Bailey & Scott's

Diagnostic Microbiology

Bailey & Scott's

Diagnostic Microbiology

Ellen Jo Baron

Ph.D., Diplomate (ABMM)

Adjunct Assistant Professor of Medicine University of California School of Medicine Los Angeles, California

Sydney M. Finegold

MD., SM(AAM), Diplomate (ABMM)

Associate Chief of Staff for Research and Development Wadsworth V.A. Medical Center Professor of Medicine and of Microbiology and Immunology University of California School of Medicine Los Angeles, California

EIGHTH EDITION

with 313 color illustrations

The C. V. Mosby Company

ST. LOUIS * BALTIMORE * PHILADELPHIA * TORONTO



Editor: Stephanie Manning
Assistant Editor: Anne Gunter
Design: Rey Umali
Editing and Production: CRACOM Corporation

EIGHTH EDITION

Copyright © 1990 by The C. V. Mosby Company

All rights reserved. No part of this publication may be reproduced, stored in a retrieval system, or transmitted, in any form or by any means, electronic, mechanical, photocopying, recording, or otherwise, without prior written permission from the publisher.

Previous editions copyrighted 1962, 1966, 1970, 1974, 1978, 1982, 1986

Printed in the United States of America

The C. V. Mosby Company 11830 Westline Industrial Drive, St. Louis, Missouri 63146

Library of Congress Cataloging-in-Publication Data

Baron, Ellen Jo.

Bailey and Scott's diagnostic microbiology.—8th ed./Ellen Jo Baron, Sydney M. Finegold.

p. cm.

Rev. ed. of: Baily and Scott's diagnostic microbiology. 7th ed./ Sydney M. Finegold, Ellen Jo Baron. 1986.

Includes bibliographical references.

ISBN 0-8016-0344-7

 Diagnostic microbiology. I. Bailey, W. Robert (William Robert), 1917- Diagnostic microbiology. II. Finegold, Sydney, M.,

1921- III. Title. IV. Title: Diagnostic microbiology.

[DNLM: 1. Microbiological Technics. QW 25 B265b] QR67.B37 1990 616'.01'028—dc20

DNLM/DLC

for Library of Congress

89-13698

CII

To
James C. Taylor,
Alma and Lee Baron,
and Mary L. Finegold
with loving appreciation for
their constant guidance
and continued understanding
again

Contributors

CHAPTER AUTHORS

O. George W. Berlin, Ph.D., M.A., MT(ASCP)
Clinical Microbiology Laboratory, Medical Center,
University of California—Los Angeles

W. Lawrence Drew, M.D., Ph.D.

Department of Microbiology and Infectious
Diseases, Mount Zion Hospital and Medical
Center, San Francisco, California; University of
California—San Francisco

Martha A.C. Edelstein, B.A., MT(ASCP) Clinical Microbiology Laboratory, Hospital of the University of Pennsylvania, Philadelphia

Lynne Shore Garcia, M.S., CLS(NCA), MT(ASCP) Clinical Microbiology Laboratory, Medical Center, University of California—Los Angeles

Glenn D. Roberts, Ph.D.

Department of Laboratory Medicine, Section of Clinical Microbiology, Mayo Medical School, Mayo Clinic and Mayo Foundation, Rochester, Minnesota

CHAPTER CONSULTANTS

Richard R. Facklam, Ph.D.

Respiratory and Special Pathogens Laboratory Branch, Centers for Disease Control, Atlanta, Georgia Gerald L. Gilardi, Ph.D.

Department of Microbiology, North General Hospital, New York, New York

Janet A. Hindler, M.S., MT(ASCP)

Clinical Microbiology Laboratory, Medical Center, University of California—Los Angeles

J. Michael Janda, Ph. D.

Microbiology Diseases Laboratory, California Department of Health Services, Berkeley, California Michael Pfaller, M.D.

Veterans Administration Medical Center; Department of Pathology, University of Iowa College of Medicine, Iowa City, Iowa

Ella M. Swierkosz, Ph.D.

Department of Diagnostic Microbiology and Virology, Cardinal Glennon Children's Hospital: St. Louis University School of Medicine, St. Louis, Missouri.

Preface

Diagnostic Microbiology presents a comprehensive view of medical microbiology from the stand-points of the organization and function of a clinical microbiology laboratory (Part One), likely agents associated with infectious syndromes (Part Three), and procedures for identification and susceptibility testing of infecting agents (Parts Two and Four). Reviews of the previous edition suggested that Diagnostic Microbiology is valuable as a textbook for students of medical technology, medical microbiology, pathology, infectious diseases, and infection control and as a laboratory reference manual and procedure book. We have strived to maintain that focus.

