Melanopsin-positive Intrinsically Photosensitive Retinal Ganglion Cells: From Form to Function

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Abstract

Melanopsin imparts an intrinsic photosensitivity to a subclass of retinal ganglion cells (ipRGCs). Generally thought of as irradiance detectors, ipRGCs target numerous brain regions involved in non-image forming vision. ipRGCs integrate their intrinsic, melanopsin-mediated light information with rod/cone signals relayed via synaptic connections to influence light dependent behaviors. Early observations indicated diversity amongst these cells and recently several specific subtypes have been identified. These subtypes differ in morphological and physiological form, controlling separate functions that range from biological rhythm via circadian photoentrainment, to protective behavioral responses including pupil constriction and light avoidance, and even image-forming vision. In this minisymposium review, we will discuss some recent findings that highlight the diversity in both form and function of these recently discovered atypical photoreceptors.

Introduction

Melanopsin-containing intrinsically photosensitive retinal ganglion cell (ipRGCs) are atypical retinal photoreceptors separate from classical rod and cone photoreceptors. The discovery of this inner retinal photoreceptor (Berson et al 2002, Hattar et al 2002, Provencio et al 1998) was motivated by attempts to explain circadian photoentrainment in otherwise blind individuals (Czeisler et al 1995) and mice lacking rods and cones (Freedman et al 1999, Lucas & Foster 1999). Circadian rhythms should be independent of light input if rods and cones were the only retinal photoreceptors, however, another, unidentified photoreceptor in these blind individuals caused alignment of their circadian rhythms to the onset and offset of light/dark cycles (photoentrainment). ipRGCs were first identified in 2002 as this third class of circadian photoreceptors, and are now known to be critical for...
mediation of not only circadian photoentrainment, but also the pupillary light reflex (PLR) and sleep (Altimus et al 2010, Berson et al 2002, Gooley et al 2001, Goz et al 2008, Guler et al 2008, Hatori et al 2008). Previously cloned in the late 1990’s (Provencio et al 2000), melanopsin soon thereafter was demonstrated to be the photopigment for ipRGCs that convey their intrinsic light response (reviewed in (Do & Yau 2010),(Berson et al 2002, Gooley et al 2001, Hattar et al 2002, Lucas et al 2003, Panda et al 2002, Ruby et al 2002). The confluence of modern molecular, transgenic, imaging and electrophysiological techniques has allowed for rapid generation of tools necessary for studying ipRGCs, and as such, this field has seen exponential growth in the decade since these cells were first discovered.

Initially identified as a single retinal cell type, morphological and physiological classifications showed that ipRGCs comprise a far more complex population than originally thought. Early studies described the dendrites of ipRGCs as covering the retina, forming a “photoreceptive net” (Provencio et al 2002), and gathering illuminance information from the entire retina. More recent characterization has shown that the cells identified by Provencio consist of two subclasses that form a mosaic covering the entire retina but that a third class of ipRGCs does not form a mosaic (Berson et al 2010, Schmidt & Kofuji 2010). In primates, ipRGCs are larger but display comparable morphologies to mice, with the additional feature that their dendrites have a minimal presence in the foveal pit, probably to avoid influencing high acuity vision (Berson et al 2010, Dacey et al 2005). Research has now suggested that the ipRGC family has up to 5 distinct cell types (called M1 to M5) with differential dendritic morphology and axonal projections, as well as some unexpected intra-retinal signaling (Barnard et al 2006, Hankins & Lucas 2002, Muller et al 2010, Sekaran et al 2003, Tu et al 2005, Viney et al 2007, Vugler et al 2007, Zhang et al 2008) (Figures 1 and 2).

Light integration occurs at the level of ipRGCs as they combine input from the highly sensitive and fast rod and cone phototransduction pathways with their own sluggish melanopsin-mediated response. This integrated information is then transmitted to numerous discrete brain regions involved in both non-image and image-forming vision (Brown et al 2010, Ecker et al 2010, Gooley et al 2003, Hattar et al 2006), including the suprachiasmatic nucleus (SCN) for circadian photoentrainment, the olivary pretectal nucleus (OPN) for the pupillary light reflex (PLR) in which bright light causes pupil constriction, and the dorsal lateral geniculate nucleus (dLGN) for image formation (Figure 3). ipRGCs project to many additional brain regions, suggesting several other as yet unidentified light-related functions.

