

T-box Transcription Regulator *Tbr2* Is Essential for the Formation and Maintenance of Opn4/Melanopsin-Expressing Intrinsically Photosensitive Retinal Ganglion Cells

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Opsin 4 (Opn4)/melanopsin-expressing intrinsically photosensitive retinal ganglion cells (ipRGCs) play a major role in non-image-forming visual system. Although advances have been made in understanding their morphological features and functions, the molecular mechanisms that regulate their formation and survival remain unknown. Previously, we found that mouse T-box brain 2 (*Tbr2*) (also known as *Eomes*), a T-box-containing transcription factor, was expressed in a subset of newborn RGCs, suggesting that it is involved in the formation of specific RGC subtypes. In this *in vivo* study, we used complex mouse genetics, single-cell dye tracing, and behavioral analyses to determine whether *Tbr2* regulates ipRGC formation and survival. Our results show the following: (1) *Opn4* is expressed exclusively in *Tbr2*-positive RGCs; (2) no ipRGCs are detected when *Tbr2* is genetically ablated before RGC specification; and (3) most ipRGCs are eliminated when *Tbr2* is deleted in established ipRGCs. The few remaining ipRGCs display abnormal dendritic morphological features and functions. In addition, some *Tbr2*-expressing RGCs can activate *Opn4* expression on the loss of native ipRGCs, suggesting that *Tbr2*-expressing RGCs may serve as a reservoir of ipRGCs to regulate the number of ipRGCs and the expression levels of *Opn4*.

Key words: circadian; *Eomes*; ipRGC; retina development; RGC subtypes; *Tbr2*

Introduction

Retinal ganglion cells (RGCs) are the sole output neurons of the retina and convey distinct features of visual information to specific nuclei in the brain (Masland, 2012). In the mouse retina,

>20 types of RGCs have been described on the basis of their distinct morphological features (Rockhill et al., 2002; Coombs et al., 2007; Völgyi et al., 2009). Each RGC subtype is postulated to have a unique genetic makeup, and the establishment of diverse RGC subtypes is likely controlled by distinct combinatorial transcriptional cohorts. Nevertheless, the molecular mechanisms that lead to such diversity remain essentially unknown.

Among these diverse types of RGCs, opsin 4 (Opn4)/melanopsin-expressing intrinsically photosensitive RGCs (ipRGCs) are the only cells with comprehensively characterized morphological features, anatomical targets, and physiological functions (Provencio et al., 1998a,b; Hattar et al., 2006; Do et al., 2009; Schmidt and Kofuji, 2009; Berson et al., 2010; Ecker et al., 2010; Xue et al., 2011; Estevez et al., 2012; Hu et al., 2013). Their primary task is to control non-image-forming visual functions, including pupillary light reflex (PLR) and circadian photo-entrainment (Provencio et al., 2000; Lucas et al., 2001; Berson et al., 2002; Hattar et al., 2002; Panda et al., 2002; Provencio et al., 2002). Attempts to address the functions of the RGC transcriptional regulators POU class 4 homeobox 2 (*Pou4f2*; also known as *Brn3b*) and *Isl1* in ipRGCs prompted more questions regarding the molecular mechanism of ipRGC formation (Chen et al., 2011; Shi et al., 2013). *Pou4f2* and *Isl1* are activated in all RGCs during the early

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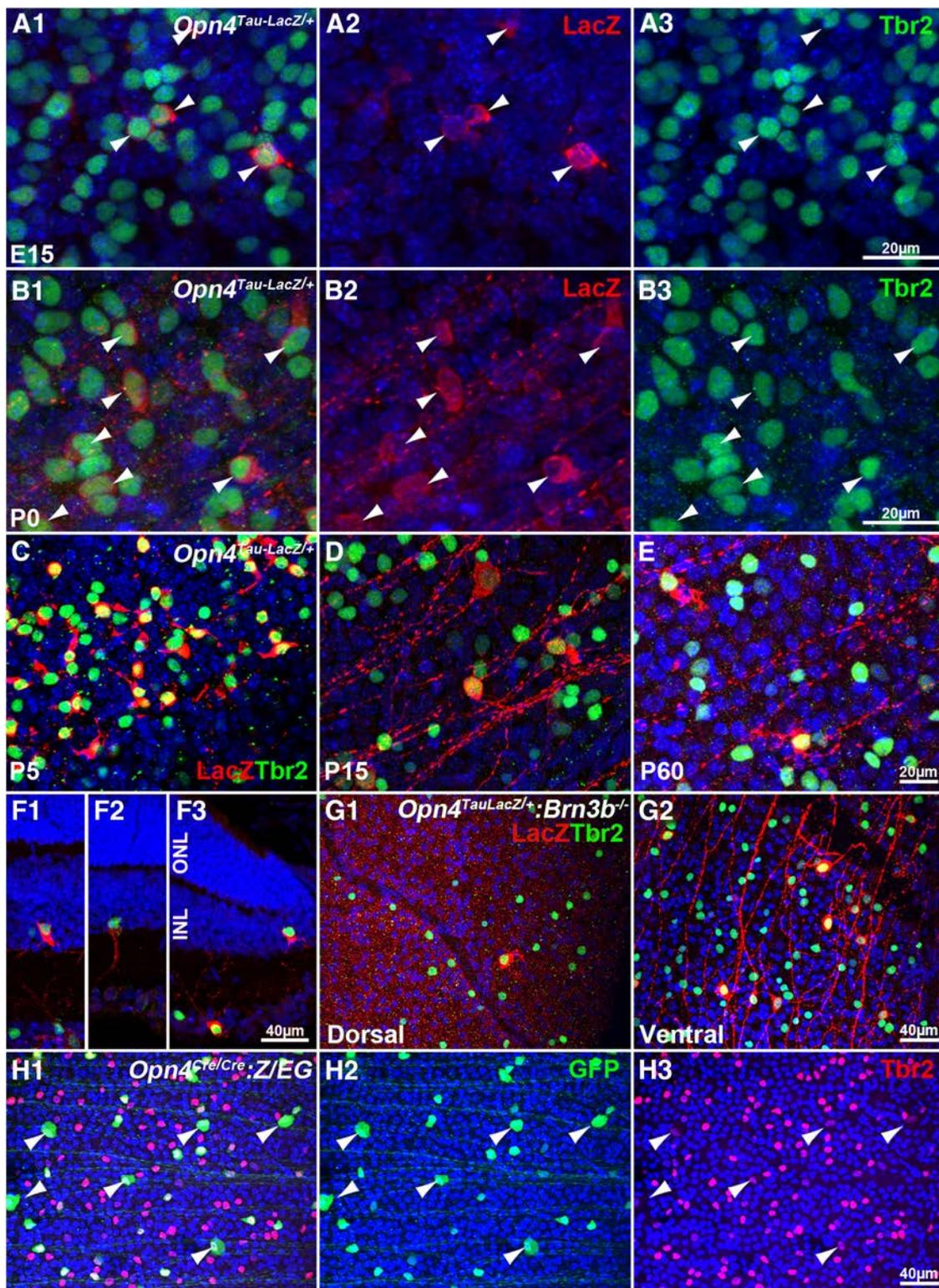


Figure 1. All ipRGCs express *Tbr2*. **A1–G2**, Confocal images of *Opn4^{Tau-LacZ/+}* retinas labeled with *Tbr2* (green) and *LacZ* (red). **A–E**, Flat-mounted *Opn4^{Tau-LacZ/+}* retinas at E15 (**A1–A3**), P0 (**B1–B3**), P5 (**C**), P15 (**D**), and P60 (**E**). **F1–F3**, Cryosections of *Opn4^{Tau-LacZ/+}* retinas at P30 that show rare displaced *LacZ*⁺ ipRGCs in the inner nuclear layer are also *Tbr2*⁺. **G**, Dorsal (**G1**) and ventral (**G2**) quadrants of flat-mounted *Opn4^{Tau-LacZ/+};Brn3b^{-/-}* retinas at P30 show more *Tbr2*⁺ cells in the ventral quadrant. **H1–H3**, Flat-mounted *Opn4^{Cre/Cre};Z/EG* retinas at P30, labeled with anti-*Tbr2* and anti-GFP antibodies, show confinement of GFP⁺ cells in the *Tbr2*⁺ cells. INL, Inner nuclear layer; ONL, outer nuclear layer. Scale bars are presented in the last image of each series. In all panels, blue is DAPI-labeled nuclei.

phase of RGC development [approximately embryonic day 12 (E12); Gan et al., 1999; Mu et al., 2008], which is much earlier than the onset of ipRGCs (approximately E15; McNeill et al., 2011). Throughout retinal development, they are not associated

with a specific RGC type (Gan et al., 1999; Mu et al., 2008). Genetically deleting either of them caused a drastic loss of total RGCs, including, but not specific to, ipRGCs (Gan et al., 1999; Lin et al., 2004; Mu et al., 2008; Shi et al., 2013), suggesting the

