SAXOCON
Co-owned by Technical University of Denmark DTU

Disinfectant hygiene products for human use (type 1)

Everyone marketing disinfectant products for use in health care (e.g. antibacterial soaps, hand and surface disinfection) is facing a paradigm shift since the Biocidal Products Regulation *EC no 528/2012 (BPR)* introduces more comprehensive legal requirements for such products.

Products containing active substances approved under the BPR, must gain product approval by the Danish EPA, a similar authority in another EU country or the European Chemicals Agency, ECHA.

If you are planning to market your biocidal products in an EU Member State you must apply for product authorisation in that country. SAXOCON provides you with everything you need to apply for authorisation, including:

- Assistance with application types and forms
- Assistance with biocide legislation and documentation requirements set by BPR
- Assistance in ECHA Guidance and information requirements and how to fulfil these
- Assistance with the European application system R4BP3
- A toxicological risk assessment of human health effects, including exposure assessments carried out in accordance with relevant BRP guidances

Why choose us?

SAXOCON services for biocide manufacturers give you access to:

- Toxicologists with experience from the Danish Environmental Protection Agency with extensive knowledge in assessing biocidal products in various product types
- Extensive experience in state-of-the-art safety evaluations according to BPR legislation

Delivery

SAXOCON compiles all documentation necessary for approval of your product in relevant procedural ECHA templates.

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