

Quality Policy

Abingdon Analytical is committed to operating and continually improving an effective Quality Management System that ensures the competence, impartiality, and consistent operation of the laboratory in accordance with ISO/IEC 17025:2017 — General requirements for the competence of testing and calibration laboratories.

The laboratory provides testing services to support the performance evaluation, verification, and validation of in vitro diagnostic (IVD) devices. These services may include analytical performance testing and other laboratory activities within the laboratory's defined scope. Activities are conducted in accordance with controlled procedures, study-specific protocols, and agreed customer requirements. Abingdon Analytical generates objective laboratory results and does not provide clinical interpretation, make diagnostic decisions, or authorise the release of IVD products. Where results are compared with predefined acceptance criteria, conclusions are limited to the agreed scope and reported in accordance with applicable procedures.

Abingdon Analytical commits to:

- Performing laboratory activities competently, impartially, and consistently
- Ensuring analytical results are technically valid, reliable, accurately reported, and suitable for the agreed testing purpose
- Using competent, trained, and appropriately authorised personnel, suitable facilities, and appropriately maintained and calibrated equipment
- Applying recognised guidance and scientifically sound methods where appropriate
- Protecting the confidentiality and integrity of customer and proprietary information
- Identifying and managing risks to impartiality, the validity of results, and the effectiveness of laboratory activities
- Complying with applicable legal, regulatory, contractual, and accreditation requirements
- Continually improving the effectiveness of the Quality Management System

Laboratory management is committed to establishing, implementing, and maintaining the Quality Management System, and to ensuring that this Quality Policy is communicated, understood, implemented, and upheld at all levels of the organisation.

Quality objectives are established, monitored, and reviewed through management review to support the continuing suitability, adequacy, and effectiveness of the management system and laboratory activities.



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Signatory Table

Action Name	User Name	Title	Signature Date
Send for Review (Written By)	Ryan Desforges	Quality Lead AA UK	06-Oct-2026 15:36
Review	Baranse Ozturk	Laboratory Manager AA UK	06-Oct-2026 16:01
Send for Approval	Ryan Desforges	Quality Lead AA UK	07-Oct-2026 11:35
Approve	Baranse Ozturk	Laboratory Manager AA UK	07-Oct-2026 13:17
QA Approval - Skip Training	Ryan Desforges	Quality Lead AA UK	07-Oct-2026 13:55

* Dates are displayed according to the system time zone: (GMT+01:00) British Summer Time (Europe/London)