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QDHL2009009987MD

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Test Report

Report No.: QDHL2009009987MD

Sample Name:	SINGLE-USE MEDICAL LATEX EXAMINATION GLOVES
Applicant:	FITONE LATEX PRODUCTS CO.,LTD.GUANGDONG
Test Type:	SUBMITTED BY CLIENT

QD 7352628

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Report No.: QDHL2009009987MD

Test Report

Sample information	Sample name	SINGLE-USE MEDICAL LATEX EXAMINATION GLOVES	Color	RUBBER
	Reviced sample quantity/ Tested sample quantity	46PCS/ 13PCS	Type/ Specifications	M SIZE/ Powdered-free gloves other than surgeon's gloves
	Lot No.	SE200819	Lot Quantity	100000PCS
	Manufacture Date	08/2020	Expiration Date	07/2025
	Material/Appearance	LATEX/FINISHED	Storage Condition	ROOM TEMPERATURE/ AVOID LIGHT
	Manufacturer	FITONE LATEX PRODUCTS CO.,LTD.GUANGDONG		

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Client information	Applicant	FITONE LATEX PRODUCTS CO.,LTD.GUANGDONG		
	Applicant address	NO.5 TONGYI ROAD, LINGBEI INDUSTRIAL ZONE, SUIXI,524338,ZHANJIANG, GUANGDONG,CHINA		
Test information	Sample Receiving Date	SEP.24,2020	Test Period Date	SEP.24,2020 TO OCT.12,2020
	Sample No.	QDHL2009009987MD (TAOHG2004309501)	Test environment	Meet requirement
	Test items	Removable Surface Powder; Proteins, leachable*		
	Testing Accordance	BS EN 455-3:2015 Medical gloves for single use – Part 3: Requirements and testing for biological evaluation clause 4.4,4.5		
Test conclusion	This report only provides the test results and individual judgment, conclusion please see follow pages. Issue date: OCT.12,2020			
Remark	/			

Approver: Jessica Gao Auditor: Jessica Gao Compiler: Lillian Diao

Date: 2020.10.12 Date: 2020.10.12 Date: 2020.10.12

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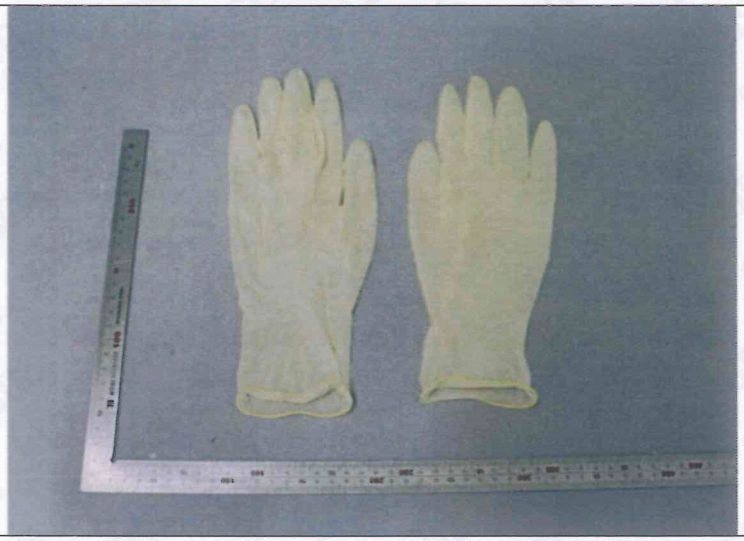
QD 7352630

Report No.: QDHL2009009987MD

Sample Photo


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QD 7352631

Report No.: QDHL2009009987MD

Test Results

Test Items	Unit	Test Method	Requirement	Test Result	Assessment
Removable Surface Powder	mg	EN ISO 21171:2006	≤2	0.08	Pass
Proteins, leachable*	µg/g	BS EN 455-3:2015	/	Not Detected (MDL: 10)	/

Remark:
1. The declaration of conformity is only based on the actual value of laboratory activity, measurement uncertainty of the results not take into account.
2. * Test items were not included in the CNAS accredited schedule for our laboratory.

End of Report

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Statement

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2. Above information and sample(s) was/were submitted and certified by the client, SGS quoted the information with no responsibility as to the accuracy, adequacy and/or completeness.

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5. Should you have any queries or objection to the test report, please contact us whthin 15 days after receiving the report.

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