The range of visual responses to which ipRGCs are known to contribute has also expanded consistently over the 10 years since their discovery. There is now convincing experimental evidence that ipRGCs ultimately modulate a multiplicity of behaviors including circadian photoentrainment, PLR, activity masking, sleep/arousal and neuroendocrine systems, anxiety, light aversion and even make a significant contribution to thalamo-cortical visual function (Brown et al 2010, Ecker et al 2010, Guler et al 2008, Hatori et al 2008, Johnson et al 2010, Thompson et al 2010, Tsai et al 2009). Furthermore, these novel functions of ipRGCs have applications in clinical conditions and diagnoses of retinal degeneration and sleep disturbances (reviewed in (La Morgia et al 2011), (Kankipati et al 2011, Kardon et al 2011, Park et al 2011)). This review will discuss recent advances in the field, describing the diversity of ipRGCs, their signaling properties, and previously unappreciated roles in image- and non-image forming vision.

**Morphological and Physiological Diversity Defines Classes of ipRGCs**

The depth at which ipRGC dendrites ramify in the inner plexiform layer (IPL) of the retina gives us the first clue as to their diversity. Dendrites can ramify near the ganglion cell layer,
in the ON-sublamina of the IPL or ramify in the outer IPL, or OFF-sublamina. ON-
stratifying RGCs receive excitatory input from ON bipolar cells, which respond to light
increments while OFF-stratifying RGCs receive excitatory input from OFF bipolar cells,
which respond to light decrements. Despite the initial view that ipRGCs consisted of a
uniform population of cells, early staining with a sensitive anti-melanopsin antibody
revealed two plexuses of melanopsin immunoreactive dendrites (Provencio et al 2002). To
examine the morphological properties of ipRGCs in detail, Schmidt et al. utilized a mouse
line expressing enhanced green fluorescent protein (eGFP) under the control of the
melanopsin (Opn4) promoter (Schmidt et al 2008). ipRGCs were identified under
epifluorescent illumination and subsequent morphological and electrophysiological analyses
were performed. These studies revealed three morphological subtypes: previously
characterized M1 cells with dendrites stratifying in the OFF sublamina of the IPL, M2 cells
with dendrites stratifying in the ON sublamina of the IPL, and M3 cells with surprising
variability in the proportion of dendrites ramifying in the ON and OFF sublaminas of the
IPL (Schmidt & Kofuji 2011, Schmidt et al 2008). These findings solidified earlier reports
of two melanopsin immunoreactive plexuses in the IPL as well as findings from retrograde
cellular labeling studies identifying three classes of ipRGCs (Provencio et al 2002, Viney et al
2007).

Morphologically, M2 cells had larger, more complex dendritic arbors and larger somas than
M1 cells (Schmidt & Kofuji 2009). Subsequent characterization of the rare M3 subtype
revealed morphological characteristics similar to M2 cells (Schmidt & Kofuji 2011). Research
from the Hattar lab has since identified the ON-stratifying M4 cells with the
largest somas and dendritic arbors of all ipRGC classes and ON-stratifying M5 cells with
small, bushy dendritic arbors (Ecker et al 2010).

When the functional properties of M1, M2 and M3 ipRGCs were examined, M1 cells were
found to have large, sensitive intrinsic light responses in comparison to M2 cells. M1 cells
were also found to have a higher input resistance, more depolarized resting membrane
potential, and to spike at lower frequencies than M2 cells (Schmidt & Kofuji 2009). Surprisingly, despite the variable morphology of M3 cells, this subtype had remarkably
homogeneous physiological characteristics that most closely resembled those of M2 cells,
arguing against M3 cells being a developmental anomaly or M1 and M2 “hybrid” somehow
left undifferentiated (Schmidt & Kofuji 2011).

