October 25, 2021

Inner Mongolia Cureguard Medical Technology Co.,Ltd.
Boyle Wang
Official Correspondent
Shanghai Truthful Information Technology Co., Ltd.
RM.1801, No.161, East Lujiazui Rd., Pudong
Shanghai, Shanghai 200120
China

Re: K212401
Trade/Device Name: Disposable Nitrile Examination Glove
Regulation Number: 21 CFR 880.6250
Regulation Name: Non-Powdered Patient Examination Glove
Regulatory Class: Class I, reserved
Product Code: LZA
Dated: July 16, 2021
Received: August 2, 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

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

Indications for Use

Form Approved: OMB No. 0910-0120
Expiration Date: 06/30/2023
See PRA Statement below.

510(k) Number *(if known)*
K212401

Device Name
Disposable Nitrile Examination Glove

Indications for Use *(Describe)*
The Disposable Nitrile Examination Glove is a non-sterile disposable device intended for medical purposes that is worn on the examiner's hands or finger 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.

This section applies only to requirements of the Paperwork Reduction Act of 1995.

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510(k) Summary

(K212401)

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

1.0 Submitter's Information

Name: Inner Mongolia Cureguard Medical Technology Co.,Ltd.
Address: Room 326, Management Committee of New Industrial Park, Tumote youqi, Baotou, Inner Mongolia Autonomous Region 014100, China.
Phone Number: +86-13485097856
Contact: Guo Hua
Date of Preparation: Oct.22,2021

Designated Submission Correspondent

Mr. Boyle Wang
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Tel: +86-21-50313932
Email: Info@truthful.com.cn

2.0 Device Information

Trade name: Disposable Nitrile 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 Examination Glove is a non-sterile disposable device intended for medical purposes that is worn on the examiner's hands or finger to prevent contamination between patient and examiner.

6.0 Device Description

The subject device is powder free nitrile patient examination gloves. The subject device is blue color and has 5 models of XS,S, M, L, XL. The design of subject device is addressing the standards as ASTM D6124, ASTM D5151, and ASTM D6319. The subject device is non-sterile.

7.0 Technological Characteristic Comparison Table

Item	**Subject Device** (K212401)	**Predicate Device (**K171422**)**	**Remark**
Product Code	LZA	LZA	Same
Regulation No.	21CFR880.6250	21CFR880.6250	Same
Class	I	I	Same
Indication for Use/ Intended Use	The Disposable Nitrile Examination Glove is a non-sterile disposable device intended for medical purposes that is worn on the examiner's hands or finger 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
Material	Nitrile	Nitrile	Same
Powdered or Powered free	Powder free	Powder free	Same
Design Feature	Ambidextrous	Ambidextrous	Same
Colorant	Blue	White/ Blue/ Black/ Pink	Similar
Labeling Information	Single-use indication, powder free, device color, device name, glove size and quantity,Non-Sterile	Single-use indication, powder free, device color, device name, glove size and quantity, Non-Sterile	Same
Dimensions(mm)	Length: XS/S:≥220； M/L/XL: ≥230； Width: XS: 70±10;	Length: XS/S:≥220；M: ≥235; L/XL: ≥245 Width: XS: 75±5;	Similar

		S: 80±10; M: 95±10; L: 105±10; XL: 115±10		S: 85±5; M: 95±5; L: 105±5; XL: 115±5		
Thickness(mm)		Finger: ≥0.05; Palm: ≥0.05		Finger: ≥0.05; Palm: ≥0.05		Same
Physical Properties	Before Aging	Tensile Strength	14MPa, min	Tensile Strength	14MPa, min	Same
		Ultimate Elongation	500% min	Ultimate Elongation	500% min	Same
	After Aging	Tensile Strength	14MPa, min	Tensile Strength	14MPa, min	Same
		Ultimate Elongation	400%min	Ultimate Elongation	400%min	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 ASTMD5151 AQL=2.5		Same
Powder Content		Meet the requirements of ASTM D6124 ＜2.0mg		Meet the requirements of ASTM D6124		Same
Biocompatibility		ISO 10993-10; Under the conditions of the study, not an irritant or a sensitizer		Comply with ISO10993-10		Same
		ISO 10993-5 Under conditions of the study, device extract is not cytotoxic		/		Similar

Analysis:
The color(blue) of the subject device is different with those (white/ blue/ black/ pink) of the predicate device, biocompatibility test has been performed on subject device and the test result can meet the requirements of ISO 10993 standards.

The physical dimensions are little different with that of the predicate, but they all meet the requirements of ASTM D6319.

Therefore, the differences will not raise any safety and effectiveness issues.

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.

Table 2: Summary of Non-clinical Testing Table

Test Methodology	Purpose	Acceptance Criteria	Results
ASTM D6319	Physical Dimensions Test	Length(mm): XS/S:≥220； M/L/XL: ≥230； Width: XS: 70±10; S: 80±10; M: 95±10; L: 105±10; XL: 115±10	Length(mm): ＞230 Width(mm): XS: 72-74; S: 80-83 M: 95-98 L: 110-114 XL: 118-121 Pass
		Thickness (mm) : Finger: ≥0.05 Palm: ≥0.05	XS: Finger: 0.07-0.10 Palm: 0.08-0.10 S: Finger: 0.08-0.11 Palm: 0.08-0.11 M: Finger: 0.08-0.12 Palm: 0.07-0.11 L: Finger: 0.08-0.12 Palm: 0.08-0.11 XL: Finger: 0.08-0.12

					Palm: 0.08-0.12 Pass
ASTM D5151	Watertightness Test for Detection of Holes	Meet the requirements of ASTM D5151 AQL 2.5			XS:2/125 leaks S:0/125 leaks M:0/125 leaks L: 1/125 leaks XL: 0/125 leaks Pass
ASTM D6124	Powder Content	Meet the requirements of ASTM D6124 < 2.0mg			XS:0.05mg S:0.06mg M:0.06mg L:0.07mg XL:0.09mg Pass
ASTM D412	Physical properties	Before Aging	Tensile Strength	≥14MPa	XS:15.4-17.3 S:15.3-16.9 M: 15.4-17.3 L:15.4-17.6 XL:15.3-17.1 Pass
			Ultimate Elongation	≥500%	XS:524-569 S:525-568 M: 525-567 L:527-566 XL:520-570 Pass
		After Aging	Tensile Strength	≥14MPa	XS:15.3-17.0 S:15.4-16.9 M:15.4-16.4 L:15.3-16.6 XL:15.2-17.2 Pass
			Ultimate Elongation	≥400%	XS:526-568 S:522-570 M:527-570 L:521-567 XL:528-565 Pass

ISO 10993-5	Cytotoxicity	Non-cytotoxic	Under conditions of the study, did not show potential toxicity to L-929 cells. Pass
ISO 10993-10	Irritation	Non-irritating	Under the conditions of the study, not an irritant. Pass
ISO 10993-10	Sensitization	Non-sensitizing	Under conditions of the study, not a sensitizer. Pass

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, Disposable Nitrile Examination Glove, is as safe, as effective, and performs as well as or better than the legally marketed predicate device.