Project Consulting in Pharma and Biotech

ASPIRAS

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We've got what it takes

. . . via external specialists for your QA-Team

Your questions:

- Audits und inspections come up and your quality-team has its work cut out. Who helps you?
- The quality system needs to be optimised or restructured. Who reorganises and sets up the new procedures?
- The regular SOP review is due and the annual PQRs need to be drafted. Who can undertake this task?

Our services:

ASPIRAS helps you compensating capacity bottlenecks and assumes tasks like e.g. composing of QM-Manuals, Site Master Files, SOPs, work instructions or Product Quality Reviews. We also assist you in processing regulatory aspects as well as in preparation and conducting of internal or external audits. The objective view on your quality system by our staff increases the potential for optimisation.

Due to their experience ASPIRAS' employees can adjust oneself flexibly on special customers concerns to develop customised solution in a team.

ASPIRAS ensures audit-proof authoring of controlled documents. As well we can take AMG responsibility for you according to §14 and provide the Qualified Person.

Your contact:

Cathrin Pauly, pharmacist, MBA

- More than 25 years experience in well-known pharmaceutical companies
- Specialisation in project management and business development, Qualified Person

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