



4367
UKAS accredited Proficiency
Testing Provider

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QUALITY POLICY

EMQN CIC is committed to helping ensure diagnostic molecular genetic laboratory test results are accurate, reliable and comparable wherever they are produced. EMQN CIC will provide a high quality and timely service which takes into account the needs and requirements of its users.

Our commitment to quality:

EMQN CIC will:

- establish and operate a quality management system designed to integrate the organisation, its processes, procedures and resources;
- set quality objectives and plans in order to implement this quality policy.
- ensure that all personnel are familiar with this policy, the quality manual and all procedures relevant to their work.
- commit to the health, safety and welfare of all its staff and visitors by providing a safe working environment.
- uphold professional values and be committed to good professional practice and conduct.

EMQN CIC will ensure compliance with the UK GDPR and Data Protection Act 2018.

- Establishing a framework to ensure EMQN meets its obligations under the regulations guided by the six data protection principles: to ensure data is processed fairly, lawfully and in a transparent manner, used only for limited, specified stated purposes and not used or disclosed in any way incompatible with those purposes, that data is adequate, relevant, limited to what is necessary, accurate and, where necessary, up to date, that data is not kept for longer than necessary and kept safe and secure.
- Establishing policy & procedures for data management, storage, processing & validation.

EMQN CIC will comply with the following standard as required by the United Kingdom Accreditation Service (UKAS):

- ISO17043

EMQN CIC may require subcontractors to comply with relevant clauses of the following standards:

- ISO 15189 / 17025 (where relevant)

EMQN is committed to:

- supporting its staff to ensure it provides a full and effective service to its users;
- the proper procurement and validation of EQA resources so as to ensure their correct performance in external laboratory examinations;
- appropriate handling and distribution of EQA resources to ensure they meet the needs of the users in a timely fashion;
- the reporting of EQA results in ways which are timely, confidential and accurate;
- the assessment of user satisfaction and internal audit in order to produce continual quality improvement.
- complying with current environmental legislation.

Signed on behalf of EMQN

Date: 22 October 2024

EMQN Director

Important note: The complete history of this document including its author, authoriser(s) and revision date, can be found on Q-Pulse

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