

DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION
 College Park, MD 20740-3835

COSMETIC PRODUCT INGREDIENT STATEMENT

(In accordance with 21 CFR 720)

Form: OMB No. 0910-0027. Expiration Date: XX/XX/XXXX
 See Burden Statement on Reverse of Part I.

TYPE OF SUBMISSION

ORIGINAL AMENDED

FOR FDA USE ONLY ON ORIGINAL SUBMISSIONS

FDA CPIS NO.

FILING DATE (MM/DD/YY)

F _____

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NOTE: This report is authorized by Public Law 21 U.S.C. 371(A); 21 CFR 720. While you are not required to respond, your cooperation is needed to make the results of this voluntary program comprehensive, accurate, and timely.

INSTRUCTIONS:

For faster processing please submit this form electronically at: <http://www.fda.gov/Cosmetics/RegistrationProgram/OnlineRegistration/default.htm>.
 Type all entries in CAPITAL LETTERS. List ingredients in the order that they are listed on the label. If CAS number is not known, leave CAS NO. blank. List additional ingredients on a separate Form FDA 2512a. Attach completed form to FORM FDA 2512. If submitting a trade secret petition, enter "confidential" next to the ingredient in your petition.

| INGRED NO. | COMMON, USUAL, OR CHEMICAL NAME | CAS NO. |
|------------|---------------------------------|---------|
| 01 | | |
| 02 | | |
| 03 | | |
| 04 | | |
| 05 | | |
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THIS FORM MUST BE SECURELY ATTACHED TO FORM FDA 2512

This section applies only to requirements of the Paperwork Reduction Act of 1995.

DO NOT SEND YOUR COMPLETED FORM TO THE PRA STAFF EMAIL ADDRESS BELOW.

The burden time for this collection of information is estimated to average 20 minutes per response, including the time to review instructions, search existing data sources, gather and maintain the data needed and complete and review the collection of information. Send comments regarding this burden estimate or any other aspect of this information collection, including suggestions for reducing this burden, to:

Department of Health and Human Services
Food and Drug Administration
Office of Chief Information Officer
Paperwork Reduction Act (PRA) Staff
PRASStaff@fda.hhs.gov

*“An agency may not conduct or sponsor,
and a person is not required to respond to, a
collection of information unless it displays a
currently valid OMB number.”*