



July 29, 2021

Siyang Threeguard Medical Supplies Co., Ltd.
% Boyle Wang
Official Correspondent
Shanghai Truthful Information Technology Co., Ltd.
RM.608, No.738, Shangcheng Rd., Pudong
Shanghai, Shanghai 200120
China

Re: K211131

Trade/Device Name: Disposable Nitrile Medical Examination Glove
Regulation Number: 21 CFR 880.6250
Regulation Name: Non-Powdered Patient Examination Glove
Regulatory Class: Class I, reserved
Product Code: LZA
Dated: June 23, 2021
Received: June 28, 2021

Dear Boyle Wang:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. Although this letter refers to your product as a device, please be aware that some cleared products may instead be combination products. The 510(k) Premarket Notification Database located at <https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfpmn/pmn.cfm> identifies combination product submissions. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal

statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803) for devices or postmarketing safety reporting (21 CFR 4, Subpart B) for combination products (see <https://www.fda.gov/combination-products/guidance-regulatory-information/postmarketing-safety-reporting-combination-products>); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820) for devices or current good manufacturing practices (21 CFR 4, Subpart A) for combination products; and, if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to <https://www.fda.gov/medical-devices/medical-device-safety/medical-device-reporting-mdr-how-report-medical-device-problems>.

For comprehensive regulatory information about medical devices and radiation-emitting products, including information about labeling regulations, please see Device Advice (<https://www.fda.gov/medical-devices/device-advice-comprehensive-regulatory-assistance>) and CDRH Learn (<https://www.fda.gov/training-and-continuing-education/cdrh-learn>). Additionally, you may contact the Division of Industry and Consumer Education (DICE) to ask a question about a specific regulatory topic. See the DICE website (<https://www.fda.gov/medical-devices/device-advice-comprehensive-regulatory-assistance/contact-us-division-industry-and-consumer-education-dice>) for more information or contact DICE by email (DICE@fda.hhs.gov) or phone (1-800-638-2041 or 301-796-7100).

Sincerely,

For Clarence W. Murray, III, Ph.D.
Assistant Director
DHT4B: Division of Infection Control
and Plastic Surgery Devices
OHT4: Office of Surgical
and Infection Control Devices
Office of Product Evaluation and Quality
Center for Devices and Radiological Health

Enclosure

Indications for Use

510(k) Number (if known)
K211131

Device Name
Disposable Nitrile Medical Examination Glove

Indications for Use (Describe)

The Disposable Nitrile Medical Examination Glove is a disposable device intended for medical purposes that is worn on the examiner's hands to prevent contamination between patient and examiner.

Type of Use (Select one or both, as applicable)

Prescription Use (Part 21 CFR 801 Subpart D)

Over-The-Counter Use (21 CFR 801 Subpart C)

CONTINUE ON A SEPARATE PAGE IF NEEDED.

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510(k) Summary

(K211131)

This summary of 510(k) is being submitted in accordance with 21 CFR 807.92.

1.0 Submitter's Information

Name: Siyang Threeguard Medical Supplies Co.,Ltd.

Address: East of Nanhai Road and South of Guilin Road,Economic
Development Zone, Siyang County,Suqian City,Jiangsu,223799
China.

Phone Number: +86-13485097856

Contact: Guo Hua

Date of Preparation: July.29,2021

Designated Submission Correspondent

Mr. Boyle Wang

Shanghai Truthful Information Technology Co., Ltd.

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Tel: +86-21-50313932

Email: Info@truthful.com.cn

2.0 Device Information

Trade name: Disposable Nitrile Medical Examination Glove

Common name: Patient Examination Gloves

Classification name: Non-powdered patient examination glove

Model(s): XS,S, M, L, XL

3.0 Classification

Production code: LZA

Regulation number: 21CFR880.6250

Classification: Class I

Panel: General Hospital

4.0 Predicate Device Information

Manufacturer: Ever Global (Vietnam) Enterprise Corp

Device: Disposable Powder Free Nitrile Examination Glove, White/
Blue/ Black/ Pink Color

510(k) number: K171422

5.0 Indication for Use

The Disposable Nitrile Medical Examination Glove is a disposable device intended for medical purposes that is worn on the examiner's hands to prevent contamination between patient and examiner.

6.0 Device Description

The subject device is powder free nitrile examination gloves. The subject device is blue. The subject device is non-sterile.

7.0 Technological Characteristic Comparison Table

Table1-General Comparison

Item	Subject Device (K211131)	Predicate Device (K171422)	Remark
Product Code	LZA	LZA	Same
Regulation No.	21CFR880.6250	21CFR880.6250	Same
Class	I	I	Same
Intended Use	Disposable Nitrile Medical Examination Glove is a disposable device intended for medical purposes that is worn on the examiner's hands to prevent contamination between patient and examiner.	The Disposable Powder Free Nitrile Examination Glove, White/ Blue/ Black/ Pink Color is a disposable device intended for medical purposes that is worn on the examiner's hands to prevent contamination between patient and examiner.	Same
Powdered or Powdered free	Powdered free	Powdered free	Same
Design Feature	Ambidextrous	Ambidextrous	Same
Labeling Information	Single-use indication, powder free, device color, device name, glove size and quantity,	Single-use indication, powder free, device color, device name, glove size and quantity, Non-Sterile	Same

	Non-Sterile	
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Table2 Device Dimensions Comparison

