

ENTRE FOR TESTING AND CERTIFICATION - MECH-TEST

Mechanical Laboratory

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Date 29.11.2016

TEST REPORT NO. *CBC-156/2016*

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Subject of testing:

Slide Sheet, Transfer gloves

Classification according to

PN-EN ISO 9999:2011: 12 30

Type / Model:

Tube slide sheet

REF.: 278011, 278012, 278013

Flat slide sheet with handles

278023 278031

One-way slide sheet Transfer gloves

278041

Manufacturer:

MOBILEX A/S

Grønlandsvej 5

DK - 8660 Skanderborg

Number of specimens: 6

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: 11.10.2016

Test finished: 29.11.2016

Approved by:

YREKTOR

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.

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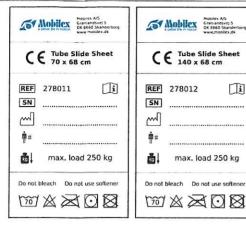
TESTING

NORMATIVE REFERENCES

PN-EN 12182:2012

Technical aids for disabled persons - General requirements and test methods

PHOTO OF PRODUCT



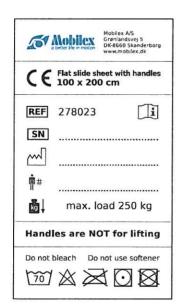




CHARAKTERISTICS OF PRODUCT Tube Slide Sheet

ArtNr.	278011, 278012, 278013	Dimensions of product	70 cm x 68 cm
Maximum load	250 kg		140 cm x 68 cm
Thickness of the fabric	0,084 mm		200 cm x 68 cm

PHOTO OF PRODUCT



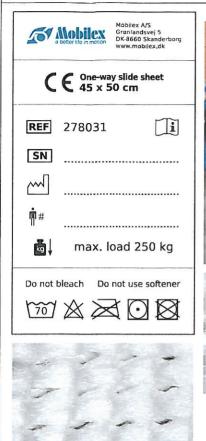


CHARAKTERISTICS OF PRODUCT Flat slide sheet with handles

ArtNr.	278023	Dimensions of product	100 cm x 200 cm
Maximum load	250 kg	Thickness of the fabric	0,084 mm



PHOTO OF PRODUCT











CHARAKTERISTICS OF PRODUCT One-way slide sheet

ArtNr.	278031	Dimensions of product	45 cm x 50 cm
Maximum load	250 kg		

PHOTO OF PRODUCT







CHARAKTERISTICS OF PRODUCT Transfer gloves

ArtNr.	278041	Dimensions of product	50 cm x 20 cm
Maximum load		Thickness of the fabric	0,084 mm

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TEST RESULTS according to PN-EN 12182: 2012

		12011	ESULTS according to r	11 1211	12102	. 2012				
Requirement s according to clause	Test method according to clause	Checked	l characteristics/assemblies/	paramet	ers	Test 50 Comme comme				
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	ation		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 con	nnections				N/A			
4.6	V/I	Boundary val	oundary values of user weight Conf. Pos.							
4.7	V/I	Immobilising	means				N/A			
	V/I, C5	Suitability of	the product for people with cognitiv	ve impairm	ent		N/T			
4.8	***	documentation	of the description in the manufacture	ırer's			N/T			
	EN (0(01 1 0	Materials				- M. A. A				
5.1	EN 60601-1-9	, ,					N/T			
5.2	V/I, B 5.2	Flammability					N/A			
5.2.2	V/I	•	arts, mattresses, bed bases and bed	ding ————			N/A			
5.2.3	V/I, EN 1021						N/A			
5.2.4	V/I, EN 597		Mattresses and bed bases				N/A			
5.2.5	V/I. EN ISO 12952						N/A			
5.2.6	V/I. EN 60695-11-1	Moulded parts	6				N/A			
5.3	EN ISO 10993-1 Annex. D		nformity and toxicity				N/T	Control of the passed to be talked the control of t		
5.4	V/I	Contaminants	and residues	111111111111111111111111111111111111111			N/A			
	V/I.,B.5.5.1	75	Cleaning				N/T			
	V/I.,B.5.5.1	iolo- ial is an natio	Disinfection				N/A	A 100 - 100		
5.5	V/I., EN ISO 22442-1 B.5.5.2	Microbiological 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 B6	Noise and vib	ration				N/A			
6.2	EN ISO 3746	Sound levels a	and frequencies of audible warning	devices			N/A			
Require ments accordi ng to clause	Test method according to clause	characteris	Checked tics/assemblies/parameters	Real value	Test result		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	Overflow, spilla	ge, leakage, and ingress of liquids		N/A					
10	V/I. Measur.	Surface tempera	ture		N/A	t ⁰ ≤ 41°C ■ requirement does not concern heat of direct solar radiation - PN-EN 12182,clause 10a ■ requirement concerns only persons with insensitiveness of skin (who do not feel heat) - PN-EN 12182,clause 10d				
11	V/I	Sterility			N/A					



