

ADVISORY DOCUMENT

INFORMATION EXCHANGE ON COSMETIC PACKAGING MATERIALS ALONG THE VALUE CHAIN



Regulatory Background



Obligations under the EU Cosmetics Regulation

- Products must be safe for the consumer
- Safety must be demonstrated via 'Safety Assessment'
- Safety assessment must
 - ➤ Be done by a qualified Safety Assessor
 - > Consider exposure and toxicological profile of ingredients
 - ➤ Include assessment of impact of impurities on safety of the product; specific reference to packaging
 - ➤ Be justified, detailed and documented
- Additional details in EU Commission implementing Guidelines



Roles and Responsibilities

- Safety is the legal responsibility of the entity placing the cosmetic product on the market (Responsible Person)
- He needs to work with a suitably qualified safety assessor
- Relevant information on the packaging material is part of the information needed
- HOWEVER, packaging suppliers have no legal obligation to provide safety information under the Cosmetics Regulation – and only limited obligations under the Chemicals legislation.
- Data exchange is a Business to Business decision.

What information to be shared? Full composition of the packaging?



- Very difficult to obtain along the supply chain
 - lack of knowledge of clients' legal obligations,
 - concerns over commercial secrecy,
 - high administrative burden,
 - lack of information from their own suppliers...
- Can be more substances than the cosmetic formulation
- Packaging safety assessment may become more complicated than the safety assessment of the cosmetic formulation

What information to be shared? Targeted, Relevant Information



EU Commission Guidelines:

- ... in direct contact with the formulation.
- ... experience with similar formulation/packaging combinations already on the market
- ... food packaging have often already been tested, so relevant information on stability and migration may be available.

Opportunity for a common sense, harmonised approach

- focus on information that is relevant for the safety assessor
- → facilitate exchange of information along the value chain



Scope of the document and principle of the approach



What it is not ...

- Not an allocation of safety responsibility to suppliers
 Cosmetic product remains the responsibility of the cosmetics
 Responsible Person
- Not Guidelines on 'packaging safety assessment'
 Rather identifies relevant information that can be transmitted along the supply chain to enable a safety assessment
- Not Guidelines on physical risks of the packaging suitability of the packaging regarding product quality
 Only addresses impact on the formulation (chemical risks)
- Not an exclusive or mandatory approach to ensure compliance with the Cosmetics Regulation
 - Other approaches may be used provided they enable an appropriate safety assessment of the cosmetic product



What it is ...

- Practical Guidance for suppliers of packaging raw materials and finished packaging items
- Tool to identify relevant packaging constituents whose presence needs to be known to the Cosmetic Safety Assessor
- Based on established practices in the area of food contact materials
- Approach that can be applied along the supply chain to aggregate relevant information



Principle approach

- Main concern is the possible migration of substances from the packaging into the formulation
 - Focus information on those components of the packaging that are in 'migrateable' contact with the formulation
- Most cosmetic products have similar physical chemical properties to typical foods/food simulants
 - Where possible, benefit from established process on information exchange on food packaging materials (based on EU legislation and Commission/industry Guidance)
- Migration exposure scenarios from food packaging can be applied to cosmetic packaging



Principle approach

- If physical/chemical properties are similar, safety information justifying food contact use can also support use as cosmetic packaging
- In addition, information needed on packaging constituents that are of specific concern for cosmetic safety and compliance (banned, restricted substances, skin sensitisers)
- The approach can be applied at any step of the supply chain. From raw material to finished packaging!



