

Quality Management Review – GAP Assessment

Zamann Pharma Support offers specialized consultancy services, aimed at evaluating the compliance level of the implemented landscape governing Quality Management Review activities. With meticulous attention to detail, we assess the effectiveness and alignment of existing processes, policies, and documentation with international regulations and industry best practices

Regulatory Compliance Evaluation

We conduct a comprehensive analysis of relevant regulations, including FDA, ICH Q10, and EMA guidelines, to identify the necessary standards and requirements for Quality Management Review.

Our experts evaluate the existing documentation and practices against these regulatory benchmarks, identifying gaps and areas for improvement.

GAP Analysis and Documentation Review

We perform a thorough GAP analysis to compare the current state of Quality Management Review processes against regulatory expectations and industry standards. Our team reviews existing documentation, including policies, SOPs, and quality records, to assess their adequacy and alignment with regulatory requirements.

Recommendations and Action Plan Development

Based on the findings from the GAP assessment, we provide actionable recommendations and develop a tailored action plan to address identified deficiencies and improve compliance. We prioritize corrective actions, establish timelines, and allocate resources effectively to enhance the Quality Management Review landscape.

