



Certificate No. 28545-6-2016

CERTIFICATE OF EXPORTABILITY [SECTION 801(e)(1)]

The Food and Drug Administration certifies that the product(s) described below is subject to its jurisdiction under the Federal Food, Drug, and Cosmetic Act (the Act). The products described below may not be sold or offered for sale in the United States. The company has certified to the Food and Drug Administration that:

- the product(s) accords to the specifications of the foreign purchaser;
- the product(s) is not in conflict with the laws of the country to which it is intended for export;
- the shipping package for the product(s) is labeled on the outside that it is intended for export; and
- the product(s) is not sold or offered for sale in the United States.

Based on the information above, the product(s) listed below may be exported pursuant to Section 801(e)(1) of the Act.

NAME OF PRODUCT

See Attached List
(3 Pages)

NAME OF COMPANY, ADDRESS

Diagnostic Automation/ Cortez
Diagnostics, Inc.
21250 Califa St, Suite 102 and 116
Woodland hills, CA 91367

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

This Certificate is valid from July 05, 2016 to July 05, 2018.

