

EC DECLARATION OF CONFORMITY

Name and address of the manufacturer: **Jiangsu Maslech Medical Technology Co.,Ltd.**
Building G39,The Third Period Factory Area,China Medical
City,Taizhou City 225300,Jiangsu Province,P.R.China

We declare under our sole responsibility that

the medical device: **Manual Suction Units**
Type: R/S/RS

of class: **Class Is**

according to annex IX of directive 93/42/EEC /

meets the provisions of the directive 93/42/EEC and its transpositions in national laws which apply to it.
The declaration is valid in connection with the "final inspection report" of the device. /

Conformity assessment procedure: / **Directive 93/42/EEC Annex II,Excluding Section 4**

Registration No.: **HD 60132442 0001**
Expiry Date:2023-06-08

Notified Body: / **TÜV Rheinland LGA Products GmbH**
Tillystraße 2
90431 Nürnberg
Deutschland

CE 0197

Authorized Representative:
SUNGO Europe B.V.
Olympisch Station 24,1076DE Amsterdam

Chen Ze (Quality Representative)

Name and function

Taizhou ,2018-09-18
Place, date