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Require ments accordi ng to clause	Test m accord clai	ling to	Checked characteristics/assemblies/parameters	Real value	Test result		Page: 5 of 8 Comments		<u> </u>
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	V/I. M	easur.	Assistive products which support or suspend users	Conf.	Pos.	1-	NO	TE 2	
17	V/I. M		Portable and mobile assistive products		N/A				
18	V/I,	B 18	Surfaces, corners, edges and protruding parts	Conf.	Pos.				
19	В		Hand held assistive products		N/A				
20	В	0.119/1	Small Parts	Conf.	Pos.				-
	V/I. M	× 2000	Stability						
21	EN 60	601-1			N/A				
22	B 22.		Forces in soft tissues of the human body		N/A				
23	V/ EN 6		Ergonomic principles		N/T	The requir	irements relate to the desi process		
Require ments accordi ng to clause	Test method accordi ng to clause		Checked characteristics/assemblies/	/paramet	ters		Real valu e	Test result	Cor
24	V/I	Requi	rements for information supplied by the manufact	urer					
24.1		Gener			74			N/T	
24.2		Instru	actions for use					N/T	
24.2.1	V/I		le Information						
	c) d) e)		formation on how to obtain the user information in a cople with visual,r eading orcognitive disabilities	format app	ropriate 1	for use by		N/T	
			l information shall as far as possible be available in Pic	togram				N/T	
			description of the intended use and the intended enviro	nment;				N/T	
			aintenance instructions, if applicable;					N/T	
		ar	an assistive product is intended to be cleaned, a de- nd suitable cleaning materials, including precautions op- pplicable;					N/T	
		f) if	an assistive product is intended to be disinfected, a ditable materials, including any precautions needed to a					N/T	
		g) th 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	assistive pro	oduct, exp	oressed in		N/T	
	V/I	53	e mass expressed in kilograms if the assistive product by removable parts that has a mass which is heavier the		smantled	or has		N/T	
24.2.1		i) if	the assistive product is supposed to be used in combin anufacturer shall state to which products, and how this	ation with		250		N/T	
		j) w of	arning about dangerous combinations of devices (e.g. decubitus ulcers often only work on correct seat sur	. cushions	for the pr	evention		N/T	
		k) a	ame resistant and non-flame resistant material; list of accessories, detachable parts and materials the etermined as being intended for use with the assistive		nufacture	er has		N/T	
		l) if pr	a programmable controller is fitted, information of cogramming, the competence required to carry out the fects on performance	n the met		the		N/T	
			perator control adjustments					N/T	
		10000	hether and how the assistive product can be folded or or transport	dismantled	to assist i	in storage		N/T	
		o) in	structions regarding transport of the assistive product ((e.g. in a ca	ır or aeroj	olane)		N/T	
		p) m	easured sound power level					N/T	



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Require ments accordi ng to clause	Test method 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 c) any adjustment or settings required before the assistive product can be used and		N/T N/T		
		d) information on how adjustments or settings affect the assistive product 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 nominał 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); i) instructions on dismantting and re-assembly of the assistive product or any		N/T		
		removable parts; j) the positions of points where the component parts can be gripped for safe moving		N/T		
		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		
		 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.	22	N/T		
		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, maintenance and calibration, including the frequency of such maintenance.		N/T		
		The service information shall provide information for the safe performance of such routine maintenance necessary to ensure the continued safe use of the assistive product.		N/T		
		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		



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Require ments accordi ng to clause	Test method accordi ng to clause	Checked characteristics/assemblies/parameters	Real valu e	Test result	Com ments		
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	1	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: • For the strength test with static load and fatigue load, it was used a probe with dimensions Ø355mm

/ R = 800mm shown in Fig. 201 104 in the standard PN- EN 60601-2-52: 2010 Medical electrical equipment - Part 2-52: Particular Requirements for basic safety and essential performance of medical beds.

- For devices REF 278011, 278012, 278013, 278023, 278041, the probe shown in Fig. 201.104 was loaded with the mass of 1000N (fig. 3). When the horizontal force of 80N was applied, the probe started to move in the horizontal direction.
- For devices REF 278011, 278012, 278013, 278023, the probe shown in Fig. 201.104 was loaded with the mass of 2500N (fig. 4). When the horizontal force of 370N was applied. The probe started to move in the horizontal direction.
- For devices REF 278011, 278012, 278013, 278023, 278041, the probe shown in Fig. 201.104 was loaded with the mass of 1000N. At the moment 4,5Nm probe began to rotate around a vertical axis.
- For devices REF 278011, 278012, 278013, 278023, 278041, the probe shown in Fig. 201.104 was loaded with the mass of 1000N. The probe of the fabric was moved forward and backward 10 000 times by 250mm. After the test, the products operate according to the manufacturer's notes.
- For the product REF 278031, the probe shown in Fig. 201.104 was loaded with the mass of 1000N. When the strength of 340N was applied in the horizontal direction to the front, the probe started to move toward horizontal direction. When the horizontal force of 250N was applied backward, the probe started to move toward horizontal direction.

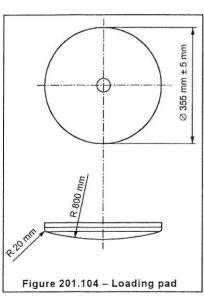




Fig. 201.104a





Fig. 3



Fig. 4





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• For devices REF 278023, the probe shown in Fig. 201.104 was loaded with the mass of 2500N (fig. 4). When the horizontal force of 370N was applied. The probe started to move in the horizontal direction.

The force appled to the ears $(2 \times 1250N)$.

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 -

