# 新語國際有限公司 Sin Yi International Corp



### SINYI ACHIEVE YOUR WISHES





#### Discover your uniqueness

**SINYI** Provide Taiwan cosmetics PIF and product registration. creates original designs that can be customized to fit your aesthetic. From small logos, to completely new product developments, we specialize in turning your product ideas into reality. Our innovation team is agile, knowledgeable, easy to communicate with, and fully respects your confidentiality.

#### Weave your dream catcher

**SINYI** can realize your innovative product concept to solution From engineering to manufacturing. From ethics to compliance Social responsibility to global product legislation Delivered with your design.

## All of SINYI

#### QUALITY

**SINYI** ensure that all the products we sell are organic, high quality and gentle on the skin.

#### EXPECT

**SINYI** focus on lower cost (tariff) of the global supply chain, but higher price goods, we can also provide them.

#### INNOVATION

**SINYI** love making fun products, be it personal care or lifestyle amenities Talk to us and let's make your Idea come true.

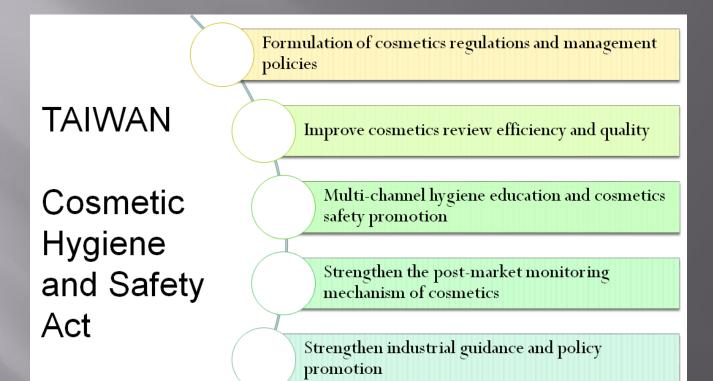


## WORLD

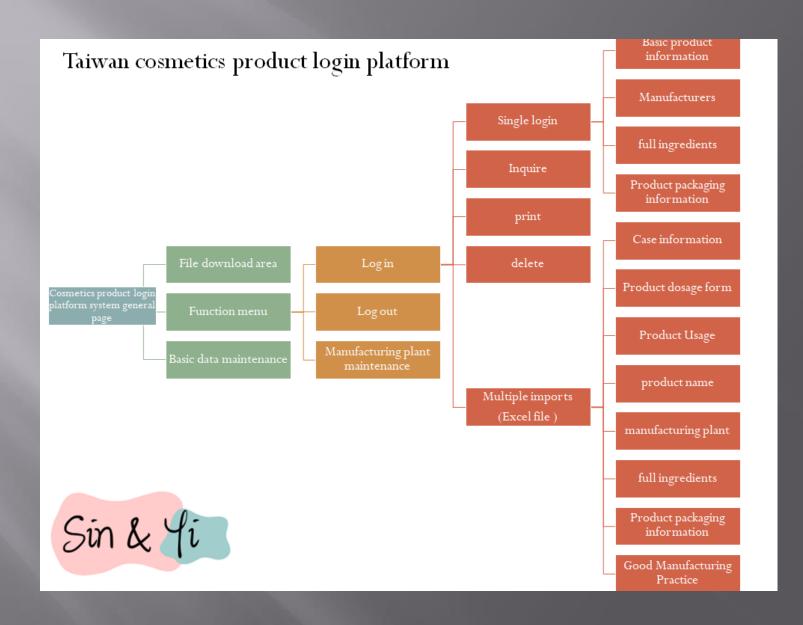
LET SINYI HELP YOUR DREAMS GO ANYWHERE.







