U.S. FOOD & DRUG ADMINISTRATION
CENTER FOR DEVICES AND RADIOLOGICAL HEALTH

U.S. Food & Drug Administration
10903 New Hampshire Avenue
Silver Spring, MD 20993
www.fda.gov

Certificate No. 6134-3-2024-1

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 Cosmetics Act (the Act). Such product(s), which is not approved for marketing in the United States, may be legally exported to foreign countries 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 USA 92618

Sincerely,

CDR Cesar A. Perez, PhD, Director
DRP2: Division of Establishment Support
Office of Regulatory Programs
Office of Product Evaluation and Quality
Center for Devices and Radiological Health
U.S. Food and Drug Administration, DHHS

This certificate is valid from March 07, 2024 to March 06, 2026.

To verify the authenticity of the information on this certificate, you may scan the QR code or visit www.access.fda.gov/fecv/CDRH.

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Certificate of Exportability Section 802 - Name of Product(s) Attachment Page 1 of 1

Manufacturing Location

TEKIA, INC.
17 Hammond
suite 414
Irvine, CA USA 92618

Name of Product(s)

Hydrophobic acrylic Intraocular Lenses. Tek-Lens III: Model 900
Tek-Clear Intraocular Lenses. Models: 500, 500S
Three piece hydrophilic acrylic Intraocular Lenses. Tek-Lens II Models: 617, 617Y, 618 618Y. MeyeLens Model: 360
Single piece hydrophilic acrylic IOLs Tek-Lens II Models: 811, 811Y, 820, 820Y, 872, 872Y, 872S, 872YS. MeyeLens: 260, 460. Ophthalmo Pro AC 3000 HD
----------END OF PRODUCT LIST----------