



FEB 3 2003

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Shanghai Huamao Gloves Company Limited
C/O Ms. Susan D. Goldstein-Falk
MDI Consultants, Incorporated
55 Northern Boulevard, Suite 200
Great Neck, New York 11021

Re: K024154

Trade/Device Name: Shanghai Huamao Powder-Free Yellow Vinyl
Examination Gloves
Regulation Number: 880.6250
Regulation Name: Patient Examination Gloves
Regulatory Class: I
Product Code: LYZ
Dated: September 9, 2002
Received: December 17, 2002

Dear Ms. Falk:

This letter corrects our substantially equivalent letter of September 9, 2002, regarding the device name.

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. 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 (PMA), 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 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); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 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), please contact the Office of Compliance at (301) 594-4618. 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 Part 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers, International and Consumer 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,



Susan Runner, DDS, MA

Interim Director

Division of Anesthesiology, General Hospital,

Infection Control and Dental Devices

Office of Device Evaluation

Center for Devices and

Radiological Health

Enclosure

SHANGHAI HUAMAO GLOVES CO., LTD

No.8 NEW INDUSTRIAL AREA, ZHU HANG ZHEN, JINSHAN COUNTY, SHANGHAI, CHINA

C/O Room C, No:201 Nanking East Rd., Sec 3, Taipei, Taiwan,

Tel:886-2-25462480 Fax:886-2-2712-5051

August 30th 2002

INDICATIONS FOR USE

Applicant: Shanghai Huamao Gloves Co., Ltd

510(k) Number (if known):* K024154

Device Name: Shanghai Huamao Gloves Co., Ltd Powder-free Yellow Vinyl Examination Glove.

Indications For Use: A powderfree patient examination glove is a disposable device made of natural rubber latex or synthetic material that is intended to be worn on the hand or fingers for medical purposes to provide a barrier against poentially infectious and other contaminants.

Sincerely



Hsieh Yi Shi

Shanghai Huamao Gloves Co., Ltd

(Please refer to Attachment #3)

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

Concurrence of CDRH Office of Device Evaluation (ODE)

Prescription Use OR Over-The-Counter

Per 21 CFR 801.109

(Optional Format 11-2-96)

* For a new submission, do NOT fill in the 510(k) number.



(Division Sign-Off)

Division of Anesthesiology, General Hospital,
Infection Control, Dental Devices

510(k) Number: K024154