



690009046



Ministero della Salute

EX DIREZIONE GENERALE DEI DISPOSITIVI MEDICI E DEL SERVIZIO FARMACEUTICO
UFFICIO 3

DGDMF/3/P/I.5.1.e.1/2024/2302

HAVING REGARD to the Legislative Decree 46/97;

HAVING REGARD to the Legislative Decree n.137/2022;

HAVING REGARD to article 120 par. 3 of the EU Regulation 2017/745 related to transitional provisions on the free movement and placing on the market of medical devices marked CE pursuant to Directive 93/42/CEE;

HAVING REGARD to article 2 par.3 of the EU Regulation 2022/2346;

HAVING REGARD the declarations of conformity CE concerning the medical devices mentioned below;

HAVING REGARD the Manufacturer's Declaration in relation to Regulation (EU) 2023/607 amending Regulation (EU) 2017/745 as regards the transitional provisions for certain medical devices;

HAVING REGARD the Notified Body Confirmation Letter in confirmation of the status of a formal application, written agreement, and appropriate surveillance in the framework of Regulation (EU) 2023/607 amending Regulation (EU) 2017/745;

HAVING REGARD to the request ref.95073- 14/11/2024, submitted by the Company **VIDA S.r.l.**, with registered office in Via Galileo Galilei 4, 56040 Crespina-Lorenzana (PI), Italy; VAT N°. 02018300505;

HAVING REGARD that the Company paid the fees required by Ministerial Decree August 06, 2021;

HAVING REGARD to the official deeds;

IT IS ATTESTED

that, the Company **VIDA S.r.l.**, with registered office in Via Galileo Galilei 4, 56040 Crespina-Lorenzana (PI), Italy, is the manufacturer and has marked CE as medical devices, according to the procedures provided by the Directive 93/42/CEE pursuant to art. 120 par. 3 of EU Regulation 2017/745, the following medical devices:

SORRENTINO MEDIUM
SORRENTINO SKINBOOSTER
SORRENTINO STRONG

The above mentioned products, according to the article 4 of Directive 93/42/CEE pursuant to article 120 par. 3 of Regulation 2017/745, can freely circulate and can be placed on the market in Italy and all over the European Union.

This document has been issued in original version upon request of the in order to export medical devices to **TUNISIA**.

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It is only allowed to show or to delivery it, upon request of the customs or Health Competent Authorities of the importing country.



The Health Manager
Dott. Marco Musella