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we will summarize the different classifications of rgc subtypes and will recapitulate the specification of some of them and describe how a genetic disease such as albinism affects neurogenesis resulting in profound visual deficits for anyone who wants to better understand glaucoma studying the retinal ganglion cells rgcs is very compelling it is known that vision loss in glaucoma results from damage to the rgcs and their axons but there is much more to learn about these important cells retinal ganglion cells rgcs convey visual information from the retina to the brain how many types of rgc exist and how they should be classified have been long standing questions electron microscopy reconstructions of retinal circuits suggest the possibility that a small proportion of midget ganglion cells might have blue off yellow on receptive fields in addition to colour discrimination midget rgcs also subserve pattern texture and stereoscopic depth perception we developed a rapid protocol for directly induced rgc irgc differentiation from human stem cells leveraging overexpression of ngn2 neuronal morphology and neurite growth were observed within 1 week of induction characteristic rgc specific gene expression confirmed identity the purpose of this study was to identify a robust representative region of interest roi for studies of retinal ganglion cell rgc soma loss in feline congenital glaucoma fcg a spontaneous large eyed glaucoma model transcriptomic and genome wide epigenetic mapping and single cell analysis confirmed rgc to astrocyte differentiation obviating neurogenesis and the gliogenic switch detailed molecular and cellular characterization experiments uncovered new mechanisms and markers for human rgcs and astrocytes retinal ganglion cells rgcs are the projection neurons of the retina and transmit visual information to postsynaptic targets in the brain while this function is shared among nearly all rgcs this class of cell is remarkably diverse comprised of multiple subtypes correlative light and electron microscopy showed that stress induced decrease in mitochondrial area length and cristae number was reversible intracellular ca² was elevated during this stage of reversible mitochondrial injury but there was no sign of mitochondrial cytochrome c release the present study investigated the morphological changes in several identified rgcs in the retinal degeneration rd1 mouse model of retinitis pigmentosa rp using a combination of viral transfection microinjection of neurobiotin and confocal microscopy the effect on retinal development and rgc differentiation was assessed by confocal microscopy of transgenic or immunolabeled embryos our results show that retinal neuroepithelial cells have an apically localized primary cilium usually protruding from the apical membrane transmission electron microscopy tem revealed alterations in mitochondrial morphology including increased membrane density and reduced mitochondrial cristae in rgcs after onc notably the ferroptosis inhibitor ferrostatin 1 fer¹ significantly promoted rgc survival

and preserved retinal function in onc and microbead induced glaucoma mouse in this chapter we review the main techniques for detecting the total population of rgcs and various of their subtypes in whole mounted retinas of pigmented and albino rats and mice four of the animal strains most studied by the scientific community in the retina field the degeneration of retinal ganglion cells rgcs leads to irreversible vision loss in a variety of pathological states here we describe a protocol to evaluate the role of a gene in protecting mouse rgcs when they sustain injuries from excitotoxicity or axonal damage we suggest that midget rgcs are not primate innovations but are descendants of evolutionarily ancient types that decreased in size and increased in number as primates evolved thereby we validated in vivo vis octf measures using both confocal microscopy of the immunostained flat mounted retina and numerical simulations vis octf for monitoring rgc axon bundle organization has the potential to bring new insight into rgc damage in optic neuropathies these new tools complement existing technologies like electron microscopy and dye filling techniques by allowing specific types of rgcs to be genetically labeled and manipulated this facilitates the search for morphological correlates of their physiological function the retinal ganglion cells rgcs are the main source of therapeutic targets for neuroprotective glaucoma treatment and evaluating rgcs is key for effective glaucoma care thus we developed a minimally invasive quick real time method to evaluate rgc death in mice background the chemokine cx3cl1 has been reported to play an important role in optic nerve protection but the underlying mechanism is still unclear cx3cr1 the only receptor of cx3cl1 is specifically expressed on retinal microglia whose activation plays a role in the pathological process of optic nerve injury this study aimed to evaluate whether cx3cl1 exerts optic neuroprotection by the macular area is important to the detection of glaucomatous retinal ganglion cell rgc damage macular thickness complementary to peripapillary retinal nerve fibre layer rnfl thickness can well reflect glaucomatous damage given that the macula contains more than 50 of the rgcs in a multilayer

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we will summarize the different classifications of rgc subtypes and will recapitulate the specification of some of them and describe how a genetic disease such as albinism affects neurogenesis resulting in profound visual deficits

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for anyone who wants to better understand glaucoma studying the retinal ganglion cells rgcs is very compelling it is known that vision loss in glaucoma results from damage to the rgcs and their axons but there is much more to learn about these important cells

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retinal ganglion cells rgcs convey visual information from the retina to the brain how many types of rgc exist and how they should be classified have been long standing questions

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electron microscopy reconstructions of retinal circuits suggest the possibility that a small proportion of midget ganglion cells might have blue off yellow on receptive fields in addition to colour discrimination midget rgcs also subserve pattern texture and stereoscopic depth perception

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transcriptomic and genome wide epigenetic mapping and single cell analysis confirmed rgc to astrocyte differentiation obviating neurogenesis and the gliogenic switch detailed molecular and cellular characterization experiments uncovered new mechanisms and markers for human rgcs and astrocytes

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correlative light and electron microscopy showed that stress induced decrease in mitochondrial area length and cristae number was reversible intracellular ca² was elevated during this stage of reversible mitochondrial injury but there was no sign of mitochondrial cytochrome c release

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in this chapter we review the main techniques for detecting the total population of rgcs and various of their subtypes in whole mounted retinas of pigmented and albino rats and mice four of the animal strains most studied by the scientific community in the retina field

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the degeneration of retinal ganglion cells rgcs leads to irreversible vision loss in a variety of pathological states here we describe a protocol to evaluate the role of a gene in protecting mouse rgcs when they sustain injuries from excitotoxicity or axonal damage

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we suggest that midget rgcs are not primate innovations but are descendants of evolutionarily ancient types that decreased in size and increased in number as primates evolved thereby

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we validated in vivo vis octf measures using both confocal microscopy of the immunostained flat mounted retina and numerical simulations vis octf for monitoring rgc axon bundle organization has the potential to bring new insight into rgc damage in optic neuropathies

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these new tools complement existing technologies like electron microscopy and dye filling techniques by allowing specific types of rgcs to be genetically labeled and manipulated this facilitates the search for morphological correlates of their physiological function

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the retinal ganglion cells rgcs are the main source of therapeutic targets for neuroprotective glaucoma treatment and evaluating rgcs is key for effective glaucoma care thus we developed a minimally invasive quick real time method to evaluate rgc death in mice

cx3cl1 cx3cr1 axis protects retinal ganglion cells by *Oct 26 2022*

background the chemokine cx3cl1 has been reported to play an important role in optic nerve protection but the underlying mechanism is still unclear cx3cr1 the only receptor of cx3cl1 is specifically expressed on retinal microglia whose activation plays a role in the pathological process of optic nerve injury this study aimed to evaluate whether cx3cl1 exerts optic neuroprotection by

macular imaging by optical coherence tomography in the *Sep 24 2022*

the macular area is important to the detection of glaucomatous retinal ganglion cell rgc damage macular thickness complementary to peripapillary retinal nerve fibre layer rnfl thickness can well reflect glaucomatous damage given that the macula contains more than 50 of the rgcs in a multilayer

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