TRISKELION

APPROVAL OF ACTIVE SUBSTANCES UNDER THE BPR

We understand that you have invested a lot in your (new) biocidal active substances. However, the process of gaining approval with the relevant authorities can be a daunting prospect. You can rely on the extensive experience of our highly specialized team to assist you with all aspects of the regulatory process. Please feel free to contact us. Our experts will be more than happy to answer any questions related to your current or future active substance application.

DUCARES B.V. | trading as TRISKELION

OUR KNOWLEDGE IS YOUR STRENGTH

www.triskelion.nl

info@triskelion.nl

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We support you in navigating through the challenges of the BPR to obtain approval as fast as possible. An early decision on the registration strategy is key, getting in touch with the competent authorities, and creating common ground and understanding for the subsequent dossier submission and evaluation process.

That is where we:

- Develop the strategy for dossier set-up, substance registration and testing
- Monitor subcontracted test work, including efficacy testing
- Prepare the IUCLID file and draft Risk Assessment Report
- Perform hazard, exposure and risk assessment
- Assess endocrine disrupting properties according to ECHA/EFSA guidance
- Submit the dossier via R4BP and interact with the authorities (eCA, ECHA)
- Participate and represent your company in registrant consortia or task forces

Our services are fully customized with short communication lines and a central point of contact dedicated to your requests.

TRISKELION offers testing and consultancy services to clients active in the food, feed, (agro)chemical and pharmaceutical industries. Our goal is to help develop products that improve the health and wellbeing of your customers.

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