

EU Declaration of Conformity

This declaration of conformity is issued under the sole responsibility of **Enraf-Nonius B.V.** Quality Management System according to EN ISO 13485:2016

Product name:	Manumed ST
Catalogue number(s):	1560000
	Delivery according to Manumed configurator 103-1560000-4A on page 2.
Basic UDI-DI:	8719992508156000038
Intended purpose:	The Manumed ST is a treatment couch, which supports a patient during treatment by a physiotherapist
Risk class:	I, according to rule 13 in Annex VIII of (EU) 2017/745
Device marking:	CE

The device covered by this declaration, in combination with its test certificate is in conformity with:

(EU) 2017/745 (Medical Device Regulation)

2006/42/EC (Machinery Directive)

2011/65/EU as amended by (EU) 2015/863 (RoHS Directive)

Any modifications to the product not authorized by Enraf-Nonius will invalidate this declaration.

Signed for and on behalf of Enraf-Nonius B.V.:

T. Doodkorte
Commercial Director

Place: Rotterdam
Date of issue: 30 April 2021

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Configuration code of Manumed ST
Appendix 103-1560000-44

T	F	XX	P	SS	G	B	W	MM	V	A	B
type	frame colour	section type	padding	upholstery colour	section width	operation	wheels	lifting mechanism	packaging	option	option

T: type

T Frame: Hydraulic
E Frame: Electric 230V/50-60Hz

F: frame colour

G grey (RAL 7035)

XX: section type

00 3 sections
01 2 sections, incl. head-section
02 2 sections, incl. back-section

P: padding

0 normal

S: upholstery colour

00 grey 013
01 sirius blue 004
02 lavender 199
03 grey 812
04 cornflower blue 169
05 red 852
09 sirius blue 204
10 grey 213
12 black 012
14 baby blue 170

G: section width

0 67 cm
1 80 cm

B: operation

X hand switch (electric)
F foot switch (electric)
J all-around (electric)
C hydraulic pump

W: wheels

Z no wheels
S small wheels

MM: lifting mechanism

44 electric motor
55 hydraulic pump

V: packaging

S packaging standard

A: options

0 no
1 paper roll holder

B: options

0 no
1 nose plug