

CSV

Computerized System Validation

The validation of computerized systems (CSV) plays a central role in pharma, biotech and medtech. With its CSV team, Valicare offers comprehensive services for digital transformation in the pharmaceutical, biotechnology and medical technology sectors. We ensure that your business and production processes and the associated computerized systems adhere to strict compliance and GxP conformity.

With our consulting and training approach, we enable your team to **seamlessly** navigate complex regulatory environments. Partner with Valicare today to **sustain** reliable and continuously validated systems and processes that will keep your business **innovative** in the evolving life science industry.



Our competence

- Extensive experience in CSV, the use of the GAMP® and the execution of CSV compliance tasks
- Decades of experiences in dealing with manifold IT applications and infrastructures
- Engineers and scientists in the use of the 2nd Edition of GAMP® 5 in pharma, biotech and medtech

Our services & support

- Preparation and realization of the validation of the computerized system and its supporting business and production processes
- Creation of CSV documentation including SOPs and Policies
- CSV training in presence and online
- CSV audits, internal and external

Your benefits

- Reducing the workload and saving time by outsourcing CSV to SMEs
- Significant and reproducible results using SOP-based and competently executed, GxP-compliant CSV methods
- GxP- and audit-compliant CSV documentation
- Valid systems and processes

Digital transformation with CSV

Transformation is an ever-evolving journey that needs to fit seamlessly into your current business and production processes. Therefore, every digital transformation with its innovative processes requires thorough planning and preparation. A customized CSV strategy for the digital transformation is indispensable.

The implementation of this strategy requires continuous evaluation to ensure that it is consistent with existing processes and maintains their integrity and effectiveness. This is of vital importance for ensuring

- **Patient safety**
- **Product quality**
- **Daten integrity**

This guarantees compliance with regulations and GxP guidelines. This responsibility extends to your entire company and concerns not only the management, but also the department heads and every single employee who is directly or indirectly involved in CSV and compliance.

Contact us if you are planning the validation of your computerized systems and their related processes. We will support you!



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