EU DECLARATION OF CONFORMITY

According to Art. 19, Annex IV of Regulation (EU) 2017/745 on Medical Devices

Manufacturer:	Ningbo Ican Machines Co., Ltd.
Trademark:	icanclave
Product Name:	Steam Sterilizer
Model	STE-8-D, STE-12-D, STE-16-C,STE-18-D, STE-23-D, STE-29-D, STE-8-K, STE-12-K, STE-18-K, STE-23-K, STE-18-E, STE-23-E, STE-3-F, STE-8-T, STE-12-T, STE-18-T,STE-23-T,STE-29-T,STE-45-T, STE-8-D Pro, STE-12-D Pro, STE-18-D Pro, STE-23-D Pro,STE-29-D Pro, STE-18-T Pro,STE-23-T Pro,STE-29-T Pro, STE-45-T Pro,STE-50-T Pro
SRN:	CN-MF-000024175
European Representative:	Icanclave Europe, S.L. Avenida Juan Ramon Jimenez, 6 46930 Quart de Poblet, Valencia, Spain
SRN:	ES-AR-000018282
EMDN code:	S9099
Basic UDI-DI:	697261510STES4
Classification acc. to MDR Ax. VIII:	Class IIa, rule 16 of MDR Annex VIII
Conformity Assessment Procedure:	Annex IX, MDR (2017/745)
CE Certificate No.:	M.2024.MDR.1039
CE marking start date:	28/05/2024
Name and ID of the Notified Body:	UDEM Adriatic d.o.o. Address: Radnička cesta 54, R3 Zagreb, Croatia Number：2696

We, the manufacturer, herewith declare under our sole responsibility that the above-mentioned products meet the provisions of the Regulation (EU) 2017/745 on Medical Devices (MDR). All supporting documentations are retained under the premises of the manufacturer.

Place: Ningbo

Position/Name/Date: Lin Hui

Management Representative

21.05.2025