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CLSI recognizes the important contributions of laboratory professionals and the health care community and applauds their efforts in the global fight against COVID-19. We've compiled a list of documents that have been identified as helpful for the laboratory community's use during the current pandemic. [View List](#)

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Clinical & Laboratory Standards Institute: CLSI Guidelines

CLSI Standards: Guidelines for Health Care Excellence. Developed by our members for use by the global laboratory community, CLSI's consensus-based medical laboratory standards are the most widely recognized resources for continually improving testing quality, safety, and efficiency.

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CLSI Style Guide for Authors and Editors

"CLSI documents VET03 and VET04 outline the standards for performing and interpreting antimicrobial susceptibility testing

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for clinically-significant aquaculture pathogens. These documents have been updated and reorganized with the end-user in mind, with input from global experts in aquatic medicine.

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M52: Verification of Commercial Microbial Identification

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- the acceptance of a document as a consensus standard or guideline. Most NCCLS documents are subject to two levels of consensus—"proposed" and "approved." Depending on the need for field evaluation or data collection, documents may also be made available for review at an

EP17-A Protocols for Determination of Limits of Detection

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CLSI C28-A3 - Defining, Establishing, and Verifying ...

CLSI C51-A Expression of Measurement Uncertainty in Laboratory Medicine; Approved Guideline. This guideline describes a practical approach to assist clinical laboratories in developing and calculating useful estimates of measurement uncertainty, and illustrates their application in maintaining and improving the quality of measured values used in patient care.

CLSI C51-A - Expression of Measurement Uncertainty in ...

CLSI C53-A Vol. 30 No. 12 Characterization and Qualification of Commutable Reference Materials for Laboratory Medicine; Approved Guideline. This guideline provides recommendations for the characterization, assessment of commutability, and assignment of analyte concentration or activity values to reference materials (RMs) that are used for calibration and trueness assessment of in vitro

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