

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

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		Date 20.05.2013
	ΤΕΩΤ ΝΕΝΔΝΤΑΙ	CPC 100/2012
	ILSI KEPURT NO	). <b>CBC-100/2013</b>
		Page 1 of 6
Subject of testing:	Ramp	Classification according to PN-EN ISO 9999:2011 : 18 30 15
Гуре / Model:	TM 300	Factory ref. no.:
Manufacturer:	MOBILEX A/S Nørskovvej 1 DK – 8660 Skanderborg	Number of specimens: 1
Applicant:	A-Net s.c. 93-469 Łódź, ul. Łaskowice174	
Kind of testing	Testing scope according to app Mechanical testing according to	
Fest started: 10.(	05.2013	
Test finished: 20.0	05.2013	
		Approved by:
	Б. -	DYREKTOR mgr inż. Andrzej Tkaczyk
Special comments /	enclosures:	
Test results refer only to Test results reported here	re are not applicable to the further modifications	o report form) s of the product affecting its structure, material or technology. e published without written consent of the Laboratory.







	anical Labo	ratory of CB			Rej	ort no.		100/2013 3 of 6	
		TEST R	ESULTS according to I	PN-EN	2182	: 2012			
Requirement s according to clause	Test method according to clause	Checked	l characteristics/assemblies/	/paramet	ers	Test result	Opinion	Comments	
4.1	4.8, 5.2, 5.4.2, 5.5, 6, 8.2.1, 9.4, 10, 22, 24 i EN 1441					<del>a d</del>	N/T		
4.2	V/I	and the second s	racteristics and technical document	ation		Conf.	Pos.		
4.3	EN ISO 14155	Clinic assessm	nent				N/T		
4.4	V/I	Technical sup	port which can be dismantled			Conf.	Pos.		
4.5	V/I	Single use con	nnections			Conf.	Pos.		
4.6	<b>V/I</b>	Boundary val	ues of user weight	197 - 197 - M		Conf.	Pos.		
4.7	V/I	Immobilising	means	alityeeneri i taalige vi			N/A		
	V/I, C5	Suitability of	the product for people with cogniti	ve impairm	ent		N/T		
4.8		documentatio	of the description in the manufactun	urer's			N/T		
		Materials							
5.1	EN 60601-1-9	8					N/T		
5.2	V/I, B 5.2	Flammability					N/A		
5.2.2	V/I		parts, mattresses, bed bases and bed	ding			N/A		
5.2.3	V/I, EN 1021	1 1	Upholstered parts				N/A		
5.2.4	V/I, EN 597		Mattresses and bed bases				N/A		
5.2.5	V/I. EN ISO 12952	Bedding	Bedding				N/A		
5.2.6	V/I.	Moulded part	Moulded parts						
	EN 60695-11-1 EN ISO 10993-1		aformity and toxicity				N/A		
5.3	Annex. D	Diological con	Biological conformity and toxicity				<i>N/T</i>		
5.4	V/I		Contaminants and residues				N/A		
ŀ	V/I.,B.5.5.1	gica and ion	Cleaning			Conf.	Pos.		
5.5	V/I.,B.5.5.1	niolo, ions ninat	Disinfection			N/A			
	V/I., EN ISO 22442-1 B.5.5.2	Microbiologica 1 infections and contamination	Animal tissue				N/A		
5.6	EN ISO 9227	Resistance to	corrosion				N/T		
6			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		Checked stics/assemblies/parameters	Real value	Test result	N/A Comments		iments	
6.3	EN ISO 3746	Feeedback			N/A			6.1. 6. 520a.v.b. 7	
7	EN 60601-1-2 7.2, 7.3, 7.4	Electromagnetic	compatibility		N/A				
8		Electrical safety			N/A			105	
	V/I	Overflow, spilla	ge, leakage, and ingress of liquids		N/A				
9		Surface tempera	ture		N/A	<ul> <li>ℓ ≤ 41°C</li> <li>requirement does not concern heat of direct solar radiation - PN-EN 12182, clause 10a</li> <li>requirement concerns only persons wi insensitiveness of skin (who do not fee)</li> </ul>			
9 10	V/I. Measur.	2				dir require	ect solar ra 12182,c ement conce	diation - PN-EN clause 10a erns only persons with	



