February 28, 2022

Chifeng Huawei Medical Science & Technology Co., Ltd
℅ Boyle Wang
Offical Correspondent
Shanhai Truthful Information Technology Co., Ltd
RM.1801,No.161 Lujiazui East Rd.,Pudong
Shanghai, Shanghai 200120
China

Re: K212833
Trade/Device Name: Disposable Nitrile Examination Gloves
Regulation Number: 21 CFR 880.6250
Regulation Name: Non-Powdered Patient Examination Glove
Regulatory Class: Class I, reserved
Product Code: LZA
Dated: January 28, 2022
Received: February 1, 2022

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, PhD
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)*
K212833

Device Name
Disposable Nitrile Examination Gloves

Indications for Use *(Describe)*
The Disposable Nitrile Examination Gloves is intended to be worn on the hands of examiners to prevent contamination between patient and examiner. This is a single-use, powder-free, non-sterile device.

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

K212833

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

1.0 Submitter's Information

Name: CHIFENG HUAWEI MEDICAL SCIENCE＆TECHNOLOGY CO.,LTD.
Address: No. 2-4, Second-stage Standardized Plant, Songshan (Anqing) Industrial Park, Chifeng, China
Phone Number: +86-18248056569
Contact: Li Xiaohong
Date of Preparation: Oct. 9th,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 Gloves
Common name: Patient Examination Gloves
Classification name: Non-powdered patient examination glove
Model(s): 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: GUANG DONG KINGFA SCI. & TECH.CO., LTD.
Device: Nitrile examination gloves
510(k) number: K203593

5.0 Indication for Use

The Disposable Nitrile Examination Gloves are intended to be worn on the hands of examiners to prevent contamination between patient and examiner. This is a single-use, powder-free, non-sterile device.

6.0 Device Description

The subject device is powder free nitrile examination gloves. The subject device is blue. It can be available in four specifications: S,M,L and XL.
The subject device is non-sterile.

7.0 Technological Characteristic Comparison Table

Table1-General Comparison

Item	Standard	Subject Device (K212833)	Predicated Device (K203593)	Comparison
Product Code	/	LZA	LZA	Same
Regulation No.	/	21CFR880.6250	21CFR880.6250	Same
Class	/	I	I	Same
Intended Use	/	The Disposable Nitrile Examination Gloves is intended to be worn on the hands of examiners to prevent contamination between patient and examiner. This is a single-use, powder-free, non-sterile device.	The nitrile examination glove is intended to be worn on the hands of examiners to prevent contamination between patient and examiner. This is a single-use, powder-free, non-sterile device.	Same
Material	/	Nitrile	Nitrile	Same
Powdered or Powered free	/	Powdered free	Powdered free	Same
Design Feature	/	Ambidextrous	Ambidextrous	Same
Colorant	/	Blue	Blue	Same
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)	As Requirements of ASTM 6319: Length: S:≥220； M/L/XL: ≥230；	Length(mm): ＞240; Width(mm): S: Average 88mm M: Average 99mm	Length: S (220mm min) M (230mm min) L (230mm min) XL (230mm min)	Similar

	Width: S: 80±10; M: 95±10; L: 110±10; XL: 120±10		L: Average 108mm XL: Average 118mm	Width: Small (80±10mm) Medium (95±10mm) Large (110±10mm) X large (120±10mm)	
Thickness (mm)	As Requirements of ASTM 6319: Finger: ≥0.05; Palm: ≥0.05		Finger: 0.10-0.12 Palm: 0.08-0.09	Palm: 0.05mm min Finger: 0.05mm min	Similar
Physical Properties	Before Aging	Tensile Strength 14MPa, min	14-32MPa	14MPa, min	Similar
		Ultimate Elongation 500% min	500-638%	500% min	Similar
	After Aging	Tensile Strength 14MPa, min	14-30MPa	14MPa, min	Similar
		Ultimate Elongation 400% min	401-609%	400%min	Similar
Freedom from Holes	ASTMD5151		Be free from holes when tested in accordance with ASTMD 5151 AQL=2.5	Be free from holes when tested in accordance with ASTMD5151 AQL=2.5	Same
Powder Content	ASTM D6124 <2.0 mg/gloves		0.16-0.23mg	<2.0 mg/gloves	Similar
Biocompatibility	ISO 10993-10		ISO 10993-10; Under the conditions of the study, not an irritant or a sensitizer	ISO 10993-10; Under the test condition of study not a sensitizer. Under the test condition of study not an irritant.	Same
	ISO 10993-11		ISO 10993-11; Under the condition of acute systemic toxicity test, the test article did not show acute systemic toxicity in vivo.	Cytotoxicity is assessed via rationale. Under the condition of acute systemic toxicity test, the test article did not show acute systemic toxicity in vivo.	Same
	ISO 10993-5		ISO 10993-5 Under conditions of the	N.A.	/

		study, device extract is cytotoxic		

Analysis:

The physical dimensions, physical properties and powder content are different with that of the predicate, but they all meet the requirements of ASTM D6319-19, so the differences do not raise any new safety or performance questions.

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 10993-5:2009 Biological evaluation of medical devices - Part 5: Tests for in vitro cytotoxicity

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

ISO 10993-11:2017, Biological evaluation of medical devices - Part 11: Tests for systemic toxicity.

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 performance testing

Test Method	Purpose	Acceptance Criteria	Results
ASTM D6319	Physical Dimensions Test	Length(mm): S:≥220; M/L/XL:≥230; Width(mm): S: 80±10; M: 95±10; L: 110±10; XL: 120±10	Length(mm): ＞240/Pass; Width(mm): S: 86-89 /Pass M: 96-101/ Pass L: 107-110/ Pass XL:116-119/ Pass
		Thickness (mm):	Thickness (mm):

		Finger: ≥0.05 Palm: ≥0.05			Finger: 0.10-0.12/Pass Palm: 0.08-0.09/Pass
ASTM D5151	Watertightness Test for Detection of Holes	Meet the requirements of ASTM D5151 AQL 2.5			0/125/Pass
ASTM D6124	Powder Content	Meet the requirements of ASTM D6124 < 2.0mg			0.16-0.23mg/Pass;
ASTM D412	Physical properties	Before Aging	Tensile Strength	≥14MPa	14-32MPa/Pass;
			Ultimate Elongation	≥500%	500-638%/Pass;
		After Aging	Tensile Strength	≥14MPa	14-30MPa/Pass;
			Ultimate Elongation	≥400%	401-609%/Pass;
ISO 10993-5	Cytotoxicity	In Vitro Cytotoxicity Test			Under conditions of the study, device extract is cytotoxic
ISO 10993-11	Cytotoxicity	Non- acute systemic toxicity			Under conditions of` the study, did not show acute systemic toxicity in vivo / 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 Gloves is as safe, as effective, and performs as well as or better than the legally marketed predicated device K203593.