January 16, 2024

Nephron Nitrile, LLC.
Lou Kennedy
CEO/President/Manager
4777 12th Street Extension
West Columbia, South Carolina 29172

Re: K233970
Trade/Device Name: Nephron Nitrile™, Nitrile Powder-Free Examination Blue Gloves (Tested For Use With Chemotherapy Drugs and Fentanyl)
Regulation Number: 21 CFR 880.6250
Regulation Name: Non-Powdered Patient Examination Glove
Regulatory Class: Class I, reserved
Product Code: LZA, LZC, OPJ, QDO
Dated: December 15, 2023
Received: December 15, 2023

Dear Lou Kennedy:

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 (the 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 available 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.

Additional information about changes that may require a new premarket notification are provided in the FDA guidance documents entitled "Deciding When to Submit a 510(k) for a Change to an Existing Device" (https://www.fda.gov/media/99812/download) and "Deciding When to Submit a 510(k) for a Software Change to an Existing Device" (https://www.fda.gov/media/99785/download).

Your device is also subject to, among other requirements, the Quality System (QS) regulation (21 CFR Part 820), which includes, but is not limited to, 21 CFR 820.30, Design controls; 21 CFR 820.90, Nonconforming product; and 21 CFR 820.100, Corrective and preventive action. Please note that regardless of whether a change requires premarket review, the QS regulation requires device manufacturers to review and approve changes to device design and production (21 CFR 820.30 and 21 CFR 820.70) and document changes and approvals in the device master record (21 CFR 820.181).

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 Part 803) for devices or postmarketing safety reporting (21 CFR Part 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 Part 4, Subpart A) for combination products; and, if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR Parts 1000-1050.

Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 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,

Bifeng Qian -S

Bifeng Qian, M.D., Ph.D.
Assistant Director
DHT4B: Division of Infection Control
and Plastic and Reconstructive 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

Form Approved: OMB No. 0910-0120
Expiration Date: 07/31/2026
See PRA Statement below.

510(k) Number *(if known)*
K233970

Device Name
Nephron Nitrile™, Nitrile Powder-Free Examination Blue Gloves (Tested For Use With Chemotherapy Drugs and Fentanyl)

Indications for Use *(Describe)*
Nephron Nitrile™, Nitrile Powder-Free Examination Blue Gloves (Tested For Use With Chemotherapy Drugs and Fentanyl) is a disposable device intended for medical purpose that is worn on the examiner's hand to prevent contamination between patient and examiner. In addition, these gloves were tested for use with chemotherapy drugs and fentanyl in accordance with ASTM D6978-05 (2019) Standard Practice for Assessment of Resistance of Medical Gloves to Permeation by Chemotherapy Drugs.

The following chemotherapy drugs and concentration had NO breakthrough detected up to 240 minutes:
Bleomycin Sulfate (15.0 mg/ml)
Busulfan (6.0 mg/ml)
Carboplatin (10.0 mg/ml)
Cisplatin (1.0 mg/ml)
Cyclophosphamide (20.0 mg/ml)
Cytarabine (100.0 mg/ml)
Dacarbazine (10.0 mg/ml)
Daunorubicin HCl (5.0 mg/ml)
Docetaxel (10.0 mg/ml)
Doxorubicin HCl (2.0 mg/ml)
Epirubicin HCl (2.0 mg/ml)
Etoposide (20.0 mg/ml)
Fludarabine (25.0 mg/ml)
Fluorouracil (50.0 mg/ml)
Gemcitabine (38.0 mg/ml)
Idarubicin HCl (1.0 mg/ml)
Ifosfamide (50.0 mg/ml)
Irinotecan (20.0 mg/ml)
Mechlorethamine HCl (1.0 mg/ml)
Melphalan (5.0 mg/ml)
Methotrexate (25.0 mg/ml)
Mitomycin C (0.5 mg/ml)
Mitoxantrone HCl (2.0 mg/ml)
Paclitaxel (6.0 mg/ml)
Rituximab (10.0 mg/ml)
Trisenox (1.0 mg/ml)
Vincristine Sulfate (1.0 mg/ml)
The following chemotherapy drugs have low permeation times:
Carmustine (3.3 mg/ml) : 33.8 minutes
Thiotepa (10.0 mg/ml) : 128.1 minutes
Warning: Not for Use with: Carmustine, Thiotepa
The tested Opioid is:
Fentanyl Citrate Injection (100 mcg/2 mL)
Permeation: no breakthrough up to 240 minutes

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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Food and Drug Administration
Office of Chief Information Officer
Paperwork Reduction Act (PRA) Staff
PRAStaff@fda.hhs.gov

"An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB number."