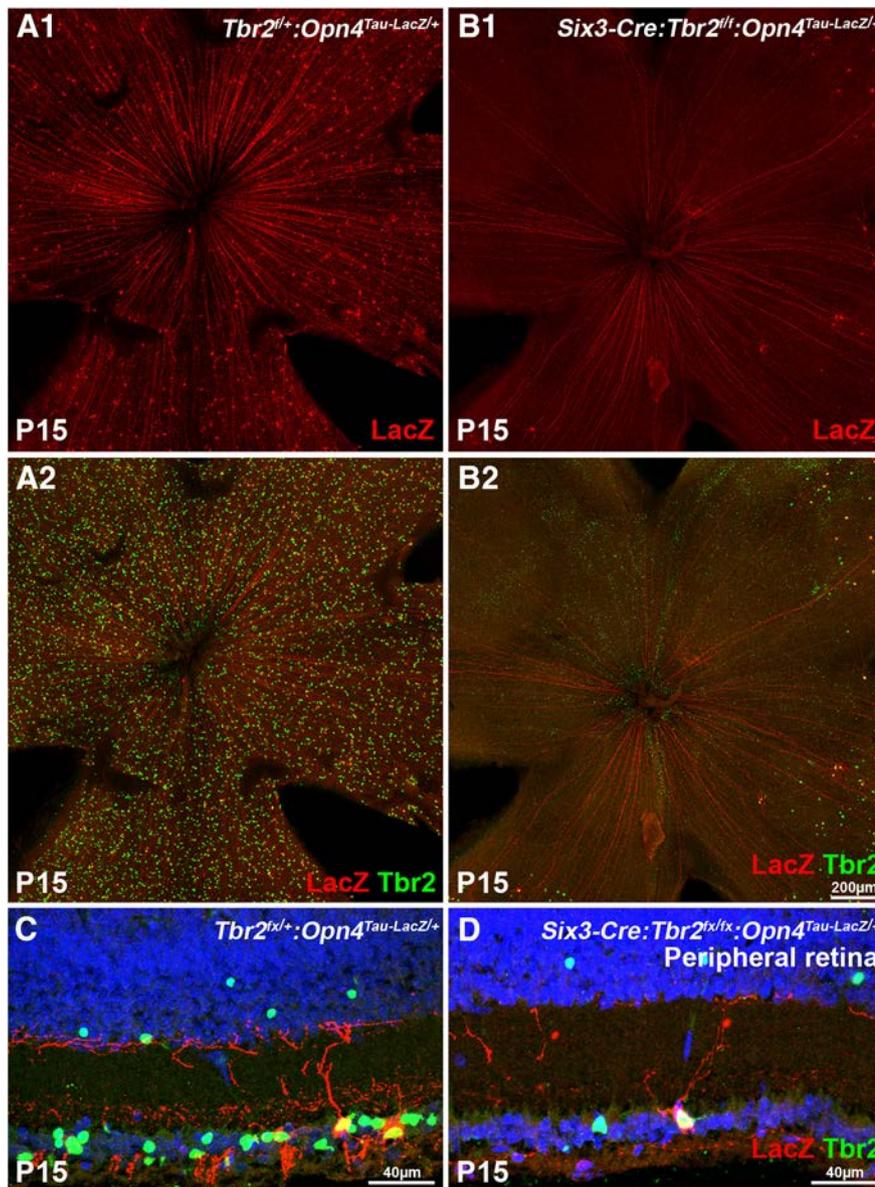


Figure 2. *Tbr2* is essential for the formation of ipRGCs. Confocal images of P15 control and *Tbr2*-deficient retinal flat mounts (**A**, **B**) and cryosections at the peripheral region (**C**, **D**) that show *Tbr2* and LacZ (reporting *Opn4*) labeling. **A1**, **A2**, **C**, The control *Tbr2*^{flax/+}:*Opn4*^{Tau-LacZ/+} retinas show normal distribution of *Tbr2*⁺ and LacZ⁺ cells. **A1** shows only the red channel from **A2**. **B1**, **B2**, **D**, The mutant *Six3-Cre:Tbr2*^{flax/flax}:*Opn4*^{Tau-LacZ/+} retinas show the absence of *Tbr2*⁺ and LacZ⁺ cells in the central retina. Only a few LacZ⁺ cells can be found in the periphery. All remaining LacZ⁺ cells are also positive for *Tbr2*, indicating that *Tbr2* is not removed by Cre in these cells. **B1** shows only the red channel from **B2**.

presence of other downstream factors that are more specific for ipRGC formation.

We observed previously that T-box brain 2 (*Tbr2*; also known as *Eomes*), a T-box-containing transcription factor, was expressed in a subset of newborn RGCs starting at E14.5, shortly preceding the onset of *Opn4* expression. Most importantly, *Tbr2* has been shown to be a common downstream factor of *Pou4f2* and *Isl1* (Mao et al., 2008; Mu et al., 2008). Single-cell transcriptome analysis also found that *Tbr2* was enriched in ipRGCs (Sieger et al., 2012). Here, we used the rich resource of genetically modified mice in *Opn4* and *Tbr2* loci to address the molecular mechanisms regulating ipRGC formation and sustaining their survival by *Tbr2*. Our results demonstrate the following: (1) no ipRGCs are detected when *Tbr2* is genetically ablated before RGC specification; (2) most ipRGCs are eliminated when *Tbr2* is de-

leted in established ipRGCs; and (3) additional ipRGCs are produced from *Tbr2*-expressing (*Tbr2*⁺) RGCs on the loss of native ipRGCs.

Materials and Methods

Animal models. The generation and genotyping of the mouse lines used in this study were described previously [*Tbr2*^{flax}, Mao et al., 2008; *SIX* homeobox 3 (*Six3*)–*Cre*, Furuta et al., 2000; *Opn4*^{Tau-LacZ}, Hattar et al., 2002; *Opn4*^{Cre}:*Z/EG*, Ecker et al., 2010; and *Brn3b*^{-/-}, Wang et al., 2000]. All mice were maintained in a mixed BL/6:129SvJ background. Both male and female mice were used in this study, and no differences were observed according to sex. All animal procedures followed the U.S. Public Health Service Policy on the Humane Care and Use of Laboratory Animals. The use of animals was approved by the local Institutional Animal Care and Use Committee and the Animal Welfare Committee at the University of Texas Health Science Center at Houston.

Immunohistochemical analysis. We used *Opn4* reporter mice to conduct *Tbr2* and *Opn4* colabeling analyses because the available antibodies against *Tbr2* and *Opn4* were both derived from rabbits. Embryos and embryonic or adult eyes were fixed, cryoembedded, and sectioned into 12–20 μm slices for immunofluorescent staining, as described by Mao et al. (2008). Retinal sections or whole retinas were placed in a 300 W microwave oven in 10 mM sodium citrate for 18 min to expose the antigen epitopes. Microwave-treated sections or whole retinas were then incubated with the following primary antibodies: rabbit anti-*Tbr2*/*Eomes* (1:1000 dilution; catalog #AB23345; Abcam), chicken anti-β-galactosidase (1:2000 dilution; catalog #9361; Abcam), chicken anti-green fluorescent protein (GFP; 1:600 dilution; catalog #A10262; Life Technologies), and rabbit anti-melanopsin/*Opn4* (1:1000 dilution; catalog #N39; Advanced Targeting Systems). Alexa Fluor-conjugated secondary antibodies were obtained from Life Technologies (1:600 dilution). DAPI (1.5 μg/ml; Vector Laboratories) was used to stain the nuclei. Images were acquired on a Zeiss 510 confocal laser scanning microscope (Carl Zeiss) and exported as .TIFF files into Adobe Photoshop (Adobe Systems). Cell counting was conducted using the cell counter plugin of NIH ImageJ. All animals used in the immunohistochemical comparisons were littermates.

Morphological analysis of ipRGCs. Mice were killed by cervical dislocation. Retinas were quickly isolated from sclera, mounted flat on black membrane filter paper, and perfused with oxygenated Ames medium (Sigma-Aldrich) in a recording chamber. The glass microelectrodes were pulled with a Brown-Flaming horizontal micropipette puller (Sutter Instruments), tip-filled with a mixture of 1% Lucifer yellow-CH (Life Technologies) and 3.5% Neurobiotin (NBT; Vector Laboratories), and backfilled with 3 M LiCl. The GFP⁺ ipRGCs were viewed with an epifluorescence microscope with a 40× water-immersion objective and randomly targeted. Cell penetration was verified within the first minute by iontophoresis of Lucifer yellow with negative current (−1 nA at 3 Hz). NBT was then injected into cells with positive current (+1 nA at 3 Hz) for 10–12 min. Tissue was fixed with 4% paraformaldehyde (PFA) in 0.1 M phosphate buffer for 1 h at ambient temperature. Dye-filled cells were

visualized by incubating with Alexa Fluor 488-conjugated streptavidin (1:1000 dilution; Life Technologies) at 4°C overnight. The injected retinal tissue was immunostained with anti-GFP antibody to confirm the correct cell targeting and colabeled with anti-Opn4 antibody to distinguish the M2, M4, and M5 ipRGC subtypes on the basis of their distinct *Opn4* expression levels. A goat polyclonal antibody for choline acetyltransferase (ChAT; 1:100 dilution; Millipore) was used to indicate the depth of dendritic stratification of the injected cells.

PLR analysis. Mice aged 3–6 months were dark adapted for >30 min, gently restrained by hand, and exposed to infrared light ($\lambda > 650$ nm). Under infrared illumination, a single eye was illuminated with white light, starting from the dimmest intensity [-7 log unattenuated light density (Io)] to the brightest (0 log Io), in 30 s, 1 log unit incremental steps. Io at the level of the mouse eye was 15 mW/cm² and was measured at the end of each experiment (IL-1700 radiometer; International Light). A series of pictures of the illuminated eye was captured with a digital camera at the end of the 30 s period, from which the percentage of pupil constriction was determined. The light source was a 150 W xenon arc lamp attenuated through neutral density filters (Lambda 10-3 and 10-2B; Sutter Instruments). To estimate the maximum pupil constriction, data were fitted to a sigmoidal dose–response curve with the lower asymptote fixed to $y = 0$ using OriginLab 8.5 software.

Wheel-running activity. Mice aged 3–6 months were individually housed in wheel-running cages (Columbus Instruments). Wheel-running activity was constantly recorded and binned in 2 min. Mice were kept in each of the different light/dark cycling conditions for 10–25 cycles. During the light phase, mice were exposed to ~650 lux of fluorescent white light. Data analysis was performed using NIH ImageJ software with the Actogram plug-in (Schmid et al., 2011) and the X^2 periodogram.

X-gal staining of the brain. To visualize the expression of β -galactosidase (LacZ) from the *Opn4^{Tau-LacZ}* allele, mice were anesthetized by intraperitoneal injection of tribromoethanol (0.2 ml/g) and arterially perfused for 20 min with 30 ml of cold 4% PFA in PBS, pH 7.3. Their brains were removed, cryoprotected in 30% sucrose in PBS, pH 7.3, overnight, and sliced coronally into 50 μ m sections. For X-gal staining, brain sections were washed twice for 10 min in wash buffer (100 mM phosphate buffer at pH 7.3, 2 mM MgCl₂, 0.01% deoxycholate, and 0.02% Nonidet P-40), and incubated in staining solution (5 mM potassium ferricyanide, 5 mM potassium ferrocyanide, and 1 mg/ml X-gal in wash buffer) at ambient temperature in a light-tight chamber overnight.

RGC axon tracing. Two microliters of cholera toxin subunit B (CTB) conjugated with Alexa Fluor-555 (1 mg/ml; Invitrogen) was injected into the vitreous of the right eye using a 35 gauge NanoFil system (World Precision Instruments) to trace the RGC axons in the suprachiasmatic nuclei (SCNs). One day after CTB injection, animals were anesthetized and killed with perfusion using 4% PFA. Whole brains were dissected and postfixed with 4% PFA. Cryopreserved brains were sectioned into consecutive 100 μ m coronal sections for additional processing.

