PFAS in Pharmaceutical and Biotech Industry

Support concerning PFAS within your company

Since 2006 Per- und Polyfluoroalkyl substances (PFAS) are subject to stricter regulations as well as partially prohibitions. Since the beginning of 2023, a general prohibition of PFAS by the European Chemical Agency (ECHA) has been planned. But these substances are today still widely used in almost all branches of industry.

The Team of Valicare supports you in the evaluation of the actual status at your site, finding solutions and the preparation of documentation concerning the use of PFAS.

Our Competence

- Many years of experience with pharmaceuticals and medical devices in compliance to regulations
- Interdisciplinary team of natural scientists (biologists, chemists, and pharmacists) and engineers

Our Services & Support

- Preparation of the risk analysis for evaluation of used PFAS
- Support by decisions for the substitution of PFAS
- Preparation on and support during audits
- Preparation of documentation for analyzing the impact of PFAS to the product
- Project management and execution

Your Benefits

- Relief in terms of personnel and time by outsourcing the activity to experienced validation experts
- Accelerated implementation based on field-proven templates and experienced staff
- Audit-compliant documentation
- GMP compliance
- Risk reduction through competence

valicare
GMP Compliance Services & Solutions

Our services concerning PFAS reaching from support for the preparation of user requirement specifications (URS), over the assessment and mitigation of risks of using PFAS at your site, to the implementation of new equipment and systems as well as their qualification or validation. If you need consulting, feel free to contact us.

Contact us if PFAS are used in your processes or at your site.
Our experts will support you!

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