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(1117) MICROBIOLOGICAL BEST LABORATORY PRACTICES

INTRODUCTION

Good laboratory practices in a microbiology laboratory consist of activities that depend on several principles: aseptic technique, control of media, control of test strains, operation and control of equipment, diligent recording and evaluation of data, and training of the laboratory staff. Because of the inherent risk of variability in microbiology data, reliability and reproducibility are dependent on the use of ac-

cepted methods and adherence to good laboratory practices.

MEDIA PREPARATION AND QUALITY CONTROL

Media Preparation

Culture media are the basis for most microbiological tests. Safeguarding the quality of the media is therefore critical to the success of the microbiology laboratory. Media preparation, proper storage, and quality control testing can ensure a consistent supply of high-quality media.

It is important to choose the correct media or components in making media based on the use of accepted sources or references for formulas. The manufacturer's formula and instructions for preparation routinely accompany dehydrated media and ready-made media. Because different media types may have different preparation requirements (e.g., heating, additives, and pH adjustment), it is important to follow these instructions to ensure preparation of acceptable media quality. A certificate of analysis describing expiration dating and recommended storage conditions accompanies ready-made media, as well as the quality control organisms used in growth-promotion and selectivity testing of that media.

Water is the universal diluent for microbiological media. Purified Water is most often used for media preparation, but in certain cases the use of deionized or distilled water may be appropriate. Water of lesser quality should not be used for microbiological media preparation. The volume of the water used should be recorded.

Consistent preparation of media requires accurate weighing of dehydrated media or media constituents. A calibrated balance with the appropriate weight range for the ingredients should be used (See Weighing on an Analytical Balance (1251)). Clean weighing containers and tools (such as spatulas) should be used to prevent foreign substances from entering the formulation. The weight of the components should be recorded.

Dehydrated media should be thoroughly dissolved in water before dispensing and sterilization. If heating is necessary to help dissolve the media, care should be taken not to overheat media, because all culture media, to a greater or lesser extent, are heat-sensitive. Equipment used in the preparation of media should be appropriate to allow for controlled heating, constant agitation, and mixing of the media. Darkening of media (Maillard-type reaction or nonenzymatic browning) is a general indication of overheating. When adding required supplements to media, adequate mixing of the medium after adding the supplement should be performed.

Preparation of media in poorly cleaned glassware can allow inhibitory substances to enter the media. Inhibitory substances can come from detergent residue after cleaning glassware or from prior materials used in the glassware. Be sure that the cleaning process removes debris and foreign matter, and that the detergent is thoroughly rinsed out with Purified Water. See *Cleaning Glass Apparatus* (1051) for additional guidance.

Sterilization of media should be performed within the parameters provided by the manufacturer or validated by the user. Commercially prepared media should provide documentation of the sterilization method used. Autoclaving by moist heat is the preferred sterilization technique, except in instances when boiling is required in order to avoid deterioration of heat-labile components of the media. Sterilization by filtration may also be appropriate for some formulations.

The effects of the sterilization method and conditions on the media should be validated by sterility and growth-promotion testing of the media. In addition, if sterilized by moist heat, the autoclave cycle should be validated to en-

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Table of Contents Usp 36 Chapter 1117

1. Understanding the eBook Usp 36 Chapter 1117
 - The Rise of Digital Reading Usp 36 Chapter 1117
 - Advantages of eBooks Over Traditional Books
2. Identifying Usp 36 Chapter 1117
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Usp 36 Chapter 1117
 - User-Friendly Interface
4. Exploring eBook Recommendations from Usp 36 Chapter 1117
 - Personalized Recommendations
 - Usp 36 Chapter 1117 User Reviews and Ratings
 - Usp 36 Chapter 1117 and Bestseller Lists
5. Accessing Usp 36 Chapter 1117 Free and Paid eBooks

- Usp 36 Chapter 1117 Public Domain eBooks
- Usp 36 Chapter 1117 eBook Subscription Services
- Usp 36 Chapter 1117 Budget-Friendly Options
- 6. Navigating Usp 36 Chapter 1117 eBook Formats
 - ePub, PDF, MOBI, and More
 - Usp 36 Chapter 1117 Compatibility with Devices
 - Usp 36 Chapter 1117 Enhanced eBook Features
- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Usp 36 Chapter 1117
 - Highlighting and Note-Taking Usp 36 Chapter 1117
 - Interactive Elements Usp 36 Chapter 1117
- 8. Staying Engaged with Usp 36 Chapter 1117
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Usp 36 Chapter 1117
- 9. Balancing eBooks and Physical Books Usp 36 Chapter 1117
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection Usp 36 Chapter 1117
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Usp 36 Chapter 1117
 - Setting Reading Goals Usp 36 Chapter 1117
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Usp 36 Chapter 1117
 - Fact-Checking eBook Content of Usp 36 Chapter 1117
 - Distinguishing Credible Sources
- 13. Promoting Lifelong Learning
 - Utilizing eBooks for Skill Development

- Exploring Educational eBooks
- 14. Embracing eBook Trends
 - Integration of Multimedia Elements
 - Interactive and Gamified eBooks

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