



Certification of Conformity

EU-MEDICAL DEVICES DIRECTIVE-93/42/EEC

Registration No.: ENC2304184GZ20

Applicant : Guangzhou Kapha Electronic Technology Co.,LTD
Applicant Address : No.21-23, Hejing Road, Liwan District, Guangzhou
Product Designation : Physio Magneto
Model Number : PM-ST1, PM-ST2, PM-ST3, PM-ST4, PM-ST5, PM-ST6, PM-ST7, PM-DST, PM-ST PRO, INTELECT WAVE, PMST, PMST-1, PMST-2, PMST-3, PMST-4, PMST-5, PMST-6, PM-ST PRO, PMST NEO, PMST-1 NEO, PMST-2 NEO, PMST-3 NEO, PMST-4 NEO, PMST-5 NEO, PMST-6 NEO, PMST NEO+, PMST-1 NEO+, PMST-2 NEO+, PMST-3 NEO+, PMST-4 NEO+, PMST-5 NEO+, PMST-6 NEO+, PMST WAVE, PMST WAVE-1, PMST WAVE-2, PMST WAVE-3, PMST VET, PMST VET-1, PMST VET-2, PMST VET-3, PEMF LOOP, PEMF LOOP-1, PEMF LOOP-2, PMST PLUS, PMST PRO, PMST DUAL, PMST THERMAL, PMST LASER
Manufacturer : Guangzhou Kapha Electronic Technology Co.,LTD
Manufacturer Address : No.21-23, Hejing Road, Liwan District, Guangzhou

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:

Directive	Applied Standards	Test Report No.
EMC Directive	EN 60601-1-2:2015+A1:2021	ENC2304184GZ20E1
	EN IEC 61000-3-2:2019+A1:2021	
	EN 61000-3-3:2013+A1:2019+A1:2021	
Safety Directive	EN 60601-1:2006+A11:2011+A1:2013+A12:2014+A2:2021	ENC2304184GZ20L1

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.



Ray Zhou / General Manager

Date of Issue: Apr. 18, 2023

East Notice Certification Service Co., Ltd.

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