中国认可
国际互认
检测
TESTING
CNAS L5772

TEST REPORT

Commissioned by:	**Shenzhen DreamCan Technology Co., Ltd**
Factory:	**Shenzhen DreamCan Technology Co., Ltd**
Name of Sample:	**Disposable Medical Mask**
Type,Specification:	**DP-A-24L-K**
Testing Purpose:	**Commission**
Testing Lab:	**Precise Testing & Certification (Guangdong) Co., Ltd. (PTC)**
Address:	**Building 1, No. 6, Tongxin Road, Dongcheng Street, Dongguan, Guangdong**

Test Report

Name of sample	Disposable Medical Mask	**Type,Specification**	DP-A-24L-K
		Trade Mark	/
Grade of sample	Type IIR	**Sample No.**	/
Testing perpose	Commission	**Batch No.**	/
Quantity of sample	150pcs	**Package**	Sealed
Sample Received Date	Oct. 29, 2020 Nov.11, 2020	**Completed Date**	Nov.06, 2020 Nov.12, 2020
Testing Environment	As requirement of testing standard		
Commissioned by	Shenzhen DreamCan Technology Co., Ltd 301/B3,Huaqiang Industrial Park, Qingfeng Ave., Baolong, Longgang District, Shenzhen, Guangdong, China		
Name & Address of Factory	Shenzhen DreamCan Technology Co., Ltd 301/B3,Huaqiang Industrial Park, Qingfeng Ave., Baolong, Longgang District, Shenzhen, Guangdong, China		
Testing Reference	EN 14683:2019+AC：2019		
Testing Item	General、Bacterial filtration efficiency、Microbial cleanliness(Bioburden)、Splash resistance、Differential pressure、Marking, labelling and packaging		
Remark	Test results are attached as below		
Note	None		

Prepare by:	Checked by：	Approved by
Anne		Tim Mo

PRECISE TESTING & CERTIFICATION (GUANGDONG) CO., LTD.

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Test Report

No.	Testing Item/ Unit		Requirements	Testing Result	Conclusion
1	General	Materials and construction	1）The medical face mask is a medical device, generally composed of a filter layer that is placed, bonded or moulded between layers of fabric. 2)The medical face mask shall not disintegrate, split or tear during intended use. 3)In the selection of the filter and layer materials, attention shall be paid to cleanliness.	Complied	PASS
		Design	1)The medical face mask shall have a means by which it can be fitted closely over the nose, mouth and chin of the wearer and which ensures that the mask fits closely at the sides 2)Medical face masks may have different shapes and constructions as well as additional features such as a face shield (to protect the wearer against splashes and droplets) with or without anti-fog function, or a nose bridge (to enhance fit by conforming to the nose contours).	Complied	PASS

Test Report

No.	Testing Item/ Unit	Requirements	Testing Result		Conclusion
2	Bacterial filtration efficiency(%）	Type I ≥95 Type II ≥98 Type IIR ≥98	99.95		Type IIR PASS
			99.84		
			100		
			100		
			99.89		
3	Microbial cleanliness (Bioburden)	Type I ≤30CFU/g Type II ≤30CFU/g Type IIR ≤30CFU/g	Sample1: (3.1g)	16CFU/g	Type IIR PASS
			Sample2: (3.1g)	13CFU/g	
			Sample3: (3.1g)	26CFU/g	
			Sample4: (3.1g)	15CFU/g	
			Sample5: (3.0g)	19CFU/g	
4	Splash resistance (Kpa)	Type IIR≥16,0	>16.0		Type IIR PASS
5	Differential pressure (Pa/cm^2)	Type I <40 Type II <40 Type IIR <60	57.80		Type IIR PASS
			56.44		
			57.18		
			57.98		
			58.51		

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No.	Testing Item/ Unit	Requirements	Testing Result	Conclusion
6	Marking, labelling and packaging	Annex I, §13, of the Medical Devices Directive (93/42/EEC) or Annex I, §23, of the Medical Device Regulation (EU) 2017/745 specifies the information that should be specified on the packaging in which the medical face mask is supplied. The following information shall be supplied: a) number of this European Standard; b) type of mask (as indicated in Table 1). EN ISO 15223-1:2016 and EN 1041:2008+A1:2013 should be considered.	Comply	PASS

Remark: This report supersedes all previous documents bearing the test report number PTC20102707901C-EN01. Report number PTC20102707901C-EN01 was invalid.

