



Certificate No. 10233-5-2017

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 Cosmetics 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(s)

See Attached List
(One Page)

Name of Company, Address

VISIONARY MEDICAL SUPPLIES, INC.
6441 Enterprise Ln Ste 216
Madison, WI USA 53719

Robin W. Newman MSN EdD CPNP
Director
Office of Compliance
Center for Devices and Radiological Health
U.S. Food and Drug Administration, DHHS

This certificate is valid from June 02, 2017 to June 01, 2019.





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Certificate of Exportability Section 801(e)(1) - Name of Product(s) Attachment Page 1 of 1

Name of Company, Address

VISIONARY MEDICAL SUPPLIES, INC.
6441 Enterprise Ln Ste 216
Madison, WI USA 53719

Name of Product(s)

VisiTryBlue (Trypan Blue Solution)
VisiTryBluePlus (Trypan Blue Solution)
VisiRing (Capsular Tension Ring)

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