Strengthen cosmetics international and cross-strait exchanges and cooperation



I. Product description						
1	Basic product information					
	1.1	Product name				
	1.2	Product category, Product dosage form				
	1.3	Purpose				
	1.4	Manufacturer name and address				
	1.5	Filling plant information, Packing plant information, Product manufacturer information (name, address and phone number)				
2	Evidentiary document of product notification completion					
3	Full ingredient name and content of specific ingredient -Ask the manufacturer to provide a complete formula sheet including Formula raw material names, INCI name, CAS No., Function for being added, Content w/w (%). Label in Chinese or English.					
4	<b>Product label, leaflet, outer packaging or container</b> -Please ask the manufacturer to provide the inner and outer packaging, labels and instructions, or present them as photos.					
5	<b>Evidentiary document or statement for cosmetics manufacturing facilities that comply with the Cosmetic</b> <b>Good Manufacturing Practice Regulations</b> -Ask the manufacturer to provide the cosmetics GMP certificate issued by the competent authority or the ISO 22716 certificate issued by the certification body					
6	Manufacturing method, procedure -The manufacturer should provide the manufacturing method of the product.					
7	Usage method, applied body part, dosage, use frequency and group -Ask the manufacturer to provide the usage method of the product, which can be explained by using words or pictures on the package.					
8	Adverse effects generated by product under normal use -It is not necessary to provide if there is no adverse reaction to the product. If there is any adverse reaction, the type of adverse reaction, treatment method and corrective and preventive measures must be stated. -Any information related to adverse reactions or serious adverse reactions should be included, and can be presented in a statistical manner, such as the number or frequency of adverse reactions per 10,000 bottles or per year.					

II. Data regarding quality				
9	<b>Physical and chemical properties of the product and its individual ingredient</b> -Ask the manufacturer to provide the quality specifications of the product and the ingredients contained in it (according to CNS 22716) or the Certificate of Analysis(COA), Safety Data Sheet (SDS).			
	9.1	Physical and chemical properties of the product		
	9.2	Physical and chemical properties of its individual ingredient		
10	<b>Toxicological information of ingredients</b> - Ingredient toxicology information mainly comes from raw material suppliers. You can learn about the characteristics, risks, hazards, hazard prevention, and toxicology of the ingredients through the provided Safety Data Sheets (SDS). Note that ingredients with the same name but come from different manufacturers may have different toxicological characteristics due to different production processes, so the SDS information must be correct and the source of the data must be listed.			
11*	The product stability test reports - Manufacturers shall provide product accelerated stability or long-term stability test data.			
12*	<b>The microbiological test reports</b> -Ask the manufacturer to provide microbiological test reports, including the specifications for microbiological control of raw materials and finished products, the test methods, and test results.			
13*	The antimicrobial effectiveness test reports - Manufacturers shall provide the antimicrobial effectiveness test data and the testing methods used. - Relevant proofs shall be provided for microbiologically low-risk products that do not require an antimicrobial effectiveness test. If the signatory for the safety report thinks that the test should be conducted, the test shall be conducted.			

III. Functional support				
14	Supporting documents of functional assessments - Products containing specific ingredients announced by the central competent authority shall comply with regulations of the limitation of quantity of ingredients and testing reports shall be provided or if there are claims for specific functions, supporting information shall be provided.			
IV. Safety Assessment Information				
15	Information about the packaging materials which have contact with the products - Manufacturers shall provide standard testing reports and quality specifications for packaging materials, and packaging material suppliers shall provide supporting information for packaging materials or packaging material test reports, stability of packaging materials, extraction substance of packaging materials or compatibility test.			
16	Product safety information			
	16.1	Safety conclusion and suggestions - Safety assessment shall cover the safety conclusion provided by the Safety Assessor for the safety report based on data analysis of chemical structures, exposure dosage, interactions of the ingredients, and toxicology research; warning labels, direction for use. The Safety Assessor shall sign name and date on the safety report.		
	16.2	Supporting qualification documents for Safety Assessor - Including the Safety Assessor's name and education; cosmetic safety evaluation training qualification certificate recognized by the central competent authority.		

### Connect With Us



Whats App:Sin Yi Cosmetic





WeChat :Sin Yi Cosmetic

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SINYI Contact us Achieve your wishes Weave your dream catcher

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