OHE-QA-REG-21049

MANUFACTURER AUTHORISATION

OMRON Healthcare Europe B.V., with its place of business at Wegalaan 73, 2132JD, Hoofddorp, the Netherlands, being the official EU representative for the Manufacturer OMRON Healthcare Co., Ltd., 53, Kunotsubo, Terado-Cho, Muko, Kyoto, 617-0002, Japan ("OMRON")

hereby declares that:

a) **CELIMED s.r.o.,** a company with its place of business at Socialni pece 3487/5A 400 11 USTI NAD LABEM, Czech Republic ("CELIMED") entered into a Distribution Agreement On 1st April 2004 ("Distribution Agreement") for the purpose of resale of certain product as identified in b) below;

b) Based on the Distribution Agreement, CELIMED is OMRON official representative and service provider in the Czech Republic of OMRON and is authorized to distribute according to Medical Device Directive 93/42/EEC the following OMRON products:

- Blood Pressure Monitors,
- Nebulizers,
- Electronic Fever Thermometers,
- Electronic Nerve Stimulators,
- Electro Cardiograph,
- Body Composition Monitors,
- Weigh Scales and Step Counters

c) Provided the Distribution Agreement between CELIMED and OMRON has not been terminated earlier in accordance with its applicable terms and conditions, in which case this Manufacturer Authorisation will automatically terminate as well, this appointment will be valid for the period from 1st January 2025 until 31st December 2025; and

d) The abovementioned appointment will automatically expire thereafter, unless OMRON agrees in writing to extend the authorization on request by CELIMED. Nevertheless, nothing contained in this Manufacturer Authorisation will barred OMRON Healthcare Europe B.V. to terminate the Distribution Agreement.

On behalf of
OMRON Healthcare Europe B.V.,

Signed by:
Karen Sanders
Signer Name: Karen Sanders
Signing Reason: I approve this document
Signing Time: December 16, 2024 | 7:50:40 AM CET
BD480D4B54464319ABAD772216332721

Karen Sanders
CEO

Date of issue: December 13th, 2024

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KVK 342.10306 **BTW** NL8132.86.281.B01