We welcome the continued participation of W. Lawrence Drew (virology), Martha A. C. Edelstein (anaerobes), Lynne Shore Garcia (parasitology), and Glenn D. Roberts (mycology), whose authoritative chapters add so much to the book. We have also received able assistance from several new contributors, George Berlin (mycobacteriology), Michael Pfaller (non-spore-forming, gram-positive bacilli), and Ella Swierkosz (chlamydiae), who helped to provide the updated information presented in this edition.

Certain chapters were evaluated extensively by Gerald Gilardi, Janet Hindler, J. Michael Janda, Richard Facklam, and Maurice White. Henry Isenberg's suggestions are always valuable. Several of the new photographs were taken by Peter Rose. We are grateful for their expert contributions. Stephanie Manning at Mosby and Mary Espenschied at CRACOM have been able editors, and we appreciate their support and expertise. As always, we welcome

input from our readers, many of whom helped us to correct errors and modify topics that were presented in the seventh edition.

The eighth edition continues the format introduced in the seventh edition, with chapter outlines listed at the beginning of each chapter. The major format change is the introduction of full color throughout, which allows all illustrations to be placed adjacent to their corresponding text. Procedures, still presented in boxes on one page or facing pages, now include a statement of the principle of the test, quality control guidelines, expected results, and performance schedule recommendations, in addition to the method itself. Chapter 4 has been retitled to reflect its current content, which has been expanded to include material on laboratory manage-

control in a final part and an a party fail to

ment and test selection. Of course, the entire book has been updated, and new identification tables have been added in certain areas. We urge readers to peruse How to Use This Book, p. xi, which explains the organization of the text and outlines the content.

The importance of microbiology continues to expand in this era of new infectious agents and difficult-to-manage infections. Although there is a definite trend toward automation and newer techniques, such as DNA probes, there will always be a place for conventional microbiology and for competent microbiologists.

Ellen Jo Baron Sydney M. Finegold

How To Use This Book

The organization of the seventh edition of this book has been retained. Part One contains material of general importance to the practice of diagnostic clinical microbiology, including laboratory safety, quality assurance, and laboratory management, as well as the role of the microbiology laboratory in infection control and epidemiologic studies. Part Two encompasses methods for specimen handling, initial observations, and procedures used to detect agents of infectious diseases. New technologies, automation, and immunologic methods are included in this section, as are methods for susceptibility testing. The pathogenesis of infectious diseases in relation to the site of infection or the organ system involved is covered in Part Three, which highlights the agents likely to be recovered from each body site or syndrome. Part Four details methods for detection and identification of specific infectious agents.

Each chapter begins with an outline of the sections within to facilitate finding a particular area and to provide a quick overview of the contents. Reference material has been reorganized into three appendixes and the glossary. Appendix A contains formulas for media and reagents, Appendix B contains formulas for commonly used stains, and Appendix C lists the names and addresses of commercial product suppliers, whose names are simply mentioned in parentheses in the text when they are cited. Even companies not cited specifically in the book are listed; Appendix C should be a valuable resource for laboratory managers. The Glossary has been expanded to include numerous new terms and the abbreviations used throughout the text are incorporated alphabetically into the Glossary. The use of boldface type in the text identifies all terms that have been defined in the Glossarv.

Bailey & scores Diagnostic
Microbiology

Contents

PART ONE Organization and Function of the Clinical Microbiology Laboratory

- Diagnostic microbiology: purpose and philosophy, 3
- 2. Laboratory safety, 8
- 3. Laboratory organization and quality assurance, 17
- 4. Managing the clinical microbiology laboratory: maximizing patient care in a cost-conscious environment, 24
- 5. Hospital epidemiology, 36

PART TWO Handling Clinical Specimens for Microbiological Studies

- 6. Selection, collection, and transport of specimens for microbiological examination, 49
- Optical methods for laboratory diagnosis of infectious diseases, 64
- 8. Cultivation and isolation of viable pathogens, 81
- 9. Conventional and rapid microbiological methods for identification of bacteria and fungi, 100
- Nontraditional methods for identification and detection of pathogens or their products, 127
- Principles of automated methods for diagnostic microbiology, 142

XH

- Diagnostic immunological principles and methods, 157
- Methods for testing antimicrobial effectiveness, 171

PART THREE

Etiologic Agents Recovered from Clinical Material

- 14. Microorganisms encountered in the blood, 197
- Microorganisms encountered in the cerebrospinal fluid, 213
- Microorganisms encountered in the respiratory tract, 223
- Microorganisms encountered in the gastrointestinal tract, 238
- Microorganisms encountered in the urinary tract, 253
- Genital and sexually transmitted pathogens, 263
- Microorganisms encountered in wounds, abscesses, skin, and soft tissue lesions, 279
- Microorganisms encountered in solid tissue, bone, bone marrow, and body fluids, 289
- 22. Infections of the head and neck, 301
- 23. Infections in the vulnerable host, 309

