



**CERTIFICATE OF FREE SALE**

1. Pursuant to the provisions of Rule 44 of the Federal Rules of Civil Procedure, I hereby certify that the copy attached (see attached list) 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.

Letter Dated

July 17, 2007

To Whom It May Concern

from, Robert J. Moore

regarding

**Antiaging Institute of California Control, 60 Softgels**  
**Antiaging Institute of California CoQ10 Red 100 mg, 60 Softgel Capsules**  
**Antiaging Institute of California Vitamin E with Selenium, 60 Softgel Capsules**  
**Antiaging Institute of California Lipid Care TT, 60 Softgels**  
**Antiaging Institute of California CoQ10 Red 30 mg, 30 Softgel Capsules**

2. In witness whereof, I have pursuant to the provisions of Title 42, United States Code, Section 3505, and 21 CFR 5.22, hereto set my hand and caused the seal of the Department of Health and Human Services to affixed this 17 day of July 2007.

Barbara Schneeman  
Director  
Office of Nutrition, Labeling  
and Dietary Supplements  
Center for Food Safety  
and Applied Nutrition

By direction of the Secretary  
of Health and Human Services

This Certificate expires on July 17, 2009.





JUL 17 2007

TO WHOM IT MAY CONCERN:

We have reviewed correspondence on behalf of:

Softgel Technologies, Inc.  
6982 Bandini Boulevard  
Los Angeles, CA 90040-3326

concerning the status of:

**Antiaging Institute of California Control, 60 Softgels**  
**Antiaging Institute of California CoQ10 Red 100 mg, 60 Softgel Capsules**  
**Antiaging Institute of California Vitamin E with Selenium, 60 Softgel Capsules**  
**Antiaging Institute of California Lipid Care TT, 60 Softgels**  
**Antiaging Institute of California CoQ10 Red 30 mg, 30 Softgel 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 (FLPA) and other related laws.

The Food and Drug Administration does not have statutory authority to approve or sanction 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 Federal Food, Drug and Cosmetic Act (FD&C Act) and related statutes. While we have not examined the specific products being offered for export or reviewed the labels, we can state that foods may be exported if they meet the specifications of section 801(e) of the FD&C Act (21 U.S.C. 318(e)).

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The issuance of this certification is for exportation of the products. Under section 801(e) of the Federal Food, Drug, and Cosmetic Act, a food or cosmetic intended for export shall not be deemed to be adulterated or misbranded if it:

- (a) accords to the specifications of the foreign government,
- (b) is not in conflict with the laws of the country to which it is intended for export,
- (c) is labeled on the outside of the shipping package that it is intended for export,
- (d) is not sold or offered for sale in domestic commerce.

Sincerely yours,

A handwritten signature in black ink, appearing to read "Robert J. Moore". The signature is fluid and cursive, written over a light blue horizontal line.

Robert J. Moore, Ph.D.  
Supervisor, Regulations Implementation Team  
Division of Dietary Supplement Programs  
Office of Nutrition, Labeling  
and Dietary Supplements  
Center for Food Safety  
and Applied Nutrition

# SOFT GEL

TECHNOLOGIES, INC.®

Date: 7/25/07

To Whom It May Concern:

The following Antiaging product(s) will be shipped in bulk quantity:

**CC-268 Antiaging Vitamin E w/Selenium**  
**CC-447 Antiaging Control**  
**CC-560 Antiaging Lipid Care TT**  
**CC-910 Antiaging CoQ10 Red 30 mg**  
**CC-930 Antiaging CoQ10 Red 100 mg**

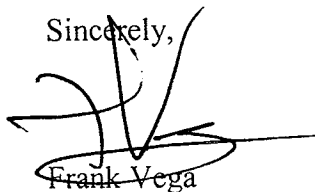
The above product(s) is/are free of gluten, yeast, sugar, preservatives, artificial color and flavor. Also the product(s) may be stored at room temperature. The expiration date is two years from date of manufacture.

This letter also certifies that the above product(s) is/are manufactured in the United States under all cGMP regulations.

With reference to BSE (Bovine Spongiform Encephalopathy), please be advised that Soft Gel Technologies, Inc. manufacturing facility only uses pharmaceutical grade gelatin in full compliance of all current United States Department of Agriculture (USDA) and the United States Food and Drug Administration (FDA) regulations. Additionally the gelatin used in the manufacturing process is derived from cattle born, raised and slaughtered in India where BSE has not been found nor expected to exist.

Soft Gel Technologies, Inc. uses gelatin that is derived only from the leg bone and shoulder bone of the Water Buffalo, no specified risk material, as defined in the July 30,1997 European Commission Decision 97/534/EC, are use to produce the gelatin.

Sincerely,



Frank Vega

Material Validation Clerk

**SOFT GEL TECHNOLOGIES**

6982 BANDINI BLVD.

LOS ANGELES, CA 90040 USA

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