



**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

June 28, 2011

To Whom It May Concern

from, Robert J. Moore

regarding

**Cyanotech Spirulina Pacifica Powder**  
**Cyanotech Spirulina Pacifica 200 mg Tablets**  
**Cyanotech Spirulina Pacifica 300 mg Tablets**  
**Cyanotech Spirulina Pacifica 400 mg Tablets**  
**Cyanotech Spirulina Pacifica 500 mg Tablets**  
**Cyanotech Spirulina Pacifica 1000 mg Tablets**  
**Cyanotech BioAstin Beadlets, Tablet Grade, 1.35% Astaxanthin**  
**Cyanotech BioAstin JointAstin Gelcaps, 3 mg Astaxanthin each**  
**Cyanotech BioAstin Gelcaps, 4 mg Astaxanthin each**  
**Cyanotech BioAstin Gelcaps, 2 mg Astaxanthin each**  
**Cyanotech BioAstin SCE5**

2. In witness whereof, I have pursuant to the provisions of Title 42, United States Code, Section 3505, and 1410.20 of the FDA Staff Manual Guide, hereto set my hand and cause the seal of the Department of Health and Human Services to be affixed this 28 day of June 2011.

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 June 28, 2013.





TO WHOM IT MAY CONCERN: JUN 28 2011

We have reviewed correspondence on behalf of:

Cyanotech Corporation  
73-4460 Queen Kaahumanu Highway, #102  
Kailua-Kona, HI 96740

concerning the status of:

**Cyanotech Spirulina Pacifica Powder**  
**Cyanotech Spirulina Pacifica 200 mg Tablets**  
**Cyanotech Spirulina Pacifica 300 mg Tablets**  
**Cyanotech Spirulina Pacifica 400 mg Tablets**  
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**Cyanotech BioAstin Gelcaps, 2 mg Astaxanthin each**  
**Cyanotech BioAstin SCE5**

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 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,

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