U.S. FOOD & DRUG ADMINISTRATION

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

Certificate No. 1578-11-2019

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 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.

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 November 07, 2019 to November 06, 2021.

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Certificate to Foreign Government - Name of Manufacturer/Distributor Attachment Page 1 of 1

Name of Owner Operator

DGH TECHNOLOGY, INC.
110 SUMMIT DR., SUITE B
EXTON, PA USA 19341

Name of Manufacturer

DGH TECHNOLOGY, INC.
110 SUMMIT DR., SUITE B
EXTON, PA USA 19341

---END OF MANUFACTURER/DISTRIBUTOR LIST---

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Name of Product(s)

DGH 6000 Ultrasonic A-Scan
Scanmate Flex Ophthalmic Ultrasound System
DGH 8000 Ultrasonic B-Scan
DGH 55B / DGH 555B / DGH 55 / DGH 555 Ultrasonic Pachymeter
----------END OF PRODUCT LIST----------