

CENTRE FOR TESTING AND CERTIFICATION - MECH-TEST **Mechanical Laboratory**

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Date 17.01.2018

TEST REPORT NO. *CBC-005/2018*

Page 1 of 8

Subject of testing:

SUPPORT BELT

Classification according to

PN-EN ISO 9999:2011: 12 30 15

Type / Model:

100 x 14 cm

117 x 14 cm 140 x 14 cm REF: 278045 278046

278047

Manufacturer:

MOBILEX A/S

Grønlandsvei 5

DK - 8660 Skanderborg

Number of specimens: 3

Applicant:

A-Net s.c.

93-469 Łódź.

ul. Łaskowice174

Kind of testing

Testing scope according to application of Client

Mechanical testing according to PN-EN 12182:2012

Test started: 2.01.2018

Test finished: 17.01.2018

Approved by:

YREKTOR mgr inż. Andrzej Tkaczyk

Special comments / enclosures:

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Test results refer only to tested units.

Test results reported here are not applicable to the further modifications of the product affecting its structure, material or technology, This test report shall be neither copied differently as in the whole nor be published without written consent of the Laboratory.

Page: 2 of 8

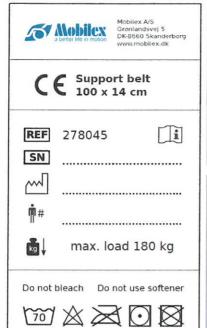
TESTING

NORMATIVE REFERENCES

PN-EN 12182:2012

Technical aids for disabled persons - General requirements and test methods

PHOTO OF PRODUCT















CHARAKTERISTICS OF PRODUCT SUPPORT BELT 278045

REF	278045	Mass of product	0,36 kg	
Maximum load	180 kg	Dimensions of product	1000 mm x 140 mm	

PHOTO OF PRODUCT



CE Support belt

REF 278046

SN

n#

max. load 180 kg

Do not bleach Do not use softener











CHARAKTERISTICS OF PRODUCT SUPPORT BELT 278046

REF	278046	Mass of product	0,44 kg	
Maximum load	180 kg	Dimensions of product	1170 mm x 140 mm	



PHOTO OF PRODUCT











CHARAKTERISTICS OF PRODUCT SUPPORT BELT 278047

REF	278047	Mass of product	0,465 kg	
Maximum load	180 kg	Dimensions of product	1400 mm x 140 mm	



Report no.: CBC-005/2018

Page: 5 of 8

TEST	RESULT	Sa	ccording	to	PN-EN	12182	:	2012	

Requirement s according to clause	Test method according to clause	Checked	l characteristics/assemblies/p	paramet	ers	Test result				
4.1	4.8, 5.2, 5.4.2, 5.5, 6, 8.2.1, 9.4 10, 22, 24 i EN 1441						N/T			
4.2	V/I	Expected char	acteristics and technical documenta	tion		Conf.	Pos.			
4.3	EN ISO 14155	Clinic assessn	nent				N/T			
4.4	V/I	Technical sup	port which can be dismantled				N/A			
4.5	V/I	Single use cor	nnections				N/A			
4.6	V/I	Boundary valu	undary values of user weight			Conf.	Pos.			
4.7	V/I	Immobilising	mmobilising means				N/A			
	V/I, C5	Suitability of	the product for people with cognitiv	e impairm	ent		N/T			
4.8		documentation	e presence of the description in the manufacturer's cumentation aterials				N/T			
F 4	EN 60601-1-9						AT/T			
5.1	V/I, B 5.2	Flammability					N/T			
5.2			ante montheres and head heave and head	•			N/A			
5.2.2	V/I		arts, mattresses, bed bases and bedd	ing			N/A			
5.2.3	V/I, EN 1021						N/A			
5.2.4	V/I, EN 597						N/A			
5.2.5	V/I. EN ISO 12952	Bedding 952					N/A			
5.2.6	V/I. EN 60695-11-1	Moulded parts	Moulded parts				N/A			
5.3	EN ISO 10993-1 Annex. D	Biological cor	formity and toxicity				N/T			
5.4	V/I	Contaminants	and residues				N/A			
	V/I.,B.5.5.1	7 5	Cleaning				N/T			
	V/I.,B.5.5.1	iolo- li ss an natio	Disinfection				N/A			
5.5	V/I., EN ISO 22442-1 B.5.5.2	Microbiolo- gical infections and contamination	Animal tissue				N/A			
5.6	EN ISO 9227	Resistance to	corrosion				N/T	NOTE 1		
6		Emitted soun	d and vibration							
6.1	EN ISO 3746	Noise and vib	ration				N/A			
6.2	B6 EN ISO 3746	Sound levels a	nd frequencies of audible warning d	levices			N/A			
Require ments according to clause	Test method according to clause	characteris	Checked tics/assemblies/parameters	Real	Test result		N/A Comments			
6.3	EN ISO 3746	Feeedback			N/A					
7	EN 60601-1-2 7.2, 7.3, 7.4	Electromagnetic	compatibility		N/A					
8		Electrical safety			N/A					
9	V/I		ge, leakage, and ingress of liquids		N/A					
10	V/I. Measur.	Surface temperat	ture		N/A	t ⁰ ≤ 41 ⁰ C ■ requirement does not concern heat direct solar radiation - PN-EN 12182,clause 10a ■ requirement concerns only persons w insensitiveness of skin (who do not fee heat) - PN-EN 12182,clause 10d				
11	V/I	Sterility		+	N/A	neut	, - 1 IV-15IV I	2102, Chuise 100		



