

CERTIFICATE

of MDD Notification

Issued date: March 12, 2020

This certificate will be automatically void if the Notification is rejected by the EU Authorities or upon termination of the EAR.

This is certify that, According to Medical Device 93/42/EEC(MDD), we accepts the appointment to be the Authorized European Representative for products which listed in the attached agreement between below manufacturer and Luxus Lebenswelt GmbH.

Sichuan AST Medical Equipment Co., Ltd
No.58 ,Jin-Peng Road C Area, Luxian Industrial Park Luzhou City Sichuan,
CHINA 646100

The Manufacturer declares that the Medical device complies with de Directive including all essential requirements.

According to Medical Device 93/42/EEC(MDD), the European Databank on Medical Devices (EUDAMED) is established as of May 1, 2011, the German Competent Authority is notified of the manufacturer's Medical Devices and has allocated registration numbers shown in:

Disposable Medical Face Mask, UMDN code: 12-530
Medical Face Shield, UMDN code: 11-961

Where the manufacturer affix the CE marking to the product listed they must ensure that all the requirements of the appropriate EU directive(s) and standards have and continue to be met.



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