ipRGCs, like conventional RGCs, receive rod and cone signals from the outer retina relayed
via bipolar and amacrine cells. Given differences in the location of dendritic stratification
and their intrinsic properties, one might predict that ipRGC subtypes receive very different
types of outer-retinal inputs. When the synaptic light response was examined in detail for
OFF-stratifying M1 and ON-stratifying M2 cells, it was found that both subtypes, perhaps
surprisingly, received a predominantly ON pathway input. Recordings from melanopsin null
animals (Opn4−/−), revealed that this ON pathway input is both larger and more influential
in shaping the light-evoked response of M2 cells compared to M1 cells which rely primarily
on melanopsin-mediated phototransduction (Schmidt & Kofuji 2010). Collectively, these
results indicate that the functional roles of the various ipRGC subtypes in behavior can be
deduced by the relative influence of melanopsin, rod and cone signaling on a given ipRGC-
mediated behavior.

**Unique Characteristics of ipRGCs as Light Sensors**

ipRGCs differ substantially from rods and cones in morphology and the molecular
mechanism of phototransduction. They also appear to signal in brighter light and over longer
durations than the rod and cone systems (Do & Yau 2010). Despite these overt
dissimilarities, common parameters govern photoreception by rods, cones, and ipRGCs including the nature of the response to each captured photon, the effectiveness of photon capture, and the threshold for generating a signal that is communicated downstream. A recent study has defined these features for ipRGCs, suggesting how these ganglion-cell photoreceptors complement rods and cones to drive the full range of mammalian vision (Do et al 2009).

All light responses represent the summed contributions of individual photopigment molecules with a linear range specific for each photopigment. Each photopigment is activated by absorption of one photon, producing the “single photon” or unitary response (Baylor et al 1979), analogous to the current flowing through a single ion channel. Because the unitary response of rods is large, relatively few rods each absorbing one photon are sufficient to elicit visual perception (Field et al 2005). Cones capture light with similar effectiveness to rods, but have a tiny unitary response and a correspondingly low sensitivity that allows them to function in daylight (Luo et al 2008). To define the unitary response of ipRGCs, electrophysiological recordings were made from cells expressing the fluorescent reporter, tdTomato under the melanopsin promoter (Do et al 2009). The smallest and brightest of these cells, which were targeted for study, were likely to be M1 cells (Do et al 2009, Ecker et al 2010, Schmidt & Kofuji 2009). ipRGCs were repeatedly given a flash of light so dim that, in the majority of trials, photons passed through the cell without consequence. On some trials, however, one melanopsin molecule was activated, producing a unitary response. The unitary response of ipRGCs appears larger than that known for any vertebrate sensory neuron, including rod photoreceptors (Do et al 2009). It is also extraordinarily prolonged, lasting nearly ten seconds, or about twenty-fold longer than rods and a hundred-fold than cones. This prolonged kinetics improves sensitivity by conferring high temporal summation, while also smoothing the response to fluctuating light levels. Thus, even at its most elementary level, the intrinsic photosensitivity of ipRGCs operates over long time scales.

Despite this large and prolonged single-photon response, ipRGCs are even less sensitive than cones (Berson et al 2002, Do et al 2009, Ecker et al 2010, Lucas et al 2003, Panda et al 2002). To determine if this low sensitivity originates from a low probability of photon capture, the melanopsin content of ipRGCs was estimated. Since melanopsin is localized to the plasma membrane (Belenky et al 2003), the number of melanopsin molecules per cell was divided by the cell’s surface area, revealing a density of ~3 molecules μm⁻² (Do et al 2009). By contrast, rods and cones express their photopigments at a density of ~25,000 molecules μm⁻², in specialized membranes that are stacked in the light path. The low melanopsin expression level may prevent interference with photon capture by rods and cones, which receive light after ipRGCs. In summary, the operation of ipRGCs in bright light is enabled by a low probability of photon capture and high amplification.

