



Certificate Unique ID: Q685-VT9S

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:
February 25, 2026
To Whom it May Concern
Regarding:

Products Listed in the Attached Product List

Lavita, 1312 17th Street, , , Denver, CO 80202

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 25th day of February, 2026.

Haijing Hu, Ph.D.
Chief, Regulatory Operations Branch
Office of Dietary Supplement Programs
Human Foods Program
By direction of the Secretary of Health and Human Services

THIS CERTIFICATE EXPIRES: February 25, 2028.





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February 25, 2026

TO WHOM IT MAY CONCERN:

We have reviewed correspondence on behalf of:

Lavita
1312 17th Street ,
Denver, CO 80202

concerning the status of:

"Certificate of Free Sale Product List"

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.
Chief, Regulatory Operations Branch
Office of Dietary Supplement Programs
Human Foods Program
U.S. Food and Drug Administration



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ATTACHMENT TO "TO WHOM IT MAY CONCERN LETTER" PRODUCT LISTING

February 25, 2026

1. LAVITA B12 500MCG (60 TABLETS)
2. LAVITA WELLNESS GUARD (30 CAPSULES)
3. LAVITA BLOOD PRESSURE SUPPORT (30 CAPSULES)
4. LAVITA COQ10 100 MG (30 CAPSULES)
5. LAVITA FULL-SPECTRUM MULTI COMPLEX (90 TABLETS)
6. LAVITA MAGNESIUM BISGLYCINATE 200 MG (30 CAPSULES)
7. LAVITA D3 10,000 IU (250 MCG) (30 SOFTGELS)
8. LAVITA ZINC 50 MG (30 TABLETS)