U.S. FOOD & DRUG ADMINISTRATION
CENTER FOR DEVICES AND RADIOLOGICAL HEALTH

U.S. Food & Drug Administration
10903 New Hampshire Avenue
Silver Spring, MD 20993
www.fda.gov

Certificate No. 8193-3-2018

CERTIFICATE TO FOREIGN GOVERNMENT

In order to allow the importation of United States products into foreign countries , the U.S. Food and Drug Administration (FDA) certifies the following information concerning the product(s) to be exported listed below:

Name of Product(s)

See Attached List

(One Page)

Name of Manufacturer/Distributor, Address

Name of Manufacturer
MEDICAL TECHNICAL PRODUCTS
1022 Fuller St
Santa Ana, CA USA 92701

The product(s) described above (and the manufacturing/distribution site(s) which produces/distributes it) is subject to the jurisdiction of the FDA under the Federal Food, Drug, and Cosmetic Act.

It is certified that the above product(s) may be marketed in, and legally exported from, the United States of America at this time. The manufacturing plant(s) in which the product(s) is produced is subject to periodic inspections. The last such inspection showed that the plant(s), at that time, appeared to be in substantial compliance with current good manufacturing practice requirements for the product(s) listed above.

CAPT Sean M. Boyd, MPH, USPHS
Deputy Director for Regulatory Affairs
Office of Compliance
Center for Devices and Radiological Health
U.S. Food and Drug Administration, DHHS

This certificate is valid from March 22, 2018 to March 21, 2020.

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Certificate to Foreign Government - Name of Product(s) Attachment Page 1 of 1

Name of Manufacturer

MEDICAL TECHNICAL PRODUCTS
1022 Fuller St
Santa Ana, CA USA 92701

Name of Product(s)

Model 2000 Ophthalmic Surgical System, Value
Model 2000 Ophthalmic Surgical System, Premium
Model 3000 Ophthalmic Surgical System
Trisonic Ophtalmic Surgical System
----------END OF PRODUCT LIST----------

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