

VERIFICATION OF CONFORMITY

No. : PTC25062613901E-EM01

Applicant : Guangzhou Ekai Electronic Technology Co., Ltd.

Address : 3/F Building E, No.81 Zijing Road, Liwan District, Guangzhou, Guangdong, China.

Manufacturer : Guangzhou Ekai Electronic Technology Co., Ltd.

Address : 3/F Building E, No.81 Zijing Road, Liwan District, Guangzhou, Guangdong, China.

Factory : Guangzhou Ekai Electronic Technology Co., Ltd.

Address : 3/F Building E, No.81 Zijing Road, Liwan District, Guangzhou, Guangdong, China.

Product : Micro-Needling System

Trade Mark : -

Model : M8S, Dr pen A6S, Dr pen A1, A6, A7, A8, A9, H2, H3, H5, M8, M7, M5, X5, E30, N2, V6, V8, V7, V9, V11, Needle Cartridge, HN20, HN25, HN64, DRS128, DRS6in1, DRS5in1, DRS4in1, DRS3in1, DRS192, DRS540, DRS600, DRS1200, DRS140A, DRS140, DRS40, ICE Roller, GTSROLLER192, MTSNeedle, A10, A11, Bio pen T6, Bio needle H24, Bio Roller G4, G5, G10, Soicy S20, S30, S40, S50, Q2, bio4, bio6, H6, H7

Ratings : 100-240V~, 50/60Hz, 0.2A

The submitted sample of the above equipment has been tested and found to comply with the following European Directive:

Electromagnetic compatibility (EMC) Directive - 2014/30/EU

The standard(s) used for showing compliance with the essential requirements:

Applicable Standard(s)	Test Report(s) Number
EN IEC 55014-1:2021 EN IEC 55014-2:2021 EN IEC 61000-3-2:2019 EN 61000-3-3:2013/A2:2021	PTC25062613901E-EM01

This verification is part of the full test report(s) and should be read in conjunction with it. This verification is based on an evaluation of one sample of above mentioned product. It does not imply assessment of the production of the product. Without the written approval of Precise Testing & Certification (Guangdong) Co., Ltd., this verification is not permitted to be reproduced, except in full. It is not permitted to use the test lab's logo. The CE marking may only be used if all the relevant and effective European Directives are applicable.



Jacky Ou

Jacky Ou
Manager
Date: 2025-07-04



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