

Declaration of Conformity

we the



P.O Box # 995 Ugoki Road,
Adalat Garh,
Sialkot-Pakistan

hereby declare under their sole responsibility
that the following medical devices

Sterile disposable products
of the class I

based on the technical documentation:

Comply with the applicable requirements of the Directive 93/42/EEC as updated directive 2007/47/EC for Class I Medical Devices. The Technical file of the products have been assessed according to the procedure of Conformity Assessment described in the Annex I and Annex VII

Products: Re-Useable, Non-Powered and Dental Instruments

The declaration of conformity is valid:

From: 11.01.2021

To: 10.01.2024

GS Instruments GmbH

Industriestraße 10 a
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www.gs-instruments.de

Stamp Signature

Geschäftsführer Ali Sakhi

Steinbach, 11.01.2021

Note

comply with the regulations, in particular the essential requirements according to Annex II (without 4) of Directive 93/42 EMG.

Effective May 26, 2020, the aforementioned directive will be replaced by EU-Regulation 2017/745.

All listed products can be found in the attached Excel document with the number:

TD-GHS-1.1.1.07