99118047058000

Notifizierung von kosmetischen Mitteln einschließlich ihrer Rahmenrezeptur sowie ggf. auch von Nanomaterialien Durchführung

Modul	Sachverhalt
Leistungsschlüssel	99118047058000
Leistungsbezeichnung I	Notifizierung von kosmetischen Mitteln einschließlich ihrer Rahmenrezeptur sowie ggf. auch von Nanomaterialien Durchführung
Leistungsbezeichnung II	Notify cosmetic products (possibly with nanomaterials) before placing them on the market
Typisierung	1 - Bund: Regelung und Vollzug
Quellredaktion	Bund
Freigabestatus Katalog	fachlich freigegeben (gold)
Freigabestatus Bibliothek	unbestimmter Freigabestatus
Begriffe im Kontext	
Leistungstyp	Leistungsobjekt mit Verrichtung
Leistungsgruppierung	

Modul	Sachverhalt
Verrichtungskennung	Durchführung (58)
SDG-Informationsbereich	nicht SDG-relevant
Lagen Portalverbund	Produkt- und Stoffzulassung (2120200)
Einheitlicher Ansprechpartner	Nein
Fachlich freigegeben am	14.12.2023
Fachlich freigegen durch	Federal Ministry for the Environment, Nature Conservation, Nuclear Safety and Consumer Protection (BMUV)
Handlungsgrundlage	https://eur-lex.europa.eu/legal-content/DE/TXT/?uri=CELEX%3A02009R1223-20221217&qid=1403001930973 https://eur-lex.europa.eu/legal-content/DE/TXT/?uri=CELEX%3A02009R1223-20221217&qid=1403001930973
Teaser	Cosmetic products that you wish to make available on the European market for the first time must be notified to the European Commission in advance via the Cosmetic Products Notification Portal (CPNP).
Volltext	Cosmetic products are not subject to authorization. However, certain components of cosmetic products such as new preservatives, colorants and UV filters are exempt from this requirement. Regardless of which cosmetic product is involved, the safety of the product must be guaranteed. If you want to make a cosmetic product available on the European market for the first time, you as the "responsible person" (usually the manufacturer or importer, and possibly also the distributor) must provide some information about your product in advance. This is done via the notification portal for cosmetic products (Cosmetic Products Notification Portal, CPNP). The CPNP is operated by the European Commission and enables uniform and centralized notification in all member states of the European Union. The CPNP makes this information available electronically to

Modul	Sachverhalt
	• competent authorities (for the purposes of market surveillance, market analysis, evaluation and consumer information) • Poison information centers or similar bodies established by European Union countries (for the purpose of medical advice) The CPNP is accessible to: • Persons responsible for cosmetic products • Distributors of cosmetic products Notification in the CPNP does not include approval or verification of data. As a responsible person, you must ensure that the cosmetic products you place on the market are safe, that their intended use is guaranteed and that they comply with the legal provisions. • The CPNP is intended for manufacturers of nanomaterials in order to be able to add detailed information on nanomaterials used in a cosmetic product to the notification of the cosmetic product. The CPNP includes 3 notification obligations that you as the responsible person must fulfill before your cosmetic product is placed on the market for the first time: • 1. information about the cosmetic product and its formulation, which will be made available to poison information centers for the purpose of providing prompt and appropriate advice in the event of adverse health effects. • 2. information on the cosmetic product without the formulation, which is made available to the competent authorities of the federal states for the purpose of monitoring. • 3. cosmetic products containing nanomaterials other than colorants, preservatives and UV filters must be notified 6 months before being placed on the market. This notification must be made in addition to the notification referred to in Article 13. This information is sent exclusively to the European Commission. If the European Commission has concerns about the safety of a nanomaterial, it may ask the Scientific Committee on Consumer Safety to carry out a risk assessment. To fulfill this obligation, you as the responsible person can

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	appoint a nanomaterial representative. In cosmetic products, "nanomaterial" refers to a natural, process-derived or manufactured material consisting of solid particles that occur either independently or as recognizable constituent particles in aggregates or agglomerates, and where at least 50 percent of these particles in the number size distribution meet at least one of the following conditions: (a) one or more external dimensions of the particles are in the size range of 1 nanometer (nm) to 100 nm; b) the particles have an elongated shape such as a rod, fiber or tube, with two outer dimensions being less than 1 nm and the other outer dimension being greater than 100 nm c) the particles have a platelet-like shape, where one outer dimension is less than 1 nm and the other outer dimension is greater than 100 nm. In addition, the national Cosmetics Ordinance must be observed. Anyone who manufactures cosmetic products in Germany must notify the competent authority responsible for monitoring the place of manufacture before placing them on the market. If cosmetic products are imported into the European Union, the person responsible for the import must notify the competent authority responsible for supervision of the place where cosmetic products are brought into the scope of this Regulation (place of import) before they are imported for the first time.
Erforderliche Unterlagen	• Information on the responsible person and the cosmetic product, including the name and quantity of the ingredients used • label • Information on nanomaterials, if applicable
Voraussetzungen	• Your company produces, has produced or imports cosmetics. • You are a "responsible person": Only cosmetic products for which a legal or natural person is named

