EC DECLARATION OF CONFORMITY

Name and address of the manufacturer: **DONGGUAN KAISER TECHNOLOGY CO., LTD.**

No.81, Sanjiang Industrial District, 523462, Hengli Town, Dongguan, Guangdong, China

Name and address of European representative **MedNet EC-REP GmbH**

Borkstrasse 10, 48163 Muenster, Germnay

We declare under our sole responsibility that

the medical device:

Product name: ***Surgical Lavage Units***
Product code: **2500 B/C/D/E /P/Z**
2500 M DP-ST01K-CT02Z
2500 K1
2500 K3
2500 K9 DPS-ST01K-CT02K-HA250
ntm-51100L
ntm-52200L
ntm-53300L
2500K1B-DPL-ST01K-CT02K-3M
2500K1B-DPL-ST01K-CT02K
2500K1B-ARCL-ST01K-CT02K
MD PL1000B
MD PL1000BE

Of class: **IIa (sterile)**

according to Rule 9 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 V+Annex VII**

Registration No.: **DD 60137545 0001**

TÜV Rheinland LGA Products GmbH

Notified Body: /

CE 0197

Dongguan, 2020.02.14

Place, date

Shawn Lu

Name

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