Results

Expression of *Tbr2* and *Opn4* in developing and mature mouse retinas suggests a stringent epistatic relationship between *Tbr2* and *Opn4*/ipRGCs

To demonstrate the link between ipRGCs and *Tbr2*, we first studied the spatiotemporal relationship between *Tbr2* and *Opn4* during retinal development using retinas from *Opn4^{Tau-LacZ/+}* mice (Hattar et al., 2002). Consistent with the findings of a previous report (McNeill et al., 2011), a few *Opn4^{Tau-LacZ/+}* RGCs were readily detectable at E15 (Fig. 1A1), and all *Opn4^{Tau-LacZ}*

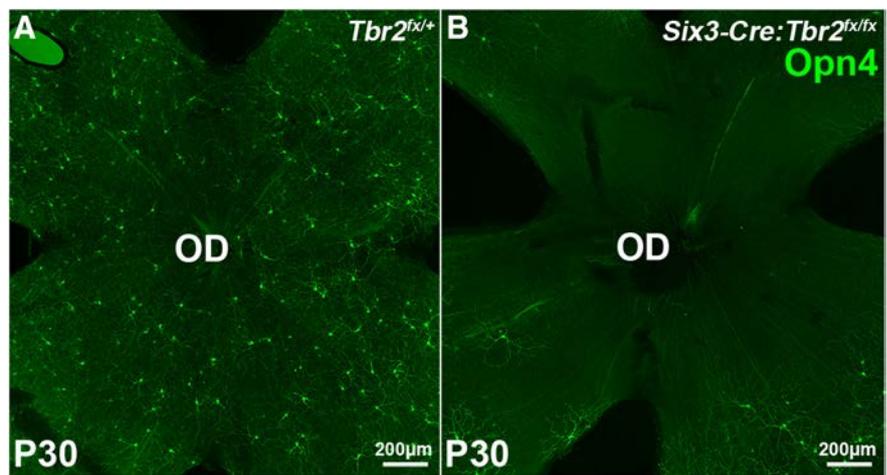


Figure 3. *Tbr2* is essential for the formation of M1–M3 ipRGCs. Confocal images of control *Tbr2^{lox/+}* (A) and mutant *Six3-Cre:Tbr2^{lox/lox}* (B) flat-mounted retinas, immunolabeled with anti-melanopsin/Opn4 antibody at P30. The central retina in B shows no Opn4⁺ cells, which is consistent with the presence of the LacZ reporter in the *Six3-Cre:Tbr2^{lox/lox};Opn4^{Tau-LacZ/+}* retina shown in Figure 2. OD: Optic disc.

expressing cells were found in *Tbr2⁺* RGCs (Fig. 1A1–A3, arrowheads; six retinas and >100 LacZ⁺ RGCs examined). At postnatal day 0 (P0), when the number of ipRGCs peaked, *Tbr2* was found in every LacZ⁺ cell (Fig. 1B1–B3, arrowheads; six retinas and >1000 LacZ⁺ RGCs examined). As the retinas matured from P5 to P60, *Opn4^{Tau-LacZ}* expression was consistently restricted to *Tbr2⁺* RGCs (Fig. 1C–E). A limited number of displaced ipRGCs was reported in the inner nuclear layer (Hattar et al., 2002); we found that these displaced ipRGCs also expressed *Tbr2* (Fig. 1F1–F3). Without exception, *Opn4* expression was restricted to *Tbr2⁺* RGCs at all stages examined.

The tight association between *Opn4* and *Tbr2* expression was further substantiated in a *Brn3b*-deficient retina that had a reduced number of RGCs and a skewed ipRGC distribution pattern. *Brn3b* is a key regulator of RGC differentiation. In *Brn3b^{-/-}* retinas, ~80% of RGC precursors die or trans-differentiate into other cell types during development, and the remaining 20% develop evenly across the retina (Gan et al., 1999; Lin et al., 2004; Qiu et al., 2008). However, *Opn4⁺* cells are concentrated at the ventral retina (Lin et al., 2004). We reasoned that any key transcription factors involved in regulating the formation of ipRGCs would likely present a similarly distorted expression pattern in the *Brn3b^{-/-}* retina. Indeed, *Tbr2⁺* cells were distributed in a dorsal-low and ventral-high pattern in *Brn3b^{-/-};Opn4^{Tau-LacZ/+}* retinas (Fig. 1G1, G2), in which every remaining ipRGC was *Tbr2⁺* (Fig. 1G1, G2).

Of the five known subtypes (M1–M5) of ipRGCs (Schmidt and Kofuji, 2009; Ecker et al., 2010; Schmidt et al., 2011b; Hu et al., 2013), the *Opn4^{Tau-LacZ}* mice, which were used to collect the data shown in Figure 1, mainly reported that the M1 subtype dominated expression (Hattar et al., 2002; Schmidt et al., 2011a). To determine whether non-M1 ipRGCs also expressed *Tbr2*, we examined retinas from *Opn4^{Cre/Cre};Z/EG* mice, in which the GFP signal reports all known ipRGC subtypes (Ecker et al., 2010; Schmidt et al., 2011a). We found that all ipRGCs were *Tbr2⁺*, but not all *Tbr2⁺* cells were ipRGCs (number of GFP⁺/number of *Tbr2⁺* = 21.1 ± 4.6%, $n = 4$) (Fig. 1H1–H3). Some GFP⁺ ipRGCs with significantly larger soma, known as the M4 subtype (Ecker et al., 2010; Estevez et al., 2012), were untraceable in *Opn4^{Tau-LacZ/+}* retinas because of their low *Opn4* levels (Ecker et al., 2010; Schmidt et al., 2011a). Correspondingly, these cells had

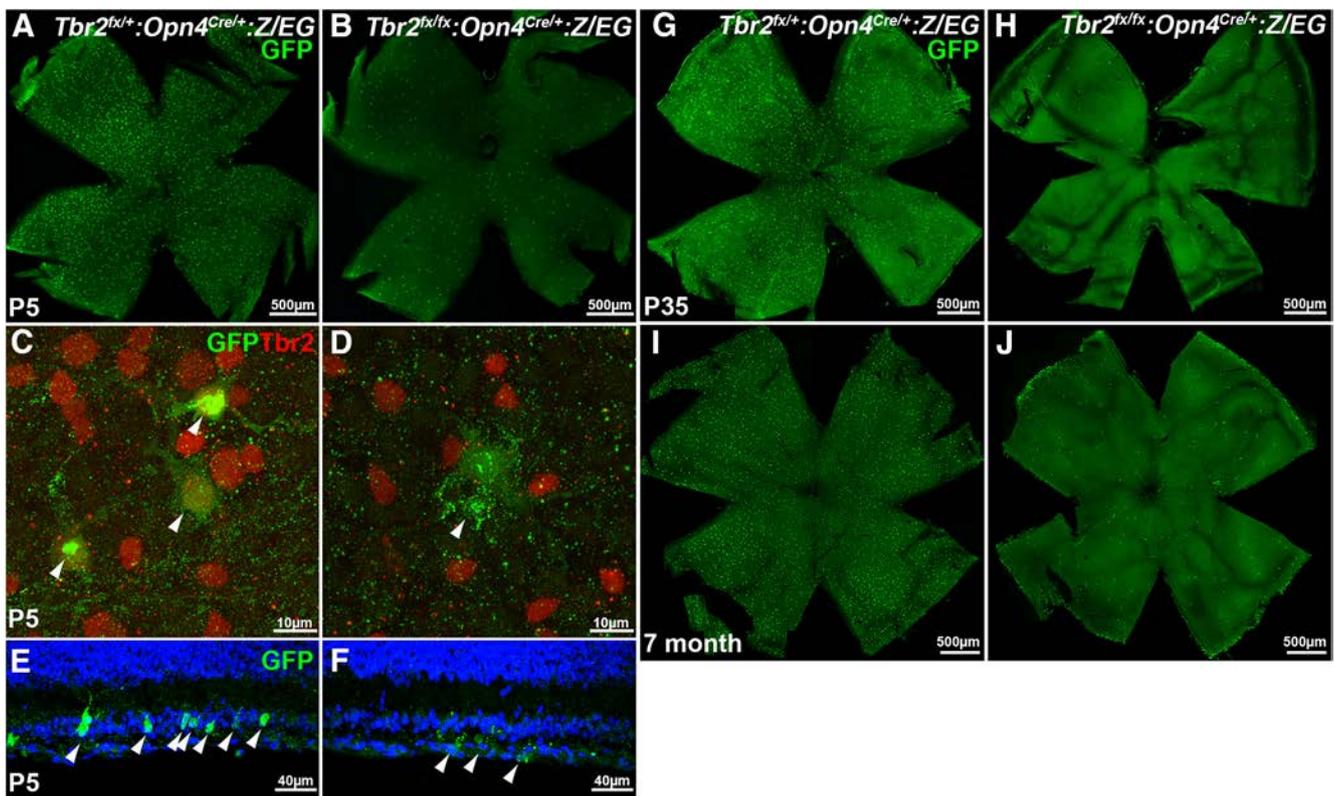


Figure 4. *Tbr2* is essential for the maintenance of ipRGCs. **A–J**, Confocal images of *Opn4*^{Cre/+}:Z/EG (**A, C, E, G, I**) and *Tbr2*^{lox/lox}:*Opn4*^{Cre/+}:Z/EG (**B, D, F, H, J**) retinas at P5 (**A–F**), P35 (**G, H**), and 7 months (**I, J**) that were labeled for GFP only (**A, B, E–J**) or GFP and *Tbr2* (**C, D**). **C–F**, Representative images show normal GFP signal in the control *Opn4*^{Cre/+}:Z/EG ipRGCs (**C, E**) and diffused abnormal GFP signal in *Tbr2*^{lox/lox}:*Opn4*^{Cre/+}:Z/EG cells (**D, F**).

lower levels of *Tbr2* (Fig. 1*H1–H3*, arrowheads). Therefore, expression levels of *Opn4* were also correlated with expression levels of *Tbr2* in ipRGCs. These results are consistent with those of a previous transcript-profiling study that showed enrichment of *Tbr2* in ipRGCs (Siegert et al., 2012). Our data show that all ipRGCs expressed *Tbr2*, indicating that a stringent epistatic relationship exists between *Tbr2* and *Opn4* and suggesting that *Tbr2* plays a significant role in regulating ipRGCs.

Tbr2 is essential for the formation of ipRGCs

To determine whether *Tbr2* is involved in the formation of ipRGCs, we deleted *Tbr2* specifically in the retinal primordia before RGC specification by crossing *Six3–Cre:Tbr2*^{lox/lox}:*Opn4*^{Tau–LacZ/+} and *Tbr2*^{lox/+} mice. The distribution of ipRGCs, as indicated by *Opn4*^{Tau–LacZ} expression, in *Tbr2*-deficient retinas (*Six3–Cre:Tbr2*^{lox/lox}:*Opn4*^{Tau–LacZ/+}) was compared with that in control littermate retinas (*Tbr2*^{lox/+}:*Opn4*^{Tau–LacZ/+}). *Six3–Cre* functions effectively in the central region of retinal primordium before neuronal formation (Furuta et al., 2000), thereby causing a near complete deletion of the *Tbr2*^{lox/lox} allele in the central retina. Although *Tbr2* and *Opn4* were expressed evenly throughout the control retinas (Fig. 2*A1,A2*), their expression disappeared concurrently in the central region of *Six3–Cre:Tbr2*^{lox/lox}:*Opn4*^{Tau–LacZ/+} retinas (Fig. 2*B1,B2*). In the peripheral region of *Six3–Cre:Tbr2*^{lox/lox}:*Opn4*^{Tau–LacZ/+} retinas, in which *Six3–Cre* was less effective (Furuta et al., 2000), some *Opn4*^{Tau–LacZ} ipRGCs remained and expressed *Tbr2* (Fig. 2*C,D*), indicating that *Six3–Cre* was not functional in these cells.

