


AUTHORISATION OF BIOCIDAL PRODUCTS IN EUROPE

The way forward



Arcadis helps companies to comply with regulations regarding biocidal products. In particular, we support our customers with upcoming BPR deadlines, obtaining an overview of and interpreting the continuously extending biocide regulatory framework .

If you manufacture, supply or use biocidal active substances or biocidal products in the EEA, you need to be aware of the EU's Biocidal Product Regulation (EU 528/2012). It is designed to make sure that the potential risk caused by biocides are in balance with their expected benefits.

Ensuring compliance with EU BPR is a resource consuming process, which requires a broad range of expertise. The Arcadis experts in Belgium and Switzerland can support you in understanding and meeting the obligations under the BPR and the corresponding legislation in Switzerland.



Is your company **placing** biocidal products on the **market**, e.g.:

- Do you want to ensure **market continuation** of your product?
- Are your **suppliers** meeting the current biocidal requirements?
- Is your **active substance** under review or already approved?
- Does your company manufacture articles that are treated with, or intentionally incorporate, one or more biocidal products (**treated articles**)?



Do you want to ensure **market continuation** of your product?



Are your **suppliers** meeting the current biocidal requirements?



Is your **active substance** under review or already approved?



Does your company manufacture articles that are treated with, or intentionally incorporate, one or more biocidal products (**treated articles**)?



Is your company **using** one of the above listed biocidal products / treated articles?



Are your employees aware of the need to use authorized biocidal products only for certain **intended biocidal uses**?



Do you have a **system** in place **tracking** all active substances, their status and the biocidal products / treated articles they are used in?

How can we support you:

Are you a manufacturer or importer of active substances or biocidal products?



Dossier preparation

- Active substance dossiers
 - Inclusion in **Annex I**?
- Product dossiers (**single** product or product **families**)
- Consortia for biocides
- Obtaining access to the dossier of the active substance (**Letter of Access**)
- Demonstrating **technical equivalence** when buying from a “new” source
- Gathering of data required to build the dossier: coordination of **tests**, writing **waivers**
- **Risk assessments** (ecotoxicological and toxicological expertise in-house)
- Putting together the **IUCLID** dossier



Dossier submission

Pre-submission meetings with the eCA

- What kind of submission?
 - National authorization
 - Mutual recognitions
 - Union authorization
- Submitting the dossier through **R4BP3**



Compliance check for the manufacturing of **treated articles**

- Check the **authorization** of the **biocidal product** for the treatment of the article
- Listing of **label requirements**

Are you a user of biocidal products / treated articles?



BPR training for users and regulatory teams for biocides



Support with the interpretation of the legislation (e.g. assessment if the products are subject to another, closely related regulation e.g. MDR, PPP, REACH)



Development and implementation of compliance strategies (e.g. via an **inventory template** for documenting compliance status of biocidal products or **process flows** for the evaluation of products used)



Act as **helpdesk** or **Trustee**

Arcadis may provide you with just the support you need!

LEGENDA



manufacturer



user

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ABOUT ARCADIS

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