

K994064

PT. BERLIAN GLOVINDO

JLN. SUTOMO UJUNG 96
MEDAN 20235 - INDONESIA

Tel : 62-61-616944
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Page Numbers 2 of 2

(5) Technological characteristic of the gloves.

a. Dimensions

Sizes	XS	S	M	L	XL
Length	240 mm	240 mm	240 mm	240 mm	240 mm
Width	80 < mm	80±10 mm	95±10 mm	111±10 mm	>111mm
Thickness					
1. Cuff (min)	0.10 mm	0.10 mm	0.10 mm	0.10 mm	0.10 mm
2. Palm (min)	0.10 mm	0.10 mm	0.10 mm	0.10 mm	0.10 mm
3. Finger Tip (min)	0.10 mm	0.10 mm	0.10 mm	0.10 mm	0.10 mm

b. Physical Properties

	Before aging	After aging at 100°C 22 hrs.
Tensile Strength	14 Mpa (min.)	14 Mpa (min.)
Ultimate Elongation	700 % (min.)	500 % (min.)

c. Performance Requirement

Characteristic	Related Defects	Inspection Level	AQL
Visible defects	Stains, Lumps, Holes etc.	S-4	2.5
Watertight	Holes	S-4	2.5
Dimensions	Width Length & Thickness	S-2	4
Physical Properties	Before and after ageing	S-2	4

d. Weight of residual powder in medium size gloves : 0.5 ± 0.2 mg

e. Bio - Compatibility

g. Test Result as per ASTM D 3578 -95

(6) Performance data is the same as mentioned immediately above.

(7) Clinical data is not needed for gloves or for most devices cleared by the 510 (K) process.

(8) Non-clinical data

Gloves meet or exceed the ASTM D 3578 Standard.

Meets FDA pin hole requirement.

Meets labeling claim.



JAN 28 2000

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Mr. Emmy Tjoeng
Official Correspondent for
PT. Berlian Glaovindo
Jln. Sutomo Ujung 96
Medan 20235 - Indonesia

Re: K994064
Trade Name: Nitrile Examination Gloves - Powder Free
Regulatory Class: I
Product Code: LZA
Dated: November 30, 1999
Received: December 1, 1999

Dear Dr. Tjoeng:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to 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). You may, therefore, market the device, subject to the general controls provisions of the Act. 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.

If your device is classified (see above) into either class II (Special Controls) or class III (Pre-market Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Good Manufacturing Practice for Medical Devices: General (GMP) regulation (21 CFR Part 820) and that, through periodic GMP inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your pre-market notification submission does not affect any

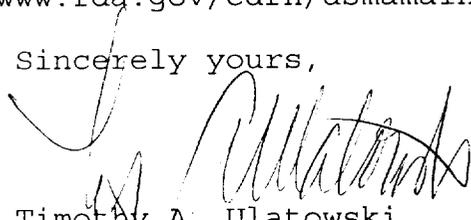
Page 2 - Mr. Tjoeng

obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4692. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "<http://www.fda.gov/cdrh/dsmamain.html>".

Sincerely yours,



Timothy A. Ulatowski
Director
Division of Dental, Infection Control
and General Hospital Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

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ANNEXURE II

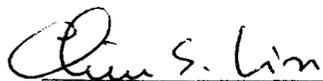
INDICATION FOR USE

Applicant : Mr. Tony Jap
Device Name : Nitrile Patient Examination Gloves Powder Free
Indication for use :

A patient examination glove is a disposable device intended for medical purpose that is worn on the examiner's hand or finger to prevent contamination between patient and examiners.

(PLEASE DO NOT WRITE BELOW THIS LINE. CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)



(Division Sign Off)
Division of Dental, Infection Control,
and General Hospital Devices
510(K) Number K994064

Prescription Use _____ OR Over-The-Counter Use X
(Per 21 CFR 801.109)