Poly Medicure Limited

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9th December 2021

EC DECLARATION OF CONFORMITY

We Poly Medicure Limited, Plot No. 104-105 & 115-116, Sector-59, HSIIDC Industrial Area, Ballabhgarh - 121004, Faridabad, INDIA.

Hereby declare and take responsibility to ensure that the following product:

NEOFLOW

S. No.	Product Description	Medical Device Class	Asept Reference codes
1	Flow Regulator without injection site	IIa	202080
2	Flow Regulator with injection site	IIa	202081
3	Flow Regulator with infusion set	IIa	202082

As per Annexure – IX of the Medical Device Directive comply with the product standards/ requirements and, meet the essential requirements according to Annexure- I of the Council Directive 93/42/EEC of 14^{th} June 1993 as amended by 2007/47/EC concerning medical devices.

Conformity Assessment Procedure was carried out according to Annexure - II excluding section 4 of the MDD and is certified by the following Notified Body.

Name, Address & No.

: TÜV SÜD Product Service GmbH, Ridlerstraße 65, 80339, Munich, Germany Notified Body Number 0123,

CE Certificate No.

: G1 041938 0007

European Authorized Representative Address

: OBELIS S.A. Boulevard Général Wahis 53, B-1030, Brussels, Belgium, mail@obelis.net

> Ramdas Sharma DGM - Quality On behalf of POLY MEDICURE LTD. Faridabad