Archived Document

This archived document is no longer being reviewed through the CLSI Consensus Document Development Process. However, this document is technically valid as of October 2022. Because of its value to the laboratory community, it is being retained in CLSI's library.



June 2004

M22-A3

Quality Control for Commercially Prepared Microbiological Culture Media; Approved Standard—Third Edition

NOTE: CLSI document M22-A3 no longer applies to US laboratories subject to the Clinical Laboratory Improvement Amendments of 1988 (CLIA). The guidance provided in this standard does not replace the need for assessment of an individualized quality control plan (IQCP). The Centers for Medicare & Medicaid Services no longer recognizes the categories of "exempt" and "nonexempt" media for the purposes of quality control but, as of January 2016, instead directs laboratories to develop an IQCP for applicable media used in their facilities or to follow the CLIA quality assurance regulations as written. M22-A3 might be applicable to international laboratories.

This document contains quality assurance procedures for manufacturers and users of prepared, ready-to-use microbiological culture media.

A standard for US application developed through the Clinical and Laboratory Standards Institute consensus process.

