Notes

XXIV International Congress of Ophthalmology

The Glaucoma Society will meet in association with the International Society for the Prevention of Blindness. The meeting is open to ophthalmologists with a special interest in glaucoma. Deadline for free paper abstracts is 1 November 1981. Details from John Hetherington Jr, MD, University of California Medical Center, 400 Parnassus Avenue, Room A-775, San Francisco, California, 94143, USA.

Contact lens technology

The Cullen Eye Institute, Baylor College of Medicine, will hold its annual course in contact lens technology on 3-5 December 1981. Details from Joseph W. Soper, Program Chairman. Contact Lens Technology Course. Department of Ophthalmology, Baylor College of Medicine, Houston, TX 77030, USA.

Research prize

The first Ludwig von Sallmann prize will be awarded at the Vth International Congress of Eye Research to be held on 3-8 October 1982 at the Koninghoff Conference Centre, Eindhoven, The Netherlands, to an individual for his outstanding achievement in ophthalmology and vision. Further information from Peter Gouras, MD, Chairman of the von Sallmann Prize Committee, Columbia University/College of Physicians and Surgeons, 630 West 168th Street, New York, New York 10032, USA.

Genetics and paediatrics

At the occasion of the International Congress of Ophthalmology in San Francisco (November 1982) there will be a combined meeting of the International Society of Ophthalmic Genetics and of the International Society of Paediatric Ophthalmology. The main topics will be functional examinations in children, malformation syndromes, the hereditary and paediatric diseases of the cornea, and the hereditary and paediatric diseases of the lens. Those who want to present a paper should write to Dr Irene Maumenee, Department of Ophthalmology, Wilmer Institute, Johns Hopkins Hospital, 601 N. Broadway, Baltimore, Md 21205, USA, for genetic papers, and to Professor M. Maione, Istituto di Oftalmologia, Via A. Gramsci 14, I-43100 Parma, Italy, for paediatric papers. Those who want to become a member of the International Society of Ophthalmic Genetics should write to Dr E. Cotlier, Yale University, Department of Ophthalmology and Visual Sciences, 310 Cedar Street, PO Box 3333, New Haven, Conn 06510, USA, and those who want to become a member of the International Society of Paediatric Ophthalmology should write to Professor M. Maione at the address given above.

Book reviews


This small booklet consists of guidelines arising out of a meeting held in San Francisco in 1979 to consider methods for assessing the magnitude and nature of avoidable blindness, particularly in those areas of the world where the major causes, such as trachoma, xerophthalmia, onchocerciasis, and untreated cataracts are particularly prevalent. The basic principles which guide the design and the conduct and interpretation of such surveys are discussed. There seems little doubt that if the criteria set out were adequately heeded in the preparation of surveys the value of the resulting material would be enhanced. The point is clearly made that, even though avoidable blindness is widely recognised in both nature and extent in many parts of the world, there are still countries where health authorities may not necessarily be aware of the presence of a serious problem within their own jurisdiction.

M. J. GILKES


This new textbook of ophthalmology from the Boston School is full of up-to-date information about clinical ophthalmology. In 346 closely packed pages the 10 authors have distilled most of the essential facts relevant to the subject. All through the emphasis is on the practical aspects — 'face to face with the patient.' There are 12 chapters, each dealing with the common ocular diseases and many of the rare ones in a systematic fashion. The text is accurate and condensed and includes brief sections on fluorescein angiography, electrodiagnosis, and ultrasound.

The practical advice on treatment is sound, whether it be on cataract, glaucoma, or squint, and there is a welcome emphasis on early operation in congenital cataract. No attempt is made to describe surgical operations in detail. Types of intraocular lenses are briefly mentioned, but in the choice of anterior chamber implants it is the corneal diameter and not the curvature that must be measured. In the section on the treatment of congenital glaucoma the corneal diameter should be included as part of the assessment of the care, and in the treatment of lime burns no mention is made of sodium versenate irrigation and prevention of symblepharon by contact lens or amnion.

These points, however, are relatively minor, and overall the impression is of a detailed practical textbook full of useful clinical information which should be of great value to anyone training in ophthalmology.

C. A. BROWN


This is an excellent and extensive account of all the factors associated with nutrition which can cause ocular pathology. Detailed accounts correlate the effects of deficiencies of animal food factors with similar deficiencies in man. The book is thus of equal value to research workers and

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Clinicians. An extensive bibliography at the end of each chapter is also of value to those who may wish to pursue a particular aspect of the subject in greater depth. In addition the author’s helpful comments in areas of dispute or deficient knowledge greatly add to the value of the book.

