declaration of conformity

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

Basic UDI-DI	401203000000000000064QX	
Device Model Basic UDI-DI	ACMA005me	
Products	seca 404	seca 405
Intended purpose	The ramp is intended to enable the clinical staff to push a patient's wheelchair, bed or stretcher onto the platform of a compatible electronic platform scale.	
Classification as a medical device	I	
Conformity assessment procedure (EU) 2017/745	Art. 52 (7)	

Regulations / Directives:

(EU) 2017/745 Regulation on medical devices

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

Single Registration Number (SRN):
DE-MF-000005469

Made in Germany
Designed in Germany

CE

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, 26 July 2024

Frederik Vogel
CEO Development & Manufacturing

17-10-09-442 rev.: -

declaration of conformity

Annex

Applied standards and specifications:

EN ISO 20417	2021
EN ISO 15223-1	2021