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
GMDN : 47890

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**
Notified Body: /	**TÜV Rheinland LGA Products GmbH** **CE 0197**

Dongguan. 2020. 02. 14

Place , date

Shawn Lu

Name

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