



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

Certificate No. 3589-7-2011

CERTIFICATE OF EXPORTABILITY (SECTION 802)

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). Such product(s), which is not approved for marketing in the United States, may be legally exported provided it meets the requirements of Section 802 of the Act.

Under Section 802 of the Act, a drug or device not approved for marketing in the United States may be exported if it is manufactured, processed, packaged, and held in substantial conformity with current good manufacturing practice requirements. 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 below. 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 802 of the Act.

Name of Product

See Attached List
(One Page)

Manufacturing Location

TEKIA, INC.
17 Hammond, Suite 414
IRVINE, CA, 92618 USA

Jennifer Medicus
Chief, Regulatory Policy and Systems Branch
Division of Risk Management Operations
Office of Compliance



This certificate expires 24 months from the date notarized.

COUNTY OF MONTGOMERY
STATE OF MARYLAND

Subscribed and sworn to before me this 31 day of July month 2011 year.

CATHRYN N. MORRIS
NOTARY PUBLIC STATE OF MARYLAND
County of Montgomery
My Commission Expires January 4, 2013



Certificate of Exportability Section 802 - Attachment (Page 1 of Page 1)

Name of Products

Manufacturing Location

- Silicone Intraocular Lenses , Model # 400 series
- Hydrophilic acrylic Intraocular Lenses ,
 - Model # 600 series,
 - Model # 600Y series,
 - Model # 800 series,
 - Model # 800Y series
- Phakic Intraocular Lenses, Model # 700 series
- Accommodative Intraocular Lenses, Model # 500 series

TEKIA, INC.
17 Hammond, suite 414
IRVINE, CA 92618 -
USA

"END OF PRODUCT LIST"