Unlike rods and cones, which signal within the retina with graded membrane voltages, ipRGCs signal to the brain using action potentials (spikes). Therefore, the number of unitary responses required to reach a spike threshold also determines their overall sensitivity. ipRGCs occasionally spike even in the absence of light, suggesting that they normally hover near spike threshold (reviewed by (Do & Yau 2010)). Because the single-photon response is so large and long-lasting, just one captured photon can increase the probability of crossing spike threshold, driving the cell to increase its firing rate (Do et al 2009). Thus, like rods, ipRGCs can signal single photon responses to the brain. The cost of this low threshold for spike generation is the noise contributed by spontaneous firing. The impact of this spontaneous noise and any effects on downstream pathways remains to be determined.
It is presently unknown whether all ipRGC subtypes have a large single photon response and low threshold for spike generation. The subtypes do appear to share phototransduction components downstream of melanopsin (reviewed in Berson 2007, Graham et al 2008, Perez-Leighton et al 2011), potentially allowing for similar single-photon responses, and at least M2 cells readily fire spikes (Schmidt 2009). Nevertheless, the M2 and M3 subtypes are less intrinsically photosensitive than M1 cells; M2 and M3 cells by ~10-fold and M4 cells by ~100-fold (Ecker et al 2010, Schmidt & Kofuji 2009, Schmidt & Kofuji 2011).

Melanopsin density, however, appears to be lower in the M2 cells and even lower in the M4 cells (Berson et al 2010, Ecker et al 2010, Schmidt & Kofuji 2009), consistent with their relative sensitivities to light (Ecker et al 2010). Thus, ipRGCs as a class may tune their sensitivities by their level of melanopsin expression.

In summary, the intrinsic photosensitivity of ipRGCs complements that of rods and cones by virtue of operating at high light intensities and over long durations. ipRGCs establish these features by pairing sparse photon capture with a highly integrating single-photon response. How the unique light responses of ipRGCs influence behavior will likely depend on the subtype of ipRGC activated and signal processing in the brain regions specifically innervated.

Specialization of Primate ipRGC-mediated Visual Function

The melanopsin-based retinal circuit appears fundamentally similar in mouse and primate, yet specializations that parallel the appearance of the unique primate fovea and trichromatic color vision suggest a more complex role in human visual processing. In primates, as in mice, the melanopsin photopigment shows a λmax of 483 nm and is expressed in two morphologically independent populations of ~3000 ganglion cells with large, sparsely branched dendritic trees that stratify at the inner and outer borders of the IPL (Dacey et al 2005). As in mouse, these ganglion cells combine input from rod and cone signaling pathways with the inherent melanopsin response and use these signals to generate an irradiance coding ON spiking response, and to drive the various components of the human and non-human primate PLR (Gamlin et al 2007).

A unique specialization of the human visual system however is the ultra-high cone and ganglion cell density of the fovea that serves peak visual acuity, color and form vision via the topographic projection to the lateral geniculate nucleus (LGN) and in turn to a greatly expanded primary visual cortex. Amazingly, the melanopsin-expressing ganglion cells of human and macaque monkey retina are present in the central retina and form a photoreceptive network of dendrites throughout the foveal specialization but with only a few processes crossing the foveal pit. These ganglion cells also project to the LGN and receive input from the spectrally distinct short (S), middle (M) and long (L) wavelength sensitive cone photoreceptor types unique to the trichromatic primate and show a rarely studied type of color sensitivity mediated by S cones (Dacey et al 2005). Irradiance detecting units have been recorded in both LGN and primary visual cortex and are implicated in the perception of surface luminance and brightness. The spectral tuning of such irradiance cells has not yet been measured. Recent evidence suggests, however, that humans lacking rods and cones can show a rudimentary conscious visual awareness (Zaidi et al 2007).

In addition to the LGN, the primate melanopsin-expressing ganglion cells project widely to the superior colliculus and the pretectum. An important question that remains to be clarified is whether, as in mouse, the M1 and M2 types in primate show distinct response properties, central projections and functional roles in the circadian, pupillary and perceptual pathways. Indeed, recent reports suggest that projections from some melanopsin cells to the dorsal
LGN in mice give rise to irradiance sensitivity in the thalamo-cortical pathway and could play a role in visual perception even in this species (Brown et al 2010, Ecker et al 2010).