Because the *Opn4*^{Tau–LacZ} line primarily reports the M1 ipRGC subtype (Hattar et al., 2002; Schmidt et al., 2011a), we also used anti-*Opn4* antibody to determine whether the formation of

M1–M3 ipRGC subtypes was affected in *Six3–Cre:Tbr2*^{lox/lox} retinas by immunostaining. We found that *Opn4* expression was completely absent in the central region of the *Six3–Cre:Tbr2*^{lox/lox} retina (Fig. 3*A,B*). These results indicate that *Tbr2* is required for the formation of at least the M1–M3 subtypes of ipRGCs. However, we could not directly determine whether *Tbr2* is required for the formation of M4 and M5 subtypes because these ipRGC subtypes have subdetectable levels of *Opn4* and can only be visualized in the *Opn4*^{Cre/+}:Z/EG line (Ecker et al., 2010; Schmidt et al., 2011a). Currently, the best known reporter line that can reveal M4 and M5 is *Opn4*^{Cre/+}:Z/EG, in which activation of enhanced GFP (eGFP) reporter relies on the activity of the *Opn4* promoter (i.e., ipRGCs have to be specified and *Opn4* has to be activated before cells can be detected). Therefore, this specific *Opn4* reporter line cannot be used to test ipRGC formation.

Tbr2 is essential for maintaining the survival and dendritic integrity of ipRGCs

To determine whether *Tbr2* is required for maintaining ipRGCs, we generated a *Tbr2*^{lox/lox}:*Opn4*^{Cre/+}:Z/EG line for simultaneously deleting the *Tbr2*^{lox} allele and activating GFP by *Opn4*^{Cre} in newly formed ipRGCs. Our results revealed that ~90% of ipRGCs had disappeared in *Tbr2*^{lox/lox}:*Opn4*^{Cre/+}:Z/EG retinas at P5 (307.3 ± 45.8 GFP⁺ ipRGCs, *n* = 3, compared with 3239 ± 170 GFP⁺ ipRGCs in *Tbr2*^{lox/+}:*Opn4*^{Cre/+}:Z/EG control retinas, *n* = 3; Fig. 4*A,B*). The few remaining GFP⁺ ipRGCs were negative for *Tbr2* (Fig. 4*C,D*), indicating that *Opn4*-driven Cre effectively removed the *Tbr2*^{lox/lox} allele and activated the Z/EG allele in ipRGC development. The GFP-marked dendritic structures and cell bodies of the residual *Tbr2*^{-/-} ipRGCs appeared to be less organized and more diffuse than did those in the control

retinas (compare the arrowheads in Fig. 4C,D and E,F), suggesting that the residual cells were unhealthy and dying.

Consistent with these findings, we observed additional ipRGC reduction at a later stage. More than 93% of ipRGCs were missing in the *Tbr2^{flox/flox}:Opn4^{Cre/+}:Z/EG* retinas at P35 (142.7 ± 7.8 GFP⁺ ipRGCs, $n = 3$, compared with 2028.7 ± 48.1 GFP⁺ ipRGCs, $n = 3$; Fig. 4G,H). This difference remained stable at 7 months of age (144 ± 9 GFP⁺ ipRGCs, $n = 3$, compared with 2677 ± 55 GFP⁺ ipRGCs, $n = 3$; Fig. 4I,J). The small number of surviving ipRGCs raised the possibility that *Tbr2* is not required for the survival of specific ipRGC subtypes. To determine whether these residual GFP⁺ cells belong to any specific ipRGC subtypes, we injected NBT tracer into the control and mutant GFP⁺ cells and systematically compared their morphological characteristics, an indicator of ipRGC subtype. In total, 52 cells from 10 control retinas (*Tbr2^{flox/+}:Opn4^{Cre/+}:Z/EG* at P21–P28) and 71 cells from 14 mutant retinas (*Tbr2^{flox/flox}:Opn4^{Cre/+}:Z/EG* at P21–P28) were dye filled and imaged for analyses. On the basis of dendritic morphological features, stratifications, and *Opn4* expression, we were able to detect five subtypes of ipRGCs [11 M1 (21.1%), 20 M2 (38.5%), 10 M3 (19.2%), 7 M4 (13.5%), and 4 M5 (7.7%) cells] from control retinas with similar features, as described previously (Schmidt and Kofuji, 2009; Ecker et al., 2010; Schmidt et al., 2011a,b; Hu et al., 2013). Among the 71 NBT-filled mutant GFP⁺ cells, two (2.8%) had M1-like OFF-lamina dendritic stratification, but their dendrites had less extensive ramifications and smaller dendritic fields (Fig. 5B, 125 μ m) than did the wild-type (WT) M1 (Fig. 5A, 281 μ m). Thirty-five (49.3%) mutant cells displayed ON-lamina dendritic stratifications. These cells contained at least two subtypes on the basis of their nuclear sizes, M4 (~14 μ m) plus M2 or M5 (9.7 ± 1.4 μ m). We could not further distinguish M2 and M5 because of the lack of *Opn4* expression for verification and the nonstandard dendritic diameter (average diameter, 290 μ m for mutant ON cells vs 266 μ m for M2, 314 μ m for M4, and 158 μ m for M5; Fig. 5C,D). Thirty-four (47.9%) mutant cells had bistratified M3-like morphological features, but their dendritic ramifications in ON and OFF laminae appeared uneven (Fig. 5E,F). In addition, their dendritic processes often traveled from

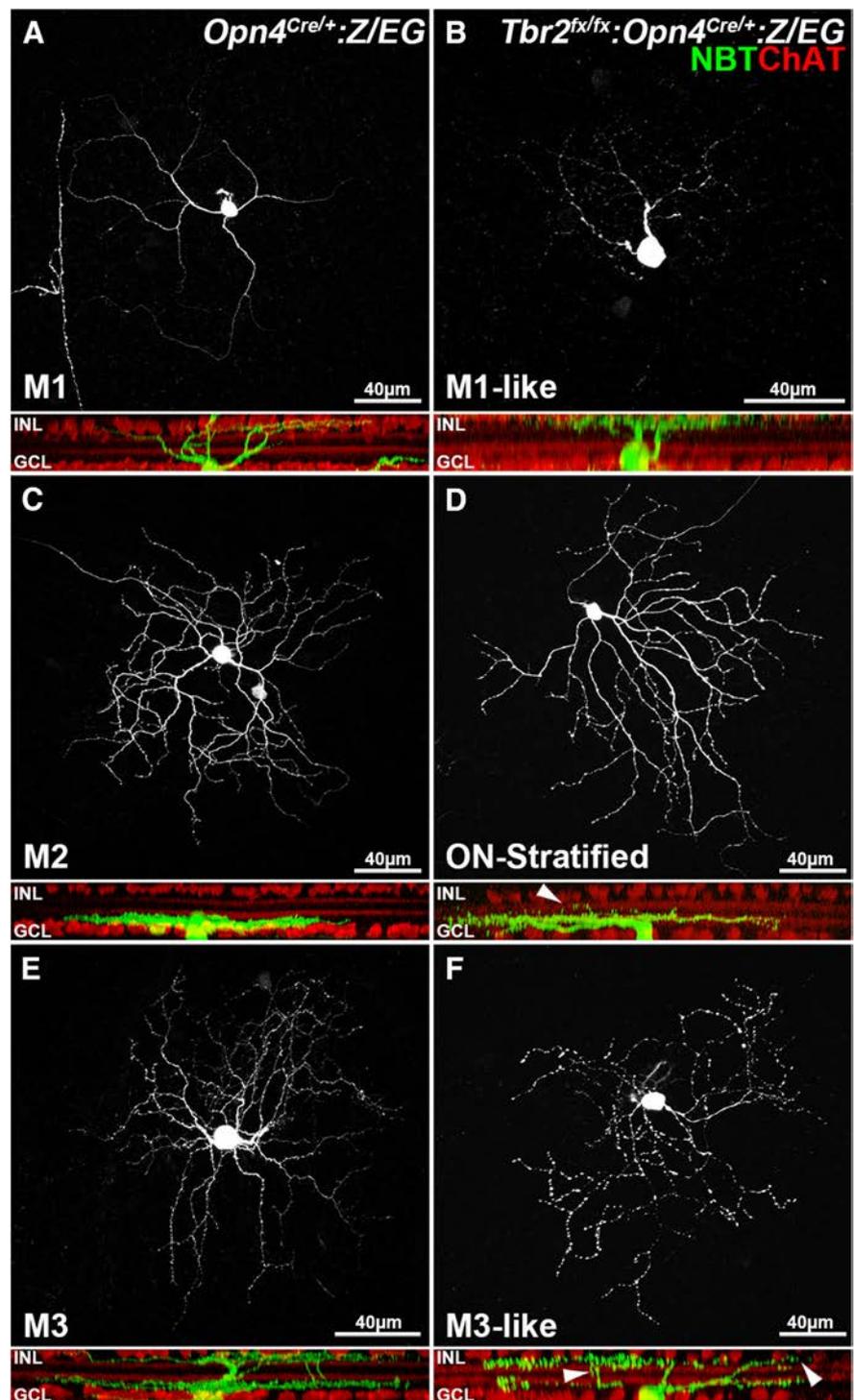


Figure 5. *Tbr2* is required for the normal dendritic morphological features of ipRGCs. **A–F**, Representative confocal images show the morphological features of NBT-injected GFP⁺ cells in *Tbr2^{flox/+}:Opn4^{Cre/+}:Z/EG* control (**A, C, E**) and *Tbr2^{flox/flox}:Opn4^{Cre/+}:Z/EG* mutant (**B, D, F**) retinas. **A**, A representative M1 cell shows sparse dendrites in a whole-mount control retina. The side view revealed its single stratification (green) distal to the OFF cholinergic (ChAT⁺) band (red). **C**, A representative M2 cell displayed more complex dendritic branches proximal to the ON cholinergic band. **E**, A representative M3 cell had bistratified dendritic arbors ramifying at both the OFF and ON sublayers. In the mutant retinas, immunostaining for *Opn4* was negative in all GFP⁺ cells. Therefore, dye-filled cells could only be categorized on the basis of their dendritic stratifications. **B**, A representative M1-like mutant cell extended its dendrites solely to the OFF sublayer. It had a smaller dendritic field and fewer branches than did the M1 cell in the control retina. **D**, A representative ON-layer-stratified mutant cell that may be M2 or M4. Although the major stratification is at the ON layer, minor runoffs can be found in the OFF layer. Thirty-five dye-filled cells (49.3%) ramified to the ON layer. Half of them displayed a small number of dendritic arbors extending to the OFF layer, as shown in the side view. The other half was relatively normal. **F**, An M3-like bistratified mutant cell. The side view shows less organized stratification. NBT, White in top view, green in side view; ChAT, red; GCL, ganglion cell layer; INL, inner nuclear layer.

