# DECLARATION OF CONFORMITY EU

*According to the Annex IX of the directive 93/42/EEC*

We, **BOW MÉDICAL**

43 Avenue d'Italie
80000 Amiens - France
SRN: FR-MF-000011517

guarantee and declare under our sole responsibility that the medical device:

**Prescription tool DIANE**

*Class I according to the Annex IX of the directive 93/42/EEC*

Complies with rule 12 of Annex IX of the directive93/42/EEC, the standard IEC 62304 « Medical device software - *Software life cycle processes »* and meets the requirements of the article 120 « transitional provisions » of the Medical Device Regulation 2017/745

*This declaration of conformity is issued under the sole responsability of the manufacturer.*

DIANE version: 4.10

DIANE Medical Device version: 3.21

Place and date: At Amiens, France, on **10/11/2022**

Managing Director: Pierre TOUTON