intertek
Total Quality. Assured.

STAY ALIGNED WITH EMA EXPECTATIONS ON GMP AUDITS

The European Medicines Agency (EMA) has recently updated its GMP Q&A to clarify the role of Third-Party for audit reports, a key concern for pharmaceutical companies and Qualified Persons (QPs). These updates reinforce the principles that have long been part of Intertek's audit methodology.

INTERTEK'S QUALITY APPROACH: BUILT FOR TRUST

When subcontracting audits to Intertek, you benefit from:

Auditor impartiality

Effective management of conflicts of interest is a key element of our process, ensuring integrity and transparency of our auditors.

Intertek auditors declare they have no conflict of interest with the supplier being audited. The signed declaration is part of the audit report.

Certified quality system

Intertek auditing process is ISO 9001 certified, ensuring the highest standards of quality and integrity in every audit we conduct.

Our auditors are qualified and trained according to a robust procedure.

Comprehensive audit reports

The quality of our audit reports is our top priority, reflecting our commitment to excellence and reliability.

All Intertek audit reports are carefully designed and reviewed by technical experts to effectively support and assist the Qualified Person in their final written assessment.

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