

CLASSIFICATION OF YOUR PRODUCT

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Are you sure that your product classifies as a cosmetic? Due to its claims, a product might sometimes classify under biocides or pharmaceuticals, instead of cosmetics.

WHAT WE DO FOR YOU:

We check your products and help you with the correct classification!



ASSIGN A RESPONSIBLE PERSON

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When imported from an extra-EU country, every cosmetic product must have a Responsible Person (RP) in the EU. The RP, among other things, accepts responsibility for the safety of a product, ensures compliance, and deals with local authorities. The Responsible Person also verifies your Product Information File (PIF), the technical documentation necessary to place a product on the EU market.

WHAT WE DO FOR YOU:

We act as a Responsible Person for your products and help create your PIF!



PRODUCT INFORMATION FILE (PIF)

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A PIF includes various mandatory documents which are needed to place your product on the EU market. Among the documents of a PIF, you can find formula, stability report, challenge test, compatibility test, safety assessment, different declarations, and others.

WHAT WE DO FOR YOU:

We guide you through the process of creating your PIF and verify the documentation!





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WHAT IS A SAFETY ASSESSMENT REPORT?

The safety assessment report is an important part of your PIF. It is a toxicological evaluation of the safety of the product and is prepared exclusively by a qualified expert.



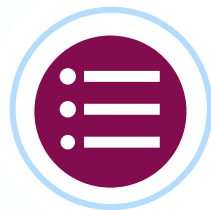
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LABELS

Labels must meet certain regulatory requirements, and claims must be substantiated with supporting documents.

WHAT WE DO FOR YOU:

We review your labels and proof of claims, and ensure they are compliant.



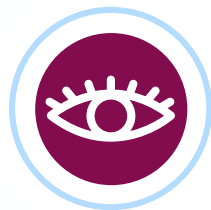
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COSMETIC PRODUCTS NOTIFICATION Portal CPNP

Once your PIF is complete, your product is ready for the EU market. Cosmetic products in the EU are notified on the Cosmetic Product Notification Portal (CPNP). Through this portal the Responsible Person submits the notification electronically to the Commission.

WHAT WE DO FOR YOU:

We help you find a Safety Assessor and check the report for you!



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POST - MARKET SURVEILLANCE

Undesirable effects might occur once the product is available on the market. In case of serious undesirable effects, the Responsible Person must notify the competent authority of the Member State where this happened and cooperate with them in order to minimize the risks.

Get a free consultation today!

SALES@OBELIS.NET



+32 27325954



obelis.net

