EU Declaration of Conformity

Manufacturer:	HANGZHOU JINLIN MEDICAL ALLPIANCES CO., LTD. M14-3-4, Hangzhou Economic &Technological Development Zone 310018 Hangzhou China
SRN:	CN-MF-000012391
European Representative:	Shanghai International Holding Corp.GmbH(Europe) Eiffestrasse 80, 20537 Hamburg Germany
Product Name:	**Manually powered suction equipments**
Models:	1311 series normal without microbial filter/ 1312 series adjustable without microbial filter/1313 series normal with microbial filter/1314 seriesadjustable with microbial filter
MDS Code:	1202
Basic UDI-DI:	69465949 MPS AS

Classification (MDR (Regulation EU MDR 2017/745) Annex Ⅷ): **Class I (rule 5, 5.1.1)**

Conformity Assessment Route: **Annexes II and III, MDR 2017/745/EU**

We herewith declare that the above mentioned products meet the transposition into national law, the provisions of the following EC Council Regulations and Standards. All supporting documentations are retained under the premises of the manufacturer. We are exclusively responsible for the Declaration of Conformity.

REGULATIONS

General applicable regulations:

EU MDR 2017/745 REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC.

Place, Date of Issue: **Hangzhou, 2021-10-28**

Signature:

Name: Fang Zhiming

Position: Management representative