μ dialysis

DECLARATION OF CONFORMITY

Manufacturer's Name: M Dialysis AB

Manufacturer's Address: Hammarby Fabriksväg 43
SE-120 30 Stockholm
SWEDEN

SRN: SE-MF-000017466

Product: Intravenous & CSF Microdialysis Catheter

Intended purpose: The Intravenous & CSF Microdialysis Catheter (67) is intended to enable microdialysis in the peripheral venous system or of the Cerebrospinal Fluid (CSF) via an External Ventricular Drain (EVD) for up to three days.

Risk class: Class IIa (Rule 7)

Basic UDI-DI: 7332699_320_004_WC

Notified Body: Intertek Medical Notified Body AB (NB 2862)

Conformity assessment: MDR 2017/745, Annex IX excluding chapter II (only those aspects related to to the reuse of the device, including cleaning, disinfections, sterilization and related instructions)

Issued certificates: EC certificate QMS # 28620170242
EC design examination certificate #TBD
ISO 13485 certificate #0106135-01

Type designation/model:

Product Name	**Catalogue number (REF)** **Basic UDI-DI**	**Risk Class**
67 IV Microdialysis Catheter 46/10	8050090 7332699_320_004_WC	IIa
67 IV Microdialysis Catheter 46/20	8050091 7332699_320_004_WC	IIa
67 IV Microdialysis Catheter 46/30	8050092 7332699_320_004_WC	IIa
67 IV Microdialysis Catheter 130/10	8050093 7332699_320_004_WC	IIa

We hereby declare that the above mentioned device(s) comply with applicable requirements in EU Medical Device Regulation (MDR) 2017/745.

A translation of this Declaration of Conformity into another EU official language is made available upon request.

This EU Declaration of conformity is issued under the sole responsibility of the manufacturer.

Signed for and on behalf of the manufacturer M Dialysis AB

Date and place of issue

Stockholm 2024-06-10

Name, position and signature

Olof Nord, CEO