Central Drugs Standard Control Organization Directorate General of Health Services Ministry of Health and Family Welfare, Government of India

NON COMPLIANCES OBSERVED DURING REVIEW PROCESS OF APPLICATIONS FOR REGISTRATION OF IMPORT OF COSMETICS

1. Covering Letter:

Not clearly mentioning the purpose (Fresh or Endorsement of Products/Pack Size/Additional Sourcing Location or Re-Registration) or details of approved Registration Certificate or Product/Category (if any)

2. Form 42:

- ➤ Incomplete Name and address of the authorized Indian agent, manufacturer & manufacturing Premises.
- ➤ If the products are to be imported in bulk, the actual pack size for import in bulk quantity is not mentioned.
- ➤ Colored scan copy of original document containing, Signature & Stamp with Name & Designation of Indian agent/Manufacturer not submitted.
- ➤ Name and address of authorized Indian agent/Manufacturer along with product name not correlated with Power of Attorney and Schedule D(III).

3. Schedule D (III):

- Name & address of the authorized Indian agent, manufacturer & manufacturing premises not correlating with Form 42.
- ➤ Name of the Cosmetic along with category applied, variant and pack size not correlating with Form 42.
- ➤ Undertaking not signed and stamped by Indian agent/ Manufacturer.
- ➤ Not mentioning list of countries where market authorization or import permission or registration was granted.
- > Incomplete information in Schedule DIII as some columns is not filled up.
- ➤ Colored scan copy of original document is not submitted.

4. Power of Attorney:

- ➤ Validity of Power of Attorney not mentioned.
- Name & address of the authorized Indian agent, manufacturer & Manufacturing premises not as per Form 42.
- Name of the Cosmetic along with category applied, variant and pack size not correlating with Form 42.
- Not co-jointly signed, stamped and dated with Name & Designation of the signatory by both Indian agent & the manufacturer.

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- ➤ Power of Attorney including product list not apostilled and not authenticated from country of origin.
- All pages of power of attorney along with product list is not signed and stamped by authorized Indian agent as well as manufacturer.

5. Fee:

- ➤ A copy of acknowledgement receipt of fee paid is not submitted.
- Amount of fee paid in USD or its equivalent in Indian Rupees is not correlating with the category of applied product as per Column 3 of the "Guidelines on Registration of Import of Cosmetics".

6. Product Composition data/Ingredient list:

- > Product composition data not submitted for all the proposed products.
- List of ingredients for all the products not submitted with exact concentration of each ingredient.
- Product composition data containing ingredients with concentration more than the prescribed limit of BIS.
- ➤ Product composition data containing substances which must not form part of the composition of cosmetic products as per Annexure A of IS 4707 (Part 1&2): 2017 of BIS.
- ➤ Product Composition data not duly signed/stamped by competent QC Person/ Person authorised from the manufacturer.

7. Free Sale Certificate:

- > Free sale Certificate not issued from country of origin/country of Manufacturer of the proposed products.
- ➤ Colored scan copy of Original/authenticated Free Sale Certificate issued by the national regulatory authority/ other competent association organizations from country of origin/country of Manufacturer not submitted for all the applied products and variants.
- > Validity of Free sale Certificate not mentioned.
- ➤ Free sale certificate does not clearly indicate that the proposed products are freely sold in the country of origin/ country of Manufacturer.
- Name of the Cosmetic along with category applied, variant and pack size not correlating with Form 42 & Power of Attorney.
- ➤ All the pages of product list not signed and stamped by issuing authority.

8. Chemical Information of Cosmetics:

➤ Test Protocol for testing of Cosmetics and specification of the product not submitted.

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- ➤ Test report including result of Mercury, Lead, Arsenic & Other Heavy metals with limits not complying BIS.
- Test report not duly signed by competent QC Person from the Manufacturer.

9. Labels/Pack insert of Proposed Products:

- ➤ Submitted Original Label/pack insert of the products which is not legible and not in English Language.
- Name of the cosmetics, Pack size & Name/address of the Manufacturer not correlating with that of Form 42.
- ➤ Label of the proposed products without word Batch no., mfg. License no., Use Before date not complying Rule 148 of Drugs & Cosmetics Rules 1945.
- Labels containing phrases that may appear to attract the definition of the drug. E.g. affects the structure or any function of the human body, Treatment of any disease/disorder, Drug Facts, Dermatologist recommended etc.
- ➤ Label does not mentions cautions and Instructions for certain ingredients (e.g. p-Phenylenediamine, fluoride content) as per requirement of Drugs & Cosmetic Rules 1945.
- > Submitted labels are not signed/ stamped by manufacturer or its authorised agent.

10. Undertaking for Heavy Metal and hexachlorophene content:

- ➤ Heavy Metal undertaking not complying with BIS, D & C Act, 1940 & Rules made thereunder and also not issued from the manufacturer.
- ➤ Declaration is not mentioning clearly that applied products are free from hexachlorophene and complies with the limit of heavy metals as per BIS.

11. Manufacturing License:

➤ Notarized Copy of Manufacturing license/Repackaging license issued by the State Licensing Authority not submitted if the proposed products are imported in bulk for repacking/relabeling.

12. Undertaking for Non Animal Testing:

- Non Animal testing Undertaking/Declaration not issued by manufacturer
- Non Animal testing declaration not duly signed/stamped by competent person/authority from the manufacturer
- **13. Others:** English translated copy of any of the above said documents not signed by the qualified translator.