ISO/IEC 17025:2017 International Standard

Services	Deliverables
• ISO/IEC 17025:2017 International Standard for Laboratories Implementation	▪ 8 Hours or 1 day. Note that this service is typically 17 - 20 days at your pace; Assessing your current QMS, defining policies and procedures, training auditors, conducting a pre-registration audit & registration assistance.

ISO/IEC 17025:2017 – Laboratory Quality Management System

Improvement and Standardization

If your company performs laboratory testing or calibration, you should consider becoming certified to the International Standard for Laboratories, ISO/IEC 17025. While it closely follows the ISO-9001 Standard, it has specific differences related to sampling of product, equipment calibration, environmental conditions and other areas. AKA will take you through the process step by step to ensure you achieve certification and can continue to serve your customers. Here's how.

AKA uses personnel who have extensive experience in implementing ISO/IEC 17025 at a variety of different types of laboratories. So even if you are starting from square one, our consultants know the standard and can apply it to your type of laboratory. They also can tell you where you don't have to apply specific elements of the standard. Our goal is to provide you with a Quality Management System that works for your company, with all the necessary documentation…and not one page more.

While the implementation effort that is needed is dependent upon size of the company, the number of laboratory processes, management support, and even multiple locations, implementing a compliant **ISO/IEC 17025 Quality Management System will typically take 17 – 20 days spread over 10 – 12 months…at your pace.** Deliverables typically include assistance with:

1. A Gap assessment of your current policies and procedures;
2. An overview of the ISO/IEC 17025 Standard for your team;
3. Development of quality policy and quality objectives;
4. Development and documentation of a quality policy manual;
5. Development and documentation of quality procedures;
6. Development and documentation of work instructions for product sampling, equipment calibration, environmental conditions, records management and other areas as needed;
7. Training of internal auditors – 2-day course;
8. Conducting a pre-registration audit;
9. Selection of a Registrar;
10. Assistance with the certification audit including addressing any corrective actions that may arise during the audit.

Give us a call today and let us walk you through the process of implementing the ISO/IEC 17025 International Standard for Laboratories.

ADVANTAGE KENTUCKY ALLIANCE