R. F. FISHER


This book is one of a series of monographs in neonatology, and its author has played a major role in achieving an understanding of the nature of retrolental fibroplasia—the retinopathy came to light in Boston 1 year previously without any true understanding of its aetiology. Furthermore the author has woven into the text personal contributions from 26 people (such as Norman Ashton, Paul Chandler, Everett Kinsey, and Arnall Patz) who in different ways played vital roles in the unravelling of the mysteries of this devastating eye condition which seemed to emerge with unusual haste.

The book is regarded by the author as a ‘modern parable’ which he has felt compelled to write because of his belief ‘that the unpleasant memory of the most dramatic epidemic of infantile blindness in recorded history was being repressed in the collective consciousness of medicine because it is too painful to recall.’

The association of the condition with prematurity was established by T. L. Terry in Boston in 1944, and before his death in 1946 he collected 117 cases. He also established the important fact that the condition in certain cases developed some time after birth, which was in conflict with the view which persisted until 1948 that it represented an inherent or acquired abnormality caused by factors which operated before birth or at the latest, immediately thereafter. In 1948, however, William and Ella Owens at the Johns Hopkins Hospital observed in a certain number of premature infants the sequence of changes in the retinal blood vessels and in the retina as a whole during the development of retrolental fibroplasia beginning some weeks after birth.

It is natural that the emergence of a new condition of such devastating proportions from a visual point of view attracted the attention of many clinical and experimental observers, and within a relatively short time it became established that retrolental fibroplasia is induced initially by an oxygen-rich atmosphere which causes a withering of the outgrowing peripheral retinal vessels in a state of prematurity. Subsequently there is a widely disorganised regrowth of these abnormal vessels on removal from the excessive oxygen leading to the formation of grey-coloured masses which represent areas of detached retina. Finally a grey membrane develops which represents the totally detached retina covered by many enlarged blood vessels.

It might be expected that the rapid recognition of the true nature of retrolental fibroplasia would have led to a disappearance of the condition by a careful control of the administration of oxygen to the premature child, but this has not been achieved to an accepted extent. However, this is natural because of the inevitable conflict between the paediatrician who has the task of maintaining life in the premature child and the ophthalmic surgeon who is aware of the risks of permanent blindness. This book discusses in detail the underlying problem, and it should be studied by all doctors who are concerned in the management of the premature infant.

Kenneth C. Wybar


This soft-back book is a review of the current state of knowledge about the retinal periphery and is a comprehensive review of that area. The first section deals with the anatomy, embryology, and physiological methods of examination and normal aspects of the retinal periphery. The book then goes on to describe pathological conditions in considerable detail.

My overall impression is that it certainly is of considerable use to the ophthalmologist in familiarising himself with many aspects of this important area of the eye. The book is a whole represents in the main a summary of our current state of knowledge and is profusely illustrated. Some of these illustrations are of somewhat inferior quality and are rather too small to be of much help. This particularly applies to the photographs. The book generously includes a large number of colour photographs, many of which are of excellent quality, but again some are not particularly satisfactory. There is an extensive bibliography from which the reader can easily refer to further work in the main subjects.

The nature of the book means that there has been a certain amount of compression of information and this does at times give some apparent imbalance in the contents. However, in general it is to be recommended, though as it is in French many English speaking ophthalmologists will find it of rather limited use.

A. H. Chignell


The first edition of this handbook was reviewed by us in 1972. At that time we expressed the view that there was a need for a publication of this nature, and the fact that it has gone into a second edition only 8 years later shows that by and large it has fulfilled this need. The second edition is an improvement on the first, and most of the more recent developments in ophthalmology find a mention in its pages.

There is a useful section on rural ophthalmology contributed by Dr W. R. Burkitt and a valuable appendix on the local production of eye drops. The index is comprehensive and accurate, and one of the few minor criticisms that can be made is that the section on the ocular complications of leprosy could usefully be expanded.

The line drawings, particularly those in which red has been used to bring out a special feature are very helpful but the black-and-white plates, although probably excellent in the original, are something of a disappointment. Although written with an understandable emphasis towards East African ophthalmology, there is no doubt that this excellent and very reasonably priced handbook will command a wide audience in developing countries wherever they may be.

D. P. Choyce