

EC Declaration of Conformity

(according to the medical device directive MDR EU 2017/745)

We,	MSP Medical Products B.V. De Sonman 7 5066 GJ Moergestel The Netherlands
Hereby declare that:	Electrically adjustable working chair Brandname "MSP Medical products"
Types :	Adam (Further referred to as working chair) With serial numbers MPADSD****** (* has a value between 0-9)

- Complies with MDR EU 2017/745 (medical device regulation) and is classified as class
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- The Working chair has been safety tested according to:
 - 🍦 EN 10535
 - 🍦 EN 12182
 - 🐓 EN 60601-1
 - Bio compatibility ISO 10993
- The Working chair is EMC tested and complies wit hall requirements in accordance with:
 - Emission EN 60601-1-2 (2007) + AC (2010)
 - Emission EN 61000-3-2 (2006) + A1 (2009) + A2 (2009)
 - Emission EN 61000-3-3 (2008)
 - Immunity EN 60601-1-2 (2007) + AC (2010)
- The internal power source is safety tested according to IEC60601-1, ANSI/AAMI ES60601-1, CAN/CSA-22.2 No 60601-1, IEC62133, UL2054.

MSP Medical Products by is not aware of any electromagnetic influences between the Working chair and other electrical products.

Technical file compiled by L.J.M.G. Dhanpat.

The Netherlands, Moergestel, 01-10-2024

L.J.M.G. Dhanpat, General manager

MSP-Medical Products B.V. Alme



Guidance and manufacturer's declaration – electromagnetic emissions

The Working chair is intended for use in the electromagnetic environment specified below. The customer or the user of the Working chair should assure that it is used in such an environment.

Emissions test	Compliance	Elektromagnetic environment – Guidance	
RF emissies CISPR11	Group 1	The working chair uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment	
RF emissies CISPR	Class B	The working chair is suitable for use in all establishments, those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.	
Harmonische emissies IEC 61000-3-2	Class A		
Voltage fluctuations Flicker emissions IEC 61000-3-3	Compliant		

Guidance and manufacturer's declaration – electromagnetic immunity

The Working chair is intended for use in the electromagnetic environment specified below. The customer or the user of the Working chair should assure that it is used in such an environment.

Immunity test	C 60601-1-2 testlevel	Compliance level	Elektromagnetic environment – guidance
Elektrostatic discharge	±6 kV contact	±6 kV contact	Floors should be wood, concrete or ceramic
(ESD)	±8 kV air	±8 kV air	tile. If floors are covered with synthetic
IEC 61000-4-2			material, the relative humidity should be at
			least 30%.
Electrical transient	±2 kV for power cables	±2 kV for power cables	Mains power quality should bet hat of a
/surges			typical commercial or hospital environment.
IEC 61000-4-4			
Overvoltage IEC	±1 kV line to line	±1 kV line to line	Mains power quality should bet hat of a
61000-4-5			typical commercial or hospital environment.
Voltage drops, short	(95% dip in Un)	(95% dip in Un)	Mains power quality should bet hat of a
power interruptions	For 0,5 cycle 40% Un	For 0,5 cycle 40% Un	typical commercial or hospital environment.
and voltage	(60% dip in Un)	(60% dip in Un)	The Working chair has an internal power
fluctuations on power	for 5 cycles 70% Un	for 5 cycles 70% Un	source.
cables	(30% dip in Un)	(30% dip in Un)	
IEC61000-4-11	for 25 cycles <5% Un	for 25 cycles <5% Un	
	(95% dip in Un)	(95% dip in Un)	
	for 5 seconds	for 5 seconds	
Mains frequency	3 A/m	3 A/m	Power frequency based magnetic fields
50/60Hz Magnetic			should be maintained at levels characteristic
field			of a typical commercial or hospital
IEC 61000-4-8			environment.