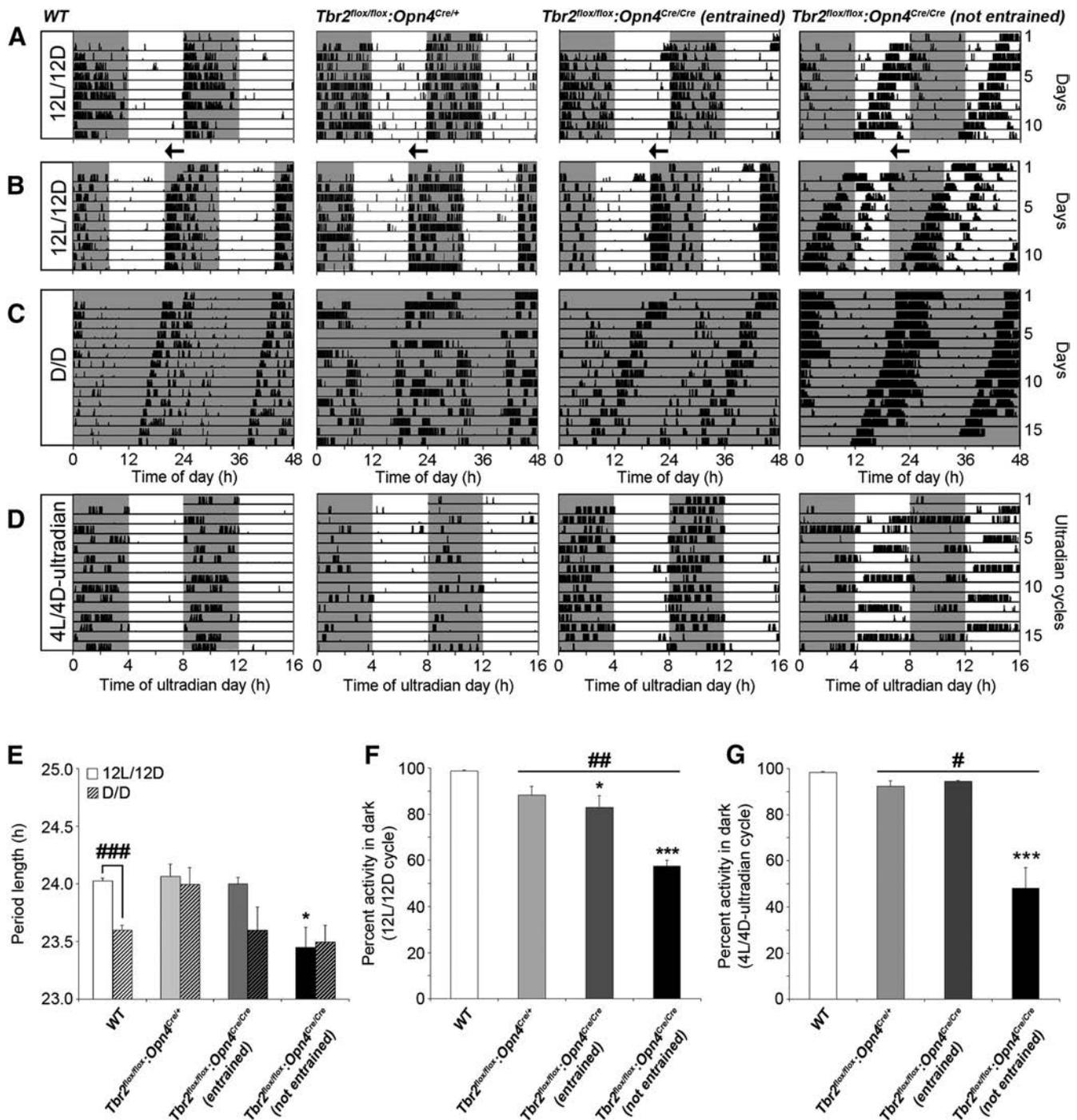


Figure 6. *Tbr2* is necessary for normal light entrainment and masking. **A–D**, Representative double-plotted profiles of wheel-running activity of a WT control (*Tbr2*^{flox/flox}:*Opn4*^{+/+}), a *Tbr2*^{flox/flox}:*Opn4*^{Cre/+} mutant, and two *Tbr2*^{flox/flox}:*Opn4*^{Cre/Cre} mutant mice, obtained under a 12L/12D cycle (**A**), including after a 4 h phase shift (jetlag paradigm; **B**), D/D (**C**), and a 4L/4D ultradian cycle (**D**). Of five *Tbr2*^{flox/flox}:*Opn4*^{Cre/Cre} mice, three showed photo-entrainment to the 12L/12D cycle (entrained to light/dark), and two were unable to photo-entrain (not entrained to light/dark). Mice from all genotypes were able to run freely under D/D conditions. **E**, Average period lengths of the wheel activity rhythm under 12L/12D and D/D conditions. A two-way ANOVA revealed significant differences between the genotypes ($F_{(3,22)} = 4.59, p < 0.01$) but not between lighting cycles ($F_{(1,22)} = 4.38, p = 0.054$) or genotype \times lighting cycles ($F_{(3,22)} = 1.42, p = 0.28$). The genotypes under 12L/12D and D/D were compared using one-way ANOVA, followed by a Tukey's *post hoc* test ($*p < 0.05$, compared with WT). A comparison between the two lighting cycles within a genotype was performed using Student's *t* test ($###p < 0.001$). **F, G**, Percentage activity in the dark under 12L/12D (**F**) and 4L/4D (**G**) cycles. Data were collected from six WT, four *Tbr2*^{flox/flox}:*Opn4*^{Cre/+}, and five *Tbr2*^{flox/flox}:*Opn4*^{Cre/Cre} mice. A comparison between WT and combined *Tbr2*^{flox/flox}:*Opn4*^{Cre/+} and *Tbr2*^{flox/flox}:*Opn4*^{Cre/Cre} animals was performed using Student's *t* test ($*p < 0.05$, $##p < 0.01$). Comparisons among the four groups of animals were performed using one-way ANOVA, followed by a Tukey's *post hoc* test ($*p < 0.05$, $***p < 0.001$, compared with WT). Error bars represent SEM. Shaded areas indicate periods of darkness. Arrows indicate the direction of the 4 h phase shift.

one sublamina into another to form small loose-end branches (Fig. 5D,F, arrowheads). These data suggest that the residual ipRGCs did not belong to a specific subtype; they also exhibited apparent abnormalities in the absence of *Tbr2*. Together, our data

demonstrate that *Tbr2* is essential for the maintenance and maturation of the dendritic structures of ipRGCs. Our data also suggest that *Tbr2* is not merely required for the expression of *Opn4* but also for the formation and survival of ipRGCs.

The remaining ipRGCs in *Tbr2*^{fllox/flox}:*Opn4*^{Cre/+} mice are sufficient to drive light entrainment and masking and the PLR

To determine the extent to which the residual ipRGCs contribute to non-image-forming visual functions, we conducted circadian photo-entrainment and PLR analyses on *Tbr2*^{fllox/flox}:*Opn4*^{Cre/+} mice at the 3–6 months of age. WT and *Tbr2*^{fllox/flox}:*Opn4*^{Cre/Cre} mice were also included, as positive and negative controls, respectively.

In the free wheel-running assay under four distinct conditions of the light/dark cycle, including the 12 h light/dark (12L/12D) cycle (Fig. 6A), a 4 h phase advance shifting (jetlag; Fig. 6B), a dark/dark (D/D) cycle (Fig. 6C), and a 4 h light/dark (4L/4D) ultradian cycle (Fig. 6D), we found that mice from all genotypes showed daily (Fig. 6A,B) and circadian (Fig. 6C) organization of their locomotor activity rhythm. We found small but significant differences between WT and *Tbr2*^{fllox/flox}:*Opn4*^{Cre/+} animals. For example, contrary to the WT controls, the endogenous period of the locomotor activity rhythm of the *Tbr2*^{fllox/flox}:*Opn4*^{Cre/+} animals recorded in a D/D cycle was not significantly different from that recorded in a 12L/12D cycle (Fig. 6C,E). In addition, mice lacking *Tbr2* in ipRGCs had a higher percentage of daily activity confined to the light period compared with the WT mice (Fig. 6D,F,G). The double knock-out mice showed greater difficulty to entrain and mask compared with the WT or the *Tbr2*^{fllox/flox}:*Opn4*^{Cre/+} mice, presumably because these animals combined a more severe reduction in the number of ipRGCs with the lack of *Opn4* (Fig. 6).

In the PLR test, the *Tbr2*^{fllox/flox}:*Opn4*^{Cre/+} mice, despite the severe loss of ipRGCs, exhibited surprisingly mild contractile defect at high irradiance levels compared with the WT (Fig. 7A,B,E). As for the locomotor activity data, *Tbr2*^{fllox/flox}:*Opn4*^{Cre/Cre} mice presented a more severe defect, likely reflecting the additional lack of *Opn4* (Fig. 7A–E), yet *Tbr2*^{fllox/flox}:*Opn4*^{Cre/Cre} mice were still able to constrict their pupil by ~40% in response to bright light (Fig. 7A,E).

Although these results are globally consistent with the requirement of ipRGCs and *Opn4* for normal non-image-forming visual functions, the deficits observed in *Tbr2*^{fllox/flox}:*Opn4*^{Cre/+} mice were surprisingly mild given the fact that the number of ipRGCs in the retinas of these animals is reduced by ~93% (Figs. 4, 5). However, the next few experiments (Figs. 8, 9) may be able to explain these mild behavioral deficits.

