June 28, 2021

Onetexx SDN BHD
% Michael Van der Woude
U.S. Agent
Emergo Global Representation
2500 Bee Cave Road, Building 1 Suite 300
Austin, Texas 78746

Re: K210587
Trade/Device Name: Violet Nitrile Powder Free Patient Examination Glove, Non-Sterile
Regulation Number: 21 CFR 880.6250
Regulation Name: Non-Powdered Patient Examination Glove
Regulatory Class: Class I, reserved
Product Code: LZA
Dated: May 23, 2021
Received: June 1, 2021

Dear Michael Van der Woude:

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,

Ryan Ortega
Acting 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)*
K210587

Device Name
VIOLET NITRILE POWDER FREE PATIENT EXAMINATION GLOVE, NON STERILE

Indications for Use *(Describe)*
A patient examination glove is a disposable device intended for medical purposes that is worn on the examiner's hand 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.

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510(k) SUMMARY - K210587

1.0 Submitter:

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Phone No.:	+60 5 8070 666
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Date of Summary Prepared: 27th May 2021

2.0 Identification of the subject device:

Trade Name	: Violet Nitrile Powder Free Patient Examination Glove, Non-Sterile
Common Name	: Patient Examination Gloves
Classification Name	: Patient Examination Gloves
Device Classification	: 1
Regulation Number	: 21 CFR 880.6250
Product Code	: LZA.

3.0 Predicate Device:

K143289
Non-Sterile, Powder Free Nitrile Examination Gloves – Violet Color -LZA
Company: YTY INDUSTRY (MANJUNG) SDN. BHD.

4.0 Description of The Device:

Violet Nitrile Powder Free Patient Examination Glove, Non-Sterile meets all requirements of ASTM standard D6319 and FDA 21 CFR 880.6250.

The powder free nitrile examination glove is manufactured from Nitrile latex. Inner surface of gloves undergoes surface treatment process to produce a smooth surface that assists the user in donning the gloves with ease without using any lubricant such as powder on the glove surface. The glove is ambidextrous, i.e., can be worn on right hand or left hand.

5.0 Indication for use:

A patient examination glove is a disposable device made of Nitrile latex intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.

6.0 Summary of the Technological Characteristics of the Device:

The Violet Nitrile Powder Free Patient Examination Glove, Non-Sterile are summarized with the following technological characteristics compared to ASTM D6319 or equivalent standards as shown in Table 1.

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510(k) SUMMARY - K210587

Table 1

CHARACTERISTICS	STANDARDS	DEVICE PERFORMANCE		COMPARISON ANALYSIS
		PREDICATE	CURRENT	
		VIOLET	VIOLET	
510(k) Number	-	K143289	K210587	
Manufacturer(s)	-	YTY Industry (Manjung) Sdn Bhd	Onetexx Sdn Bhd	Same
Material	ASTM D6319	Nitrile	Nitrile	Same
Color	-	Violet	Violet	Same
Product Code		LZA	LZA	Same
Physical Properties <u>Before Aging</u> Tensile Strength: Ultimate Elongation: <u>After Aging</u> Tensile Strength: Ultimate Elongation:	ASTM D6319	 23.22-29.11Mpa 520-580% 27.39-30.82 440-500%	 32.35Mpa 568% 36.10Mpa 551%	 Different but within the ASTM standard Different but within the ASTM standard
Thickness: - Finger - Palm - Finger	ASTM D6319	 0.09-0.11mm 0.06-0.07mm 0.05-0.06mm	 0.10mm 0.07mm	 Different but within the ASTM standard
Length:	ASTM D6319	240 – 244mm	246 – 255mm	Different but within the ASTM standard
Powder Free	ASTM D6124	0.14mg/glove	0.24 mg/glove	Different but within the ASTM standard

