

20ZCTS0323013SP



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TEST REPORT EN 14126

Protective clothing - Performance requirements and tests methods for protective clothing

against infective agents

20ZCTS0323013SP Report Reference No.

Tested by (+ signature):

Kevin Yang

Approved by (+ signature) :

King Hu

Date of issue: March 12, 2020

Shenzhen ZCT Technology Co.,Ltd Testing laboratory.....

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Xixiang Street, Bao'an District, Shenzhen, Guangdong, China.

Applicant's name Hunan Mingyu Nonwovens Co.Ltd

Address NO.333 DESHAN AVENUE, CHANGDE ECONOMIC AND

TECHNOLOGY DEVELOPMENT DISTRICT, HUNAN

PROVINCE, CHINA

Manufacturer's name.....: Same as applicant

Address....:

Same as applicant Factory's name:

Address....:

Test specification:

X EN 14126:2003 Standard....::

Test procedure: Commission test

Non-standard test method..... N/A

Test Report Form No.:

TRF Originator:: ZCT

Master TRF: Dated 2019-01

Test item description:: Disposable isolation clothin

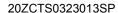
Trade Mark....::

Model/Type reference: MY003

Ratings....: Type 4-B Class 2









Test item particulars:	
Test case verdicts:	
Test case does not apply to the test object:	N/A
Test object does meet the requirement:	Pass (P)
Test object does not meet the requirement:	Fail (F)
Testing:	
Date of receipt of test item	Mar. 12, 2020
Date(s) of performance of test	Mar. 13, 2020 to Mar. 24, 2020
General remarks:	
The test results presented in this report relate o	nly to the item(s) tested.

This report shall not be reproduced, except in full, without the written approval of the testing laboratory. "(see remark #)" refers to a remark appended to the report.

"(see Annex #)" refers to an annex appended to the report.

"(see appended table)" refers to a table in the Test Report.

Throughout this report a comma (point) is used as the decimal separator.

No Marking





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	EN 14126		
Clause	Requirement - Test	Result - Remark	Verdict
4	Requirements		-
4.1	Materials requirements		-
4.1.1	General		-
	If the care instructions indicate that the clothing can be cleaned and reprocessed at least five times, protective clothing materials shall be submitted to five cleaning and reprocessing cycles according to the manufacturer's care instructions before testing.		Р
	If the care instructions specify a lower number of cleaning/reprocessing cycles, then materials shall be submitted to the number of cleaning/reprocessing cycles indicated.		Р
-	Unless otherwise stated in the relevant test procedure, the specimens shall be conditioned for at least 24 h in an atmosphere of (20 ± 2) °C and (65 ± 5) % relative humidity before testing.		Р
	Tests shall be carried out in the same atmosphere or within 5 min of removing the sample from the conditioning atmosphere.		Р
4.1.2	Mechanical and flammability requirements		-
	The materials shall be tested and classified in accordance with the test methods and performance classification system specified in the relevant clauses of prEN 14325.		Р
4.1.3	Chemical requirements		-
	If protection against chemicals is claimed, the materials shall be tested and classified in accordance with the test methods and performance classification system specified in the relevant clauses of prEN 14325.		Р
4.1.4	Performance requirements against penetration by infective agents		-
4.1.4.1	Resistance to penetration by contaminated liquids under hydrostatic pressure		Р
	When tested in accordance with ISO/FDIS 16603 and ISO/FDIS 16604 the material shall be classified according to the levels of performance given in Table 1, as obtained in the bacteriophage		Р





	EN 14126			
Clause	Requirement - Test	Result - Remark	Verdict	
	test (ISO/FDIS 16604).			
	NOTE The synthetic blood test (ISO/FDIS 16603)		-	
	is used for screening purposes, i.e. to predict the			
	level where a strikethrough can be expected when			
	performing the bacteriophage test (ISO/FDIS			
	16604)			

Table 1 - Classification of resistance to penetration by contaminated liquids under hydrostatic pressure (ISO/FDIS 16604)

Class	Hydrostatic pressure at which the	Result	Verdict
	material passes the test		
6	20 kPa	1	1
5	14 kPa	1	1
4	7 kPa	1	1
3	3,5 kPa	1	1
2	1,75 kPa	2,3 kPa	Р
1	0 kPa a	1	1

а this means that the material is only exposed to the hydrostatic pressure of the liquid in the test cell

4.1.4.2	Resistance to penetration by infective agents	Р
	due to mechanical contact with substances	
	containing contaminated liquids.	
	When tested in accordance with Annex A the	Р
	material shall be classified according to the levels	
	of performance given in Table 2.	