K233970 - 510(K) SUMMARY

AS REQUIRED BY: 21CFR§807.92

A. APPLICANT INFORMATION

510(K) Owner's Name	Nephron Nitrile, LLC.
Address	4777 12th Street Extension, West Columbia, SC 29172.
Phone	844-937-3888
Fax	1-803-926-9853
E-mail	lkennedy@nephronpharm.com NitrileRegulatory@nephronnitrile.com
Contact Person	Lou Kennedy
Designation	Chief Executive Officer
Contact Number	1-803-569-3110
Contact Email	lkennedy@nephronpharm.com NitrileRegulatory@nephronnitrile.com
Preparation date	28 September 2023
Date Submitted	15 December 2023

B. DEVICE IDENTIFICATION

Name of the device	Nephron Nitrile™, Nitrile Powder-Free Examination Blue Gloves (Tested For Use With Chemotherapy Drugs and Fentanyl)
510(k) Number	K233970
Product proprietary or trade name	Nephron Nitrile™
Common or usual name	Nitrile Examination Glove (Tested For Use With Chemotherapy Drugs and Fentanyl)
Classification name	Non-Powdered Patient Examination Glove, Specialty
Device Classification	Class-1
Product Code	LZA, LZC, OPJ, QDO
Regulation Number	21 CFR 880.6250
Review Panel	General Hospital

C. PREDICATE DEVICE

Predicate Device	Nephron Nitrile Powder-Free Nitrile Examination Gloves (Tested For Use With Chemotherapy Drugs and Fentanyl)
510(k) Number	K231349
Regulatory Class	Class-1
Product code	LZA, LZC, OPJ, QDO
Manufacturer	Nephron Nitrile, LLC.

D. DESCRIPTION OF THE DEVICE:

Nephron Nitrile™, Nitrile Powder-Free Examination Blue Gloves (Tested For Use With Chemotherapy Drugs and Fentanyl) are Class I patient examination gloves, bearing the product codes LZA, LZC, OPJ, QDO (21 CFR 880.6250). They meet all the current specifications listed under the ASTM D6319-19, *Standard Specification for Nitrile Examination Gloves for Medical Application* and also complies with requirements for *Standard Practice for Assessment of Resistance of Medical Gloves to Permeation by Chemotherapy Drugs* as per ASTM D6978-05 (2019). The powder-free gloves are made from Nitrile (NBR) latex and are blue in color. The product is non-sterile, fingertip textured, ambidextrous with beaded cuff, and single use only.

Nephron Nitrile™, Nitrile Powder-Free Examination Blue Gloves (Tested For Use With Chemotherapy Drugs and Fentanyl) include glove sizes: X-Small, Small, Medium, Large, X-Large and XX-Large.

E. INDICATION FOR USE OF THE DEVICE:

Nephron Nitrile™, Nitrile Powder-Free Examination Blue Gloves (Tested For Use With Chemotherapy Drugs and Fentanyl) is a disposable device intended for medical purpose that is worn on the examiner's hand to prevent contamination between patient and examiner. In addition, these gloves were tested for use with chemotherapy drugs and fentanyl in accordance with ASTM D6978-05 (2019) Standard Practice for Assessment of Resistance of Medical Gloves to Permeation by Chemotherapy Drugs.

