CERTIFICATE

Anbotek
Product Safety

CERTIFICATE

Of Conformity
EU Council Directive 2014/30/EU
Electromagnetic Compatibility

Registration No.: AT18250EC300055

Report No.: 18250EC30005501

Applicant	:	**Shanghai Smart Industrial Co., Ltd.**		
		Lane 1118, Jinyuan 1st Road, Jiangqiao Town, Shanghai, China		
Product Identification	:	**Derma Rolling System (Derma pen/ Hydra pen /Needles)**		
	:	**Test Model No.**	:	**H2**
		Reference Model No.	:	**Dr.pen derma pen, A1, A3, A5, A6, A6S, A7, A8, A9, A10, M5, M6, M7, M8, MYM, N2, E30, X5, H2, M8S, S3, MOLE PEN & Dr.pen NEEDLE CARTRIDGE,MRF NEEDLE, V6, V7, V8, V9, V11, HR64, HN20, HN25, H24, MR16, MR18, DRS derma roller, LESCOLTON, G4, DRS140A, NEATCELL**
		Trade Mark	:	
		Rating	:	**DC 5V, 2A(Battery: 5V, 2000mA)**
Test Standards	:	**EN IEC 55014-1: 2021**		
		EN IEC 61000-3-2: 2019+A1:2021		
		EN 61000-3-3: 2013+A1:2019+A2:2021		
		EN IEC 55014-2: 2021		

The certificate of conformity is based on an evaluation of a sample of the above-mentioned product. Technical report and documentation are at the applicant's disposal. This is to certify that the tested sample is in conformity with all provisions of Annex II of Council Directive 2014/30/EU, in its latest amended version, referred to EMC Directive. The certificate does not imply assessment of the production and does not permit the use of Lab's logo. The applicant of the certificate is authorized to use this certificate in connection with EU declaration of conformity to Article 15 of the Directive.

Anbotek (SZ) Compliance Laboratory Limited
Anbotek
Approved

Certified by

KingKong Jin

Feb. 23, 2023
Date

CE **The CE Marking may only be used if all relevant and effective EU Directives are complied with** CE

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