Elevated Intraocular Pressure, Optic Nerve Atrophy, and Impaired Retinal Development in ODAG Transgenic Mice

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PURPOSE. In an earlier study, a cDNA was cloned that showed abundant expression in the eye at postnatal day (P2) but was downregulated at P10; it was named ODAG (ocular development-associated gene). Its biological function was examined by generating and analyzing transgenic mice overexpressing ODAG (ODAG Tg) in the eye and by identifying ODAG-binding proteins.

METHODS. Transgenic mice were generated by using the mouse Crx promoter. EGFP was designed to be coexpressed with transgenic ODAG, to identify transgene-expressing cells. Overexpression of ODAG was confirmed by Northern and Western blot analysis. IOP was measured with a microneedle technique. The eyes were macroscopically examined and histologically analyzed. EGFP expression was detected by confocal microscopy. Proteins associated with ODAG were isolated by pull-down assay in conjugation with mass spectrometry.

RESULTS. Macroscopically, ODAG Tg exhibited gradual protrusion of the eyeballs. The mean IOP of ODAG Tg was significantly higher than that of wild-type (WT) littermates. Histologic analysis exhibited optic nerve atrophy and impaired retinal development in the ODAG Tg eye. EGFP was expressed highly in the presumptive outer nuclear layer and weakly in the presumptive inner nuclear layer in the ODAG Tg retina. Rab6-GTPase-activating protein (Rab6-GAP) and its substrate, Rab6, were identified as ODAG-binding proteins.

CONCLUSIONS. Deregulated expression of ODAG in the eye induces elevated intraocular pressure and optic nerve atrophy and impairs retinal development, possibly by interfering with the Rab6/Rab6-GAP-mediated signaling pathway. These results provide new insights into the mechanisms regulating ocular development, and ODAG Tg would be a novel animal model for human diseases caused by ocular hypertension. (Invest Ophthalmol Vis Sci. 2009;50:242–248) DOI:10.1167/iovs.08-2206

Ocular development is a complex process, involving several genes with expression that is strictly controlled in a spatial and temporal manner. Although several genes, including Pax6, Rx, and Crx, are essential for normal ocular formation, 1–5 the molecular mechanism(s) governing eye development has not been fully elucidated.

To identify genes that are preferentially expressed in the developing eye, we performed a differential display using mRNAs extracted from postnatal day (P2) and P10 mouse eyes. 4 As a result, we isolated a cDNA fragment that had high expression in the eye at P2 but was downregulated at P10 and named it ODAG (ocular development-associated gene). 4 The ODAG gene product is composed of 266 amino acids and contains a putative zinc finger domain at the N-terminus. 4 In situ hybridization in the eye has revealed that ODAG mRNA is expressed in various regions, including the lens, ciliary body, retina, sclera, and conjunctiva, but its expression level in the retina attenuates with growth. 4 At P2, ODAG was highly expressed in all the retinal layers (presumptive outer nuclear layer [ONL], presumptive inner nuclear layer [INL], and ganglion cell layer [GCL]), but at P7, its expression decreases, especially in the GCL, and at P14, no apparent expression is detected. 4 These results strongly suggest that physiologically controlled ODAG expression in the eye would play an important role in normal ocular development, but its biological function remains unknown.

To investigate, we generated transgenic mice overexpressing ODAG (ODAG Tg). The mouse Crx promoter, which directs transgene expression in photoreceptors, 5,6 was chosen as a regulatory element, and IRES/EGFP was attached to the ODAG coding region as a reporter gene, to detect transgenic ODAG-expressing cells. Macroscopically, ODAG Tg exhibited gradual eye protrusion with increased intraocular pressure. Histologic examination of the ODAG Tg eye revealed optic nerve atrophy and retinal dysplasia. Analysis of EGFP expression demonstrated that transgenic ODAG was expressed highly in the presumptive ONL and weakly in the presumptive INL in the developing retina. In addition, by a pull-down assay using GST-ODAG conjugated with mass spectrometry, we identified Rab6-GAP and its substrate, Rab6, as ODAG-binding proteins. These results demonstrated that physiologically regulated ODAG expression is essential for normal eye development, possibly through a Rab6/Rab6-GAP-mediated pathway.
MATERIALS AND METHODS