Mech	ianical	Labo	ratory of CBC		керо	rt no.: C		05/2018 e: <i>6 of a</i>	
Require ments accordi ng to clause	Test m accord clau	ling to	Checked characteristics/assemblies/parameters	Real value	Test result			ments	
12	V/I. M	easur.	Safety of moving parts		N/T				
13	V/I. M	easur.	Prevention of traps for parts of the human body		N/T				
14	V	/I	Folding and adjusting mechanisms		N/T				
15	V/I. M	easur.	Carrying handles		N/T			16:	
16	V/I. M	easur.	Assistive products which support or suspend users	Conf.	Pos.		NO.	TE 2	
17	V/I. M	easur.	Portable and mobile assistive products		N/A				
18	V/I,	B 18	Surfaces, corners, edges and protruding parts	Conf.	Pos.				
19	В	19	Hand held assistive products	 	N/A				
20	В	20	Small Parts	Conf.	Pos.				
21	V/I. M		Stability		N/A				
	EN 60								
22	B 22,		Forces in soft tissues of the human body		N/A	TI :			1 1 .
23	V/ EN 6		Ergonomic principles		N/T	The requir	rements i pro		ne aesi
Require ments accordi ng to clause	Test method accordi ng to clause		Checked characteristics/assemblies/	/paramet	ers		Real valu	Test result	Con
24	V/I	Requi	rements for information supplied by the manufacti	ırer					
24.1		Gener						N/T	
24.2		Instru	ctions for use					N/T	
24.2.1	V/I Pre-sale Information								
		pe	formation on how to obtain the user information in a cople with visual,r eading orcognitive disabilities		ropriate f	for use by		N/T	
			l information shall as far as possible be available in Pic					N/T	
			description of the intended use and the intended environaintenance instructions, if applicable;	nment;				N/T	
		e) if	an assistive product is intended to be cleaned, a dead suitable cleaning materials, including precautions replicable;					N/T N/T	
	f) if		an assistive product is intended to be disinfected, a ditable materials, including any precautions needed to a	avoid corro	sion, if ap	plicable;		N/T	
		m	e overall dimensions (width, length and height) of the a illimetres, and its mass, expressed in kilograms, when oplicable, when it is folded or dismantled					N/T	
	V/I	an	e mass expressed in kilograms if the assistive productly removable parts that has a mass which is heavier that	an 10 kg;				N/T	
24.2.1		m	the assistive product is supposed to be used in combin anufacturer shall state to which products, and how this	can be don	e in a saf	e way;		N/T	
		of fla	arning about dangerous combinations of devices (e.g. decubitus ulcers often only work on correct seat surame resistant and non-flame resistant material;	face) and	combinat	ions of		N/T	
		de	list of accessories, detachable parts and materials the stermined as being intended for use with the assistive	product		r has		N/T	
		pr	a programmable controller is fitted, information or ogramming, the competence required to carry out the fects on performance			the		N/T	
		m) op	perator control adjustments					N/T	
		1000	nether and how the assistive product can be folded or or transport	lismantled	to assist i	n storage		N/T	
			structions regarding transport of the assistive product (e.g. in a ca	r or aerop	olane)		N/T	
		p) me	easured sound power level					N/T	



