EC Declaration of Conformity

Manufacturer:

Chison Medical Technologies Co.,Ltd.

No.3 Changjiang South Road,
Xinwu District, Wuxi, 214028
Jiangsu, P.R. China
No.9, Xinhuihuan Road, Xinwu District,
Wuxi, Jiangsu, China 214028

whose single Authorized Representative:

Shanghai International Holding
Corp.GmbH(Europe)
Eiffestrasse 80,20537Hamburg,Germany
DIMDI NO.:DE/0000040627

We, the manufacturer, herewith declare that the products
Ultrasound Diagnostic Systems
Model: i3,i8,Q5,Q9,QBit 5,ECO 1,ECO 2,ECO 3,ECO 5,ECO 6,ECO 3 EXPERT.
Probe: D3C60L,D7L40L,D12L40L,D7L60L,D6C12L,D7C10L,D3P64L,D6P64L,D3C20L,D5C20L, D6C15L, V4C40L,D7L50L,D5P64L,D6C15L,D3C20L,M8L40L,M8L50L,D10L30L,V4C40L, D7L40L-REC,C3-A,MC3-A ,MC5V-A,MC6-A,L7M-A,L7S-A,P3-A ,V6-A,R7-A,L7V-A.

UMDNS-Code: **15976**

meet the provisions of Directive 93/42/EEC which apply to them.
The medical device has been assigned to class IIa according to Annex II of the Directive 93/42/EEC. It bears the mark

CE 0197

The product concerned has been manufactured under a quality management system according to Annex II of Directive 93/42/EEC.
Compliance of the designated product with the Directive 93/42/EEC has been assessed and certified by the Notified Body

TÜV Rheinland LGA Products GmbH
Tillystraße 2, 90431, Nürnberg, Germany
Certificate No.: HD 60147775 0001
Issue date: 03,04.2020
Expiry date: 26.05.2024

following the procedure relating to the EC Declaration of Conformity set out in Annex II of Directive 93/42/EEC.

09.06.2022

Liu Qifei

CHISON MEDICAL TECHNOLOGIES CO.,LTD. 无锡祥生医疗科技股份有限公司 国际部 INTERNATIONAL DEPT.