Require ments accordi ng to clause	Test m accord clau	ing to	Checked characteristics/assemblies/parameters	Real value	Test result	Page: 4 of 6 Comments			
12	V/I. M	easur.	Safety of moving parts	Conf.	Pos.				
13	V/I. M	easur.	Prevention of traps for parts of the human body	Conf.	Pos.				
14	V	/1	Folding and adjusting mechanisms	Conf.	Pos.	Not	e in ser	vice manu	al
15	V/I. M		Carrying handles	Conf.	Pos.		loading	ct below 16,96kg	
16	V/I. M	easur.	Assistive products which support or suspend users	Conf.*)	Pos.	loading 187,5 kg, 70sek./ 1pi 1000 cycles, 125kg / 1 piec			
17	V/I. M	easur.	Portable and mobile assistive products		N/A	2 - 500 GE 441			
18	<b>V/I,</b> ]	B 18	Surfaces, corners, edges and protruding parts	Conf.	Pos.				455 FM 11
19	<b>B</b> 1	19	Hand held assistive products		N/A			- K	
20	B 2	20	Small Parts	Conf.	Pos.				
21	V/I. M EN 60	601-1	Stability		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		relate to t. cess	he desig
*) Elasti			2mm, sustained deformation – 1mm				pro		
Require ments accordi ng to clause	Test method accordi ng to clause		Checked characteristics/assemblies	/paramet	ers		Real valu e	Test result	Com ment
24	V/I	Requi	rements for information supplied by the manufact	urer					
24.1		Gener	al			1.0779-07-07-07 H		N/T	
24.2 24.2.1	V/I	A second s	ictions for use le Information					<i>N/T</i>	
	<ul> <li>a) information on how to obtain the user information in a format appeople with visual, r eading orcognitive disabilities</li> <li>b) all information shall as far as possible be available in Pictogram</li> <li>c) a description of the intended use and the intended environment;</li> <li>d) maintenance instructions, if applicable;</li> <li>e) if an assistive product is intended to be cleaned, a description and suitable cleaning materials, including precautions needed to applicable;</li> <li>f) if an assistive product is intended to be disinfected, a description</li> </ul>				of the m	ethod osion, if		N/T           N/T           N/T           N/T           N/T	
	<ul> <li>suitable materials, including any precautions needed to avoid corrosion, if applicable;</li> <li>g) the overall dimensions (width, length and height) of the assistive product, expressed in millimetres, and its mass, expressed in kilograms, when it is ready for use and, if applicable, when it is folded or dismantled</li> </ul>							N/T N/T	
24.2.1	<ul> <li>V/I h) the mass expressed in kilograms if the assistive product can be dismantled or has any removable parts that has a mass which is heavier than 10 kg;</li> </ul>							N/T	
27.2.1		m	the assistive product is supposed to be used in combin anufacturer shall state to which products, and how this arning about dangerous combinations of devices (e.g.	s can be don	e in a saf	e way;		N/T	
	of decubitus ulcers often only work on correct seat surface) and combinations of flame resistant and non-flame resistant material;							<i>N/T</i>	
1	<ul> <li>k) a list of accessories, detachable parts and materials that the manufacturer has determined as being intended for use with the assistive product</li> <li>if a programmable controller is fitted, information on the method of</li> </ul>							<i>N/T</i>	
		programming, the competence required to carry out the programming and the effects on performance						N/T	
		pi ef	fects on performance	he program					
		pr ef m) or	fects on performance perator control adjustments		-			N/T	
		m) op n) w	fects on performance	dismantled	to assist	in storage			



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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
24.2.2	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	
		<ul> <li>d) information on adjustment possibilities and the competence required to carry out these adjustments</li> </ul>		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	
		<ul> <li>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);</li> </ul>		N/T	
		<ul> <li>instructions on dismantting and re-assembly of the assistive product or any removable parts;</li> </ul>		N/T	
		j) 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	
		<ul> <li>k) a warning if surface temperatures can increase / decrease when exposed to external sources of heat or cold (e.g. sunlight, outdoor environment);</li> </ul>		N/T	
		<ol> <li>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.);</li> </ol>		N/T	
		<ul> <li>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);</li> </ul>		N/T	
		<ul> <li>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;</li> </ul>		N/T	
		<ul> <li>o) 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;</li> </ul>		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 solve 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		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.			
		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.		<i>N/T</i>	
		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	
			20 a		



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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		N/T	Note 1
$\partial \alpha = nc$	anitima:	Neg pagative: N/T not togtad: N/A not applicable: N/D not applied N/O not applied N/O	a - 20 000 <b>1</b>	1	L

NOTE 1: 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 2: Conformity assessment of product according to standard requirements refer to the scope of mechanical tests ordered by client

NOTE 3 : During visual inspection before testing any visible defects that can have an effect on test results were not stated.

NOTE 4: Sample/object for testing was delivered to the Laboratory by the Orderer.

## **CONCLUSIONS:**

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

- END –

