



June 24, 2021

Careplus (M) SDN BHD
Lim Shyan
CEO/Managing Director
Lot 120 & 121, Jalan Senawang 3, Senawang Industrial Estate
Seremban, Negeri Sembilan 70450
Malaysia

Re: K210656

Trade/Device Name: ENCORE Latex Ortho, Powder Free Latex Surgical Gloves with Protein Content Labeling Claim ($<50\mu\text{g}/\text{dm}^2$ per glove of total aqueous extractable protein) and Tested for use with Chemotherapy drugs
GAMMEX Latex Ortho, Powder Free Latex Surgical Gloves with Protein Content Labeling Claim ($<50\mu\text{g}/\text{dm}^2$ per glove of total aqueous extractable protein) and Tested for use with Chemotherapy drugs

Regulation Number: 21 CFR 878.4460

Regulation Name: Non-Powdered Surgeon's Glove

Regulatory Class: Class I, reserved

Product Code: KGO

Dated: May 25, 2021

Received: May 27, 2021

Dear Lim Shyan:

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 and Part 809); 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, Ph.D.
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

Indications for Use

510(k) Number
k210656

Device Name

GAMMEX® Latex Ortho, Powder free Latex Surgical Gloves with Protein Content Labeling Claim (< 50 µg/dm² per glove of total aqueous extractable protein) and Tested for use with Chemotherapy drugs, ENCORE® Latex Ortho, Powder free Latex Surgical Gloves with Protein Content Labeling Claim (< 50 µg/dm² per glove of total aqueous extractable protein) and Tested for use with Chemotherapy drugs

Indications for Use (*Describe*)

Powder Free Surgical gloves are sterile disposable devices intended to be worn by operating room personnel to protect a surgical wound from contamination.

These gloves were tested for use with Chemotherapy Drugs as per ASTM D6978 Standard Practice for Assessment of Medical Gloves to Permeation by Chemotherapy:

Test Chemotherapy Drug (Concentration)	Minimum Breakthrough Detection Time in minutes
Fluorouracil (50.0mg/ml)	>240
Etoposide (20.0mg/ml)	>240
Cyclophosphamide (20.0mg/ml)	>240
Carmustine (3.3mg/ml)	22.0
Thiotepa (10.0mg/ml)	24.2
Paclitaxel (6.0mg/ml)	>240
Doxorubicin Hydrochloride (2.0mg/ml)	>240
Methotrexate (25.0mg/ml)	>240
Vincristine Sulfate (1.0mg/ml)	>24

"WARNING: Do not use with Carmustine and Thiotepa"

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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