DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
College Park, MD 20740

Certificate Unique ID: ZP8N-5NEE

Application Number: 2015-13048

18 March 2015

CERTIFICATE

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

To Whom it May Concern letter dated

March 18, 2015

From Jon Hicks

Regarding

One (1) Product
Listed in the Attached Product List

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 to the Director, Office of Cosmetics and Colors, Center for Food Safety and Applied Nutrition, hereto set my hand and cause the seal of the Department of Health and Human Services to be affixed this March 18, 2015.

Linda M. Katz, M.D., MPH
Director, Office of Cosmetics and Colors
Center for Food Safety
and Applied Nutrition

By direction of the Secretary of
Health and Human Services

This Certificate expires on March 17, 2017.

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TO WHOM IT MAY CONCERN

We have received correspondence on behalf of:

Ms. Jessica Rodriguez
Hentek Products Factory, Inc.
13157 Flores St
Santa Fe Springs, CA 90670

concerning the status of cosmetic products that are intended to be exported from the United States.

If you are a foreign government official authorized by your government to accept export certificates, you may verify the authenticity of this export certificate by creating an online account at: http://www.access.fda.gov/oaa. Please note FDA does not require the authentication of cosmetic export certificates. This is an optional service to foreign governments that may require authentication. Once you have signed up for an account, to activate your account you will need to submit a document verifying your position, credentials, or authority to accept export certificates in an email to CAP-OCAC-CFSAN@fda.hhs.gov. You will be notified by email once your account has been activated. Once activated, login to http://www.access.fda.gov/oaa, access the Certificate Application Processing module, and enter the unique certificate ID located at the top left hand corner of the certificate and below to verify the certificate authenticity.

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Cosmetic products marketed in this country 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, as appropriate, the Fair Packaging and Labeling Act (FPLA). The FDA does not have the statutory authority to approve or sanction any cosmetic product or any manufacturer or distributor of such product. It is the responsibility of the manufacturer and/or distributor to market a safe and properly labeled product (i.e., one that is neither adulterated nor misbranded within the meaning of the FD&C Act and/or the FPLA, as applicable).

A cosmetic product may be exported if it meets the specifications of section 801(e) of the FD&C Act (21 U.S.C. 381(e)). In accordance with the provisions of section 801(e), a cosmetic product intended for export shall not be deemed to be adulterated or misbranded under the FD&C Act if it-

(a) accords to the specifications of the foreign purchaser,

(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, and

(d) is not sold or offered for sale in domestic commerce.

Cosmetic products may be exported in accordance with the provisions of section 801(e) of the FD&C Act but, when sold in domestic commerce, must comply with all applicable provisions of and regulations promulgated under the FD&C Act and/or the FPLA.

The certification of this document does not suggest or imply that FDA approves or sanctions the labels and labeling of the firm's products or that the firm's products are in compliance with the requirements of the FD&C Act and/or FPLA. Further, certification of this document does not preclude the Agency from taking regulatory action against the firm's products if such action is warranted.

Sincerely,

Jon Hicks
Office of Cosmetics and Colors
Center for Food Safety
and Applied Nutrition
Food and Drug Administration

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PRODUCT LIST

1. SOSORO (500 Milliliters (ml))

----------END OF PRODUCT LIST----------