

New progress in research on retinal diseases

The glaucomas are a group of optic neuropathies characterized by progressive degeneration of retinal ganglion cells. Glaucoma affects more than 70 million people worldwide, making it the leading cause of irreversible blindness in the world. Glaucoma can remain asymptomatic until it is severe, resulting in a high likelihood that the number of affected individuals is much higher than the number known to have it. Glaucomas can be classified into 2 broad categories: open-angle glaucoma and angle-closure glaucoma. Both open-angle and angle-closure glaucoma can be primary diseases. Secondary glaucoma can result from trauma, certain medications such as corticosteroids, inflammation, tumor, or conditions such as pigment dispersion or pseudo-exfoliation. Currently available treatments cannot reverse glaucomatous damage to the visual system; however, early diagnosis and treatment can prevent progression of the disease. Recently, multiple studies have reported on glaucoma, which may provide assistance for the prevention and treatment of this disease.

1. Large-scale multitrait genome-wide association analyses identify hundreds of glaucoma risk loci

Glaucoma is a highly heritable human disease. Since much of glaucoma heritability remains unexplained, Xikun Han, Statistical Genetics Lab, QIMR Berghofer Medical Research Institute, Australia, and his team conducted a large-scale multitrait genome-wide association study in participants of European ancestry combining primary open-angle glaucoma (POAG) and its two associated traits to substantially improve genetic discovery power^[1]. They further increased our power by then employing a multiancestry approach, which increased the number of independent risk loci to 312. Leveraging multiomics datasets, they identified many potential druggable genes (table 1), including neuro-protection targets likely to act via the optic nerve, a key advance for glaucoma because all existing drugs only target intraocular pressure. They also identified novel associations of POAG with other complex traits, including numerous immune-related diseases. These findings provide insights into the pathogenesis of POAG and enable new drug development for this common cause of blindness.

Progress In Retinal Research

Shasha Hu



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Chalupa, 2013-10-25 A comprehensive review of contemporary research in the vision sciences reflecting the rapid advances of recent years Visual science is the model system for neuroscience its findings relevant to all other areas This essential reference to contemporary visual neuroscience covers the extraordinary range of the field today from molecules and cell assemblies to systems and therapies It provides a state of the art companion to the earlier book The Visual Neurosciences MIT Press 2003 This volume covers the dramatic advances made in the last decade offering new topics new authors and new chapters The New Visual Neurosciences assembles groundbreaking research written by international authorities Many of the 112 chapters treat seminal topics not included in the earlier book These new topics include retinal feature detection cortical connectomics new approaches to mid level vision and spatiotemporal perception the latest understanding of how multimodal integration contributes to visual perception new theoretical work on the role of neural oscillations in information processing and new molecular and genetic techniques for understanding visual system development An entirely new section covers invertebrate vision reflecting the importance of this research in understanding fundamental principles of visual processing Another new section treats translational visual neuroscience covering recent progress in novel treatment modalities for optic nerve disorders macular degeneration and retinal cell replacement The New Visual Neurosciences is an indispensable reference for students teachers researchers clinicians and anyone interested in contemporary neuroscience Associate Editors Marie Burns Joy Geng Mark Goldman James Handa Andrew Ishida George R Mangun Kimberley McAllister Bruno Olshausen Gregg Recanzone Mandyam Srinivasan W Martin Usrey Michael Webster David Whitney Sections Retinal Mechanisms and Processes Organization of Visual Pathways Subcortical Processing Processing in Primary Visual Cortex Brightness and Color Pattern Surface and Shape Objects and Scenes Time Motion and Depth Eye Movements Cortical Mechanisms of Attention Cognition and Multimodal Integration Invertebrate Vision Theoretical Perspectives Molecular and Developmental Processes Translational Visual Neuroscience Progress in Retinal Research editor. Osborne (Neville), editor. Chader (Gerald), 1983

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Neurobiology of the Inner Retina Reto Weiler, Neville N. Osborne, 2013-06-29 The relatively simple stratified nature of the retina and its specialized use in the visual process has long made it an inviting tissue to study both for its own sake and as a model for the more complex processes of the brain. For these dual purposes, the retina can be thought of as basically consisting of two functional parts. First, the outer retina comprised of the photoreceptor cells and attendant pigment epithelium serves to capture the photic energy and convert it into a neurochemical response. Second, the inner layers of the retina, mainly bipolar, amacrine, and ganglion cells and their attendant Müller cells, function more clearly as a typical part of the CNS, transmitting the photic signals to the brain. Between the 8th and 12th of August 1988, more than seventy scientists from all over the world gathered in Oldenburg, Federal Republic of Germany, for a meeting. The neurobiology of the inner

retina which was devoted entirely to the neural mechanism of the inner synaptic layer of the vertebrate retina. The meeting comprised twenty three separate lectures and four specially arranged discussion groups. In addition a number of posters were displayed and a period was allotted specifically for the discussion of these posters. The articles contained in this book will serve as a record of the papers delivered at the Oldenburg Meeting and illustrate the advances made in trying to understand the importance of the diversity of amacrine cell morphology and physiology in retinal function. **Veterinary**

Ophthalmology Kirk N. Gelatt, Gil Ben-Shlomo, Brian C. Gilger, Diane V. H. Hendrix, Thomas J. Kern, Caryn E.

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