Construction of the Transgene and Generation of Transgenic Mice
The mouse Crx promoter (~12 kb) was excised from a pCrx12K-transgenic cassette with NotI and Pmel and subcloned into the NotI-EcoRV sites of a phagemid (pBluescript; pCrx12K/pBS; Stratagene, Tokyo, Japan). The coding region of mouse ODAG cDNA was amplified with 5′ (5′-TCTTTCAAAAGTGCCCCTCA-3′) and 3′ (5′-CCGAGAAGGATTGATCCCCT-3′) primers by using mouse ODAG cDNA, as described elsewhere.4,7 A DNA fragment of the expected size was subcloned into the PCR-Blunt II-TOPO cloning vector (Invitrogen, Invitrogen, Tokyo, Japan), and correct amplification was verified by sequencing. The ODAG coding region was excised with XhoI and SpeI, blunt-ended by Klenow enzyme (Takara Shuzo, Kyoto, Japan), and subcloned into the EcoRV site of an IRES/EGFP/pA vector, which contains IRES (internal ribosomal entry site)/EGFP (enhanced green fluorescent protein) derived from pIRE2-EGFP; BD-Clontech Laboratories, Inc., Palo Alto, CA) and SV40 splicing and poly(A)(pA) signals downstream of the cloning sites. A fragment containing the ODAG coding region, IRES/EGFP, and SV40 splicing and pA signals was excised with XhoI and SacII, blunt-ended by T4 DNA polymerase (Takara Shuzo), and subcloned into the Hincll site of pCrx12K/pBS, which was located downstream of the mouse Crx promoter. A NotI-digested fragment containing the mouse Crx promoter, mouse ODAG coding region, IRES/EGFP, and SV40 splicing and pA signals was excised, purified, and microinjected into the pronuclei of fertilized eggs of BDF2 mice, as previously described.8 Founder mice and their transgenic progeny were identified by Southern blot analysis, using ODAG cDNA as a probe.8 All the mice were kept according to the guidelines of the Institute of Laboratory Animal Science, Hiroshima University, and the protocol complied with the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research.

Histologic Examination and Detection of EGFP-Expressing Cells
Histologic analysis was performed essentially as previously described.8 In brief, mouse eyeballs from anesthetized mice were fixed in 10% formalin, cut in 4-μm sections, stained with hematoxylin and eosin (HE), and examined under a light microscope. For detecting EGFP expression in the retina, retinas dissected from eyeballs were placed in PBS and were examined under a stereomicroscope (SteREO Lumar.V12 microscope; Carl Zeiss Meditec, Tokyo, Japan). To identify EGFP-expressing cells in the retinal layers, the retinas were fixed in 4% paraformaldehyde in 0.1 M phosphate buffer (pH 7.4), rinsed in phosphate buffer, and equilibrated in 30% sucrose/PBS overnight at 4°C. They were then embedded in optimal cutting temperature (OCT) compound and frozen at ~20°C. Frozen blocks were sectioned at 20-μm thickness with a cryostat (Cryostar HM 560 MV; Carl Zeiss Meditec), washed in PBS, and mounted with mounting medium. Digital images were acquired on a confocal microscope (Axiovert 100M; Carl Zeiss Meditec) and processed by graphics software (LSM 510 ver. 3.0 software; Carl Zeiss Meditec).

Intraocular Pressure Measurement
IOP was measured by using a microneedle technique.9 In brief, after mice were anesthetized with 100 μg/kg of ketamine and 9 μg/kg of xylazine, the tip of a microneedle was inserted into the anterior chamber through the cornea, and the released pressure was transferred to a transducer that measured the IOP. The mean IOPs of ODAG Tg and WT were compared by using Student’s t test.

