









Good Manufacturing Practices for Cosmetics (Second Draft for Comments)

化妆品生产质量管理规范(征求意见稿)

National Medical Products Administration

Release Date: Sep. 8, 2021 Implementation Date: XXXX-XX-XX



You can now catch up with the latest updates of China CSAR on ChemLinked:



Stay tuned for more translations:

We will provide English, Japanese and Korean translations of CSAR and supporting rules soon.



Watch our webinars:

Interpretation of China's Cosmetic Supervision and Administration Regulation



Check out the breaking news:

China Finalises the Long-Awaited Cosmetic Supervision and Administration Regulation



Visit the featured page for more information:

Featured — China Cosmetic Regulatory Reform Tracking



Disclaimer

This is an unofficial document provided by **ChemLinked** (chemlinked.com), a platform of REACH24H Consulting Group, as an informational service to assist non-Chinese companies to better understand Asia Pacific especially China Chemical, Cosmetic, Food and Agrochemical regulatory issues.

This document should only be used as a reference and in case of any discrepancy between the English and original versions the original version shall prevail.

Nondisclosure:

You may not disclose this document to anyone else without the written permission of ChemLinked.

For further clarification and questions, you can read our <u>Privacy Polices</u> or contact us at <u>cosmetic@chemlinked.com</u>



Attachment 1

Good Manufacturing Practices for Cosmetics (Second Draft for Comments)

Chapter I General Provisions

Article 1 Purpose and Basis In order to regulate the quality management of cosmetics production, these Practices are hereby formulated in accordance with the *Cosmetics Supervision and Administration Regulations* (CSAR), *Supervision and Administration Measures on Cosmetics Manufacture and Operation* and other regulations and rules.

Article 2 Scope of Application These Practices are the basic requirements for quality management of cosmetics production. All cosmetics registrants, notifiers and entrusted production enterprises within the territory of the People's Republic of China shall abide by these Practices.

Article 3 General Requirements Cosmetics registrants, notifiers and entrusted production enterprises shall be honest and self-disciplined and establish a production quality management system to realize the control and traceability of the entire process of cosmetics material purchase, production, testing, storage, sales and recall, so as to ensure continuous and stable production of cosmetics that meet the quality and safety requirements.

Chapter II Institutions and Personnel

Article 4 Organizational Structure Cosmetics registrants, notifiers and entrusted production enterprises (hereinafter referred to as the "production enterprises") engaged in cosmetics production activities shall establish an organizational structure that is compatible with the type, quantity, and production license items, etc., of the cosmetics they respectively produce, and define the responsibilities and authorities of quality management, production and other departments. They shall also be equipped with the management personnel, technical personnel and testing personnel that are compatible with the type, quantity, and production license items, etc., of the cosmetics.

The quality management department of production enterprises shall be set up independently, perform quality assurance and quality control responsibilities, and participate in all activities related to quality management.

Article 5 Quality and Safety Responsibility System Production enterprises shall establish a cosmetics quality and safety responsibility system, specifying the

CHEMLINKED

responsibilities of the enterprises' legal representative (or the person chiefly in charge, the same below), person in charge of quality and safety, person in charge of the quality management department, person in charge of the production department, and the posts related to cosmetics quality and safety. The personnel in each post shall be responsible for cosmetics quality and safety level by level, and implement the main responsibility for cosmetics quality and safety.

Article 6 Legal Representative Legal representatives shall be fully responsible for the quality and safety of cosmetics, and shall be in charge of providing necessary resources, rationally formulating and organizing the implementation of quality policies, and ensuring the realization of quality objectives.

Article 7 Person in Charge of Quality and Safety Production enterprises shall assign a person in charge of quality and safety who shall master professional knowledge related to cosmetics quality and safety such as cosmetics, chemistry, chemical engineering, biology, medicine, pharmacy, food, public health or legal science, *etc.*, and be familiar with relevant laws and regulations, mandatory national standards, technical specifications, and have more than 5 years of experience in cosmetics production or quality management.

The person in charge of quality and safety shall assist the legal representative to undertake the following corresponding product quality and safety management and product release responsibilities:

- (1) Establishing and organizing the implementation of the enterprise's production quality management system, implementing quality and safety management responsibilities, and regularly reporting the operation of the quality management system to the legal representative;
- (2) Making decisions on product quality and safety issues and issuing relevant documents;
- (3) Review and management of product formulas, production techniques, material suppliers, product labels, *etc.*;
- (4) Material release management and product release;
- (5) Monitoring and management of cosmetics adverse reactions;
- (6) Product recall management.

The person in charge of quality and safety shall perform his responsibilities independently and shall not be interfered by other personnel of the enterprise.

