



CERTIFICATE OF FDA REGISTRATION

REGISTRATION & LISTING OF COSMETIC PRODUCT FACILITIES

THIS CERTIFICATE ONLY REPRESENTS THAT THE REGISTERED AGENT HAS REGISTERED THE
FDA FOR THE FOLLOWING COMPANY, AND DOES NOT REPRESENT THE FDA ENDORSEMENT

Enterprise Name : Guangzhou Bangya Cosmetics Co.Ltd
Address : Guangdong Guangzhou Baiyun 13A12-13A13, Building 7, No.
50 Juyuan Street, Shicha Road, Baiyun District, Guangzhou
City

HUAX Testing has been registered with the US Food and Drug Administration (FDA) for the aforementioned company.
HUAX Testing has registered these facilities with the US Food and Drug Administration under the 2024 Cosmetic
Modernization Act (MoCRA) and completed the marketing of cosmetic facilities.

U.S. FDA does not issue or recognize Certificates of Registration.HUAX Testing is not affiliated with the U.S. US
Food and Drug Administration (FDA).

Facility FEI Number : 3032194482

Brand Name : Bangya

According to the requirements of the Federal Food, Drug, and Cosmetic (FD&C) Act: Manufacturers and processors
must register their facilities with the FDA and renew their registration every two years, including any updates to the
product and its ingredients If the FDA determines that cosmetics manufactured or processed by a registered facility
has the potential for series adverse health sequences, the FDA has the authority to suspend the facility's FDA
registration.

Note :Under Section 607 (c) (5) of the FD&C Act , product updates are required every year ,including ingredient
updates and discontinuation updates .

Under Section 607 (a) (2) of the FD&C Act , registered facilities must be renewed every two years

This certificate only represents that the registered
agent has registered the FDA for the above company,
and does not represent the FDA endorsement.
Shenzhen huaxiang Testing Technology Co., Ltd Tel:0755-23010432

Kevin su / Registered Agent
Valid from - until
Sep. 12, 2024 - Sep. 12, 2026
Email: Kevin_su@hua-x.com