PART FOUR

Methods for Identification of Etiologic Agents of Infectious Disease

- Micrococcaceae: staphylococci, micrococci, and stomatococci, 323
- 25. Streptococci and related genera, 333
- Aerobic gram-negative cocci (Neisseria and Branhamella), 353
- 27. Enterobacteriaceae, 363
 Budvicia, Buttiauxella, Cedecea, Citrobacter,
 Edwardsiella, Enterobacter, Erwinia,
 Escherichia, Ewingella, Hafnia, Klebsiella,
 Kluyvera, Koserella, Leclercia, Leminorella,
 Moellerella, Morganella, Obesumbacterium,
 Proteus, Providencia, Rhanella, Salmonella,
 Serratia, Shigella, Tatumella, Xenorhabdus,
 Yersinia

- Nonfermentative gram-negative bacilli and coccobacilli, 386
 Pseudomonas, Alcaligenes, Acinetobacter, Moraxella, Oligella, Eikenella, Flavobacterium, Sphingobacterium, Weeksella, Agrobacterium
- Gram-negative facultatively anaerobic bacilli and aerobic coccobacilli, 408
 Francisella, Brucella, Bordetella, Haemophilus, Actinobacillus, Pasteurella, Kingella, Capnocytophaga, Cardiobacterium, HB-5
- Vibrio and related species, Aeromonas, Plesiomonas, Campylobacter, and others, 431
- Spirochetes and other spiral-shaped organisms, 445
 Treponema, Borrelia, Leptospira, Brachyspira
- Aerobic or facultative spore-forming rods (Bacillus species), 451
- Aerobic, non-spore-forming, gram-positive bacilli, 457
 Listeria, Erysipelothrix, Corynebacterium, Rhodococcus, Arcanobacterium, Actinomyces pyogenes, Nocardia, Nocardiopsis, Actinomadura, Dermatophilus, Streptomyces, Rothia, Kurthia, Oerskovia, Lactobacillus
- Processing clinical specimens for anaerobic bacteria: isolation and identification procedures, 477
 Martha A. C. Edelstein
- Anaerobic gram-positive bacilli, 508
 Clostridium, Actinomyces, Bifidobacterium, Eubacterium, Lactobacillus, Propionibacterium
 Martha A. C. Edelstein
- Anaerobic gram-negative bacilli, 529
 Bacteroides, Porphyromonas, Fusobacterium, anaerobic vibrios, and curved rods
 Martha A. C. Edelstein
- 37. Anaerobic cocci, 549

 Peptococcus, Streptococcus,
 Peptostreptococcus, Veillonella,
 Acidaminococcus, Megasphaera,
 Ruminococcus, Coprococcus
 Martha A. C. Edelstein
- 38. Chlamydia, Mycoplasma, and Rickettsia, 558

- 39. Unclassified or unusual but easily cultivated etiologic agents of infectious disease, 572
 Bartonella, Chromobacterium,
 Dermatophilus, Gardnerella, Legionella,
 Prototheca, Simonsiella, Stomatococcus,
 Streptobacillus, thermophilic bacteria,
 unnamed bacteria
- New, controversial, difficult-to-cultivate, or noncultivatable etiologic agents of disease, 592
- Mycobacteria, 597
 George W. Berlin

- Laboratory methods in basic virology, 641
 W. Lawrence Drew
- Laboratory methods in basic mycology, 681
 Glenn D. Roberts
- 44. Laboratory methods for diagnosis of parasitic infections, 776

Lynne Shore Garcia

Appendix A, A-1

Appendix B, A-35

Appendix C, A-50 Glossary, G-1

Part One

Organization and Function of the Clinical Microbiology Laboratory



Lagration and Function is a second or Street Street

Constitute of this nature should be not freed to the

Diagnostic Microbiology: Purpose and Philosophy

sub-to-season, out all companie in mentions

1.1 Purpose of Diagnostic Microbiology

1.2 Responsibility to the Clinician and to the Patient

the periods viewed who was nation before on the

week out by iterated by low return of a specimen

1.3 Specimen Collection and Transport

Noncultural Methods of Diagnosis 1.4

Rejection of Specimens 1.5

Extent of Identification Required

1.7 Quantitation of Results

1.8 Expediting Results

1.9 Interaction with the Clinician

Clinical microbiologists are part of the health team and serve an important role in the diagnosis, management, and prevention of infections in patients. Microbiologists should take pride in this role and should feel responsibility commensurate with it.