CHARACTERISTICS	STANDARDS	DEVICE PERFORMANCE		COMPARISON ANALYSIS
		PREDICATE	CURRENT	
		VIOLET	VIOLET	
Biocompatibility	Primary Skin Irritation – ISO 10993-10:2010 (E) & Consumer Product Safety Commission Title 16. Chapter II, Part 1500	Under the condition of the study the device is non-irritant and non-sensitizer	Under the conditions of this study the device did not cause an irritant response. The Primary Irritant Response Category is deemed 'Negligible'	Similar
	Dermal Sensitization- ISO 10993-10: 2010 (E) & Consumer Product Safety Commission, Title 16, Chapter II, Part 1500.3 (c) (4)	Under the condition of the study the device is non-irritant and non-sensitizer	Under the conditions of this study the device did not produce a skin sensitization effect in the guinea pigs.	Similar
	Acute Systemic Toxicity, ISO 10993-11:2017 (E)	Not Tested	Under conditions of this study the device did not induce any systemic toxicity.	Different

CHARACTERISTICS	STANDARDS	DEVICE PERFORMANCE		COMPARISON ANALYSIS
		PREDICATE	CURRENT	
		VIOLET	VIOLET	
Watertight (1000ml)	ASTM D5151:2019	AQL 2.5 Result: 0	Gloves passed AQL 1.5	Different but within the ASTM standard
Indication for Use	-	A patient examination glove is a disposable device intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.	A patient examination glove is a disposable device made of synthetic rubber latex intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.	Same
Size	Medical Glove Guidance Manual – Labeling	Medium	Extra Small Small Medium Large Extra Large	Similar
Single use	Medical Glove Guidance Manual – Labeling	Single Use	Single Use	Same

There are no significant differences between the two products and are identical in terms of intended use, materials design, physical properties, color, thickness and biocompatibility.

7.0 Summary of Non-Clinical Testing

The performance test data of the non-clinical test for this powder free nitrile examination glove is summarized as per below.

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510(k) SUMMARY - K210587

Test Method	Standard	Purpose of Testing	Acceptance Criteria			Results		Status
			[redacted]	Before aging	After aging	Before aging	After aging	
Physical Properties	ASTM D412 (Standard Test Method for Vulcanized Rubber and Thermoplastic Elastomers-Tension)	To evaluate the tensile (tension) properties of glove.	Tensile strength	Min 14.0 MPa	Min 14.0 MPa	32.35Mpa	36.10Mpa	Pass
			Ultimate elongation	Min 500%	Min 400%	568%	551%	Pass

Test Method	Standard	Purpose of Testing	Acceptance Criteria		Results		Status
Dimension	ASTM D3767 Standard Practice for Rubber—Measurement of Dimensions	To measure the length, width and thickness of glove	Length	Min 240 mm	Length	249 mm	Pass
			Width	95 ± 10 mm	Width	98 mm	Pass
			Thickness	Finger – min 0.05mm Palm – min 0.05mm	Thickness	0.10mm 0.07mm	Pass

Test Method	Standard	Purpose of Testing	Acceptance Criteria	Results	Status
Watertight	ASTM D5151 (Standard Test Method for Detection of Holes in Medical Gloves)	To detect holes that leak water and thereby compromise the usefulness of the glove.	Sample size: 500 pcs Inspection level: G1 AQL: 1.5, Acceptance No. 10	The batch size for this sampling is 150,001 to 500,000. Hence, according to the single sampling plan GI, the sample to be drawn is under code M equivalent to 315 pieces with accept 10 and reject 11 to be accepted under AQL 1.5. During the test, 0 piece was found with leaks. Hence it falls within the acceptance criteria.	Pass

Test Method	Standard	Purpose of Testing	Acceptance Criteria	Results	Status
Residual Powder	ASTM D6124 (Standard Test Method for Residual Powder on Medical Gloves)	To determine the amount of residual powder and non-powder solids found on gloves.	Less than 2 mg per glove	Sample size : 5 pcs Requirement : <2mg/glove Result :0.24mg/glove	Pass

8.0 Summary of Clinical Testing:

No clinical study is included in this submission.

9.0 Conclusion

The conclusion drawn from the non-clinical tests demonstrate that the subject Violet Nitrile Powder Free Patient Examination Glove, Non-Sterile is as safe, as effective, and performs as well as or better than the legally marketed predicate devices K143289.