Northern Blot Analysis
Total RNA was prepared from whole eyes (Isogen; Invitrogen), electrophoresed in 1% agarose gel containing 3% formaldehyde, transferred to a membrane (Hybond-N; GE Healthcare, Tokyo, Japan), and hybridized with 32P-dCTP-labeled ODAG cDNA or GAPDH as a probe. Positive signals were detected by a phosphoimage analyzer (Fuji X Bas2000; Fujifilm, Tokyo, Japan).

Western Blot Analysis
Western blot analysis was performed as previously described.10 In brief, cell lysates were extracted in lysis buffer (150 mM NaCl, 20 mM Tris-HCl [pH 7.4], 1% Triton-X, 0.5% deoxycholate, 1 mM phenylmethylsulfonyl fluoride, 10 mM NaF, and 20 mg/mL aprotinin). Protein aliquots were separated by SDS-PAGE, transferred to a nitrocellulose membrane (Hybond-C; GE Healthcare), blocked with 5% skim milk in TBS-T buffer (10 mM Tris-HCl [pH 7.4], 150 mM NaCl, and 0.1% Tween-20), and incubated with an anti-ODAG polyclonal antibody that was raised against a peptide, WTHVGPITATIKETVANHL, that corresponds to the C-terminal region of the mouse ODAG protein (1:1000; Kurabo Co., Ltd., Osaka, Japan) or an anti-Rab6 antibody (1:500; Santa Cruz Biotechnology, Santa Cruz, CA). The blots were developed by enhanced chemiluminescence (ECL; GE Healthcare).

GST Pull-Down Assay
To purify a recombinant ODAG protein, we inserted the coding region of ODAG cDNA into the EcoRI-XhoI sites of the pGEX4T1 vector (BD Biosciences, Piscataway, NJ) and introduced into the BL21 bacterial cells. Expression of the glutathione-Transferase (GST)-ODAG fusion protein (GST-ODAG) or GST alone was induced by addition of 1 mM isopropyl-β-thiogalactopyranoside (IPTG; Sigma-Aldrich, St. Louis, MO). Bacterial cells were lysed in PBS containing 1% Triton X-100, and the recombinant protein was collected using Sepharose 4B (GE Healthcare) and eluted with elution buffer (10 mM glutathione and 50 mM Tris-HCl [pH 8.0]). Whole-cell extracts of R28 retinal precursor cells were incubated with GST-ODAG or GST alone, and the bound proteins were collected on glutathione-Sepharose, as previously described.11

Two-Dimensional Electrophoresis and Protein Identification
Two-dimensional electrophoresis and protein identification were performed as previously described.12 In brief, proteins precipitated with GST-ODAG and GST alone were solubilized in sample buffer (8 M urea, 4% CHAPS, 0.5% dithiothreitol [DTT], and IPG [pH 3–10]). Samples were applied by ingel rehydration to a strip with immobilized pH gradient (pH 3–10, nonlinear gradient, 18 cm; GE Healthcare). First-dimensional isoelectrofocusing was performed by PIPephor (GE Healthcare). A strip was equilibrated and placed on 10% SDS-PAGE. After SDS-PAGE electrophoresis, the gels were stained with silver nitrate. Silver nitrate-stained protein spots were scanned (Epson, Nagano, Japan) and analyzed (PD-Quest software; Bio-Rad, Tokyo, Japan). Protein spots of interest were directly excised and subjected to in-gel digestion with trypsin. Spectra were obtained (MALDI-TOF-MS; Bruker Daltonics, Billerica, MA) and a database search (NCBIInr, Matrix Science Ltd., Tokyo, Japan; using the Profound protein search engine available at http://prowl.rockefeller.edu/prowl-cgi/profound.exe; developed by W. Zhang and B. T. Chait and provided in the public domain by Rockefeller University, New York, NY) was used for identification of protein names.