<table>
<tr><td colspan="3">The following chemotherapy drugs and concentration had NO breakthrough detected up to 240 minutes:</td><td>The tested Opioid is:</td></tr>
<tr><td>Bleomycin Sulfate (15.0 mg/ml)
Busulfan (6.0 mg/ml)
Carboplatin (10.0 mg/ml)
Cisplatin (1.0 mg/ml)
Cyclophosphamide (20.0 mg/ml)
Cytarabine (100.0 mg/ml)
Dacarbazine (10.0 mg/ml)
Daunorubicin HCl (5.0 mg/ml)
Docetaxel (10.0 mg/ml)</td><td>Doxorubicin HCl (2.0 mg/ml)
Epirubicin HCl (2.0 mg/ml)
Etoposide (20.0 mg/ml)
Fludarabine (25.0 mg/ml)
Fluorouracil (50.0 mg/ml)
Gemcitabine (38.0 mg/ml)
Idarubicin HCl (1.0 mg/ml)
Ifosfamide (50.0 mg/ml)
Irinotecan (20.0 mg/ml)</td><td>Mechlorethamine HCl (1.0 mg/ml)
Melphalan (5.0 mg/ml)
Methotrexate (25.0 mg/ml)
Mitomycin C (0.5 mg/ml)
Mitoxantrone HCl (2.0 mg/ml)
Paclitaxel (6.0 mg/ml)
Rituximab (10.0 mg/ml)
Trisenox (1.0 mg/ml)
Vincristine Sulfate (1.0 mg/ml)</td><td rowspan="3">Fentanyl Citrate Injection
(100 mcg/2 mL)

Permeation: no breakthrough up to 240 minutes</td></tr>
<tr><td colspan="3">The following chemotherapy drugs have low permeation times:
Carmustine (3.3 mg/ml) : 33.8 minutes Thiotepa (10.0 mg/ml) : 128.1 minutes</td></tr>
<tr><td colspan="3">Warning: Not for Use with: Carmustine, Thiotepa</td></tr>
</table>

F. SUMMARY OF THE TECHNOLOGICAL CHARACTERISTICS OF THE DEVICE COMPARED TO THE PREDICATE DEVICE

CHARACTERISTICS	STANDARDS	DEVICE PERFORMANCE		Comparison
		PREDICATE	SUBJECT	
510(K) Number	---	K231349	K23970	
Name of device	---	Nephron Nitrile Powder-Free Nitrile Examination Gloves (Tested For Use With Chemotherapy Drugs and Fentanyl)	Nephron Nitrile™, Nitrile Powder-Free Examination Blue Gloves (Tested For Use With Chemotherapy Drugs and Fentanyl)	**Similar**
Product Code	---	LZA, LZC, OPJ, QDO	LZA, LZC, OPJ, QDO	**Identical**
Indication for use	---	Nephron Nitrile Powder-Free Nitrile Examination Gloves (Tested For Use With Chemotherapy Drugs and Fentanyl) is a disposable device intended for medical purpose that is worn on the examiner's hand to prevent contamination between patient and examiner. In addition, these gloves were tested for use with chemotherapy drugs and fentanyl in accordance with ASTM D6978-05 (2019) Standard Practice for Assessment of Resistance of Medical Gloves to Permeation by Chemotherapy Drugs.	Nephron Nitrile™, Nitrile Powder-Free Examination Blue Gloves (Tested For Use With Chemotherapy Drugs and Fentanyl) is a disposable device intended for medical purpose that is worn on the examiner's hand to prevent contamination between patient and examiner. In addition, these gloves were tested for use with chemotherapy drugs and fentanyl in accordance with ASTM D6978-05 (2019) Standard Practice for Assessment of Resistance of Medical Gloves to Permeation by Chemotherapy Drugs.	**Identical**
Regulation Number	---	21 CFR 880.6250	21 CFR 880.6250	**Identical**
Material	---	Nitrile	Nitrile	**Identical**
Color	---	Blue Gloves	Blue	**Identical**
Size	---	M, L, XL, XXL	XS, S, M, L, XL, XXL	**Similar** Adding size XS and S
Single Use	---	Single-use	Single-use	**Identical**
Sterile/Non-sterile	---	Non-Sterile	Non-Sterile	**Identical**
Rx Only or OTC	---	OTC	OTC	**Identical**