Modul	Sachverhalt
	as the "responsible person" on the label within the European Union (EU) or a state of the European Economic Area (EEA) may be placed on the market (usually manufacturer or importer, possibly also distributor). • They have access to the CPNP as the "responsible person".
Kosten	Abgabe: Es fallen keine Kosten an There are no costs.
Verfahrensablauf	The required data is transmitted by you as the responsible person via online forms of the CPNP system of the European Commission and can also be corrected or updated later. Registration in the CPNP: In order to be able to use the CPNP, you must complete two primary registration steps: • One-time personal registration in EU Login: this provides you with the user name and determines the access password. • This is followed by the registration of a company or the application of an existing company via the (SANTE Authorization System, SAAS). The CPNP can only be used once these registration steps have been carried out and the applications have been completed in accordance with the user profile. Notification in the CPNP: You must enter the following data in the CPNP if you are making a cosmetic product available on the European/German market for the first time: • the category of the cosmetic product and its name or names by which the specific identification is possible It is not sufficient to merely state the name of a cosmetic series under which various products are marketed. • the name and address of the person responsible where the product information file is made easily accessible

Modul	Sachverhalt
	• the country of origin in the case of imports Country of origin does not mean the last country in a supply chain, but rather the country in which the cosmetic product was actually manufactured. • the Member State in which the cosmetic product is placed on the market, i.e. made available for the first time • the information that makes it possible to contact a natural person if necessary (e.g. first and last name, address, telephone number and e-mail address); information on a legal person alone is not sufficient. • the presence of substances in the form of nanomaterials and their identification, including the chemical name • the reasonably foreseeable exposure conditions • the name and Chemicals Abstracts Service (CAS) or EC number of the substances classified as carcinogenic, mutagenic or toxic for reproduction (CMR) • the frame formulation to enable rapid and appropriate medical treatment in the event of difficult incidents All data you submit to the CPNP will be treated as confidential information and can be corrected and updated at any time.
Bearbeitungsdauer	none
Frist	6 Monat(e) Cosmetic products that contain nanomaterials other than colorants, preservatives and UV filters and are not otherwise restricted are subject to an additional procedure. They require a separate notification to the CPNP 6 months before being placed on the market.
weiterführende Informationen	https://www.bvl.bund.de/cpnp https://www.bvl.bund.de/SharedDocs/Downloads/03_Verbraucherprodukte/Kosmetik/CPNP_Benutzerhandbuch.html?nn=11019868 https://webgate.ec.europa.eu/cpnp https://webgate.ec.europa.eu/cas/eim/external/register.cgi https://webgate.ec.europa.eu/saas https://webgate.ec.europa.eu/cpnp/faq/?event=faq.show

Modul	Sachverhalt
Hinweise	
Rechtsbehelf	• There are no legal remedies against the BVL.
Kurztext	• Notification of cosmetic products including their master formulation and, if applicable, nanomaterials Implementation • Notification required for cosmetic products that are to be made available on the European market for the first time • Transmission ("notification") takes place exclusively electronically via the European Commission's Cosmetic Products Notification Portal (CPNP) • Objectives: Consumer protection and product safety • CPNP provides information electronically for Competent authorities (for the purposes of market surveillance, market analysis, evaluation and consumer information) Poison information centers or similar facilities established by EU countries (for the purpose of medical treatment) • CPNP is accessible for: persons responsible for cosmetic products distributors of cosmetic products Raw material suppliers of nanomaterials • CPNP contains a separate module for cosmetic products containing nanomaterials • Notification that nanomaterials are present must be made in addition to the notification of the product • if the European Commission has concerns about the safety of a nanomaterial, it can call in the Scientific Committee on Consumer Safety for a risk assessment • Costs: none • Responsible: competent national contact point: Federal Office of Consumer Protection and Food Safety (BVL)
Ansprechpunkt	
Zuständige Stelle	
Formulare	
Ursprungsportal	Notifizierung von kosmetischen Mitteln einschließlich ihrer Rahmenrezeptur sowie ggf. auch von Nanomaterialien Durchführung, Notifizierung von kosmetischen Mitteln einschließlich ihrer Rahmenrezeptur sowie ggf. auch von Nanomaterialien

Modul	Sachverhalt
	Durchführung