



Certificate No. 5908-2-2021

## 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)

Name of Manufacturer/Distributor, Address

See Attached List

See Attached List

(One Page)

(One Page)

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 device product(s) may be marketed in, and legally exported from, the United States of America at this time. While the manufacturing plant(s) in which the device product(s) is produced is subject to inspection, FDA does not routinely inspect manufacturing firms that only make Class 1 medical devices. However, the firm has certified that it is currently operating in substantial compliance with current good manufacturing practice requirements for the device product(s) listed above.

Sincerely,

CDR Cesar A. Perez, PhD, Director DRP2: Division of Establishment Support

Office of Regulatory Programs

Office of Product Evaluation and Quality Center for Devices and Radiological Health U.S. Food and Drug Administration, DHHS

This certificate is valid from February 18, 2021 to February 17, 2023.





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

## Certificate No. 5908-2-2021 Certificate to Foreign Government - Name of Product(s) Attachment Page 1 of 1 Name of Corporate Headquarters

Biomiga, Inc. 10637 Roselle Street, Suite C San Diego, CA USA 92121

## Name of Manufacturer

BIOMIGA, INC. 10637 Roselle Street, Suite C San Diego, CA USA 92121

## Name of Product(s)