Article 8 Person in Charge of Quality Management Department The person in charge of the quality management department shall master professional knowledge related to cosmetics quality and safety such as cosmetics, chemistry, chemical engineering, biology,



附件 1

化妆品生产质量管理规范 (征求意见稿)

第一章总则

第一条【目的及依据】 为规范化妆品生产质量管理,根据《化妆品监督管理条例》 《化妆品生产经营监督管理办法》等法规、规章,制定本规范。

第二条【适用范围】 本规范是化妆品生产质量管理的基本要求, 在中华人民共和 国境内的化妆品注册人、备案人、受托生产企业应当遵守本规范。

第三条【总体要求】 化妆品注册人、备案人、受托生产企业应当诚信自律,建立 生产质量管理体系,实现对化妆品物料采购、生产、检验、贮存、销售和召回等全过程 的控制和追溯,确保持续稳定地生产出符合质量安全要求的化妆品。

第二章 机构与人员

第四条【组织机构】 从事化妆品生产活动的化妆品注册人、备案人、受托生产企 业(以下简称"生产企业")应当建立与生产的化妆品品种、数量和生产许可项目等相 适应的组织机构,明确质量管理、生产等部门的职责和权限。配备与化妆品品种、数量 和生产许可项目等相适应的管理人员、技术人员和检验人员。

生产企业的质量管理部门应当独立设置,履行质量保证和质量控制职责,参与所有 与质量管理有关的活动。

第五条【质量安全责任制】 生产企业应当建立化妆品质量安全责任制,明确企业 法定代表人(或者主要负责人,下同)、质量安全负责人、质量管理部门负责人、生产 部门负责人以及与化妆品质量安全相关的岗位职责,各岗位人员应当逐级对化妆品质量 安全负责,落实化妆品质量安全主体责任。

第六条【法定代表人】 法定代表人对化妆品质量安全工作全面负责,应当负责提 供必要的资源,合理制定并组织实施质量方针,确保实现质量目标。

第七条【质量安全负责人】 生产企业应当设质量安全负责人,质量安全负责人应 当具备化妆品、化学、化工、生物、医学、药学、食品、公共卫生或者法学等化妆品质



量安全相关专业知识, 熟悉相关法律法规、强制性国家标准、技术规范, 并具有 5 年以上化妆品生产或者质量管理经验。

质量安全负责人应当协助法定代表人承担下列相应的产品质量安全管理和产品放 行职责:

- (一)建立并组织实施本企业生产质量管理体系,落实质量安全管理责任,定期向 法定代表人报告质量管理体系运行情况;
 - (二) 产品质量安全问题的决策及有关文件的签发;
 - (三) 产品配方、生产工艺、物料供应商、产品标签等的审核管理;
 - (四)物料放行管理和产品放行;
 - (五) 化妆品不良反应监测管理;
 - (六) 产品召回管理。

质量安全负责人应当独立履行职责,不受企业其他人员的干扰。

第八条【质量管理部门负责人】 质量管理部门负责人应当具备化妆品、化学、化工、生物、医学、药学、食品、公共卫生或者法学等化妆品质量安全相关专业知识,并具有化妆品生产或者质量管理经验。质量管理部门负责人应当承担下列职责:

- (一) 所有产品质量有关文件的审核;
- (二)组织与产品质量相关的变更、自查、不合格品管理、不良反应监测、召回等活动;
 - (三) 保证质量标准、检验方法和其他质量管理规程有效实施;
 - (四) 保证完成必要的验证工作, 审核和批准验证方案和报告;
 - (五) 物料和产品放行的具体审核;
 - (六) 评价物料供应商;
- (七)制定并实施生产质量管理相关的培训计划,保证员工经过与其岗位要求相适应的培训;
 - (八) 负责其他与产品质量有关的活动。

第九条【生产部门负责人】 生产部门负责人应当具备化妆品、化学、化工、生物、 医学、药学、食品、公共卫生或者法学等化妆品质量安全相关专业知识,并具有化妆品 生产或者质量安全管理经验。生产部门负责人应当承担下列职责:

Make Sure Your Products Entering Global Market with Compliance

What we can do for you?



- + Responsible agent service
- + Cosmetic formula and label compliance review
- + Product registration and filing
- + New cosmetic ingredient registration



- + Cosmetic formula and label compliance review
- + EDI Notification
- + Cosmetic QC test service
- + Functional cosmetic registration
- + Ingredients registration by ICID



- + Marketing license application
- + Cosmetics notification
- + Ingredients analysis and full ingredient list translation
- + Cosmetic test service
- + Label & advertisement review



- + Responsible Person (RP) service in the EU & UK
- + Cosmetic Product Notification Portal (CPNP)
- + Cosmetic Product Safety Report (CPSR)
- + Preparation of Product Information File (PIF)



- + Cosmetic formula and label compliance review
- + FDA Voluntary Cosmetic Registration Program (VCRP)
- + California Safe Cosmetics Program (CSCP)
- + Toxicological Risk Assessment (TRA)
- + INCl application

What makes us unique?



REMOVE COSMETIC REGULATORY BARRIERS **EXPEDITE ASIA MARKET ACCESS**

- · In-time regulatory information and market access requirements
- · Supported by local and experienced technically adept technical team
- · Strategic partnerships with competent authorities and industrial associations worldwide
- · Customized consulting service for complete compliance solutions.

OUR SERVICES



Information

News, Alerts



Knowledge

Pedia-articles,

Regulatory Analysis, Market Insights, Reports



Databases

Regulatory Database,

English Translations, Lists/Inventories, Q&A



Training

Webinars, Online courses, Offline events



Solutions

Translation, Consultancy, Advertising, Tailored Regulatory Report, Business-matching services, Tailored online and offline training



14th Floor, Building 3, Haichuang Technology Center, 1288 West Wen Yi Road, Hangzhou, China (311121)