K233970 - 510(K) SUMMARY

AS REQUIRED BY: 21CFR§807.92

CHARACTERISTICS	STANDARDS	DEVICE PERFORMANCE		Comparison
		PREDICATE	SUBJECT	
510(K) Number	---	K231349	K23970	
Dimensions - Length	ASTM D6319-19	Minimum 230 mm (sizes M – XXL)	Minimum 220 mm (Sizes XS-S) Minimum 230 mm (sizes M – XXL)	**Similar** Adding size XS and S
Dimensions - Width	ASTM D6319-19	M: 95±10 mm L: 110±10 mm XL: 120±10 mm XXL: 130±10 mm	XS: 70±10 mm S: 80±10 mm M: 95±10 mm L: 110±10 mm XL: 120±10 mm XXL: 130±10 mm	**Similar** Adding size XS and S
Physical Properties- Tensile Strength	ASTM D6319-19	Before aging 14MPa, min	Before aging 14MPa, min	**Identical**
		After aging 14MPa, min	After aging 14MPa, min	**Identical**
Physical Properties- Ultimate Elongation	ASTM D6319-19	Before aging 500%, min	Before aging 500%, min	**Identical**
		After aging 400%, min	After aging 400%, min	**Identical**
Thickness	ASTM D6319-19	Palm: Minimum 0.05 mm Finger: Minimum 0.05 mm	Palm: Minimum 0.05 mm Finger: Minimum 0.05 mm	**Identical**
Powder Free Residue	ASTM D6319-19	≤ 2 mg per glove	≤ 2 mg per glove	**Identical**
Freedom from holes	ASTM D5151-2019	In accordance with ASTM D 5151-19, following ASTM D6319- 19, G-I, AQL 2.5	In accordance with ASTM D 5151-19, following ASTM D6319- 19, G-I, AQL 2.5	**Identical**
Chemotherapy Drugs Tested with Minimum Breakthrough Detection Time	ASTM D6978-05 (2019)	Bleomycin Sulfate 15 mg/ml (15,000 ppm) >240 Minutes	Bleomycin Sulfate 15 mg/ml (15,000 ppm) >240 Minutes	**Identical**
		Busulfan 6 mg/ml (6,000 ppm) >240 Minutes	Busulfan 6 mg/ml (6,000 ppm) >240 Minutes	**Identical**
		Carboplatin 10 mg/ml (10,000 ppm) >240 Minutes	Carboplatin 10 mg/ml (10,000 ppm) >240 Minutes	**Identical**
		Carmustine 3.3 mg/ml (3,300 ppm) 33.8 Minutes	Carmustine 3.3 mg/ml (3,300 ppm) 33.8 Minutes	**Identical**
		Cisplatin 1 mg/ml (1,000 ppm) >240 Minutes	Cisplatin 1 mg/ml (1,000 ppm) >240 Minutes	**Identical**
		Cyclophosphamide 20 mg/ml (20,000 ppm) >240 Minutes	Cyclophosphamide 20 mg/ml (20,000 ppm) >240 Minutes	**Identical**

