U.S. FOOD & DRUG ADMINISTRATION
CENTER FOR FOOD SAFETY & APPLIED NUTRITION

Certificate Unique ID: YA9A-A3DK

CERTIFICATE OF FREE SALE

1. Pursuant to the Provisions of Rule 44 of the Federal Rules of Civil Procedure, I hereby certify that the attached letter (and product list, if applicable), as described below, is a true copy of material on file in the Food and Drug Administration, Department of Health and Human Services and is a part of the official records of said Administration and Department.

Attachment Dated:
August 12, 2024
To Whom it May Concern
Regarding:
Higo AKG Women (60)

Higo Corp, 9660 Flair Dr, , , ,, El Monte, CA 91731

2. In witness whereof, I have pursuant to the provisions of Title 42, United States Code, Section 3505, and the authority delegated by the Commissioner of Food and Drugs, hereto set my hand and cause the seal of the Department of Health and Human Services to be affixed this 12th day of August, 2024.

Haijing Hu, Ph.D.
Director, Regulatory Implementation Staff
Office of Dietary Supplement Programs
Center for Food Safety and Applied Nutrition
By direction of the Secretary of Health and Human Services

THIS CERTIFICATE EXPIRES: August 12, 2026.

Certificate Unique ID: YA9A-A3DK

August 12, 2024

TO WHOM IT MAY CONCERN:

We have reviewed correspondence on behalf of:

Higo Corp
9660 Flair Dr , , ,
El Monte, CA 91731

concerning the status of:

Higo AKG Women (60)

This product is regulated by the Food and Drug Administration (FDA) pursuant to the requirements of the Federal Food, Drug, and Cosmetic Act (FD&C Act) and the Fair Packaging and Labeling Act (FPLA) and other related laws.

The Food and Drug Administration does not have statutory authority to approve any food or any food manufacturer or distributor of such products.

The above referenced product is under the jurisdiction of the Food and Drug Administration which has primary responsibility for the administration and enforcement of the FD&C Act and the FPLA and other related laws. We have not examined the specific product being offered for export or reviewed the label. Under the FD&C Act, such a product may be exported if:

1. It is not adulterated or misbranded and it meets the other requirements of the FD&C Act for marketing in the U.S.; or

2. It cannot be lawfully marketed in the U.S. but meets the requirements of section 801(e) of the FD&C Act (21 U.S.C. 381(e)).

Sincerely yours,

Haijing Hu, Ph.D.
Director, Regulatory Implementation Staff
Office of Dietary Supplement Programs
Center for Food Safety and Applied Nutrition
U.S. Food and Drug Administration