The Role of ipRGCs in Vision

Convergent evidence from recent studies indicates that ipRGCs project to distinct brain regions involved in image-forming vision. In 2005, Dacey et al back labeled ipRGCs from the primate LGN (Dacey et al 2005), suggesting that the projection pattern of these photoreceptive ganglion cells extends to the visual thalamus. A discrepancy arose, however, when extensive analysis of the Opn4^{taulacZ} knock-in mouse, in which histological stains can be used to label both cell bodies and axons of melanopsin-expressing cells, implied that any such input to thalamo-cortical projection neurons in the dorsal LGN (dLGN) would be extremely limited (Hattar et al 2006). The subsequent discovery that the Opn4^{taulacZ} reporter selectively labels the M1 class of ipRGC raised renewed uncertainty regarding the central projection pattern of ipRGCs. This controversy was resolved using Cre-recombinase-based reporter mouse lines, which more fully labeled the entire ipRGC population, and revealed extensive innervation of the dLGN (Brown et al 2010, Ecker et al 2010), showing species conservation of ipRGC projections.

Recent experiments from the Lucas laboratory directly assessed the contribution of melanopsin photoreceptors to visual activity in the mouse dLGN (Brown et al 2010). Using a combination of extracellular recording electrodes and intrinsic optical imaging, widespread light-evoked activity was observed in both the dLGN and visual cortex of rodless/coneless mice. The thalamic light response in these mice recapitulated the low sensitivity and slow response kinetics reported for melanopsin phototransduction.

The contribution of melanopsin input to the mouse dLGN was not merely a feature of the retinally degenerate preparation. In mice with an intact visual system, 60s stimuli at relatively high intensity elicited a persistent increase in firing in a large number of neurons in the dLGN. Two lines of evidence suggest that, under these conditions, melanopsin contributes to this persistent firing. Firstly, it was recorded only in response to wavelengths within the melanopsin-sensitivity range. Secondly, it was much reduced in amplitude in melanopsin knockout mice. Further study of melanopsin knockout mice indicated that melanopsin functioned not only to sustain firing of these dLGN neurons under steady illumination, but was also critical in allowing this part of the brain to encode stimulus irradiance. Thus, in visually intact animals, the firing rate of many dLGN cells tracked irradiance over at least 6 orders of magnitude. This ability was substantially impaired in melanopsin knockout mice.

These data imply that melanopsin plays an important role in providing information about spatial brightness to the thalamo-cortical visual projection. Experiments in visually intact, retinally degenerate and melanopsin knockout mice all indicate that this signal appears in a large proportion of dLGN neurons, approximately 40% of all light-responsive units detected (Brown et al 2010). As all of those units also showed rod/cone-dependent transient responses to lights ‘on’ and ‘off’, the melanopsin-dependent signal of illuminance is likely super-imposed upon other visual codes at the level of the thalamus. Future work will be required to determine melanopsin’s significance for brightness perception and, perhaps, other aspects of vision.

M1 and Non-M1 Specification of Central Function

Previous studies have indicated that rods, cones and melanopsin all influence circadian photoentrainment (Lall et al 2010) and PLR (McDougal & Gamlin 2010) via ipRGCs (Guler et al 2008, Hatori et al 2008). Rods have an unparalleled sensitivity while the kinetics of

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cone opsin and melanopsin signaling suggests that cones mediate an initial fast response across a range of light levels and that the sustained response of melanopsin mediates a longer-lasting physiological response (Brown et al 2010). This simple model highlights complementary roles for each photoreceptive system in non-image forming behaviors but an additional layer of complexity arises if differential synaptic input occurs. Research from the Hattar lab indicates that circadian photoentrainment is preferentially dependent on rod photoreceptors with cones providing little input (Altimus et al 2010) but also see (Lall et al 2010). To the contrary, cones seem to provide strong input to the PLR (Lall et al 2010). These data provide the intriguing possibility that M1 ipRGCs projecting to the OPN are different than the M1 ipRGCs projecting to the SCN, hinting at the possibility of subpopulations within the M1 cells and adding another layer of complexity to this narrative.