Require ments	Test method			e: 7 of 8	
accordi ng to clause	accordi ng to clause	Checked characteristics/assemblies/parameters	Real valu e	Test result	Com
24.2.2	V/I	User information			
		User information shall be provided by the manufacturer with each assistive product. Information shall contain all pre-sale warnings and informations and the following as applicable for each assistive product:		N/T	
		a) the location and the type of identification number/word on the assistive product shall be given for the unique identification number of the assistive product		N/T	
		b) the intended user		N/T	
		c) any adjustment or settings required before the assistive product can be used and information on how adjustments or settings affect the assistive product		N/T	
		d) information on adjustment possibilities and the competence required to carry out these adjustments		N/T	
		e) instructions on operation of all controls		N/T	
		f) the battery type and nominal vottage		N/T	
		g) instructions for battery maintenance		N/T	
		h) instructions for operating the battery charger, including warnings regarding any			
		potential safety hazards (e.g. a possibility of gas accumulating in the charging area);		N/T	
		 i) instructions on dismantting and re-assembly of the assistive product or any removable parts; 		N/T	
		 the positions of points where the component parts can be gripped for safe moving and handling and/or a method for handling during dismantling, assembly or carrying; 		N/T	
		k) a warning if surface temperatures can increase / decrease when exposed to external sources of heat or cold (e.g. sunlight, outdoor environment);		N/T	
		a warning if the assistive product might disturb the operation of devices in its environment that emit electromagnetic fields (e.g. alarm systems of shops, automatic doors, etc.);		N/T	
		m) a warning if the performance of the assistive product can be influenced by electromagnetic fields {e.g. those emitted by portable telephones, electricity generators or high power sources);		N/T	
		n) if the intended purpose of an assistive product cannot be met without a hazard {e.g, holes, V-shaped opening), a warning and instructions on howto operatethe assistive product safely;		N/T	
		 if the intended purpose of an assistive product cannot be met without a hazard due to moving parts such as squeezing, a warning and instructions on how to operate the assistive product safely; 		N/T	
		p) the level of resistance to ignition of materials and assemblies;		N/T	
24.2.2	V/I	q) information on the recycling of used batteries and other parts of the assistive product;		N/T	
		r) expected lifetime of the assistive product.		N/T	
		- It is recommended to include instructions on how to sotve simple problems for the ease of use.		N/T	
24.2.3	V/I	Service information			
		The service information shall contain all the pre-sale information, user information and instructions necessary for the maintenance, adjustment and repair of the assistive product and for the replacement of parts.		N/T	
		and for the replacement of parts. The service information shall contain all the pre-sale information and the user information.		N/T	
		The service information shall be sufficiently detailed concerning preventive inspection,		N/T	
		maintenance and calibration, including the frequency of such maintenance. The service information shall provide information for the safe performance of such		N/T	
		routine maintenance necessary to ensure the continued safe use of the assistive product. Additionalty, the service information shall identify the parts on which preventive inspection and maintenance shall be performed by service personnel, including the periods to be applied and details about the actual performance of such maintenance.		N/T	



Mech	nanical	Laboratory of CBC Report no.: C	BC-005/2018			
			Page	}		
Require ments accordi ng to clause	Test method accordi ng to clause	Checked characteristics/assemblies/parameters	Real valu e	Test result	Comments	
24.3	V/I	Labelling		N/T		
		- year of production for the product		N/T		
		- Detachabfe parts of an assistive product with a mass of more than 10 kilograms shall be marked with the actual mass on the part.		N/T		
		- Symbole for use in the labelling of medical devices shall be in accordance with EN 980		N/T		
25	V/I	Packaging		N/T	Note 3	

Pos. – positive; Neg – negative; N/T – not tested; N/A – not applicable; N/R – not required , N/O – not occurred , V/I. – visual inspection, Conf. - conformed.

- *NOTE 1: The risk of corrosion that affects the safety of the users should be evaluated in the risk analysis.*
- NOTE2: Handles are loaded with 210 kg

Test result positive.

After all the tests, the products operate according to the manufacturer's notes.

• The buckle was loaded with a force of 90 kg. Test result positive.

After all the tests, the products operate according to the manufacturer's notes.

- NOTE 3: Assessment of package, clause 25 concerns risk of threats caused by improper protection against damage, fall or impurity during storage and transport to place of use
- NOTE 4: Conformity assessment of product according to standard requirements refer to the scope of mechanical tests ordered by client
- NOTE 5: During visual inspection before testing any visible defects that can have an effect on test results were not stated.
- NOTE 6: Sample/object for testing was delivered to the Laboratory by the Orderer.

CONCLUSIONS:

Testing object conforming with requirements of PN-EN 12182:2012

END -