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Photo(s) of Sample:

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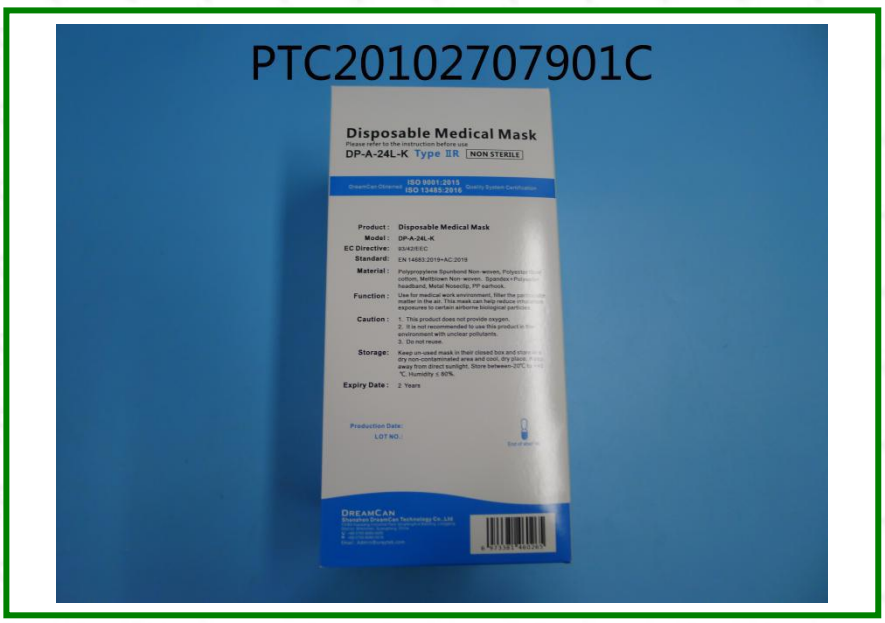

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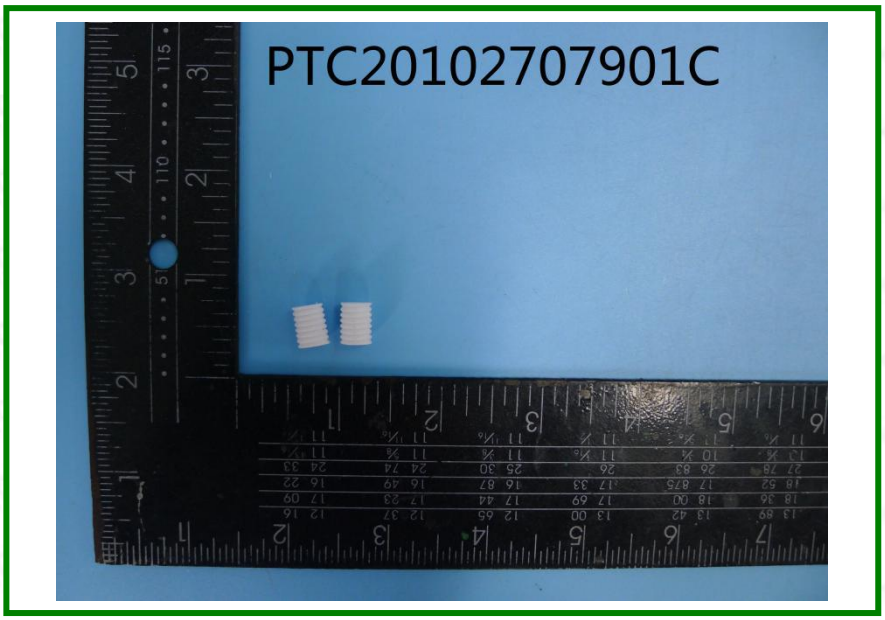

End of Report

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