



U.S. Pharmacopeia
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April 27, 2011

Tom Nguyen
President & CEO
Robinson Pharma Inc.
3330 S. Harbor Blvd.
Santa Ana, CA 92704

DSVP Ref. No.: 1083100401

Subject: Fish Oil 30% EPA / DHA MEG-3 USP Softgels – Authorization to Use USP Verified Mark

Dear Mr. Nguyen:

I am pleased to inform you that, as of the date of this letter, the subject article, Fish Oil 30% EPA / DHA MEG-3 USP Softgels, submitted by Robinson Pharma, was found to be in compliance with the requirements of the USP Dietary Supplement Verification Program. This determination was based on our good manufacturing practices audits of Robinson Pharma's facilities located in Costa Mesa, CA and Santa Ana, CA; review of manufacturing, quality control and quality assurance documents; and testing of the following product lots:

Headquarters

12601 Twinbrook Parkway
Rockville, Maryland 20852
+1-301-881-0666

Europe/Middle East/Africa

Münchensteinerstrasse 41
CH-4052 Basel, Switzerland
+41 (0)61 316 30 10

USP-India Private Limited

ICICI Knowledge Park
Genome Valley
Labs 7-10, Phase III
Turkapally, Shameerpet
Ranga Reddy District
Hyderabad 500 078, A.P., India
+91-40-2348-0088

USP-China

Building 11
Lane 67 Libing Road
Zhangjiang Hi-Tech Park
Shanghai, 201203, China
+86-21-51370600

USP-Brazil

Avenida Ceci, 1600 - Tamboré
Barueri/SP, Brazil
06460-120
+55-11-3245-6400

Product Name	Product Code	Sample Lot # Bulk / Finished	Manufacture Date	Expiration Date
Fish Oil 30% EPA / DHA MEG-3 USP Softgels	RP-USPFO-1000	S8D128 / NA	June 2008	June 2010
		S8D129 / NA	June 2008	June 2010
		S8D130 / NA	June 2008	June 2010

Accordingly, during the term of the License Agreement, between Robinson Pharma and USP, Robinson Pharma may place the USP Verified Mark on the following approved dietary supplements, manufactured at Robinson Pharma located at 3330 S. Harbor Blvd., Santa Ana, CA 92704; and 1683 Sun Flower Ave., Costa Mesa, CA 92628.

Product Name	Product Code
Fish Oil 30% EPA / DHA MEG-3 USP Softgels	RP-USPFO-1000

This authorization is subject to the USP Verification Mark's usage requirements, described in the Verification Mark Usage Manual.

The USP Verified Mark may also be placed on private label brands of Robinson Pharma's customers, if the subject products are manufactured under the same conditions and at the same facility. Such use requires specific written authorization from USP. Please notify us in writing with your intent to utilize the USP Verified Mark on such products.

As a reminder, USP requires submission of artwork for pre-approval for product labels, advertising, promotional, collateral, or other materials that include the USP Verified Mark. The artwork may be submitted to the undersigned when ready.

Any major changes to a product's specifications, process control data, raw material source, manufacturing site change, testing or other criteria deemed by you to be essential or significant, must be reported in writing to USP. If necessary, USP may require the product to be re-evaluated and a facility audit to be conducted.

USP may also perform periodic "off-the-shelf" evaluations of the approved product to ensure that it continues to meet the criteria to carry the USP Verified Mark.

We are pleased to be able to verify the subject articles and look forward to continue working with you and your staff closely in this verification process.

Sincerely,



V. Srinivasan, Ph.D.
Head, International Sites and Verification Programs

Cc: Tracy Hunt, Senior Financial Analyst, USP