Tbr2⁺ RGCs activate *Opn4* on the loss of ipRGCs

The distinct behavioral outcomes in *Tbr2*^{fllox/flox}:*Opn4*^{Cre/+} and *Tbr2*^{fllox/flox}:*Opn4*^{Cre/Cre} mice indicate that there is a small, but relatively stable (compared with the *Tbr2*^{fllox/flox}:*Opn4*^{Cre/Cre} retina), number of ipRGCs in the *Tbr2*^{fllox/flox}:*Opn4*^{Cre/+} retina. To investigate the underlying mechanism responsible for the presence of these cells, we examined *Tbr2*^{fllox/+}:*Opn4*^{Cre/+}:*Z/EG* (control) and *Tbr2*^{fllox/flox}:*Opn4*^{Cre/+}:*Z/EG* (mutant) retinas at P30 to compare their *Opn4* and GFP expression. In the control retinas, the *Opn4*-expressing cells that were positive for GFP (Fig. 8A,C, yellow arrowheads) were M1–M3 subtypes (Ecker et al., 2010). Many of the GFP⁺ ipRGCs that were *Opn4* negative (*Opn4*⁻;

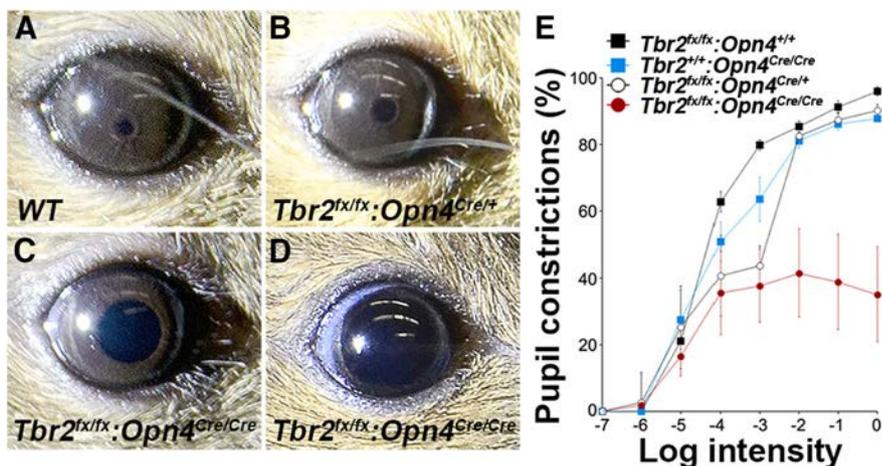


Figure 7. *Tbr2* is required for normal PLR. **A–D**, Representative PLR images of WT control (**A**), *Tbr2*^{fllox/flox}:*Opn4*^{Cre/+} (**B**), and *Tbr2*^{fllox/flox}:*Opn4*^{Cre/Cre} (**C, D**) pupils in response to a 30 s bright light stimulation (0 log lo). Some *Tbr2*^{fllox/flox}:*Opn4*^{Cre/+} mice (3 of 5) showed an ~60% pupil constriction (**C**), and others (2 of 5) were unable to maintain pupil constriction (**D**). **E**, Quantification of the PLR as a function of light intensity. Data were collected from five animals of each genotype. A two-way ANOVA revealed significant light intensity ($F_{(7,119)} = 58.74, p < 0.001$), phenotype ($F_{(2,119)} = 29.64, p < 0.001$), and light intensity \times phenotype ($F_{(14,119)} = 4.75, p < 0.001$) effects. The maximum constriction was $96.03 \pm 1.36\%$ in the WT control mice, $90.22 \pm 1.94\%$ in the *Tbr2*^{fllox/flox}:*Opn4*^{Cre/+} mice, $38.35 \pm 5.41\%$ in the *Tbr2*^{fllox/flox}:*Opn4*^{Cre/Cre} mice, and $87.85 \pm 0.89\%$ in *Opn4*^{Cre/Cre} mice. $I_0 = 15$ mW \cdot cm $^{-2}$ white light. Error bars represent SEM.

Fig. 8A, C, green cells not marked by arrowheads) were defined as M4 or M5 ipRGC subtypes (Ecker et al., 2010). We consistently found that a small percentage of *Opn4*⁺ cells were GFP⁻ (1.5 ± 0.6 per 0.1 mm², $n = 4$; Fig. 8A, C, white arrowheads). As illustrated in Figure 8E and reasoned below, these *Opn4*⁺ but GFP⁻ cells were designated as “newly activated” ipRGCs with recently activated *Opn4* but no activated GFP. In the *Tbr2*^{fllox/+}:*Opn4*^{Cre/+}:*Z/EG* control retinas, *Tbr2* activated *Opn4*^{Cre/+}, which produced Cre and *Opn4*. Over time, the accumulated Cre activity removed *loxP*-flanked β geo (a LacZ and Neo fusion cassette) from the *Z/EG* allele to allow eGFP expression. An important determinant of Cre efficiency is *Opn4* promoter activity, which is variable in different ipRGCs. Therefore, a transitional time window of variable lengths exists between *Opn4* activation and GFP visualization. *Opn4*⁺/GFP⁻ cells were detected in the transitional time when *Opn4*-driven Cre was accumulating to an effective level to activate the *Z/EG* allele.

The few remaining GFP⁺ cells in the *Tbr2*^{fllox/flox}:*Opn4*^{Cre/+}:*Z/EG* mutant retina did not contain *Opn4*, as we predicted, because of the *Opn4*^{Cre}-mediated *Tbr2* deletion (Fig. 8B,D,F). Nonetheless, an extensive number of newly activated *Opn4*⁺/GFP⁻ cells (11.8 ± 3.1 per 0.1 mm², $n = 4$; Fig. 8B,D) was detected throughout the entire retina, suggesting that some RGCs begin to turn on *Opn4* expression in the absence of ipRGCs. Given the stringent requirement for *Tbr2* in the formation and maintenance of ipRGCs (results from Figs. 2–4), it is conceivable that these *Opn4*⁺ cells arose from other *Tbr2*⁺ RGCs. As illustrated in Figure 8F, in the mutant *Tbr2*^{fllox/flox}:*Opn4*^{Cre/+}:*Z/EG* retina, *Tbr2* activates *Opn4*^{Cre/+} in ipRGCs. Both *Opn4* and Cre are presumably initially produced using the same kinetics. The accumulated Cre recombinase eventually excises the floxed *Tbr2*, leading to *Opn4* inactivation and ultimately cell death. Other *Tbr2*⁺ RGCs sense the loss of ipRGCs and activate *Opn4*^{Cre/+}. Over time, the accumulated Cre activity removed *loxP*-flanked β geo from the *Z/EG* allele to allow eGFP expression. Similar to the control (Fig. 8E), a transitional time window of unknown

