

## Book reviews

**Molecular Methods for Virus Detection.** Wiedbrauk D L, Farkas D H, eds. (£46.00.) Academic Press. 1995. ISBN 012 748920 7.

**PCR: Protocols for Diagnosis of Human and Animal Viruses.** Becker Y, Darai G, eds. (DM168.00.) Springer Verlag. 1995. ISBN 354 058899 X.

*Molecular methods for virus detection* has an all American cast and only an occasional reference from beyond those shores. One of the interesting features is the author list and the numbers of individuals who manage to undertake their research and development of molecular methods within private and profit making institutions. It is this expansion of molecular methods that has fuelled the biotechnology industry and now provides a powerful diagnostic arm for virologists.

A number of us who have become partly conversant with enzyme and DNA technology look for reliable recipe books for the laboratory. Recent requirements in the United Kingdom for laboratory accreditation have encouraged formal recording of results in the virology laboratory. Indeed, many techniques that have been passed from senior to junior technicians over several generations is formalised as methodology on every bench. This book supplies basic and well described formal methods and those who wish to be more creative may mix and develop from these protocols. There is enough insight into the techniques both to stimulate interest in the underlying mechanisms involved and to meet one's own requirements.

The book covers all basic nucleic acid detection techniques in an interesting first chapter. The second chapter addresses the problems of quality assurance: which is a constant worry in a diagnostic laboratory and one that enters a new dimension with the exquisitely sensitive amplification techniques that we have to hand. There are a number of overviews within the chapters which are especially valuable for those of us who move from laboratory to clinic and back again. These demonstrate the value of frequently revising and updating our techniques.

One chapter describing the technique proposed for assessing antiviral susceptibility in a wild virus isolate was of particular interest. The "prophets of doom" tell us that we will see an escalation in antiviral drug resistance with the increasing use of virus directed pharmaceuticals. Many of us may see this in the (at least) hundreds of distraught young people who will be afflicted with drug resistant genital herpes. They may no longer achieve remission and relief from the excellent antiherpes drugs currently available. The technique proposed in this chapter marries a home-grown and well established plaque assay with a commercial kit assaying antiviral susceptibility with a DNA/DNA hybridisation. I am slightly worried about the use of a radioactive probe in the technique recommended: surely we may move on from this marker? While the method proposed for a commercial kit evaluation of viral sensitivity (they supply even the cells in the United States) is interesting, comments in the proce-

dural notes leave me uneasy: "If the virus inoculum is too weak (and) too much time is required to achieve the appropriate CPE (then) the antiviral agent may overwhelm the isolate, giving a false impression of the effectiveness of the drug." The reader should consider this carefully. The logic used is a bit shaky. Do we have a resistant isolate or are we selecting even as we culture in vitro?

In the chapter on in situ PCR there are good instructions for a technique which we are finding increasingly useful, especially for unexplained pathology where tissue is available. The demonstration of viral nucleic acid within a cell is support for a theory of viral pathology. But each case should be interpreted with caution. We carry many viral passengers, sometimes for life, and others for a short time. One PCR followed by a hybridisation in situ does not confirm a cause and effect diagnosis; it merely tips the balance of evidence. I have to admit a certain worry about the descriptions under the black and white photomicrographs of autoradiographs within the chapter which look less selective and convincing than the captions would have us believe.

It is a book to stimulate interest and, I suspect, will be a primer for laboratory staff at all levels. I would not recommend it as a sole book on molecular techniques but as one of several good reference guides and a constantly available reminder about the underlying mechanisms that we are addressing. Making things work in the laboratory is one pleasure; developing mechanisms and teasing away at the methods for refinement and the creation of new techniques is another.

*PCR: protocols for diagnosis of human and animal viruses* is a manual which enters a different dimension from the previous book. It comes with its own MS-DOS disk (although my fear of virus infection has prevented me loading it until my Dr Solomon's virus checker is updated). Part 1, as the preface tells us, contains 49 useful protocols in 13 sections which deal with the diagnosis and typing of disease causing viruses in humans. The 12 chapters of part II deal with viruses from animals which cause disease. Spumaviruses are not the everyday story of (even) country folk and their role in the aetiology of human disease is not, to my knowledge, certain, although a number of aetiological associations have been suggested; these include Graves' disease, kidney disease, and encephalopathy and myasthenia gravis. This is a very well researched, fascinating, and useful book for all of us who seek to use and develop modern technology and science for diagnostic purposes. However, I detect that there has been no close liaison between the scientist and the clinician in formulating this published material. The physician quickly becomes sceptical when the scientist is seen to sell his expertise but lacks the particular diagnostic perspective that years of grinding through outpatient clinic provides. Nonetheless, it provides a stimulus. It contains a description of applying PCR to poorly diagnosed but common illnesses such as influenza, respiratory syncytial virus, and even Rhinoviruses. Whether a diagnosis of *Molluscum contagiosum* by PCR is ever necessary seems doubtful as it is clinically obvious in its effects. Furthermore, the sequencing for smallpox and of primers for PCR will I hope always be an academic exercise. All the chapters on animal viruses are interesting, although I wonder whether the sales team who allowed this juxtaposition thought the world

to be filled with virologists with such refreshingly broad interests? I wonder for whom this book is written. Smallpox (chapter 33) will, we hope, remain historical but hanta virus and the pulmonary syndrome in the USA (chapter 45) was an exercise in diagnosis and cooperation with good laboratory techniques, sound epidemiology, and inquisitive physicians from which we all may learn. I will want to own this book for the insight that all this excellent information provides.

COLIN G FINK

**Molecular Diagnostics for the Clinical Laboratorian.** Coleman W B, Tsongalis G J, eds. (Pp 390; £60.00.) Humana Press. 1996. ISBN 089 603373 2.

This multi-author text consists of four sections that: introduce molecular biological principles; describe the basic molecular techniques applicable to laboratory medicine; discuss application of these techniques to clinical problems; and assess the routine laboratory implementation of such technology. One useful aspect of this structure is the inclusion of a wide range of techniques, together with the principles of molecular biology, in one volume specifically designed for clinical molecular diagnostics. However, it is inconsistent in places, with some chapters representing overviews of technical principles and others including practical protocols. Moreover, the protocols that are included are incomplete, assuming some degree of knowledge of the procedures involved. Although to some extent sacrificing depth for breadth, the chapters addressing clinical application of the techniques described are useful as they are generally restricted to abnormalities of clinical relevance.

Given the assumption that this book was not intended as a practical manual, and in view of the editors' stated wish particularly to address the interpretation and limitations of molecular diagnostic results, this text gives a good theoretical background for those with little experience in this field. It is readable, comprehensive, and relatively up-to-date with some references from 1996. However, those directly involved in molecular diagnostics are likely to require a supplementary practical manual.

C S HERRINGTON

**Molecular Biology of Cancer.** Macdonald F, Ford C H F. (Pp 218; £19.95.) BIOS Scientific Publishers Ltd. 1996. ISBN 185 996225 4.

This book is an overview of the molecular events involved in the process of tumour development. As such, it is competing in a market awash with similar texts designed to make some sense of the morass of information available to those interested or involved in this field. In this regard, the book succeeds, largely as a result of the way in which it is structured. The initial section, dealing with general principles, gives a distillation of the basic information required to understand the second section, which comprises specific chapters dealing with oncogenes, tumour suppressor genes, and the cell cycle control molecules and mismatch repair genes. However, it is not essential to read and assimilate all of this information to be able to understand the chapters in the third and fourth sections that deal with specific tumour types and possible clinical molecular strategies. Thus, it is possible to take from the



## Molecular Methods for Virus Detection

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