CHARACTERISTICS	STANDARDS	DEVICE PERFORMANCE		Comparison
		PREDICATE	SUBJECT	
510(K) Number	---	K231349	K23970	
		Cytarabine 100 mg/ml (100,000 ppm) >240 Minutes	Cytarabine 100 mg/ml (100,000 ppm) >240 Minutes	**Identical**
		Dacarbazine 10 mg/ml (10,000 ppm) >240 Minutes	Dacarbazine 10 mg/ml (10,000 ppm) >240 Minutes	**Identical**
		Daunorubicin HCl 5 mg/ml (5,000 ppm) >240 Minutes	Daunorubicin HCl 5 mg/ml (5,000 ppm) >240 Minutes	**Identical**
		Docetaxel 10 mg/ml (10,000 ppm) >240 Minutes	Docetaxel 10 mg/ml (10,000 ppm) >240 Minutes	**Identical**
		Doxorubicin HCl 2 mg/ml (2,000 ppm) >240 Minutes	Doxorubicin HCl 2 mg/ml (2,000 ppm) >240 Minutes	**Identical**
		Epirubicin HCl 2 mg/ml (2,000 ppm) >240 Minutes	Epirubicin HCl 2 mg/ml (2,000 ppm) >240 Minutes	**Identical**
		Etoposide 20 mg/ml (20,000 ppm) >240 Minutes	Etoposide 20 mg/ml (20,000 ppm) >240 Minutes	**Identical**
		Fludarabine 25 mg/ml (25,000 ppm) >240 Minutes	Fludarabine 25 mg/ml (25,000 ppm) >240 Minutes	**Identical**
		Fluorouracil 50 mg/ml (50,000 ppm) >240 Minutes	Fluorouracil 50 mg/ml (50,000 ppm) >240 Minutes	**Identical**
		Gemcitabine 38 mg/ml (38,000 ppm) >240 Minutes	Gemcitabine 38 mg/ml (38,000 ppm) >240 Minutes	**Identical**
		Idarubicin HCl 1 mg/ml (1,000 ppm) >240 Minutes	Idarubicin HCl 1 mg/ml (1,000 ppm) >240 Minutes	**Identical**
		Ifosfamide 50 mg/ml (50,000 ppm) >240 Minutes	Ifosfamide 50 mg/ml (50,000 ppm) >240 Minutes	**Identical**
		Irinotecan 20 mg/ml (20,000 ppm) >240 Minutes	Irinotecan 20 mg/ml (20,000 ppm) >240 Minutes	**Identical**
		Mechlorethamine HCl 1 mg/ml (1,000 ppm) >240 Minutes	Mechlorethamine HCl 1 mg/ml (1,000 ppm) >240 Minutes	**Identical**
		Melphalan 5 mg/ml (5,000 ppm) >240 Minutes	Melphalan 5 mg/ml (5,000 ppm) >240 Minutes	**Identical**
		Methotrexate 25 mg/ml (25,000 ppm) >240 Minutes	Methotrexate 25 mg/ml (25,000 ppm) >240 Minutes	**Identical**
		Mitomycin C 0.5 mg/ml (500 ppm) >240 Minutes	Mitomycin C 0.5 mg/ml (500 ppm) >240 Minutes	**Identical**

CHARACTERISTICS	STANDARDS	DEVICE PERFORMANCE		Comparison
		PREDICATE	SUBJECT	
510(K) Number	---	K231349	K23970	
		Mitoxantrone HCl 2 mg/ml (2,000 ppm) >240 Minutes	Mitoxantrone HCl 2 mg/ml (2,000 ppm) >240 Minutes	**Identical**
		Paclitaxel 6 mg/ml (6,000 ppm) >240 Minutes	Paclitaxel 6 mg/ml (6,000 ppm) >240 Minutes	**Identical**
		Rituximab 10 mg/ml (10,000 ppm) >240 Minutes	Rituximab 10 mg/ml (10,000 ppm) >240 Minutes	**Identical**
		Thiotepa 10 mg/ml (10,000 ppm) 128.1 Minutes	Thiotepa 10 mg/ml (10,000 ppm) 128.1 Minutes	**Identical**
		Trisenox 1 mg/ml (1,000ppm) >240 Minutes	Trisenox 1 mg/ml (1,000ppm) >240 Minutes	**Identical**
		Vincristine Sulfate 1 mg/ml (1,000 ppm) >240 Minutes	Vincristine Sulfate 1 mg/ml (1,000 ppm) >240 Minutes	**Identical**
Opioid Drugs Tested with Minimum Breakthrough Detection Time	ASTM D6978-05 (2019)	Fentanyl Citrate Injection (100mcg/2mL) >240 Minutes	Fentanyl Citrate Injection (100mcg/2mL) >240 Minutes	**Identical**
Biocompatibility	Primary Skin Irritation- ISO 10993-23: First Edition 2021-01	Under the conditions of the study, the test article met the requirements of the test.	Under the conditions of the study, the test article met the requirements of the test.	**Identical**
	Dermal Sensitization- ISO 10993-10: Fourth Edition 2021-11	Under the conditions of the study, the test article was not considered a sensitizer.	Under the conditions of the study, the test article was not considered a sensitizer.	**Identical**
	In vitro cytotoxicity- ISO 10993-5: Third Edition 2009-06-01	Under the Conditions of the study, the undiluted test article extract and 50% test article extract dilution did not meet the requirements of the test and the 25%, 12.5%, 6.25%, and 3.13% test article extract dilutions met the requirements of the test.	Under the conditions of the study, the undiluted test article extract and 50% test article extract dilution did not meet the requirements of the test and the 25%, 12.5%, 6.25%, and 3.13% test article extract dilutions met the requirements of the test.	**Identical**
	Acute Systemic Toxicity- ISO 10993-11: Third Edition 2017-09	Under the conditions of the study, there was no mortality or evidence of systemic toxicity	Under the conditions of the study, there was no mortality or evidence of systemic toxicity	**Identical**

