Declaration of the EU conformity of Prontomed cosmetic products.

The EU Cosmetics Regulation (EG) No. 1223/2009 ensures a high level of protection of the human health. It primarily affects manufacturers of cosmetics, but retailers are also affected, because some of their information obligations and advertising regulations are of concern.

Cosmetic products must be safe for human health under normal or reasonably foreseeable use.

Safety assessment, notification and product information file

Prontomed has performed a safety assessment by a qualified person for all cosmetic products. Each product has been submitted to the EU Commission by notification in the CPNP.

For ten years, from the date of placing the product on the market, we keep production information files for the cosmetic products, which are updated regularly. Hygiene monitoring and batch documentation are just as much a part of our comprehensive quality checks as the review of instructions for use and packaging.

Quality standards at Prontomed GmbH

Prontomed ensures that all cosmetic products are manufactured in accordance with the requirements of "Good Manufacturing Practice" (GMP) according to harmonized standard EN ISO 22716:2007.

Production, filling and labeling are performed in cooperation with certified contract manufacturers. Testing of raw materials, batch, packaging materials and finished products is done at accredited laboratories.

As a manufacturer, Prontomed fulfills all the requirements that must be met as a responsible person under the EU Cosmetics Regulation (EG) No. 1223/2009.

Hiddenhausen, 12.04.2021

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