To discriminate between OPN- and SCN-projecting M1 ipRGCs, identifying distinct markers for individual subtypes of ipRGCs was the critical next step. Functionally distinct subpopulations of M1 ipRGCs were molecularly defined by expression of the Brn3b transcription factor (Chen et al 2011) with Brn3b expressed in all non-M1 ipRGCs but only in a fraction of M1 cells. Coincidentally, Brn3b-positive M1 ipRGCs project to all known M1 targets except the SCN. Consistent with this innervation pattern, ablation of Brn3b-positive ipRGCs severely impairs the PLR, but does not affect circadian photoentrainment. These data support the classification of the M1 subtype into two distinct sub-populations and provide a potential mechanistic explanation for the differential influence from rods and cones on circadian photoentrainment and the PLR, although the electrophysiological profile of M1 cells suggests a singular classification. The behavioral responses of these two M1 subpopulations could be controlled by integration of signals at their axonal targets but could also be explained by differential rod versus cone input at their dendritic arbors. Cone-mediated input predominates control of the PLR, presumably via Brn3b-positive M1 cells, suggesting that Brn3b-positive M1 cells either receive more input via cone signaling pathways or that their axonal targets weight this input more heavily. This raises the intriguing question of the interaction between the molecular identity, dendritic connectivity and axonal targeting of ipRGCs. The transcription factor(s) responsible for specifying Brn3b-negative ipRGCs and differential targeting of M1 sub-populations to specific brain regions remain to be identified. Similar genetic strategies using cell-type specific tools will eventually uncover the circuitry and specialized functions of all ipRGC subtypes.

It is now well established that M1 ipRGCs are predominantly responsible for circadian photoentrainment and PLR (Guler et al 2008, Hatori et al 2008). The M1 subclass is the best characterized and exemplifies how its design subserves its role. With the highest melanopsin levels and a resting membrane potential sitting near threshold for firing, it is the most light sensitive of the characterized subclasses thereby contributing the most robust melanopsin component to their related behaviors. The non-M1 cells are likely to mediate behaviors that rely predominantly on rod and cone signaling, and less so on melanopsin photoreception. Thus understanding the relative influence of all three photoreceptive systems on ipRGC signaling is key to understanding how each subclass of ipRGC mediates specific behaviors. The non-M1 subclasses have only now begun to be studied for their functional role in behavior. These non-M1 ipRGCs may be responsible for image forming behaviors via projections to the dLGN (Ecker et al 2010) and projections to numerous other regions including the periaqueductual gray and amygdala, which suggests contributions to the pain, fear and anxiety circuitry (Hattar et al 2006, Maren & Fanselow 1996).

Photo-allodynia (photophobia) is a condition in which even low levels of illumination result in mild discomfort to extreme ocular pain. Photo-allodynia affects up to 80% of migraineurs (Choi et al 2009, Robbins & Lipton 2010), approximately 50% of patients with mild
traumatic brain injury (Craig et al. 2008), and also occurs from ocular inflammation or abrasion.

Similar to circadian photoentrainment, photo-allodynia is even experienced by patients who are visually blind (Amini et al. 2006). Furthermore, people with photo-allodynia are maximally sensitive to blue light and experience pain throughout the duration of light exposure, consistent with a role for ipRGCs in this condition (Amini et al. 2006, Newman et al. 2003, Noseda et al. 2010). To directly test this hypothesis, researchers used OPN4
dta
mice, in which all ipRGCs are ablated, in a light-aversion behavioral assay. Other groups had used naïve mice in a light/dark box behavioral test that compounds light aversion with light-dependent anxiety (Recobé et al. 2009, Semo et al. 2010, Thompson et al. 2010), an equally important consideration for photo-allodynia as pain and anxiety are inter-related. To assess light aversion separately from anxiety, mice were first habituated to a light/dark box chamber until their anxiety subsided. They were then tested for aversion to bright lights as an endophenotype for photo-allodynia. Matynia et al. showed that mice with intact ipRGCs avoided bright lights but mice lacking ipRGCs showed very little aversion to these bright lights, suggesting that ipRGCs are the primary circuit for light aversion and by extension, photo-allodynia (Matynia and Gorin, unpublished results).

Under normal conditions, ipRGCs control light aversion behavior, however, rod/cone photoreceptors may also provide input in pathological states of light sensitivity. Following administration of a sub-analgesic dose of morphine, mice normally show enhanced light aversive behavior that is μ opioid receptor-dependent (Matynia and Gorin, unpublished observations). To test if this behavior is also mediated by ipRGCs, researchers also gave sub-analgesic doses of morphine to OPN4
dta
mice lacking ipRGCs. Interestingly, these mice exhibit the same level of opiate-induced light aversion, indicating that morphine-induced light aversion is mediated by the classical rod/cone photoreceptors through conventional RGCs (Matynia and Gorin, unpublished observations). These data implicate two distinct circuits for light avoidance behavior under normal (ipRGC-dependent) and pathological (ipRGC-independent) conditions. Future studies will help determine the specific brain regions that receive ipRGC projections to mediate light aversion in normal and pathological models, helping to define the neural circuitry associated with photo-allodynia in people.