Predicate Device(K171422)	Designation	Size					Tolerance	
		XS	S	M	L	XL		
	Length, mm	230	230	230	230	230	min	
	Width, mm	75	85	95	105	115	±5	
Thickness, mm:								
	Finger	0.05					min	
	Palm	0.05					min	
Subject Device (K211131)	Designation	Size					Tolerance	
		XS	S	M	L	XL		
		Length, mm	230	230	230	230	230	min
		Width, mm	75	85	95	105	115	±5
	Thickness, mm:							
		Finger	0.05					min
	Palm	0.05					min	
Remark	Same							

Table3 Performance Comparison

Item			Subject device (K211131)	Predicate device (K171422)	Remark
Colorant			Blue	White/ Blue/ Black/ Pink	Same
Physical Properties	Before Aging	Tensile Strength	14MPa, min	14MPa, min	Same
		Ultimate Elongation	500% min	500% min	Same
	After Aging	Tensile Strength	14MPa, min	14MPa, min	Same
		Ultimate Elongation	400%min	400%min	Same
	Comply with ASTM D6319			Comply with ASTM D6319	Same
Freedom from Holes			Be free from holes when tested in accordance with ASTM D5151 AQL=2.5	Be free from holes when tested in accordance with ASTM D5151 AQL=2.5	Same

Powder Content	Meet the requirements of ASTM D6124 < 2.0mg	Meet the requirements of ASTM D6124	Same
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Table4 Safety Comparison

Item		Subject device (K211131)	Predicate device (K171422)	Remark
Material		Nitrile	Nitrile	Same
Biocompatibility	Irritation (ISO 10993-10:2010 Biological Evaluation of Medical Devices - Part 10: Tests For Irritation And Skin Sensitization)	Under the conditions of the study, not an irritant	Comply with ISO10993-10	Same
	Sensitization (ISO 10993-10:2010 Biological Evaluation of Medical Devices - Part 10: Tests For Irritation And Skin Sensitization)	Under conditions of the study, not a sensitizer.		
	Cytotoxicity (ISO 10993-5:2009 Biological Evaluation of Medical Devices - Part 5: Tests For In Vitro Cytotoxicity)	Under conditions of the study, device extract is not cytotoxic	/	Similar

8.0 Summary of Non-clinical Testing

Non-clinical tests were conducted to verify that the proposed device met all design specifications. The test results demonstrated that the proposed device complies with the following standards:

ISO 10993-10:2010 Biological Evaluation of Medical Devices - Part 10: Tests For Irritation And Skin Sensitization.

ISO 10993-5:2009 Biological Evaluation of Medical Devices - Part 5: Tests For In Vitro Cytotoxicity

ASTM D6124-06 (Reapproved 2017), Standard Test Method for Residual Powder on Medical Gloves

ASTM D5151-19, Standard Test Method for Detection of Holes in Medical Gloves.

ASTM D6319-19, Standard Specification for Nitrile Examination Gloves for Medical Application.

Test Methodology	Purpose	Acceptance Criteria	Results
ASTM D6319	Physical Dimensions Test	Length(mm): ≥230; Width(mm): XS: 75±5; S: 85±5; M: 95±5; L: 105±5; XL: 115±5	Length: > 230 Width: XS: 77-79; S: 82-86 M: 95-97 L: 104-106 XL: 115-117 <u>Pass</u>
		Thickness (mm): Finger: ≥0.05 Palm: ≥0.05	XS: Finger: 0.10-0.12 Palm: 0.06-0.10 S: Finger: 0.09-0.12 Palm: 0.07-0.09 M: Finger: 0.08-0.12 Palm: 0.06-0.09 L: Finger: 0.09-0.13 Palm: 0.06-0.09 XL: Finger: 0.11-0.12 Palm: 0.06-0.08 <u>Pass</u>
ASTM D5151	Watertightness Test for Detection of Holes	Meet the requirements of ASTM D5151 AQL 2.5	XS: 0/125 leaks S:1/125 leaks M:0/125 leaks

					L:2/125 leaks XL:2/125 leaks <u>Pass</u>
ASTM D6124	Powder Content	Meet the requirements of ASTM D6124 < 2.0mg			XS:0.01mg/glove; S:0.02mg/glove; M:0.03mg/glove; L:0.01mg/glove; XL:0.02mg/glove; <u>Pass</u>
ASTM D412	Physical properties	Before Aging	Tensile Strength	≥14MPa	XS:15.3-16.3 S:15.6-19.2 M:15.9-17.3 L:15.3-17.2 XL:15.3-17.1 <u>Pass</u>
			Ultimate Elongation	≥500%	XS: 525-579 S:518-571 M:515-575 L:521-562 XL:523-568 <u>Pass</u>
		After Aging	Tensile Strength	≥14MPa	XS: 15.6-16.9 S:15.8-17.0 M:15.2-17.7 L:15.3-17.1 XL:15.0-18.7 <u>Pass</u>
			Ultimate Elongation	≥400%	XS:520-564 S:516-570 M:511-554 L:523-564 XL:528-570 <u>Pass</u>
ISO 10993-5	Cytotoxicity	Non-cytotoxic			Under conditions of the study, did not show potential toxicity to L-929 cells. <u>Pass</u>

ISO 10993-10	Irritation	Non-irritating	Under the conditions of the study, not an irritant. <u>Pass</u>
ISO 10993-10	Sensitization	Non-sensitizing	Under conditions of the study, not a sensitizer. <u>Pass</u>

9.0 Summary of Clinical Testing

Clinical testing is not needed for this device.

10.0 Conclusion

The conclusions drawn from the nonclinical tests demonstrate that the subject device is as safe, as effective, and performs as well as or better than the legally marketed predicated device under K171422.