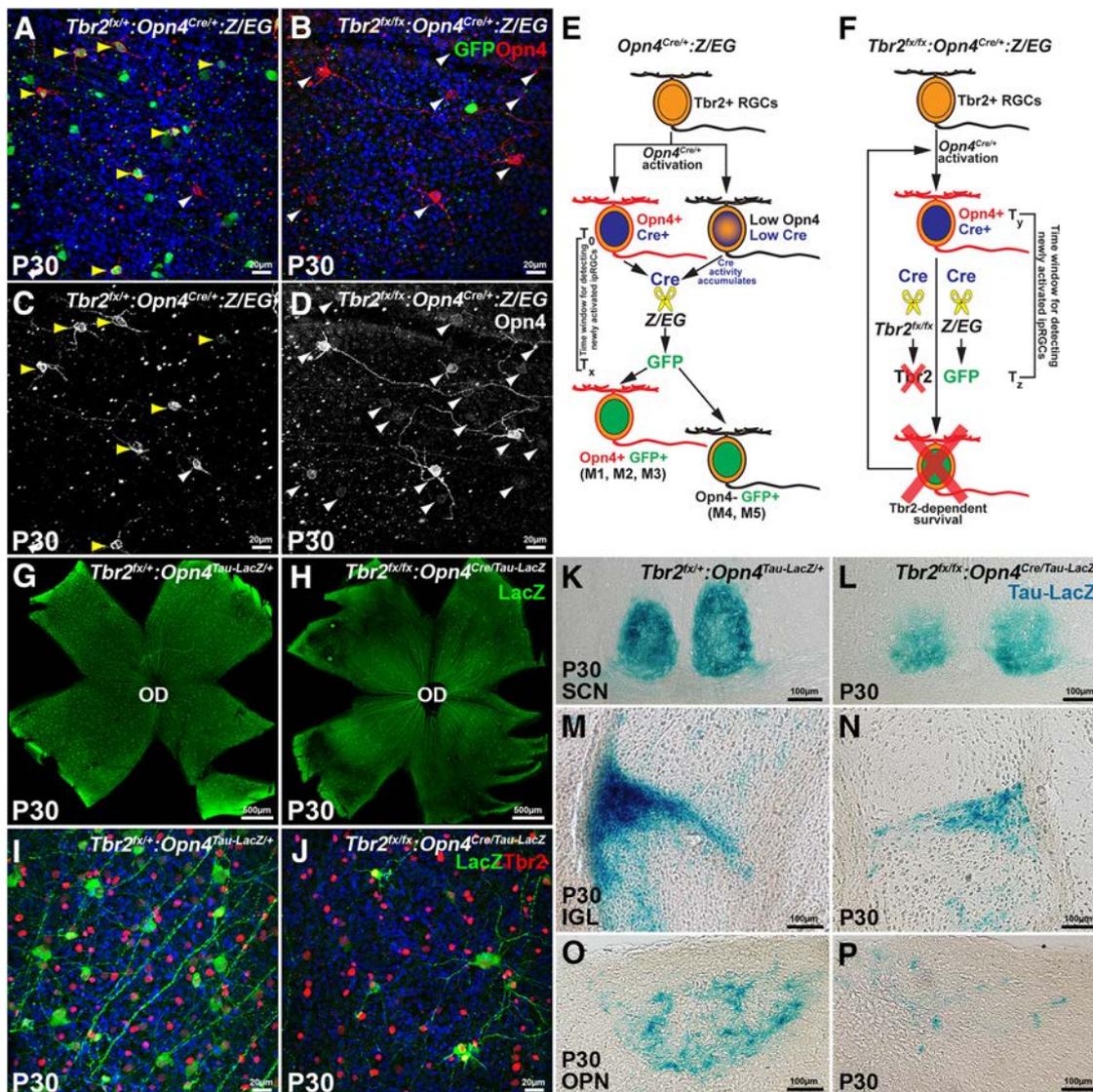


Figure 8. *Tbr2*⁺ RGCs activate *Opn4* on the loss of ipRGCs. **A–D**, Confocal images of randomly selected regions from *Tbr2^{flx/+} :Opn4^{Cre/+} :Z/EG* control retinas (**A**) and *Tbr2^{flx/flx} :Opn4^{Cre/+} :Z/EG* mutant retinas (**B**) at P30, labeled for GFP (green) and *Opn4* (red). In the *Tbr2^{flx/+} :Opn4^{Cre/+} :Z/EG* retina, all *Opn4*⁺ RGCs were negative for GFP (white arrowheads in **B**), suggesting that they are new ipRGCs. **C, D**, Same images as in **A** and **B**, respectively, but showing only the red (*Opn4*) channel. Note that only two new ipRGCs can be detected in the control retinas (white arrowheads in **C**), yet many more new ipRGCs with weak *Opn4* expression can be seen in the *Tbr2*-deficient retinas (white arrowheads in **D**). **E**, An illustration of the molecular events in ipRGCs in the *Opn4^{Cre/+} :Z/EG* retinas, which are commonly used as WT to trace the morphological features of M1–M5 ipRGC subtypes. In this WT control retina, *Tbr2* activates *Opn4^{Cre/+}*, which produces Cre and *Opn4*. Over time, the accumulated Cre activity removes *loxP*-flanked β geo (a LacZ and Neo fusion cassette) from the *Z/EG* allele to allow eGFP expression. A transitional time window of variable lengths (represented by the bracket between *T₀* and *T_x*) exists between *Opn4* activation and GFP visualization. *Opn4*⁺/*GFP*[−] cells are detected in the transitional time and can be identified as “new ipRGCs” because their *Opn4* has been activated but their GFP has not. *T₀*, the time point when *Tbr2* activates *Opn4* and Cre in the new ipRGCs; *T_x*, the time point when the newly produced GFP becomes visible. **F**, An illustration showing the dynamics of *Tbr2*-activated new ipRGCs in the mutant *Tbr2^{flx/flx} :Opn4^{Cre/+} :Z/EG* retina. *Tbr2* activates *Opn4^{Cre/+}* in ipRGCs. Both *Opn4* and Cre are presumably initially produced using the same kinetics. The accumulated Cre recombinase eventually excises the floxed *Tbr2*, leading to *Opn4* inactivation and ultimately cell death. Similar to the control, a transitional time window of unknown length (between *T₀* and *T_z*) should be present to detect *Opn4*⁺/*GFP*[−] new ipRGCs. **G–J**, Confocal images of *Tbr2^{flx/+} :Opn4^{Tau-LacZ/+}* retinas (**G, I**) and *Tbr2^{flx/flx} :Opn4^{Cre/Tau-LacZ}* retinas (**H, J**) at P30 had been labeled for LacZ (**G, H**) or LacZ and *Tbr2* (**I, J**). The mutant retinas show many LacZ⁺ cells throughout the retinas. This finding is in contrast to the *Six3*–Cre-active region in Figure 2*B*, which has no LacZ⁺ cells. **K–P**, *Opn4^{Tau-LacZ/+}* ipRGCs from the control *Tbr2^{flx/+} :Opn4^{Tau-LacZ/+}* and the mutant *Tbr2^{flx/flx} :Opn4^{Cre/Tau-LacZ}* retinas project to identical brain regions in the SCN (**K, L**), IGL (**M, N**), and olivary pretectal nuclei (OPN; **O, P**). OD: Optic Disc.

length (between *T₀* and *T_z*) should be present to detect *Opn4*⁺/*GFP*[−] newly activated ipRGCs.

To trace whether these newly activated *Opn4*⁺ cells project axons to the same brain areas as native ipRGCs, we created a *Tbr2^{flx/flx} :Opn4^{Cre/Tau-LacZ}* line on the basis of genetic logic similar to that used to develop the *Tbr2^{flx/flx} :Opn4^{Cre/+} :Z/EG* allele. In brief, *Tbr2*-dependent *Opn4^{Cre/Tau-LacZ}* activation produces Cre and Tau–LacZ. Accumulated Cre excises the floxed *Tbr2* allele, while Tau–LacZ aggregates in cell bodies and processes. *Tbr2* excision leads to ipRGC death. New rounds of *Opn4^{Cre/Tau-LacZ}*

are activated in other *Tbr2*⁺ cells, which leads to additional cell death and *Opn4^{Cre/Tau-LacZ}* activation. We learned from the *Tbr2^{flx/flx} :Opn4^{Cre/+} :Z/EG* retina that most *Opn4* active cells are newly activated ipRGCs at P30. Although differences exist between *Opn4^{Cre/+}* and *Opn4^{Cre/Tau-LacZ}*, it is known that a lack of *Opn4* does not cause ipRGC death (Lucas et al., 2003). Therefore, Tau–LacZ⁺ cells found in the *Tbr2^{flx/flx} :Opn4^{Cre/Tau-LacZ}* retina represent the equivalent population of *Opn4*⁺ cells in the *Tbr2^{flx/flx} :Opn4^{Cre/+} :Z/EG* retina at the same age described in Figure 8, **B** and **D**. Our results showed that many Tau–LacZ⁺

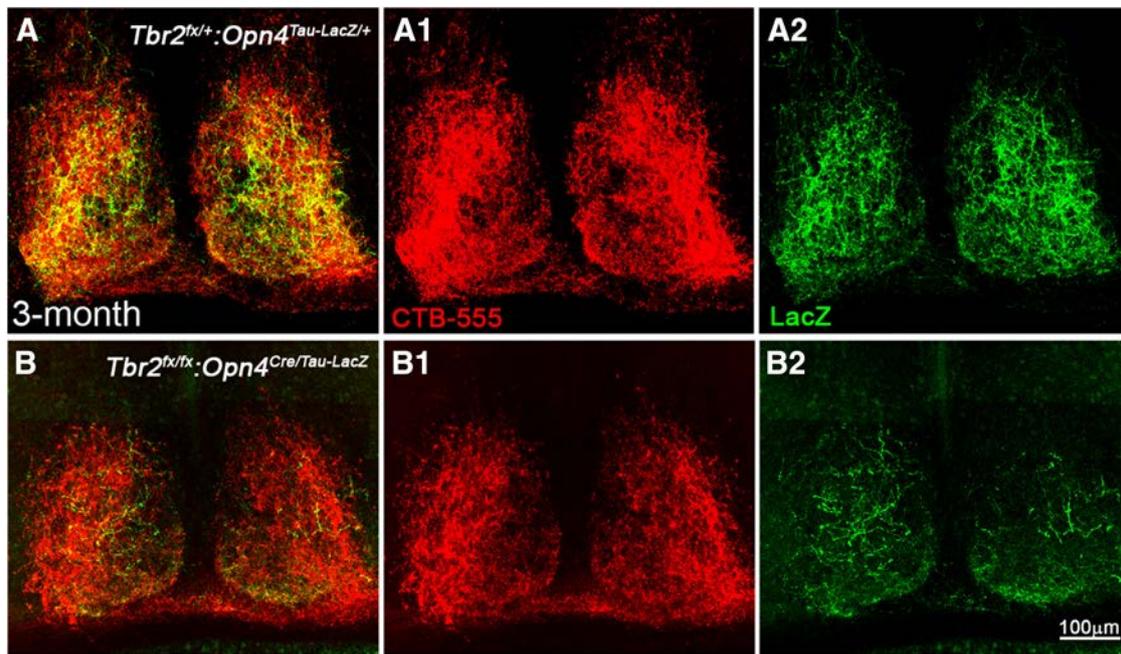


Figure 9. Anterograde RGC axon tracing reveals more axon coverage in the SCN than reported by *Opn4^{Tau-LacZ}*. **A**, A section of center SCN from a *Tbr2^{lox/+}:Opn4^{Tau-LacZ/+}* control mouse that received 2 μ l of CTB-555 in the right eye. **A1** shows CTB-555 labeling, and **A2** shows LacZ labeling. **B**, A section of center SCN from a *Tbr2^{lox/lox}:Opn4^{Cre/Tau-LacZ}* mutant mouse received the same CTB-555 treatment as the control. **B1** shows CTB-555 labeling, and **B2** shows LacZ labeling.

cells were present throughout P30 *Tbr2^{lox/lox}:Opn4^{Cre/Tau-LacZ}* retinas (Fig. 8*G,H*); as predicted, they were all positive for *Tbr2* (Fig. 8*I,J*), supporting the hypothesis that additional *Tbr2⁺* RGCs activate *Opn4* in response to the loss of native ipRGCs. More importantly, these *Tbr2⁺/Tau-LacZ⁺* cells projected to central brain targets that are identical to those of native ipRGCs, including SCNs, intergeniculate leaflet (IGL), and olivary pretectal nuclei (Fig. 8*K–P*); thus, these cells are indeed ipRGCs (henceforth termed new ipRGCs vs native ipRGCs).

However, it was noticeable that the LacZ signal in the brain targets of a *Tbr2^{lox/lox}:Opn4^{Cre/Tau-LacZ}* mouse is less intense than those in a control mouse. We believe, in addition to cell loss, one important reason for such low LacZ signal is that most new ipRGCs have very low *Opn4* promoter activity as indicated by extremely weak *Opn4* labeling in Figure 8, *B* and *D*. If this is true, an anterograde label of ipRGCs should reveal axons that are not detected using the Tau-LacZ reporter. Therefore, we injected Alexa Fluor-555-tagged CTB (CTB-555) into the eyes of 3-month-old mutant (*Tbr2^{lox/lox}:Opn4^{Cre/Tau-LacZ}*) and control (*Tbr2^{lox/+}:Opn4^{Tau-LacZ/+}*) mice ($n = 3$) and compared their RGC projections in the SCN. Results showed that CTB-555 signal covered a much bigger area than did the LacZ reporter in both mutant and control SCN (Fig. 9), indicating that more RGCs target to SCNs than those revealed by LacZ reporter. The extra SCNs coverage revealed by CTB-555 anterograde tracing may include ipRGCs that have low *Opn4* promoter activity and other *Tbr2⁺* RGCs that have projected to the same area.

To further substantiate that the new ipRGCs were pre-routed to their brain targets but not produced through *de novo* genesis in the mature retinas, we conducted birthdating experiments by injecting BrdU into P23 animals (one injection per day for 5 consecutive days) and concluded that the new ipRGCs were not born in the adult retina (negative data not shown). Our cumulative results indicate that some *Tbr2⁺* RGCs that were initially lacking *Opn4* but projected to the same regions as ipRGCs began to activate *Opn4* expression on the loss of native ipRGCs.