Table 2 - Classification of resistance to penetration by infective agents due to mechanical contact with substances containing contaminated liquids

Class	Breakthrough time, t min	Result	Verdict
6	t > 75		1
5	60 < t ≤ 75		1
4	45 < t ≤ 60		1
3	30 < t ≤ 45		1
2	15 < t ≤ 30	>28 min	Р
1	≤ 15 min		1

4.1.4.3	Resistance to penetration by contaminated	-
	liquid aerosols	
	When tested in accordance with ISO/DIS 22611	Р
	the material shall be classified according to the	
	levels of performance given in Table 3.	





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Clause	Requirement - Test		Result - Remark	Verdict

Table 3 - Classification of resistance to penetration by contaminated liquid aerosols

Class	Penetration ratio (log)	Result	Verdict
3	log > 5		1
2	3 < log ≤ 5	4	Р
1	1 < log ≤ 3		1

4.1.4.4	Resistance to penetration by contaminated	
	solid particles.	
	When tested in accordance with ISO/DIS 22612	Р
	the material shall be classified according to the	
	levels of performance given in Table 4.	

Table 4 - Classification of resistance to penetration by contaminated solid particles

Class	Penetration (log cfu)	Result	Verdict
3	≤1	1	1
2	1 < log cfu ≤ 2	1	Р
1	2 < log cfu ≤ 3	1	1

4.2	Performance requirements for seams, joins	-
	and assemblages	
	Seams, joins and assemblages of protective	Р
	clothing against infective agents shall fulfil the	
	requirements specified in the relevant clauses of	
	prEN 14325 Seam strength shall be classified	
	according to 5.5 of prEN 14325:2001.	
4.3	Whole suit requirements	-
	Protective clothing against infective agents shall	Р
	fulfil the relevant requirements of EN 340 and the	
	whole suit requirements specified in the relevant	
	standard for chemical protective clothing (see	
	Table 5).	
	The materials and design used shall not cause	Р
	skin irritation nor have any adverse effect to	
	health.	
	NOTE The suit should be as light and as flexible	-
	as possible in order to ensure the comfort of the	
	wearer, not to hinder movements and still provide	
	at the same time effective protection.	

Table 5 - Types of protective clothing against infective agents

Type of clothing	Relevant standard
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Add: 3/F., Building 5, Hongsheng Industrial Zone, Bao'an Road, Xixiang

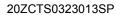


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Clause	Requirement - Test		Result - Remark	Verdict
type 1a, 1b, 1c, 2		EN 943-1 (EN 943-2 for ET suits)		
type 3		EN 466		
type 4 EN 465				
type 5 prEN ISO 13982-1				
type 6		prEN 13034		
partial body	protection	EN 467		

5	Marking		-
	The clothing shall be marked in accordance with		Р
	the applicable requirements of the relevant		
	standard for chemical protective clothing.		
	The marking of protective clothing against		Р
	infective agents shall contain the following		
	additional information:		
	a) the number of this European Standard;	EN 14126	Р
	b) the type of protective clothing, as specified in	Type 4-B	Р
	Table 5, with the suffix "-B", e.g. type 3-B;		
	c) the pictogram "protection against biological		Р
	(\$2)		
	hazard"		
6	Information supplied by the manufacturer		-
	The information for the user shall be worded		Р
	clearly and unambiguously and be understandable		
	by a trained person.		
	The information for the user of protective clothing		Р
	against infective agents shall contain all the		
	information required by EN 340 and by the		
	relevant standard for that specific type of chemical		
	protective clothing. In addition it shall contain the		
	following information:		
	a) the number of this European Standard;	EN 14126	Р
	b) the type designation, e.g. type 3-B;	Type 4B	Р
	c) the biological agents against which the		Р
	protective clothing has been tested. This		
	information shall be expressed as performance		
	levels, as specified in 4.1.4.1 to 4.1.4.4 for the		
	relevant types of biological challenge;		
	d) all other relevant information on performance		Р







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Clause	Requirement - Test	Result - Remark	Verdict
	levels, preferably as a Table;		
	e) the information necessary for trained persons about:		Р
	 application and limitations of use (temperature range, etc.); 		Р
	if relevant, checks to be carried out by the wearer before use;		Р
	 fitting and adjustments, and any accessories needed to provide the claimed level of protection; 		Р
	- use;		Р
	- maintenance, cleaning and disinfection;		Р
	- storage;		Р
	 if relevant, a warning against problems likely to be encountered; 		Р
	if relevant, illustrations, part numbers and marking of spare parts, etc.		Р
	– disposal after use.	5.	Р







Annex I: Photo documentation

Type of equipment, model: Disposable isolation clothin, MY003

Details of: MY003



- End of Test Report -

