


TECHNICAL DOCUMENTATION		
Product: Medical face mask air20 PRO		
Version 01, drafted on 22 Jan. 2021	Author: Lisa Drießen	
IT: Konformitätserklärung air20 PRO Klasse I Typ I 01[37144]_eng-GB	Page 1 of 2	

EC Declaration of Conformity

pursuant to EEC Council Directive 93/42/EEC of 14 June 1993 concerning medical devices

The product concerned is a Class I medical device.

MEDICAL face masks Type I

The device is designed and produced in compliance with Directive 93/42/EEC on the sole responsibility of:

Manufac- sauer product GmbH
turer: Frankfurter Str. 73
 D-64807 Dieburg

The complete scope of technical documentation including risk analysis is available. This declaration shall be valid until 25 May 2021.

The device meets the fundamental requirements of **Annex I** to Directive 93/42 EEC. The conformity assessment procedure has been carried out in accordance with **Annex VII** to Directive 03/42/EEC.

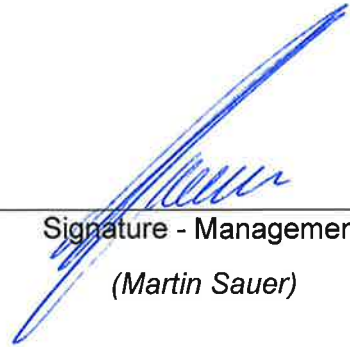
The device complies with the applicable standards which are contained in the list of standards in the available Technical Documentation.


Dieburg, 26 February 2021

Place, date

Signature - Management

(Martin Sauer)



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Product list

REF	Product designation	GMDN	UMDNS
502.827	F500 (Filter Sawascreen 8607)	37713	15-230
201.707.00.WR	Medical face mask air20 pro white/rosy	37713	15-230
201.707.00.WW	Medical face mask air20 pro white/white	37713	15-230
201.707.00.DA	Medical face mask air20 pro dark/anthracite	37713	15-230
201.707.00.DS	Medical face mask air20 pro dark/blue sea	37713	15-230
201.707.00.DB	Medical face mask air20 pro dark/burgundy	37713	15-230
201.707.00.DD	Medical face mask air20 pro dark/dark green	37713	15-230
201.707.00.DN	Medical face mask air20 pro dark/neon green	37713	15-230
201.707.00.WB	Medical face mask air20 pro dark/blue sea	37713	15-230
201.707.00.WC	Medical face mask air20 pro white/celeste	37713	15-230