



DECLARATION OF CONFORMITY

LINAK A/S
Smedevænget 8
DK - 6430 Nordborg

Hereby declares that LINAK Actuator

31*****0**
31*****1**
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31*****A*_*_*_*_*_*_*_*_*_*00
31*****B*_*_*_*_*_*_*_*_*_*00
31*****D*_*_*_*_*_*_*_*_*_*00
31*****E*_*_*_*_*_*_*_*_*_*00
31*****F*_*_*_*_*_*_*_*_*_*00

(The * in the product description can either be a character or a number, thereby defining the variation of the product)

complies with the EMC Directive 2014/30/EU according to following standards:
EN 55011:2016+A1+A2
EN 61000-4-2:2009, EN IEC 61000-4-3:2020, EN 61000-4-4:2012, EN 61000-4-5:2014+A1,
EN 61000-4-6:2014, EN 61000-4-8:2010, EN IEC 61000-4-11:2020

complies with RoHS2 Directive 2011/65/EU according to the standard:
EN IEC 63000:2018

Additional information:

The product does also comply with the standard:

EN 60601-1-2: 2015, Medical electrical equipment - Part 1-2: General requirements for basic safety and essential performance - Collateral standard: Electromagnetic compatibility - Requirements and tests

EMC parts of:

EN 60601-2-52:2010 Medical electrical equipment - Part 2-52: Particular requirements for basic safety and essential performance of medical beds.

Electromagnetic immunity performance is evaluated on system level.

Nordborg, 2022-07-08

LINAK A/S

John Kling, B.Sc.E.E.
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Authorized to compile the relevant technical documentation

This declaration of conformity is issued under the sole responsibility of the manufacturer.
Original Declaration