
used for manufacture and packaging of medical device products**

This certificate is valid from 26 March 2021 until 26 March 2024
and remains valid subject to satisfactory surveillance audits.

Re certification audit due before 15 November 2023

Issue 3. Certified since 26 March 2018

Multiple certificates have been issued for this scope

The main certificate is numbered US18/81827348.00

Authorised by

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