

declaration of conformity



We, the manufacturer, declare in sole responsibility that the products mentioned below are in compliance with the respective regulations of the following directives.

Category	Non-automatic electronic multifunctional scales							
Products	635	645	657	665	667	675	677	685
EU-type examination certificate	D00-09-010							
Accuracy class	III							
Conformity assessment procedure for non-automatic weighing instruments	The notified body Physikalisch-Technische Bundesanstalt (0102) has verified conformity in accordance with Module D of Directive 2014/31/EU and has issued the following certificate: DE-M-AQ-PTB123							
Classification as a medical device	Class I with measuring function							
Conformity assessment procedure for medical devices	In accordance with Annex VI of Medical Devices Directive 93/42/EEC							

Directives:

- 2014/31/EU** Directive relating to the making available on the market of non-automatic weighing instruments
- 93/42/EEC** Directive concerning medical devices
- 2011/65/EU** Directive on the restriction of the use of certain hazardous substances in electrical and electronic equipment

Manufacturer: seca gmbh & co. kg
Hammer Steindamm 3-25
22089 Hamburg, Germany

Made in Germany

Notified bodies:

2014/31/EU: Physikalisch-Technische Bundesanstalt (PTB) Bundesallee 100 38116 Braunschweig, Germany Reference number: 0102	93/42/EEC: TÜV SÜD Product Service GmbH Ridlerstrasse 65 80339 München, Germany Reference number: 0123
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This declaration of conformity is valid from the date of signature until a revised declaration of conformity is issued due to modification of the above-mentioned products.

Hamburg, 10 / 08 / 2021

Frederik Vogel
CEO Development & Manufacturing

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Annex

Applied standards and specifications:

EN 45501	Metrological aspects of non-automatic weighing instruments
EN 60601-1	Medical electrical equipment – General requirements for basic safety
EN 60601-1-2	Medical electrical equipment – Electromagnetic compatibility