* Identical except for the expansion of glove sizes to include XS and S while maintaining existent M to XXL, which is the subject of this submission.

There are no significant differences between the products and are identical in terms of intended use, materials, design and manufacturing methods. The devices meet the ASTM standard D6319-19 and D6978-05 (2019).

G. NON-CLINICAL TESTING SUMMARY

BENCH TEST DATA

TEST METHOD	PURPOSE	ACCEPTANCE CRITERIA	RESULT		
ASTM D6319-19 Standard Specification for Nitrile Examination Gloves for Medical Application.	To determine the length of the gloves	Extra-Small: 220 mm min Small: 220 mm min Medium : 230 mm min Large : 230 mm min X-Large : 230 mm min XX-Large : 230 mm min	Extra-Small: 242 mm Small: 243 mm Medium : 235 mm* Large : 237 mm* X-Large : 250 mm* XX-Large : 238 mm*		
ASTM D6319-19 Standard Specification for Nitrile Examination Gloves for Medical Application.	To determine the width of the gloves	XS: 70±10 mm S: 80±10 mm M: 95±10 mm L: 110±10 mm XL: 120±10 mm XXL: 130±10 mm	Extra-Small: 74 mm Small: 85 mm Medium : 95 mm* Large: 113 mm* X-Large : 121 mm* XX-Large : 129 mm*		
ASTM D6319-19 Standard Specification for Nitrile Examination Gloves for Medical Application.	To determine the thickness of the gloves	Palm: 0.05 mm min for all sizes Finger: 0.05 mm min for all sizes	**Size** Extra-Small Small Medium* Large* X-Large* XX-Large*	**Palm** 0.112 mm 0.095 mm 0.077 mm 0.106 mm 0.089 mm 0.113 mm	**Finger** 0.145 mm 0.115 mm 0.111 mm 0.109 mm 0.115 mm 0.107 mm
ASTM D6319-19 Standard Specification for Nitrile Examination Gloves for Medical Application.	To determine the physical properties-Tensile strength	**Before Ageing** Tensile Strength 14MPa min for all sizes **After Ageing** Tensile Strength 14MPa min for all sizes	**Size** Extra-Small Small	**Before ageing** 33.74MPa 32.35MPa	**After ageing** 35.99MPa 32.14MPa
			The results for Medium, Large, X-Large and XX-Large are the same as the predicate.		
	To determine the physical properties-Ultimate Elongation	**Before Ageing** Ultimate Elongation 500% min for all sizes **After Ageing** Ultimate Elongation 400% min for all sizes	**Size** Extra-Small Small	**Before ageing** 565% 538%	**After ageing** 514% 435%
			The results for Medium, Large, X-Large and XX-Large are the same as the predicate.		

ASTM D5151-19 Standard Test Method for Detection of Holes in Medical Gloves	To determine the holes in the gloves	AQL 2.5	Gloves Passes AQL 2.5
ASTM D6124-06 (Reapproved 2017) Standard Test Method for Residual Powder on Medical Gloves	To determine the residual powder in the gloves	≤ 2 mg/glove	Extra-Small: 0.822 mg/glove Small: 0.147 mg/glove
			The results for Medium, Large, X-Large and XX-Large are the same as the predicate.

