ISO/IEC 17025:2017 Pre-Assessment

Services	Deliverables
• ISO/IEC 17025:2017 - Pre-Assessment	▪ Up to 16 hours on-site Document and Process Review

Pre-assessment is used to identify any last steps necessary for registration

ISO/IEC 17025:2017 – QMS Pre-Assessment

(This service is needed to determine readiness for registration)

This is an audit/review of your Laboratory's Quality Management System and its processes prior to the actual ISO/IEC 17025 assessment by a Registrar. The objective is to determine your company's readiness for registration by determining if all the requirements found in the ISO/IEC 17025 International Standard for Laboratories have been met and are effective.

Objectives:

- The primary objective of a pre-assessment is to determine the company's readiness for registration by ensuring all the requirements found in the ISO/IEC 17025 International Standard for Laboratories have been met and are effective.
- This audit is very similar to an actual certification audit by a Registrar.
- Findings will be documented and corrective actions will be discussed for correction.
- **The pre-assessment audit activities can be used as one of the company's required internal audits.**

Upon completion, the company will;

- Be assured they are ready for ISO/IEC 17025 registration;
- Have a complete audit of company's QMS processes against the International Standard;
- Have any non-conformances documented with corrective actions determined; and,
- Incorporate the pre-assessment audit into company's internal audit program.

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