

TYPE OF SUBMISSION: ORIGINAL AMENDED

COSMETIC PRODUCT INGREDIENT STATEMENT

(In accordance with 21 CFR 720)

FOR FDA USE ONLY ON ORIGINAL SUBMISSIONS

FDA CPIS NO. **F** _____ FILING DATE _____

Read Instruction Booklet Before Completing. Type entries in CAPITAL LETTERS.

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.

INGRED NO.	1. COMMON, USUAL, OR CHEMICAL NAME							
01	2. 9 - DIGIT CAS NO.		3. TYPE OF ACTION		4. DATE OF ACTION		5. CONF	
	6. BASE CPIS NO. F _____ B _____		7. BASE NAME OR TRADE NAME MATERIAL			8. COMPANY NAME		
	1. COMMON, USUAL, OR CHEMICAL NAME							
02	2. 9 - DIGIT CAS NO.		3. TYPE OF ACTION		4. DATE OF ACTION		5. CONF	
	6. BASE CPIS NO. F _____ B _____		7. BASE NAME OR TRADE NAME MATERIAL			8. COMPANY NAME		
	1. COMMON, USUAL, OR CHEMICAL NAME							
03	2. 9 - DIGIT CAS NO.		3. TYPE OF ACTION		4. DATE OF ACTION		5. CONF	
	6. BASE CPIS NO. F _____ B _____		7. BASE NAME OR TRADE NAME MATERIAL			8. COMPANY NAME		
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04	2. 9 - DIGIT CAS NO.		3. TYPE OF ACTION		4. DATE OF ACTION		5. CONF	
	6. BASE CPIS NO. F _____ B _____		7. BASE NAME OR TRADE NAME MATERIAL			8. COMPANY NAME		
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05	2. 9 - DIGIT CAS NO.		3. TYPE OF ACTION		4. DATE OF ACTION		5. CONF	
	6. BASE CPIS NO. F _____ B _____		7. BASE NAME OR TRADE NAME MATERIAL			8. COMPANY NAME		
	1. COMMON, USUAL, OR CHEMICAL NAME							
06	2. 9 - DIGIT CAS NO.		3. TYPE OF ACTION		4. DATE OF ACTION		5. CONF	
	6. BASE CPIS NO. F _____ B _____		7. BASE NAME OR TRADE NAME MATERIAL			8. COMPANY NAME		
	1. COMMON, USUAL, OR CHEMICAL NAME							
07	2. 9 - DIGIT CAS NO.		3. TYPE OF ACTION		4. DATE OF ACTION		5. CONF	
	6. BASE CPIS NO. F _____ B _____		7. BASE NAME OR TRADE NAME MATERIAL			8. COMPANY NAME		
	1. COMMON, USUAL, OR CHEMICAL NAME							
08	2. 9 - DIGIT CAS NO.		3. TYPE OF ACTION		4. DATE OF ACTION		5. CONF	
	6. BASE CPIS NO. F _____ B _____		7. BASE NAME OR TRADE NAME MATERIAL			8. COMPANY NAME		
	1. COMMON, USUAL, OR CHEMICAL NAME							
09	2. 9 - DIGIT CAS NO.		3. TYPE OF ACTION		4. DATE OF ACTION		5. CONF	
	6. BASE CPIS NO. F _____ B _____		7. BASE NAME OR TRADE NAME MATERIAL			8. COMPANY NAME		
	1. COMMON, USUAL, OR CHEMICAL NAME							
10	2. 9 - DIGIT CAS NO.		3. TYPE OF ACTION		4. DATE OF ACTION		5. CONF	
	6. BASE CPIS NO. F _____ B _____		7. BASE NAME OR TRADE NAME MATERIAL			8. COMPANY NAME		
	1. COMMON, USUAL, OR CHEMICAL NAME							

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

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and a person is not required to respond to, a
collection of information unless it displays a
currently valid OMB number.”*