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

Certificate Unique ID: ZPZ4-M79R

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:
January 21, 2021
To Whom it May Concern
Regarding:

Products Listed in the Attached Product List

MedHija LLC, 316 River Rd N, Wappingers Falls, NY 12590

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 21st day of January, 2021.

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: January 21, 2023.

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Certificate of Free Sale Product List

January 21, 2021

1. Pro-listem Stage One (90 Capsules)
2. Pro-listem Stage Two (90 Capsules)
3. Pro-listem Stage Three (90 Capsules)

NutraScience Labs, 70 Carolyn Blvd, Farmingdale, NY 11735

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January 21, 2021

TO WHOM IT MAY CONCERN:

We have reviewed correspondence on behalf of:

MedHija LLC
316 River Rd N
Wappingers Falls, NY 12590

concerning the status of:

Pro-listem Stage One (90 Capsules), Pro-listem Stage Two (90 Capsules), and Pro-listem Stage Three (90 Capsules)

These products are 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 products are 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 products being offered for export or reviewed the labels. Under the FD&C Act, such products 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