

## S.I.I.T. S.r.1. UNIPERSONALE

Innovative Healthcare Products Contract Development & Manufacturing

## **DECLARATION OF CONFORMITY CE**

The company S.I.I.T srl, located at Trezzano sul Naviglio, via Ariosto 50/60, as Manufacturer of the Medical device

## Bei Durchfall

declares under its own responsibility that the Medical device object of this declaration satisfies the dispositions of Law Decree 24 February 1997, N° 46, (Directive 93/42/CEE on Medical devices) and subsequent amendments and additions (Implementation of Directive 93/42/EEC and subsequent amendments concerning the CE marking Medical Devices)

## We declare that:

- The Medical device satisfies the requisites specified in Enclose I of Law Decree 24 February 1997, N° 46
- The Medical device is classified as Class IIa, invasive in natural orifice, of short-term use, following criteria of classification specified in Enclose IX, rule 5 - Law Decree 24 February 1997, N° 46
- Destination of use: treatment of acute and chronic diarrhea and painful symptoms related to gastrointestinal disorders
- The Medical device is commercialized in NON STERILE packaging
- The Medical device is NOT MEASURE DEVICE
- The Medical device is not to be used for CLINICAL INVESTIGATIONS
- The Medical device is commercialized with the mark CE as specified in art. 16 of Law Decree 24 February 1997, N° 46 and is manufactured according to the Quality Management System that meets the requirements of Annex V and VII to Directive 93/42/EEC and subsequent amendments, in accordance with a certificate issued by ITALCERT (ON 0426) in force at the date of this declaration

Trezzano s/N, printed on 8 April 2022

Person Responsible of Regulatory Compliance

Sara Gianellini

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