TEST METHOD	PURPOSE	ACCEPTANCE CRITERIA	RESULT
ASTM D6978-05 (Reapproved 2019) Standard Practice for Assessment of Resistance of Medical Gloves to Permeation by Chemotherapy Drugs.	To determine the breakthrough detection time of chemotherapy drugs	Bleomycin Sulfate 15mg/ml (15,000 ppm) >240 Minutes	Bleomycin Sulfate 15 mg/ml (15,000 ppm) >240 Minutes
		Busulfan 6 mg/ml (6,000 ppm) >240 Minutes	Busulfan 6 mg/ml (6,000 ppm) >240 Minutes
		Carboplatin 10 mg/ml (10,000 ppm) >240 Minutes	Carboplatin 10 mg/ml (10,000 ppm) >240 Minutes
		Carmustine 3.3 mg/ml (3,300 ppm) >240 Minutes	Carmustine 3.3 mg/ml (3,300 ppm) >240 Minutes
		Cisplatin 1 mg/ml (1,000 ppm) >240 Minutes	Cisplatin 1 mg/ml (1,000 ppm) >240 Minutes
		Cyclophosphamide 20 mg/ml (20,000 ppm) >240 Minutes	Cyclophosphamide 20 mg/ml (20,000 ppm) >240 Minutes
		Cytarabine 100 mg/ml (100,000 ppm) >240 Minutes	Cytarabine 100 mg/ml (100,000 ppm) >240 Minutes
		Dacarbazine 10 mg/ml (10,000 ppm) >240 Minutes	Dacarbazine 10 mg/ml (10,000 ppm) >240 Minutes
		Daunorubicin HCl 5 mg/ml (5,000 ppm) >240 Minutes	Daunorubicin HCl 5 mg/ml (5,000 ppm) >240 Minutes
		Docetaxel 10 mg/ml (10,000 ppm) >240 Minutes	Docetaxel 10 mg/ml (10,000 ppm) >240 Minutes
		Doxorubicin HCl 2 mg/ml (2,000 ppm) >240 Minutes	Doxorubicin HCl 2 mg/ml (2,000 ppm) >240 Minutes
		Epirubicin HCl 2 mg/ml (2,000 ppm) >240 Minutes	Epirubicin HCl 2 mg/ml (2,000 ppm) >240 Minutes
		Etoposide 20 mg/ml (20,000 ppm) >240 Minutes	Etoposide 20 mg/ml (20,000 ppm) >240 Minutes
		Fludarabine 25 mg/ml (25,000 ppm) >240 Minutes	Fludarabine 25 mg/ml (25,000 ppm) >240 Minutes
		Fluorouracil 50 mg/ml (50,000 ppm) >240 Minutes	Fluorouracil 50 mg/ml (50,000 ppm) >240 Minutes

		Gemcitabine 38 mg/ml (38,000 ppm) >240 Minutes	Gemcitabine 38 mg/ml (38,000 ppm) >240 Minutes
		Idarubicin HCl 1 mg/ml (1,000 ppm) >240 Minutes	Idarubicin HCl 1 mg/ml (1,000 ppm) >240 Minutes
		Ifosfamide 50 mg/ml (50,000 ppm) >240 Minutes	Ifosfamide 50 mg/ml (50,000 ppm) >240 Minutes
		Irinotecan 20 mg/ml (20,000 ppm) >240 Minutes	Irinotecan 20 mg/ml (20,000 ppm) >240 Minutes
		Mechlorethamine HCl 1 mg/ml (1,000 ppm) >240 Minutes	Mechlorethamine HCl 1 mg/ml (1,000 ppm) >240 Minutes
		Melphalan 5 mg/ml (5,000 ppm) >240 Minutes	Melphalan 5 mg/ml (5,000 ppm) >240 Minutes
		Methotrexate 25 mg/ml (25,000 ppm) >240 Minutes	Methotrexate 25 mg/ml (25,000 ppm) >240 Minutes
		Mitomycin C 0.5 mg/ml (500 ppm) >240 Minutes	Mitomycin C 0.5 mg/ml (500 ppm) >240 Minutes
		Mitoxantrone HCl 2 mg/ml (2,000 ppm) >240 Minutes	Mitoxantrone HCl 2 mg/ml (2,000 ppm) >240 Minutes
		Paclitaxel 6 mg/ml (6,000 ppm) >240 Minutes	Paclitaxel 6 mg/ml (6,000 ppm) >240 Minutes
		Rituximab 10 mg/ml (10,000 ppm) >240 Minutes	Rituximab 10 mg/ml (10,000 ppm) >240 Minutes
		Thiotepa 10 mg/ml (10,000 ppm) >240 Minutes	Thiotepa 10 mg/ml (10,000 ppm) >240 Minutes
		Trisenox 1 mg/ml (1,000 ppm) >240 Minutes	Trisenox 1 mg/ml (1,000 ppm) >240 Minutes
		Vincristine Sulfate 1 mg/ml (1,000 ppm) >240 Minutes	Vincristine Sulfate 1 mg/ml (1,000 ppm) >240 Minutes
ASTM D6978-05 (Reapproved 2019) Standard Practice for Assessment of Resistance of Medical Gloves to Permeation by Chemotherapy Drugs.	To determine the breakthrough detection time of Opioid drugs	Fentanyl Citrate Injection (100mcg/2mL) >240 Minutes	Fentanyl Citrate Injection (100mcg/2mL) >240 Minutes