Raw Material vs Finished Packaging Item

Approach can be used for all steps in the supply chain:

Raw material → compounded raw materials → packaging components → final packaging

Mixture

Article



Step-by-Step Excel Macro

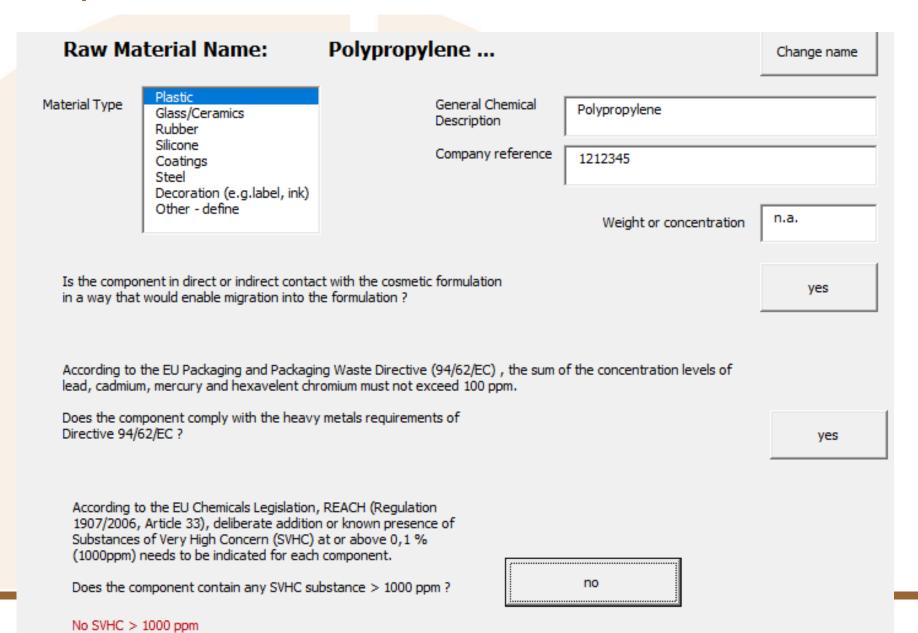
- Interactive questionnaire that allows to build a packaging statement according to the guidelines.
- No obligation to use, but helpful for some companies



Step 1 – Identification

Section I) Document Scope		×
In this section you are asked to define the s	scope of the document and identify the Packagi	ng Rawmaterial
Name of supplier company	Company ABC	
Identification of the Packaging Rawmaterial	Polypropylene	
Supplier reference code	12.12345	
Client reference code	AB.ABCDE	
If this document is covered by a confidentialty agreement, fill in the reference :	n.a.	
		ок

Step 2 – General Chemical information





Step 3 – Compliance with Food Contact Legislation (1)

Compliance with EU Food Contact Legislati	ompliance	with EU Food	d Contact L	egislatio.
---	-----------	--------------	-------------	------------

The composition of the Raw Material is compliant with the requirements of EU Food Contact Framework Regulation 1935/2004 EC, and in particular with Regulation 10/2011 EC.

Conditions for which compliance is valid (eg. types of food / or simulants , specific conditions)

oil, fresh meat, ...

Migration

Type of information available on migration (e.g. test with food, test with simulant, modeling, worst case asumptions, supplier information, ...)

Test with food simulant (see attached)



Step 3 – Compliance with Food Contact Legislation (2)

Presence of substances restricted under Food Contact Legislation (Regulation 10/2011 EC) by a specific migration limit (SML), a maximum concentration (QM) in the plastic or a "no detectable migration" requirement at a certain detection limit (DL) - with a residual quantity greater or equal to 1/10 of the SML / QM

Substance XXXX: CAS 1234-05-6

yes

GMP

Component manufactured according to EU Good Manufacturing Practice (EC) No 2023/2006 ?

yes



Substance XXXX	
CAS N°	1234-05-6
Concentration in the migrating levelin a re	. 0.23 /0
Reason for reportir	banned/restricted in Annex II / III of the EU Cosmetics Regulation (incl. CMR) Sensitiser Cat 1/Cat 1B
	Sensitiser Cat 1A
EU Cosmetics Regu	et(s) was used to identify the substances (e.g. CLP regulation, lation, Cross Industry Guidance Document (Appendix 1),?

Cross industry guidance listi



Small tangent: Why do we need information beyond Food contact compliance?



