

Declaration of conformity no. MF12/1/23/A

Manufacturer: *Elektronika i Elektromedycyna M.Lewandowski Sp.J.*
Address: ul.Zaciszna 2, 05-402 Otwock, Poland

Product: Low frequency magnetic field therapy device with accessories

Model: Magnetronic MF-12

Product class: IIa (rule 9)

Concerns devices' SN: 23001 to 23100

We herewith declare that the above mentioned product(s) meet the provisions of the EC Directive 93/42/EEC which apply to them and stated in annex I of this Directive.

The conformity assessment was performed according to annex II of the EC Directive 93/42/EEC (excluding p.4) under the supervision of Notified Body:

TÜV Rheinland LGA Products GmbH, Notified Body ID. No. 0197
Tillystraße 2, D-90431 Nürnberg, Germany,

Place, date: Otwock, Poland, 2.01.2023

Signature:

Adam Lewandowski
Co-owner

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