Annex 14

Ingredient Safety Related Information

[Ingredient commodity name, ingredient basic information and brief description of ingredient production process]

*Mandatory Fields

Name of the ingredient manufacturer*				
(Chinese name, for a domestic enterprise) (English name, for an overseas enterprise)				
Ingredient brand name*				
NMPA Registration	YES, please add the info below and skip to the last page for signature		NO, please complete the remaining sections	
Number Available?*	NMPA n.			
	Component	Component Chinese name The name of the component should be consistent with that specified in the Inventory of Existing Cosmetic Ingredients in China (IECIC)	Component INCI name	Range of percentages
	1			
Ingredient composition	2			
(including compound components)	3			
		e of addition of ingredients in cosmetics should be determined from the perspective ddition, it should be indicated separately.	e of safety or efficacy. If there are differences in non-rinse off cosmetics	and rinse-off cosmetics in terms of th
Recommended maximum addition amount in cosmetics				
Ingredient use restrictions	use restrictions, incompatibility and labeling requirements for warning language of the ingredient in cosmetics, etc.			
(if any)				

	Color	
Ingredient traits	Odor	
	State	
Description of physical and chemical properties		
	The type of production process shall be specified: physical crushing; physical pressing; extraction with water or other kinds of solvents; chemical synthesis; biological fermentation. The processes of stirring, heating, distillation, filtration, drying and packaging shall be briefly summarized. For natural ingredient directly derived from animals and plants, the species information, Latin name, and extraction parts shall be clear (as reference for algae and macrofungi). For biotechnology source ingredients ¹ , the necessary information such as the gene source, vector construction, engineering bacteria information, donor organisms, recipient organisms, modified microorganisms, <i>etc.</i> used in the production shall be specified.	
Description of production		
process type		

(If the ingredient composition cannot be provided, it may not be filled in, but relevant explanatory documents shall be submitted)

[Quality control requirements]

Serial number	Indicator name	Molecular formula or structural formula (if can be specified)	CAS number (if can be specified)	Name of test method	Quantitative range
1					
2					
3					
4					

¹ Biotechnology source ingredients include hydrolyzed plant components

(If unavailable, it may not be filled in, but relevant explanatory documents shall be submitted)

- For a single ingredient with a clear chemical structure, purity requirements shall be provided
- For an ingredient with unclear chemical structure, quantitative requirements for index components or total components, evaporation residue/solid content, loss on drying/moisture, residue on ignition, typical physical and chemical indexes etc. shall be provided
- For polymer ingredients, the degree of polymerization and average molecular weight shall be clarified simultaneously, ifany
- For oligopeptides, the amino acid sequence shall also be specified

1. Identification method

(If cannot be provided, relevant explanatory materials shall be submitted)

- 2. Test method of quantitative control indicators/characteristic indicators (one-to-one correspondence with the indicators in the above table) (For the testing method, only the method name is required)
- 3. Microbial indicator (if applicable)

there is any	describe it briefly and the authoritative organizati	ion name shall be attached		
unere is arry,	describe it briefly and the authoritative organizati	on name shall be attached		
ief descrip	otion of requirements for use in other in	ndustries]		
equiremen	ts for limit of substances with risks] (if	any)		
Serial number	Name of substances with risks	CAS number	Requirements for limit	Notes
1				
2				
Risk of pe	etal indicators (if applicable): esticide residues: (only for ingredients fron pollution control: (only for ingredients fron f host pathogenic and toxic ingredients: (o	n biotechnology sources, if appl		
ther issue	s that need explanation]			

Company seal of ingredient manufacturer or authorized enterprise filling out this form*

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	Date [Month] [Day] [Year]*