

EC Declaration of Conformity

Manufacturer:

Jawon Medical co., Ltd.
29, Gongdan 4-ro, Jillyang-eup, Gyeongsan-si, Gyeongsangbuk-do, 34109 Republic of Korea

whose single Authorized Representative:

VITAKO Sp. z o.o.
UL. MALEJ SYRENKI 2 71-790 SZCZECIN,
POLAND

We, the manufacturer, herewith declare that the products

Body Composition Analyzer

Model: ioi 353

UMDNS-Code: 17417

meet the provisions of Directive 93/42/EEC which apply to them.

The medical device has been assigned to class IIa (body composition analyzer) according to Annex IX of the Directive 93/42/EEC. It bears the mark



The product concerned has been manufactured under a quality management system according to Annex V of Directive 93/42/EEC, and the essential requirement of Annex I pertaining to medical devices

Compliance of the designated product with the Directive 93/42/EEC has been assessed and certified by the Notified Body

TÜV Rheinland LGA Products GmbH

Tillystraße 2

90431 Nürnberg

Country: Germany

Certificate No.: **DD 60105723 0001**

Expiry date: **25.11.2020**

following the procedure relating to the EC Declaration of Conformity set out in Annex V of Directive 93/42/EEC.

This Declaration of conformity is valid in connection with the release document for the respective batch of produced devices

The above mentioned declaration of conformity is exclusively under the responsibility of

Company: Jawon Medical co., Ltd.

Address: 29, Gongdan 4-ro, Jillyang-eup, Gyeongsan-si, Gyeongsangbuk-do, 712-837 Republic of Korea

Factory: 29, Gongdan 4-ro, Jillyang-eup, Gyeongsan-si, Gyeongsangbuk-do, 712-837 Republic of Korea

Gyungsan-si, 03, Feb, 2016

Place, date

A handwritten signature in blue ink, appearing to read 'Jin-Gyu Beak'.

Jin-Gyu Beak/QMR

Legally binding signature, Function