

EC CERTIFICATE

Full Quality Assurance System

Certificate no.:
10400-2017-CE-RGC-NA-PS

Initial certification date:
11 July 2017

Valid Until:
07 July 2023

This is to certify that the management system of
Sturdy Industrial Co., Ltd.
NO. 168, Sec. 1, Zhongxing Rd., WuGu District, 248, New Taipei City, -, Taiwan

For design, production and final product inspection/testing of:
Sterilizers /Autoclaves

has been assessed and found to comply with respect to:
**the conformity assessment procedure described in Annex II excluding
section 4 of Council Directive 93/42/EEC on Medical Devices, as
amended**

Place and date:
Høvik, 21 May 2021



For the issuing office:
**DNV Product Assurance AS - Notified Body
2460
Veritasveien 3, 1363 Høvik, Norway**

Björg Synnøve Nesgård
Operations Manager Medical

Jurisdiction

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as "Forskrift om Medisinsk Utstyr" by the Norwegian Ministry of Health and Care Services.

Certificate History			
Revision	Description	Issued Date	
0.0	Replace the certificate 39067-2008-CE-NOR 5.0 (NB 0434) following the transfer of Notified Body functions to DNV GL NEMKO Presafe AS (NB 2460) issued	2017-07-11	
1.0	Re-certification	2018-11-06	
2.0	Editorial Changes and blockchain information	2021-05-21	
Products covered by this Certificate:			
Product Description	Type	Product Name	Class
Small Sterilizers	B	SA-260MB, SA-300MB, SA-302MB	IIb
	S	SA-230MA-R, SA-260MA-R, SA-300MA-R, SA-300VMA-R	
	N	SA-230M-R, SA-260M-R, SA-300M-R, SA-300VM-R	
	Others	SA-230MA, SA-260MA, SA-300MA, SA-300VMA, SA-230M, SA-260M, SA-300M, SA-300VM	
	N	SA-230FA-R, SA-260FA-R, SA-300FA-R, SA-300VFA-R SA-202N, SA-232N, SA-252N, SA-300N	
	Others	SA-230FA, SA-260FA, SA-300FA, SA-300VFA	
	Others	SA-202, SA-230, SA-232, SA-232V, SA-202X, SA-232X, SA-300E, SA-302E,	
	Others	SA-252F, SA-300H, SA-300VF, SA-300VL, SA-300VLA	
Large Sterilizers	-----	SAT-450HP, SAT-450HPD, SAT-500HP, SAT-500HPD, SAT-600HP, SAT-600HPD	

Sites covered by this certificate	
Site Name	Site Address
Sturdy Industrial Co., Ltd.	No. 168, Sec. 1, Zhongxing Rd., Wugu District, New Taipei City 24872, Taiwan

EU Representative

APEX MEDICAL S.L. Elcano 9, 6a planta 48008 Bilbao. Vizcaya SPAIN



Terms and conditions

The certificate is subject to the following terms and conditions:

- Any producer (see 2001/95/EC for a precise definition) is liable for damage caused by a defect in his product(s), in accordance with directive 85/374/EEC, as amended, concerning liability of defective products.
- The certificate is only valid for the products and/or manufacturing premises listed above.
- The Manufacturer shall fulfil the obligations arising out of the quality system as approved and uphold it so that it remains adequate and efficient.
- The Manufacturer shall inform the Notified Body of any intended updating of the quality system and the Notified Body will assess the changes and decide if the certificate remains valid.
- Periodical audits will be held, in order to verify that the Manufacturer maintains and applies the quality system. Notified Body reserves the right, on a spot basis or based on suspicion, to pay unannounced visits.

The following may render this Certificate invalid:

- Changes in the quality system affecting production.
- Periodical audits not held within the allowed time window

Conformity declaration and marking of product

When meeting with the terms and conditions above, the producer may draw up an EC declaration of conformity and legally affix the CE mark followed by the Notified Body identification number.