BIOCOMPATIBILITY DATA

TEST METHOD	PURPOSE	ACCEPTANCE CRITERIA	RESULT
ISO 10993-23 First edition 2021-01 Biological Evaluation of Medical Devices - Part 23, Tests for Irritation.	To evaluate the local dermal irritation of a test article extract following intracutaneous injection in rabbits.	Under the condition of study not an irritant	Under the conditions of the study, the test article met the requirements of the test
10993-10 Fourth edition 2021-11 Biological Evaluation of Medical Devices - Part 10, Tests for Skin Sensitization.	To evaluate the test item, for the skin sensitization in Guinea pigs by maximization test.	Under the conditions of the study, not a sensitizer	Under the conditions of the study, the test article was not considered a sensitizer
ISO 10993-5 Third Edition 2009-06-01 Biological Evaluation of Medical Devices - Part 5, Tests for In Vitro Cytotoxicity.	To determine the potential of a test article to cause cytotoxicity	Under the conditions of the study, non-cytotoxic	The undiluted test articles extract and 50% test article extract dilution did not meet the requirements of the test and the 25%, 12.5%, 6.25%, and 3.13% test article extract dilutions met the requirements of the test. Cytotoxicity concern was addressed by acute systemic toxicity testing.
ISO 10993-11 Third edition 2017-09 Biological Evaluation of Medical Devices - Part 11, Tests for Systemic Toxicity.	To evaluate the acute systemic toxicity of a test article, extract following injection in mice.	Under the conditions of study, the device extracts do not pose a systemic toxicity concern	Under the conditions of study, there was no mortality or evidence of systemic toxicity

The following non-clinical tests were performed:

- ASTM D6319-19 *Standard Specification for Nitrile examination Gloves for Medical Application.*
- ASTM D5151-19 *Standard Test Method for Detection of Holes in Medical Gloves.*
- ASTM D6124-06 (Reapproved 2017) *Standard Test Method for Residual Powder on Medical Gloves.*
- ASTM D6978-05 (Reapproved 2019) *Standard Practice for Assessment of Resistance of Medical Gloves to Permeation by Chemotherapy Drugs.*
- ISO 10993-23 First Edition 2021-01 *Biological Evaluation of Medical Devices - Part 23, Tests for Irritation.*
- ISO 10993-10 Fourth Edition 2021-11 *Biological Evaluation of Medical Devices - Part 10, Tests for Skin Sensitization.*
- ISO 10993-5 Third Edition 2009-06-01 Biological Evaluation of Medical Devices - Part 5, Tests for In Vitro Cytotoxicity.
- ISO 10993-11 Third Edition 2017-09 *Biological Evaluation of Medical Devices - Part 11, Tests for Systemic Toxicity.*

H. CLINICAL TESTING SUMMARY

Not applicable - Clinical data is not needed for gloves.

I. CONCLUSION

The conclusions drawn from the non-clinical test demonstrate that the subject device in 510(K) submission K233970, Nephron Nitrile™, Nitrile Powder-Free Examination Blue Gloves (Tested For Use With Chemotherapy Drugs and Fentanyl), is as safe, as effective, and performs as well as or better than the legally marketed predicate device K231349.