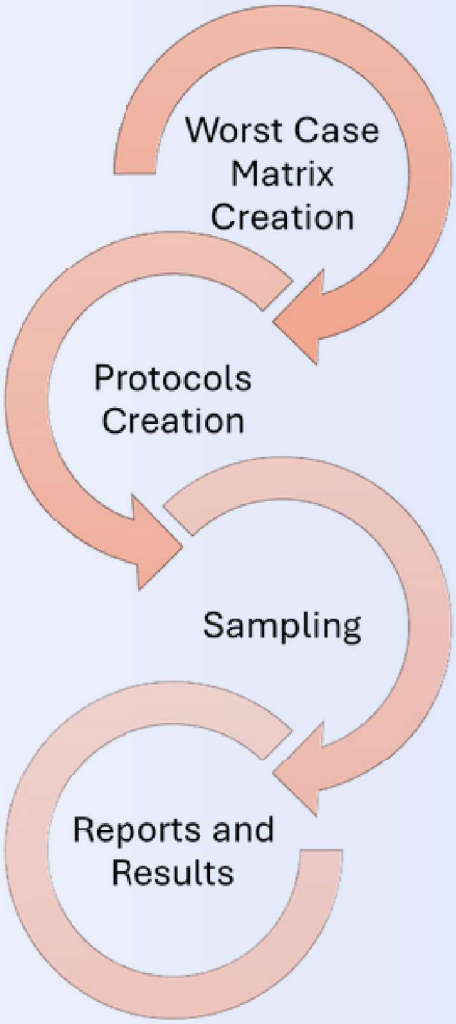
Cleaning Validation – Operational Documents Creation

Zamann Pharma Support offers consultancy services, specializing in the development of essential operational documents required for executing the Cleaning Validation process. Our focus is on creating comprehensive documents such as worst-case matrices, validation protocols, reports, and risk assessments tailored to the client's specific needs and regulatory standards

Worst Case Matrix Development

We identify worst-case scenarios for Cleaning Validation, considering factors such as equipment design, product characteristics, and cleaning procedures. Our team develops detailed worst-case matrices that systematically outline the most challenging cleaning conditions to be addressed during validation testing.



Protocols Creation

Zamann assists clients in drafting validation protocols that define the scope, objectives, methodologies, acceptance criteria, and testing procedures for cleaning validation activities.
We ensure that protocols are comprehensive, clear, and aligned with requirements, facilitating efficient execution and documentation of validation tests.

Reports Creation and Results Compilation

Our consultancy service includes compiling validation results and generating comprehensive reports summarizing the findings.
We meticulously document test outcomes, observations, deviations, and conclusions, ensuring clarity and accuracy in reporting the outcomes of cleaning validation activities.

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- Quality Consulting and Risk Management
- Qualification and Validation
- Personnel Qualification
- Customer-specific provision of QMS

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