



Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

OCT 20 2000

Mr. James H. Lale
Chief Executive Officer
Biopro (M) SDN BHD
Lot 14, PT. 4204 Lingkarn Sultan
Hishamuddin, North Port Industrial Estate
Port Klang
MALAYSIA

Re: K994276
Trade Name: Biopro, Nitrilon, Correct Touch - nitrile
Powdered Examination Gloves
Regulatory Class: I
Product Code: LZA
Dated: October 4, 2000
Received: October 6, 2000

Dear Mr. Lale:

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 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). 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 Current Good Manufacturing Practice requirements, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic QS 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:

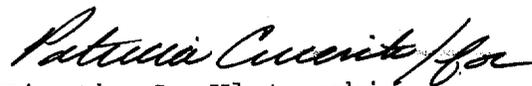
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this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

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" (21CFR 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/dsma/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

510(k) Submission - *Nitrile Powdered Examination Gloves (Green, Purple)*

Submitted by Biopro (M) Sdn Bhd Tel (603) 376-1390 Fax (603) 376-1787

3.0 Indications for Use Statement:

Applicant: Biopro (M) Sdn Bhd

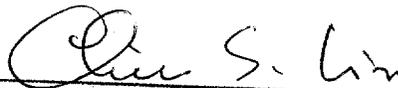
510(k) Number (if known): K994276

Device Name: Biopro™, Nitrilon™, Correct Touch^R or any other private labeling Nitrile Powdered Examination Gloves (Green, Purple)

Indications For Use: Biopro™, Nitrilon™, Correct Touch^R (Powdered) is a disposable patient examination glove which is worn on the hand of healthcare and similar personnel to prevent contamination between patient and examiner.

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

Concurrence of CDRH, Office of Devices Evaluation (ODE)



(Division Sign-Off)
Division of Dental, Infection Control,
and General Hospital Devices
510(k) Number K994276

Prescription Use _____
(Per 21 CFR 801.109)

OR

Over-The Counter Use X
(Optional Format 1-2-96)