Concluding Remarks

The research discussed here demonstrates that understanding a complex neuronal system requires convergent molecular, physiological, genetic, circuitry and behavioral approaches. The ipRGC population consists of several subtypes, each with distinct morphological features including dendritic field and soma size, as well as level of dendritic stratification. Each subtype has unique responses to light and other electrophysiological characteristics. Furthermore, the subtypes are distinguished by their axonal projection patterns. These distinct features suggest possible specific functional roles for each subtype in non-image forming and image-forming behavior. However, given the lack of molecular markers for each ipRGC subtype, study of their individual roles in behavior is restricted. Molecular markers and development of additional genetic tools will be necessary to delineate their unique functions.

ipRGCs have transcended their earliest functional designations and their influence is now seen in image-forming vision, retinal development, migraine research, photo-allodynia, sleep disorders, anxiety and depression. Furthermore, ipRGCs are now also implicated in numerous visually guided behaviors, suggesting that ipRGCs could one day be a therapeutic target in degenerative retinal diseases. It has already been shown that ectopic melanopsin expression in retinal ganglion cells normally devoid of melanopsin improved vision in mice.
lacking classic photoreceptors (Lin et al 2008). This research has also made its mark in various clinical settings, as assessment of retinal degeneration via examination of the relative contribution of rods, cones and ipRGCs to pupil constriction is now being applied as a diagnostic tool (Kankipati et al 2011, Kardon et al 2011, Park et al 2011). Initially classified as a single cell type regulating a small number of behaviors, ipRGCs have greatly exceeded expectations in the last decade, repeatedly pushing back the boundaries of retinal dogma. Future research will undoubtedly continue to illuminate the diverse characteristics, from form to function, of these atypical ganglion cell photoreceptors.

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Figure 1.
ipRGC subtypes differ in morphology and site of dendritic fields
The Outer Nuclear Layer (ONL) of the mammalian retina consists of rod and cone photoreceptors, which have synaptic connections to their respective rod (R) or cone (C) bipolar cells in the Outer Plexiform Layer (OPL). The soma of bipolar, amacrine (DA, MA and II) and horizontal (not depicted) cells are localized to the Inner Nuclear Layer (INL). The INL and GCL (Ganglion Cell Layer) are separated by the Inner Plexiform Layer (IPL), another region dendritic-axonal interactions. The GCL contains retinal ganglion cells (not depicted) and the melanopsin-positive class of retinal ganglion cells, the ipRGCs. The classification of ipRGCs, (M1, M2, M3, M4 and M5) is based on morphology, and somatic and dendritic localization. Figure modified from (Do & Yau 2010).
Figure 2.
Morphology of M1, M2 and M3 ipRGCs and their physiological responses to light
Top panel: M1 (A), M2 (B) and M3 (C) ipRGCs are identified by their morphological characteristics and location of their dendritic arbors. M1 cells have dendritic ramifications in the OFF sublamina of the IPL whereas M2 cells ramify in the ON sublamina. M3 cells have dendrites bistratifying in both the ON and OFF sublaminas of the IPL. Bottom panel: M1 cells have the largest intrinsic, melanopsin-mediated light-evoked response, while M2 and M3 cells have smaller light-evoked responses. Scale = 10mV. Figure modified from (Schmidt & Kofuji 2011).
Figure 3.
ipRGC project to numerous central regions, and play a role in shaping behavior
Central projections of M1 cells include the OPN, controlling pupil constriction, and the
SCN, controlling circadian photoentrainment. Non-M1 cells, which currently include M2-
M5 cells, send projections to the LGN and are involved in a rudimentary, low-acuity visual
function. Additional sites of axonal terminations are located throughout the brain,
influencing activity, sleep/wake states, nociception, and areas implicating novel functions of
these cells.