1.1. Purpose of Diagnostic Microbiology

The purpose of clinical microbiology is to work closely with clinicians and other health team members to provide diagnosis and optimum management of infectious disease in patients and to prevent the spread of infection to other individuals. In general, it is important to document the presence of infection. to determine its specific nature, and to provide appropriate therapy early in the course of the illness. Early diagnosis and treatment are usually more urgent with infectious diseases than with any other type of disease process. In many infections, speedy treatment is crucial. Both mortality and morbidity may be reduced significantly if proper treatment is provided early. In others, hospitalization may be avoided or shortened and surgery may be avoided if specific antimicrobial therapy is provided early. While the responsibility is large, the satisfaction that may be achieved in successfully diagnosing and treating patients with infectious diseases can be great. It is very gratifying to see rapid improvement in a critically ill patient; all members of the health team share in this gratification.

1.2. Responsibility to the Clinician and to the Patient

Microbiology laboratory service must be available at all times. This means that microbiologists must be on call if it is not reasonable to have the laboratory staffed at all times. Others may be able to set up certain types of specimens in the absence of the microbiologist, but the expertise of the microbiologist may be required to properly interpret a Gram stain or colonial morphology late at night. In assessing early findings, one should provide as much detail as possible and one should not hesitate to indicate the various possibilities as to the microbiological diagnosis. The clinician sets up his or her own differential diagnosis based on the clinical picture, roentgenograms, blood studies, and so on. The diagnostic possibilities, as seen by a competent microbiologist, are of paramount importance to the clinician in further narrowing down or substantiating a diagnosis. The microbiologist must be alert to unusual findings. As suggested earlier, speed is critical in serious illness caused by infectious agents. Good microbiologists can usually find one or more ways to expedite presumptive identification of microorganisms in clinical specimens. If the health team is to function with maximum efficiency, the members of that team must communicate freely and frequently, particularly when dealing with seriously ill patients. The physician should lead in this respect, but if he or she does not, the others on the team should initiate communication when it may be helpful. Our primary concern always must be the patient. The clinician should supply the microbiologist with pertinent historical and other data regarding the patient's illness; the microbiologist should be able to utilize this information.

The microbiologist can also be of service to the clinician by instructing him or her concerning the importance of proper specimen collection and transport and regarding specific techniques to be used. An interested physician can be taught how to judge the quality of a sputum specimen and how to interpret a Gram stain properly.

Microbiologists also have a responsibility to notify infection control personnel or epidemiologists and public health authorities. Urgent matters should be communicated by telephone.

The microbiologist also has an obligation to the patient aside from considerations of rapid, accurate diagnosis. Professionalism demands that microbiologists, as well as other members of the health team, not hesitate to deal with certain specimens or patients and that they respect the patient's rights and feelings with regard to sensitive information. At times the microbiologist may be pressured to release

information on a patient to the patient, his or her family, or others; such pressure must be resisted. Demands of this nature should be referred to the physician.

1.3. Specimen Collection and Transport

Specimen collection and transportation are urgent considerations because the quality of a laboratory's work may be limited by the nature of a specimen and its condition on arrival in the laboratory. Chapter 6 includes a detailed discussion of correct collection and transportation methods.

Specimens should be obtained so as to preclude or minimize the possibility of introducing extraneous microorganisms that are not involved in the infectious process. The problem is greatest when the specimen may become contaminated with elements of the normal (or colonizing) flora that may serve as pathogens in the entity being studied (e.g., Klebsiella as an oral cavity colonizer in a patient with pneumonia). Even normally sterile body fluids may be contaminated during specimen collection in such a way as to cause significant difficulty in interpretation. For example, a patient suspected of having bacterial endocarditis, although he does not really have the disease, has one or more blood cultures that have become contaminated with coagulase-negative staphylococci during venipuncture. Since this organism can cause endocarditis in numerous clinical settings, this patient's contaminated cultures may lead to additional testing, administration of unnecessary antibiotics, and a prolonged hospital stay for the patient. Careful skin preparation before procedures such as blood cultures and spinal taps and techniques for bypassing areas of normal flora when this is important and feasible (e.g., percutaneous transtracheal aspiration in critically ill patients with pneumonia) will prevent many problems. Personnel must be instructed to label all specimens submitted to the laboratory with data such as the patient's name, hospital number, date and time, and exact nature and source of the specimen, as discussed in Chapter 3.

1.4. Noncultural Methods of Diagnosis

As noted previously, the physician depends on data from many sources in arriving at tentative and definitive diagnoses. In addition to the usual medical history and physical examination, the competent clinician explores epidemiologic information (travel history; exposure to animals, ticks, or other vectors;