



Announcement of the National Medical Products Administration on Matters Concerning Cosmetics Registration and Notification (No. 70 of 2026)

国家药监局关于化妆品注册备案有关事项的公告
(2026年第70号)

National Medical Products Administration

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Announcement of the National Medical Products Administration (NMPA) on Matters Concerning Cosmetics Registration and Notification (No. 70 of 2026)

Date of Publication: July 29, 2026

To implement the *NMPA Opinions on Deepening the Reform of Cosmetic Administration and Promoting the High-Quality Development of the Industry* (Guo-Yao-Jian-Zhuang [2025] No. 18), continuously advance reform of cosmetics review and approval, and promote high-quality industry development, the following announcement is hereby issued regarding the optimization of cosmetics registration and notification management:

I. Encouraging the first launch of new cosmetics in China

By aligning with high-standard international economic and trade rules and fostering a first-launch economy in China's cosmetics sector, for new international cosmetics first launched in China or simultaneously launched in China and other countries (or regions), registrants and notifiers may provide a declaration of commitment regarding the product's first launch in China during the registration or notification process, and be exempt from submitting proof of sales in the country (or region) where the registrant or notifier is located, or the country (or region) of manufacture. For the above-mentioned new international cosmetics, the sales packaging submitted during the registration or notification may be a design draft.

II. Exemption from or reduction of animal testing data for cosmetics

For special cosmetics, including hair perm products, non-oxidative hair dye products, and skin-whitening products with only a physical covering effect, as well as general cosmetics using new cosmetic ingredients (excluding children's cosmetics), where the manufacturer has obtained qualification documents related to its production quality management system issued by the competent government authority of the country (or region) where it is located, and the product safety risk assessment results are sufficient to confirm product safety, the toxicological test report for the product shall be exempt from submission during product registration or notification. Based on advances in scientific research, NMPA's cosmetic technical review department may adjust the scope of products eligible for exemption from or reduction of animal testing data in a timely manner through the formulation of technical guidelines.

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- + Responsible person (RP) service in China
- + Cosmetic formula and label compliance review
- + 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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国家药监局关于化妆品注册备案有关事项的公告

(2026 年第 70 号)

发布时间：2026-07-29

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，注册人、备案人可以提供产品在中国首发上市的承诺性声明，免于提交在注册人、备案人所在国或者生产国（地区）已经上市销售的证明文件。上述国际化妆品新品在注册备案时提交的的生产国（地区）销售包装可以是设计图。

二、关于减免化妆品动物试验资料

特殊化妆品中烫发产品、非氧化型染发产品、仅具物理遮盖作用的祛斑美白产品，以及使用了新原料的普通化妆品（儿童化妆品除外），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。根据科学研究的发展，国家药监局化妆品技术审评部门可以通过制定技术指导原则，适时调整减免动物试验产品范围。

三、关于调整化妆品（含牙膏）原料安全相关信息为企业存档备查

产品注册备案时，注册人、备案人无需填写产品所使用原料的安全信息文件和原料报送码，仅需填写原料生产商名称，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、备案人应当在产品配方栏或者安全评估资料中提交原料的质量规格或者检验报告。国家药监局不再公开原料报送码。已经注册备案的产品，所使用原料的生产商、原料质量规格增加或者改变的，按照《化妆品注册备案资料管理规定》要求进行更新维护或者申请变更。



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