



**Certificate No. 2161-11-2016**

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/Distributor**

LABOMED, INC.  
2728 S LA CIENEGA BLVD  
LOS ANGELES, CA USA 90034

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.

Carl Fischer, Ph.D.  
Director  
Division of International Compliance Operations  
Office of Compliance  
Center for Devices and Radiological Health

**This certificate is valid from November 23, 2016 to November 22, 2018.**





Food and Drug Administration  
10903 New Hampshire Avenue  
Silver Spring, MD 20993

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

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**Name of Product(s)**

UV-VIS SPECTROPHOTOMETERS  
MICROSCOPES  
DIAGNOSTIC INSTRUMENTS

-----END OF PRODUCT LIST-----

