Expiration Date: April 30, 2014. Form Approved: OMB No. 0910-0027. DEPARTMENT OF HEALTH AND HUMAN SERVICES See Burden Statement on Reverse of Part I. FOOD AND DRUG ADMINISTRATION COLLEGE PARK, MD 20740-3835 TYPE OF SUBMISSION: ☐ ORIGINAL ☐ AMENDED ☐ DISC ☐ BASE COSMETIC PRODUCT INGREDIENT STATEMENT FOR FDA USE ONLY ON ORIGINAL SUBMISSIONS FDA CPIS NO. FILING DATE (In accordance with 21 CFR 720) Read Instruction Booklet Before Completing. Type entries in CAPITAL LETTERS. F 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. 01. NAME OF MANUFACTURER / PACKER / DISTRIBUTOR (On Label) 11. NAME OF MANUFACTURER / PACKER (Private Labeler) 02. KIND OF BUSINESS ☐ MFR □ PKR ☐ DISTR 03. NAME OF PARENT COMPANY (If any) 12. NAME OF PARENT COMPANY (If any) 04. COMPLETE MAILING ADDRESS: 13. COMPLETE MAILING ADDRESS: 15. PRODUCT CATEGORY CODE: 14. IS THIS STATEMENT FILED BY COMPANY 01 OR COMPANY 11? (Please check one) ☐ COMPANY 01 ☐ COMPANY 11 18. DATE OF ACTION BRAND NO. 16. BRAND NAME OF COSMETIC PRODUCT 17. TYPE OF ACTION 01 02 03 04

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19. TYPE NAME AND TITLE OF AUTHORIZED INDIVIDUAL 20. TELEPHONE NO. 21. SIGNATURE AND DATE

Public reporting burden for this collection of information is estimated to average 20 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to:

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OFFICIAL RECEIPT	FOR FDA USE ONLY
DEPARTMENT OF HEALTH AND HUMAN SERVICES FOOD AND DRUG ADMINISTRATION COLLEGE PARK, MD 20740-3835	
BRAND NAME OF COSMETIC PRODUCT	
COSMETIC PRODUCT INGREDIENT STATEMENT	
TO:	FDA CPIS NO. ¹ F
	FILING DATE
	THIS STATEMENT IS
	THIS STATEMENT IS
	☐ COMPLETE ☐ INCOMPLETE

misleading. 21 CFR 720.9

FORM FDA 2512 (9/11)