



**Certificate No. 6414-6-2016**

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 Corporate Headquarters**

Cesca Therapeutics Inc. dba Thermogenesis Corp.  
2711 CITRUS RD. --  
Rancho Cordova, CA USA 95742

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.

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

**This certificate is valid from June 22, 2016 to June 21, 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)**

MXP® Device  
MXP® Startup Kit  
XpressTRAK software  
MXP® Docking Station.

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