Tbr2⁺ RGCs serve as a reservoir for ipRGCs

Overall, our data support a model in which many *Tbr2⁺* RGCs have the capacity to activate *Opn4* expression and can serve as a reservoir for ipRGCs. They are pre-established during retinogenesis and project to the same regions as native ipRGCs. In *Tbr2^{lox/lox}:Opn4^{Cre/+}* mutant mice, most native ipRGCs die shortly after the *Opn4^{Cre}*-mediated deletion of *Tbr2*, triggering the activation of *Opn4* in these reservoir *Tbr2⁺* RGCs (as illustrated in Fig. 10*A,B*). If this model is correct, we predict that the number of *Tbr2⁺* RGCs will decline in older *Tbr2^{lox/lox}:Opn4^{Cre/+}* mutant retinas when the pool of *Tbr2⁺* cells decreases because of recurring *Tbr2* deletion (Fig. 10*B*). To test this, we examined retinas from 7-month-old mice and found that the density of *Tbr2⁺* cells dropped significantly in the mutant mice ($8.55 \pm 2.28\%$ as normalized by DAPI⁺ cells, $n = 3$; Fig. 10*D*) compared with that in the control mice ($13.65 \pm 2.29\%$ as normalized by DAPI⁺ cells, $n = 3$; Fig. 10*C*). Furthermore, <30 *Opn4⁺* cells were found in the entire mutant retina at 7 months of age (data not shown), suggesting that *Tbr2⁺* RGCs, which can activate *Opn4*, were nearly exhausted in the aging mutant retinas.

Discussion

We present an integrated study that combines sophisticated mouse genetics, single-cell dye filling, and behavior analyses to demonstrate that *Tbr2* is required to regulate the formation and survival of ipRGCs; this represents the first report of this molecular regulator in conjunction with any known RGC subtypes. Our study also reveals a previously unknown mechanism whereby some *Tbr2⁺* RGCs, which are pre-established during developmental stages, can compensate for the loss of native ipRGCs by activating *Opn4*. It is known that there is no neurogenesis in the WT adult retina (Kiyama et al., 2012). However, the number of GFP⁺ ipRGCs in *Opn4^{Cre/+}:Z/EG* control mice increases significantly with age [2028 ± 48.1 at P35 (Fig. 3*G*); 2677 ± 55 at 7 months; 2870 at 10 months], suggesting that the number of ipRGCs can be modulated and that *Tbr2⁺* RGCs can

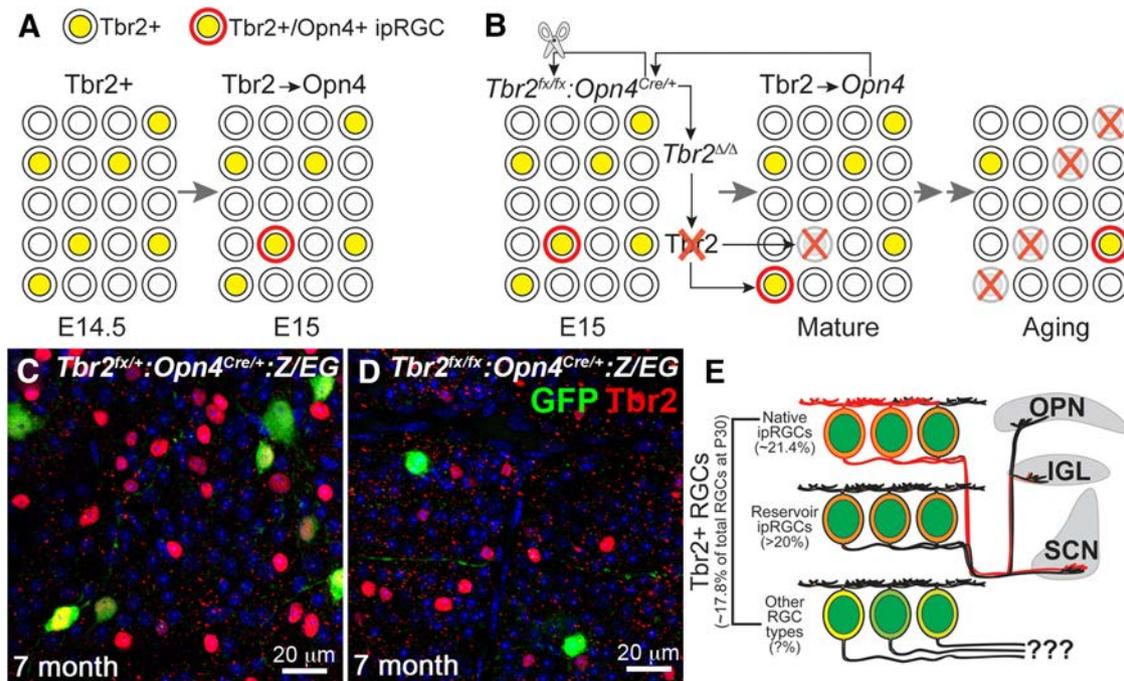


Figure 10. A working model of *Tbr2*⁺ RGCs as a reservoir for ipRGCs. **A**, In normal development, *Tbr2*⁺ RGCs (represented by yellow circles) appear at approximately E14.5. Many of them begin to project to SCN, IGL, and olivary pretectal nuclei (OPN) from E15 onward. A few *Opn4*⁺ ipRGCs (represented by yellow circles with red rings) emerge from these *Tbr2*⁺ RGCs. *Tbr2*–*Opn4* regulation continues through developmental stages and is maintained in adulthood. **B**, In *Tbr2*^{lox/lox};*Opn4*^{Cre/+} retinas, *Opn4*-driven Cre deletes *Tbr2*^{lox/lox} alleles and *Tbr2* protein slowly diminishes, leading to inactivation of *Opn4* and finally the death of native ipRGCs. Through an as-yet unclear mechanism, this event triggers activation of *Opn4* in other *Tbr2*⁺ RGCs that have projected to the same brain regions during development. These new ipRGCs with newly activated *Opn4* repeat the *Opn4*^{Cre}-mediated deletion of the *Tbr2*^{lox/lox} cycle until the *Tbr2*⁺ RGC source is exhausted, as found in older animals. **C**, **D**, Confocal images of control *Tbr2*^{lox/lox};*Opn4*^{Cre/+};*Z/EG* (**C**) and mutant *Tbr2*^{lox/lox};*Opn4*^{Cre/+};*Z/EG* (**D**) retinas labeled for GFP (green) and *Tbr2* (red) at 7 months of age show a drastically reduced density of *Tbr2*⁺ cells in the mutant retina. This result supports the model that *Tbr2*⁺ RGCs can serve as a reservoir for ipRGCs. **E**, *Tbr2*⁺ RGC population in a P30 WT mouse retina. *Tbr2*⁺ RGCs occupy ~17.8% of all RGCs. Among them, ~21.4% (4–5% of total RGCs) are native ipRGCs that are detectable by the *Opn4*^{Cre};*Z/EG* reporting system. A currently unknown number of other *Tbr2*⁺ RGCs that project to identical regions in the brain as do the native ipRGCs can serve as reservoir, or buffer, ipRGCs. Additional populations of *Tbr2*⁺ RGCs with yet-to-be characterized morphological features, functions, and central projections (labeled by question marks) may also be present.

serve as a reservoir for this task (details described in Fig. 10E). The number of reservoir ipRGCs is currently unknown. The *Opn4*-expressing new ipRGCs in P30 *Tbr2*^{lox/lox};*Opn4*^{Cre/+} retinas is ~1840 (Fig. 8D, 11.8 per 0.1 mm² × 15.6 mm², total area of mouse retina; derived from Remtulla and Hallett, 1985), which accounts for ~20% of *Tbr2*⁺ RGCs. Because the upregulation of *Opn4* in reservoir ipRGCs is a continuous process in *Tbr2*^{lox/lox};*Opn4*^{Cre/+} retinas, more RGCs may also have the potential to activate *Opn4* expression. Supporting this view is the finding that the SCN of 3-month-old *Tbr2*^{lox/lox};*Opn4*^{Cre/Tau-LacZ} animals still received extensive RGC axon projection (Fig. 9B). Whether these SCN-projected RGCs are all *Tbr2*⁺ and whether they all have the potential to activate *Opn4* expression remain to be studied.

The novel concept of reservoir ipRGCs provides mechanistic explanations of why ipRGCs are more resistant to injury and mitochondrial optic neuropathies (La Morgia et al., 2010; Sexton et al., 2012), why the SCN receives projections from RGCs that do not express *Opn4* in rats and hamsters (Gooley et al., 2003; Morin et al., 2003; Sollars et al., 2003), and why the number of ipRGCs and *Opn4* level can be modulated by light/dark conditions (Hannibal et al., 2005; Hannibal, 2006). In addition, the increased number of GFP⁺ cells detected in older *Opn4*^{Cre/+};*Z/EG* retinas suggest that *Opn4* expression in the ipRGC reservoir likely fluctuates at subdetectable levels. In this reporting system, the fluctuating *Opn4*^{Cre} in each ipRGC may activate the *Z/EG* allele to turn on GFP when it reaches the threshold level. Subsequently, the ipRGC will be registered as a GFP⁺ ipRGC in its lifetime, regardless of its current *Opn4* expression level. A plausible expla-

nation for rodents (and perhaps other mammals) developing a reservoir of ipRGCs is to cope with the need to adapt to the distinct light/dark regimens in different geographic environments and seasons and thus reduce the sensory noise for reliable irradiance detection (Hannibal, 2006; Peirson et al., 2009). Despite our speculation, whether this ipRGC buffering mechanism is used by normal retinas to respond to changes in environmental lighting conditions remains to be scrutinized.

Consistent with our findings, a recent study using *Tbr2*–GFP bacterial artificial chromosome transgenic mice showed that *Tbr2*⁺ RGCs primarily project to non-image-forming visual brain targets (Marsh et al., 2014). Our results are in line with those of all existing reports except one (eGFP-expressing pseudorabies virus was used for retrograde labeling in mouse SCN; all SCN-afferent RGCs expressed *Opn4*; Baver et al., 2008). The difference is likely attributable to the sensitivity of techniques for detecting *Opn4* expression in separate studies or other unknown mechanisms that trigger *Opn4* expression in virus-infected SCN afferent RGCs because pseudorabies virus can cause aberrant gene expression in other regions of the CNS (Taharaguchi et al., 2003; Ono et al., 2007).

The results of the present study not only demonstrate that *Tbr2* is a key regulator of the formation and survival of ipRGC but also reveal that retinal neurons have a novel compensatory response to the loss of native ipRGCs. In addition to ipRGCs, Sweeney et al. (2014) reported that several subsets of non-ipRGC exist in *Tbr2*⁺ RGCs, including CART⁺ (cocaine- and amphetamine-regulated transcript-expressing) ON–OFF direction-

selective RGCs, *Cdh3*-GFP-labeled RGCs, and *Unc5d*⁺ (*Unc-5* homolog D-expressing) RGCs. Future studies of how *Tbr2* specifies and regulates different aspects of ipRGCs and other RGC types will provide novel insights into the genetic regulatory networks of RGC subtype formation.

Whether or not the *Tbr2*⁺ RGCs are hardwired during developmental stages and serve only as a reservoir for ipRGCs, how *Tbr2*⁺ RGCs activate *Opn4* expression in distinct lighting conditions, and whether similar mechanisms are adopted by other neurons in the CNS are all pressing questions that remain to be elucidated.

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