



General Technical Requirements for Biotechnology-derived Cosmetic Ingredients 《化妆品用生物技术来源原料技术要求通则》

National Medical Products Administration

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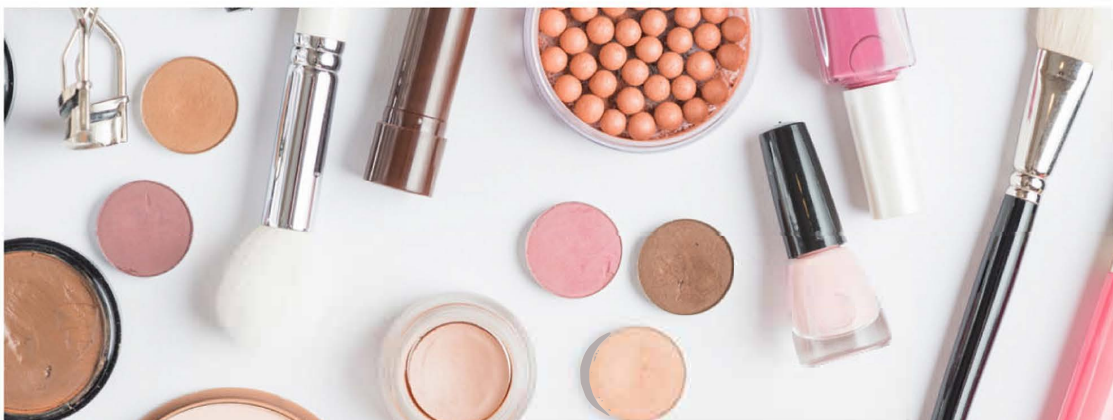
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Foreword

This document was drafted in accordance with the provisions of GB/T 1.1-2020 *Directives for Standardization - Part 1: Rules for the Structure and Drafting of Standardizing Documents*.

Attention is drawn to the possibility that some of the contents of this document may be the subject of patent rights. The issuing body of this document shall not be held responsible for identifying any or all such patent rights.

This document was proposed by the National Medical Products Administration (NMPA).

This document is under the jurisdiction of the Sub-Technical Committee on Ingredients and Packaging Materials of the Technical Committee on Cosmetic Standardization under the NMPA.

The drafting units of this document are as follows: Jiangnan University, Shenzhen Institute for Drug Control, Shandong Institute for Food and Drug Control, and National Institutes for Food and Drug Control.

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- + Cosmetic formula and label compliance review
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- + Cosmetic Product Safety Report (CPSR)
- + Preparation of Product Information File (PIF)



- + Cosmetic formula and label compliance review
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- + Toxicological Risk Assessment (TRA)
- + INCI application

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化妆品用生物技术来源原料技术要求通则

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前 言

本文件按照GB/T1.1—2020《标准化工作导则第1部分：标准化文件的结构和起草规则》的规定起草。

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本文件由国家药品监督管理局提出。

本文件由国家药品监督管理局化妆品标准化技术委员会原料和包装材料分技术委员会归口。

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