Ignorance is not bliss but a liability





CPR Banned Substances

- Annex II of the Cosmetics Regulation bans the presence of +/- 1600 Substances
- Most substances have very low likelihood to be actively used as cosmetic ingredient
 - > 1100 CMR Substances (classified as Carcinogenic, Mutagenic, Reprotoxic)
 - Active drug ingredients, antibiotics
 - Certain natural and synthetic toxins
 - BSE Risk materials
 - Hormones
 - Psychotropic drugs
 - Certain potential cosmetic ingredients



Restricted Substances

- Annex III of the Cosmetics Regulation restricts the max. concentration of +/- 300 substances
- Most substances have very high likelihood to be actively used as cosmetic ingredient
- Easy to control the actively added concentration BUT difficult to manage additional amounts from impurities



Skin Sensitisers

- Skin compatibility is major concern for cosmetics (direct prolonged contact)
- Skin reactions in allergic consumers can be elicited at low concentrations
- Presence of skin allergens may not always be avoidable – but safety can be achieved through concentration/formulation



Substances of concern to the Cosmetic Safety Assessor

- Cosmetic companies can manage direct addition but need to know impurities from raw materials and packaging !!!
- Otherwise risk with regard to safety, regulatory compliance and public perception



Step 5 – Additional Information

afety when used with the cosmetic formulation.In particular, provide Documents of Compliance (DoC) as well as references/summaries of migration valuation (OML, SML) in relation to the components or its raw material :							migration

Please provide any additional relevant information on the packaging Rawmaterial that may help the Cosmetic Product Safety Assessor to evaluate its



Step 6 – Date and Contact

12.09.2019 Date: John Doe Contact: OK

Information ultimately available for the safety assessor (1)

- Description of packaging and its components, incl. their chemical nature and weight
- For each component :
 - identity and concentration of SVHC substances > 0,1 %
 - Heavy metals > 100 ppm
 - Identification if component is in 'migrateable' contact with the formulation

Information ultimately available for the safety assessor (2)

- For each component in 'migrateable contact':
 - Statement on compliance with Food Contact Legislation (incl. applicable foods/simulants)
 OR
 - Reason of non compliance. (If linked to a specific substance: Identification and concentration*)
 - Identification and concentration * of SML substances
 - Identification and concentration* of substances of specific concern to the cosmetic safety assessor (banned/restricted under Cosmetics Regulation & skin sensitisers)

Information ultimately available for the safety assessor (2)

- For each component in 'migrateable contact':
 - Statement on compliance with Food Contact Legislation (incl. applicable foods/simulants)
 OR
 - Reason of non compliance. (If linked to a specific substance: Identification and concentration*)
 - Identification and concentration * of SML substances
 - Identification and concentration* of substances of specific concern to the cosmetic safety assessor (banned/restricted under Cosmetics Regulation & skin sensitisers)



Challenges



Practical Issues

- Reporting levels for CMR substances 1 ppm too low
 - → aligned to 10 ppm with other substances of concern due to changed approach of Commission on CMR in cosmetics
- Expection by clients that final cosmetic packaging must be fully food contact compliant
 - → clarified that formal food compliance is not mandatory. The guidance approach can also be applied for non-compliant packaging. However, in this case more information may need to be provided on the reasons of non-compliance
- Many final cosmetic packaging respect food contact legislation composition, but do not do specific migration tests
 - → Clarified that relevant information on food contact compliance can be passed on regarding raw materials/compounded raw materials



Practical Issues

- Impossible to systematically analyse for 2000 substances of concern to cosmetic safety assessor (CPR Annex II, Annex III and CLP sensitisers)
 - Requires common sense approach
 - Most of the substances of concern would never reasonably expected to occur in packaging
 - Expert team reviewed the lists and identified +/- 400 substances whose presence in food contact packaging cannot be realistically excluded



Future Challenges

- Implement guidance into daily practice along all steps in the supply chain
- Expectation management of competent authorities
- Update and further refine list of reportable substances of concern
- Revision of the Cosmetics Regulation (+/-2025):
 - Ban of Endocrine Disruptors (similar to CMR) → many more reportable substances ?
- Revision of Packaging/Packaging Waste Directive:
 - Mandatory recycled content in packaging
 - Need to evolve the apporach to recycled materials
 - Competition for 'food grade' recycled materials



Thank you for your attention!