



Announcement on Matters Related to Cosmetic Registration and Notification (Draft for Comments)

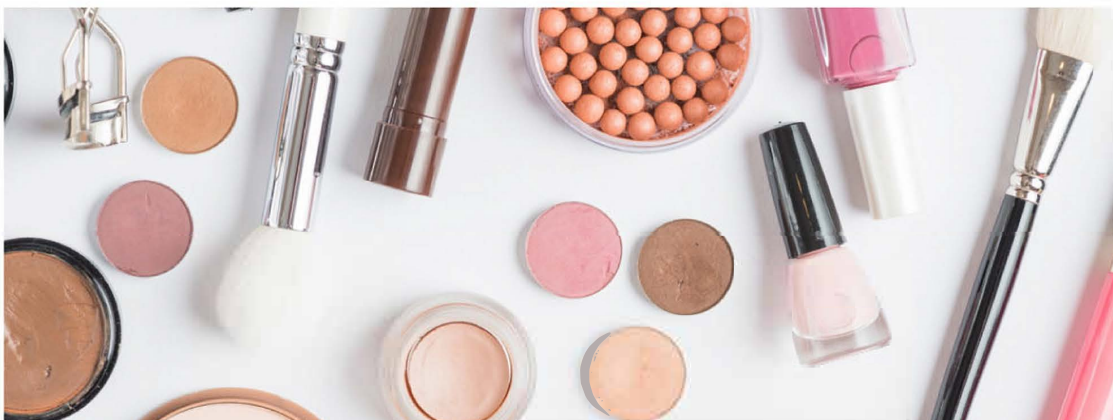
关于化妆品注册备案有关事项的公告（征求意见稿）

National Medical Products Administration

Release Date: Mar. 31, 2026

Implementation Date: /

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Attachment 1

Announcement on Matters Related to Cosmetic Registration and Notification (Draft for Comments)

In order to implement the *Opinions of the NMPA on Deepening the Reform of Cosmetic Administration and Promoting the High-quality Development of the Industry* (No. 18, 2025 of the NMPA), continue to advance the reform of cosmetic review and approval, and promote the high-quality development of the industry, the relevant matters for optimizing the registration and notification management of cosmetics are hereby announced as follows:

I. Regarding Encouragement of the First Launch of New Cosmetic Products in China

To align with high-standard international economic and trade rules and cultivate the first launch economy in China's cosmetic field, when registering and notifying an international new cosmetic product that is to be first launched in China or simultaneously launched in China and other countries/regions, the cosmetic registrant/notifier shall provide a description on the innovation of the product formula, technique and other related aspects, along with a commitment statement for the first launch of the product in China, and is exempt from submitting the certificate that the product has already been marketed in the country/region of production.

II. Regarding Exemption from Submitting Animal Test Data for Cosmetics

For hair perming, non-oxidative hair dyeing, freckle-removing and whitening cosmetics with only physical covering effects among special cosmetics, as well as general cosmetics and the aforementioned special cosmetics that contain a new cosmetic ingredient still under safety monitoring, if the production enterprise has obtained the relevant qualification

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- + Product registration and notification
- + New cosmetic ingredient (NCI) registration and notification
- + Ingredient safety information submission
- + Cosmetic efficacy evaluation



- + 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)
- + INCI application

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附件 1

关于化妆品注册备案有关事项的公告 (征求意见稿)

为贯彻落实《国家药监局关于深化化妆品监管改革促进产业高质量发展的意见》(国药监妆〔2025〕18号)，持续推进化妆品审评审批改革，促进产业高质量发展，现就优化化妆品注册备案管理有关事项公告如下：

一、关于鼓励化妆品新品在中国首发

对标国际高标准经贸规则，培育我国化妆品领域首发经济，对国际化妆品新品在中国首发上市或者在中国以及其他国家(地区)同步上市的，产品注册备案时，化妆品注册人、备案人应当提供产品配方、工艺等方面的创新性说明，同时提供产品在中国首发上市的承诺性声明，免于提交在生产国(地区)已经上市销售的证明文件。

二、关于减免化妆品动物试验资料

特殊化妆品中烫发、非氧化型染发和仅具物理遮盖作用的祛斑美白化妆品，以及使用了尚处于安全监测期中化妆品新原料的普通化妆品和前述特殊化妆品，其生产企业已取得所在国家(地区)政府主管部门出具的生产质量管理体系相关资质证明，且产品安全风险评估结果能够充分确认产品安全性的，产品注册备案时，免于提交该产品的毒理学试验报告。

三、关于调整原料安全相关信息为企业存档备查

产品(含牙膏)注册备案时，注册人、备案人无需填写产品所使用原料的安全信息文件，相关资料由企业自行存档备查。《化妆品安全技术规范》等技术文件对原料质量规格有相关要求的，注册人、备案人应当在提交安全评估资料时，一并提交原料的质量规格或者检验报告。国家药监局不再提供原料安全信息服务平台各项服务，不再公开原料报送码相关信息。

四、关于共用配方体系近似产品安全性技术资料

(一) 同一化妆品注册人、备案人拟注册备案同一品牌多个产品(含牙膏)的，注册人、备案人可以选择其中一个具有代表性的产品，开展微生物与理化检验、毒理学试

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