

Certification of Conformity

EU-MEDICAL DEVICES DIRECTIVE-93/42/EEC

Registration No.: ENC1910281GZ79

Applicant : Shijiazhuang DragonHawk Industry Co., Ltd.

Applicant Address : No. 1-1-3001, Tianshan Jiufeng, Jiantong Street, Yuhua District,

Shijiazhuang, Hebei, China

Product Designation : WQ Tattoo Machines

Model Number : WQ-series; Edge Cartridge: 1RL, 3RL, 5RL, 7RL, 9RL, 11RL, 14RL,

3RS, 5RS, 7RS, 9RS, 11RS, 14RS, 7M, 9M, 11M, 13M, 15M, 23M, 25M, 27M, 7RM, 9RM, 11RM, 13RM, 15RM, 23RM, 25RM

Brand Name : Dragonhawk; Edge

Classification of Medical Devices: Class I

Manufacturer : Shijiazhuang DragonHawk Industry Co., Ltd.

Manufacturer Address : No.1 Third floor East, Cang Sheng Road, Cang Xing Street

South, YuHua district, Shijiazhuang City, Hebei Province, China

The submitted products have been tested by us with the listed standard and found in compliance with the Medical Devices Directive 93/42/EEC and following European Standards:

Requirement	Applied Standards	Test Report No.
DELAYED-TYPE HYPERSENSIVITY	EN ISO 10993-1:2009	ENC1910281GZ79E1
	EN ISO 10993-10:2013	
	EN ISO 10993-12:2012	

Test result: WQ Tattoo Machines did not cause skin irritation or allergic sensitization.

This certificate is based on an evaluation of a sample of the above mentioned product. Technical report and documentation are at the licence applicant's disposal. This certificate does not imply assessment of the series-production of the product. The CE markings as shown below can be affixed on the product after preparation of necessary technical documentation.

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Ray Zhou / General Manager Date of Issue: Nov: 92/2019

East Notice Certification Service Co., Ltd.

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