

# Declaration of Conformity



MANUFACTURER:

Alicn Medical(Shenzhen) Co., Ltd.  
3/F, No4 Building, Niucheng industrial Park, XiLi, Nanshan District,  
518055 Shenzhen, GuangDong Province, China

MEDICAL DEVICE:

MAGNETO THERAPY DEVICE, *POCKET EMAVIT*

ACCESSORIES:

SOLENOIDS AC0008008, MATTRESS AC0008008,  
PILLOW AC0009030, BELT AC0009031

CLASSIFICATION - ANNEX IX:

CLASS IIA, RULE 9

CONFORMITY ASSESSMENT ROUTE:

ANNEX V+ ANNEX VII

WE, THE MANUFACTURER, HEREWITH DECLARE THAT THE STATED MEDICAL DEVICES  
MEET THE TRANSPOSITION INTO NATIONAL LAW, THE PROVISIONS OF COUNCIL DIRECTIVE 93/42/EEC  
AMENDED BY 2007/47/EC CONCERNING MEDICAL DEVICES;  
ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURER.

STANDARDS APPLIED: (HARMONISED - EN) STANDARDS FOR WHICH DOCUMENTED EVIDENCE OF COMPLIANCE  
CAN BE PROVIDED.

- |                   |                     |                   |
|-------------------|---------------------|-------------------|
| (1) EN 60601-1;   | (2) EN 60601-1-2;   | (3) IEC62304;     |
| (4) ISO 10993-1;  | (5) ISO 10993-5;    | (6) ISO 10993-10; |
| (7) IEC60601-1-6; | (8) IEC 60601-1-11; |                   |

NOTIFIED BODY:

TÜV SÜD PRODUCT SERVICE GMBH  
RIDLERSTR 65, D-80339 MÜNCHEN, GERMANY

IDENTIFICATION NUMBER

**CE** 0123

(EC) CERTIFICATE(S):



EUROPEAN REPRESENTATIVE:

Renault-Petersen Limited  
Couching House, Couching Street, Watlington, Oxfordshire OX495PX

UK.

START OF CE-MARKING:

PLACE, DATE OF DECLARATION: SHENZHEN, 2018-03-20

SIGNATURE:

NAME: MEISONG FANG POSITION: (GM)

