

EUDAMED - European Database on Medical Devices

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Manufacturer, CN-MF-000056687, Shenzhen Ruochuang Intelligent Manufacturing Co. Ltd. - [China]

Actor information

Version 1 (Current)  Last update date: 2026-06-25

Actor identification

Actor ID/SRN	CN-MF-000056687
Role	Manufacturer
Country	China
Actor/Organisation name	Shenzhen Ruochuang Intelligent Manufacturing Co. Ltd. [EN]
Abbreviated name	-
VAT number	-
EORI	-
National trade register	-
Last confirmation date of actor data accuracy	-

Actor address

Street name	Room 802, Building 1, Jia'an Science Park, No. 2 Longchang Road, Area 67, Xing Xin'an Street, Bao'an District
Street number	-
Address line 2	-
PO box	-
City name	Shenzhen City, Guangdong
Postal Code	518101
Country	China
Latitude	-
Longitude	-

Actor contact details

Email	certification@strutt.inc
Telephone number	-

Web site		-
Person Responsible for Regulatory Compliance		
yang john		^
First name	john	
Last name	yang	
Email	certification@strutt.inc	
Telephone number	+86 18691878180	
Responsible for	-	

Street name	Room 802, Building 1, Jia'an Science Park, No. 2 Longchang Road, Area 67, Xing Xin'an Street, Bao'an District
Street number	-
Address line 2	-
PO box	-
City name	Shenzhen City, Guangdong
Postal Code	518101
Country	China
Latitude	-
Longitude	-

Authorised Representative(s) mandates

NL-AR-000000247		^
Actor ID/SRN	NL-AR-000000247 	
Role	Authorised Representative	
Country	Netherlands	
Organisation name	SUNGO Europe B.V. [NL]	
Address	Fascinatio Boulevard 522, Unit 1.7, 2909 VA, Capelle aan den IJssel	
Telephone number	-	
Email	ec.rep@sungoglobal.com	
Mandate validity dates	From 2026-06-10 To 2031-06-09	

Competent Authority

Name	Ministerie van Volksgezondheid, Welzijn en Sport
Address	P.O. Box, 20350, The Hague
Country	Netherlands

Email	medische_hulpmiddelen@minvws.nl
Telephone number	+31 70 340 79 11

Devices

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Economic Operator devices

The search for Devices and Systems/procedure Packs allows you to search and retrieve all records that contain the search terms you enter. At least one search criterion is mandatory.

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 Search criteria

Actor ID/SRN: CN-MF-000056687 × Status: On the EU market × [New search](#)

1 records found.

(Master) UDI-DI / EUDAMED ID	Version	Basic UDI-DI/ EUDAMED DI	Trade name	Risk class	Manufacturer/Producer (and Auth orised Representative) name	Actor ID/SRN	View detail
06980226800018	1 (Current)	698022680SEVS7		Class I	Shenzhen Ruochuang Intelligent M anufacturing Co. Ltd. (SUNGO Europe B.V.)	CN-MF- 000056687 (NL-AR- 000000247)	

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DECLARATION OF CONFORMITY

ACCORDING TO (EU) 2017/745 MEDICAL DEVICE REGULATION

EU Representative

Name: SUNGO Europe B.V.
Address: Fascinatio Boulevard 522, Unit 1.7,
2909VA Capelle aan den IJssel, The
Netherlands
SRN: NL-AR-000000247

Conformity Assessment

Conformity Assessment Procedure
Annex II+III of Regulation (EU) 2017/745
Applicable Standards
EN ISO 14971:2019+A11:2021
EN ISO 15223-1:2021+A1:2025
EN ISO 20417:2026、EN ISO 10993-1:2025
EN ISO 10993-5:2009+A11:2025
EN ISO 10993-10:2023
EN ISO 10993-23:2021+A1:2025
EN 60601-1-2:2015+A1:2021、EN 12184:2022
ISO 7176-1:2014、ISO 7176-2:2017
ISO 7176-3:2012、ISO 7176-4:2008
ISO 7176-5:2008、ISO 7176-6:2018
ISO 7176-7:1998、ISO 7176-8:2014
ISO 7176-9:2009、ISO 7176-10:2008
ISO 7176-11:2012、ISO 7176-13:1989
ISO 7176-14:2022/Amd1:2025
ISO 7176-15:1996、ISO 7176-21:2025
ISO 7176-22:2014、ISO 7176-31:2023
ISO 16840-10:2021/Amd1:2024

Remark

The declaration of conformity is valid in connection with the release technical document MDR-TCF-001. All the supporting documentation is retained at the premises of the manufacturer. The Declaration of Conformity is exclusively under the sole responsibility of the manufacturer.
File No.: MDR-TCF-001-08

Manufacturer

Name: Shenzhen Ruochuang Intelligent Manufacturing Co.Ltd.
Address: Room 802, Building 1, Jia'an Science Park, No. 2 Longchang Road, Area 67, Xingdong Community, Xin'an Street, Bao'an District, Shenzhen City, Guangdong, the PRC
SRN:CN-MF-000056687

Product Information

Name: Smart Everyday Vehicle
Model: Strutt ev¹c, Strutt ev¹e
EMDN: Y1299
Basic UDI-DI: 698022680SEVS7
Classification: Class I, According to Rule 13, Annex VIII, Regulation (EU) 2017/745

Declaration

We herewith declare that the above-mentioned products meet the requirements of Medical Device Regulation (EU) 2017/745 and the applicable standards above.

Signature: 
Position: Management Representative
Date: 2026-06-25
Place: Shenzhen, China