

CENTRE FOR TESTING AND CERTIFICATION - MECH-TEST

Mechanical Laboratory

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Date 28.05.2021

TEST REPORT NO. *CBC-111/2021*

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Subject of testing:

Ramp

Classification according to

PN-EN ISO 9999:2017-02: 18 30 15

Type / Model:

Rubberramp, grev.

without alue

SN .: --

Item nr.: 306482

Number of specimens: 1.

Manufacturer:

MOBIL FX A/S

Grønlandsvei 5

DK-8660 Skanderborg

Applicant:

MOBILEX A/S

Grønlandsvei 5

DK-8660 Skanderborg

Kind of testing

Testing scope according to application of Client

Mechanical testing according to PN-EN 12182:2012:

Test started: 21.05.2021

Test finished: 28.05.2021

Approved by:

DYREKTOR 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.



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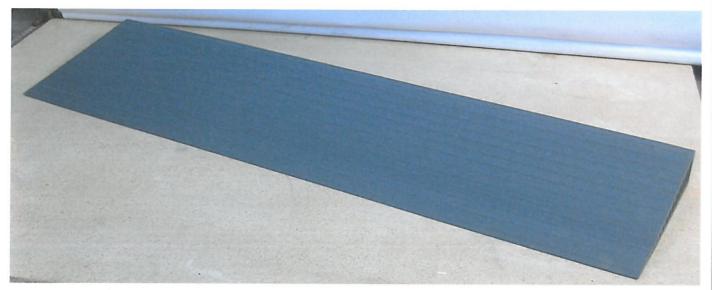
TESTING

NORMATIVE REFERENCES

PN-EN 12182:2012

Technical aids for disabled persons - General requirements and test methods

PHOTO OF PRODUCT





C€MOBILEX A/S - WWW.MOBILEX.DK - 48MM - PROD. 15-03-21

CHARAKTERISTICS OF PRODUCT

Name of product: Rubberramp, grey, without glue

SN: --

Max. loading: 350kg

Mass of product: **4,1 kg**

Dimensions:

1000 mm x 250 mm x 48 mm

Material:

rubber



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TEST RESULT	S according to	PN-EN	12182:2012
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Requirement s according to clause	Test according of the characteristics assemblies parameters Checked characteristics assemblies parameters					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	1, 9.4, i EN							
4.2	V/I	Expected char	acteristics and technical documenta	tion		Conf.	Pos.		
4.3	EN ISO 14155	Clinic assessm	nent				N/T		
4.4	V/I	Technical sup	port which can be dismantled				N/A		
4.5	V/I	Single use con	nections				N/A		
4.6	V/I	Boundary valu	es of user weight	The second second		Conf.	Pos.		
4.7	V/I	Immobilising	means				N/A		
	V/I, C5	Suitability of t	he product for people with cognitiv	e impairm	ent		N/T	***************************************	
4.8		documentation	of the description in the manufactur	er's			N/T		
		Materials							
5.1	EN 60601-1-9	Recycling					N/T		
5.2	V/I, B 5.2	Flammability	Flammability					A CONTRACTOR OF THE CONTRACTOR	
5.2.2	V/I	Upholstered p	Upholstered parts, mattresses, bed bases and bedding				N/A		
5.2.3	V/I, EN 1021	Upholstered pa	Upholstered parts				N/A		
5.2.4	V/I, EN 597	Mattresses and	Mattresses and bed bases				N/A		
5.2.5	V/I. EN ISO 12952	Bedding	Bedding				N/A		
5.2.6	V/I. EN 60695-11-10	Moulded parts					N/A		
5.3	EN ISO 10993-1 Annex. D	Biological con	formity and toxicity				N/T		
5.4	V/I	Contaminants	and residues				N/A		
	V/I.,B.5.5.1	nd nd	Cleaning			Conf.	Pos.		
	V/I.,B.5.5.1	ologi nns an natio	Disinfection	-10-1912			N/A		
5.5	V/I., EN ISO 22442-1 B.5.5.2	Microbiologica I infections and contamination	Animal tissue				N/A		
5.6	EN ISO 9227	Resistance to corrosion					N/T	(V. 0-100-100-100-100-100-100-100-100-100-1	
6		Emitted sound and vibration							
6.1	EN ISO 3746 B6	Noise and vibration					N/A		
6.2	EN ISO 3746	Sound levels a	nd frequencies of audible warning of	levices		N/A			
Require ments	Test method according to		Checked	Real	Test		Comments		

ments accordi ng to clause	Test method according to clause	Checked characteristics/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, spillage, leakage, and ingress of liquids		N/A	
10	V/I. Measur.	Surface temperature		N/A	t ⁰ ≤ 41 ⁰ 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 clau	ing to	Checked characteristics/assemblies/parameters	Real value	Test result	Comments			
12	V/I. M	easur.	Safety of moving parts		N/A				
13	V/I. M	easur.	Prevention of traps for parts of the human body	Conf.	Pos.	Not	te in service manual		
14	V	/I	Folding and adjusting mechanisms		N/A			55-21	
15	V/I. M	easur.	Carrying handles		N/A				
16	V/I. M		Assistive products which support or suspend users	Conf.	Pos.		1,5x350i vcles, 3	kg=525kg 50kg	, 70se
17	V/I. M	The second contract of the second	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		10 10 10 10 10 10 10 10 10 10 10 10 10 1		
20	B	20	Small Parts	Conf.	Pos.				1000000
21	V/I. M EN 60		Stability		N/A				
22	B 22,	, V/I	Forces in soft tissues of the human body		N/A				
23	V/ EN 6		Ergonomic principles		N/T	The requirements relate to the process			he desi
Require ments accordi ng to clause	Test method accordi ng to clause		Checked characteristics/assemblies/	/paramet	ters		Test result	Comer	
24	V/I	Requi	irements for information supplied by the manufact	urer			e		
24.1		Gener						N/T	
24.2		Instru	actions for use					N/T	
24.2.1	V/I		le Information						
		p	aformation on how to obtain the user information in a eople with visual,r eading orcognitive disabilities		ropriate	for use by		N/T	
			Il information shall as far as possible be available in Pic					N/T 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 dond suitable cleaning materials, including precautions					N/T	
		f) if	opplicable; an assistive product is intended to be disinfected, a cuitable materials, including any precautions needed to					N/T	
		m	ne overall dimensions (width, length and height) of the audillimetres, and its mass, expressed in kilograms, when opplicable, when it is folded or dismantled					N/T	
2421	V/I	aı	ne mass expressed in kilograms if the assistive producing removable parts that has a mass which is heavier the	an 10 kg;				N/T	
24.2.1		m	the assistive product is supposed to be used in combinanufacturer shall state to which products, and how this	can be dor	ne in a sat	è way;		N/T	
		0	rarning about dangerous combinations of devices (e.g. f decubitus ulcers often only work on correct seat sur ame resistant and non-flame resistant material;					N/T	
		k) a list of accessories, detachable parts and materials that the manufacturer has determined as being intended for use with the assistive product						N/T	
		p	a programmable controller is fitted, information or orgramming, the competence required to carry out the ffects on performance			the		N/T	
								AT/T	
			perator control adjustments	W 400 C C				N/T	
		n) w	hether and how the assistive product can be folded or a transport					N/T	
		n) w on o) in	hether and how the assistive product can be folded or						



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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		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		
		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		
		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 sgueezing, 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		
		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	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

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: 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