Generation of ODAG-EGFP Fusion Protein
To generate the ODAG-EGFP fusion construct, the mouse ODAG coding region was amplified with 5′ (5′-GAATTCCGATATCCAGTTGGCCTGAGC-3′) and 3′ (5′-GGATCCGAACTGTTGCGACTTC-3′) primers, where the bold nucleotide was substituted to match Kozak’s rule and the italic nucleotides were added to create restriction enzyme sites. The amplified fragment was digested with EcoRI and BamHI (Takara Shuzo) and subcloned into the EcoRI-BamHI sites of an expression vector, pEGFP-N1 (BD-Clontech), where the ODAG coding region and EGFP cDNA were fused in-frame and placed downstream of the cytomegalovirus promoter. Correct ampli-
fication of the ODAG coding region and its in-frame fusion to EGFP cDNA were verified by sequencing.

DNA Transfection

COS7 cells were seeded at a density of \(3 \times 10^5\) in a 60-mm tissue culture dish at least 24 hours before transfection. On the next day, when the cells reached 50% to 60% confluence, they were transfected with 2 \(\mu\)g of ODAG-EGFP expression plasmid (4 \(\mu\)L of FuGENE 6; Roche Molecular Biochemicals, Indianapolis, IN) according to the manufacturer’s instructions. Twelve to 48 hours after transfection, living cells were observed under a fluorescence microscope (BioZero BZ-8000; Keyence, Osaka, Japan).

RESULTS

Generation of ODAG Tg and Overexpression of ODAG in the Transgenic Eye

We planned to generate transgenic mice overexpressing ODAG in the eye. For this purpose, we used mouse Crx promoter that directs transgene expression in photoreceptor cells\(^5,6\) as a regulatory element and to detect transgene-expressing cells, IRES/EGFP was placed after the ODAG coding region (Fig. 1A). From this transgene, a single mRNA containing the ODAG coding region and IRES/EGFP (larger than the endogenous ODAG message) was generated by the Crx promoter, whereas two different proteins, ODAG (the same molecular weight as the endogenous one) and EGFP, were produced by the Crx promoter in combination with the IRES (Fig. 1A). Three independent transgenic lines containing approximately 5 to 10 copies of the transgene were established, and all the lines showed similar transgene expression levels and almost the same phenotype (data not shown).

To examine the overexpression of ODAG in the transgenic eye, we subjected RNAs extracted from eyes of ODAG Tg and WT littermates to Northern blot analysis with ODAG cDNA used as a probe. The results show that the transgenic eye expressed a high amount of exogenous ODAG mRNA (Fig. 1B, arrow) in addition to endogenous ODAG mRNA (Fig. 1B, arrowhead). To confirm the overexpression of ODAG at the protein level, we generated a polyclonal antibody against C-terminal peptides of ODAG and performed a Western blot analysis. ODAG protein was abundantly expressed in the transgenic eye compared with the WT eye (Fig. 1C, arrow). These results indicate that ODAG Tg overexpressed ODAG mRNA and protein in the eye.

Eye Protrusion with High IOP in ODAG Tg

At birth, the eyes of the ODAG Tg were macroscopically indistinguishable from those of WT. However, at 7 to 8 weeks of age, exophthalmos appeared in all the ODAG Tg and progressed and developed into megalocornea-like buphthalmos (Fig. 2A, arrows, top right). The surfaces of the extended corneas of ODAG Tg became turbid and opaque, and infiltration of neovessels into the cornea was occasionally observed (Fig. 2A, bottom right, arrows and an arrowhead).

We then measured IOPs in the ODAG Tg and compared them to those of the WT (7–11 weeks of age). The results indicate that the IOPs of the ODAG Tg were significantly higher than those in the WT. The IOPs of the ODAG Tg \((n = 30\) eyes) were 24.02 ± 7.35 mm Hg, with maximum and minimum IOPs being 41.78 and 8.77 mm Hg, respectively. Among them, 5 mice exhibited extremely high IOP (>30 mm Hg), 10 mice showed high IOP (>20 mm Hg), and only 1 mouse had normal IOP (<20 mm Hg). In contrast, IOPs of WT mice \((n = 30\) eyes) were 15.87 ± 1.45 mm Hg, with maximum and minimum IOPs being 19.22 and 13.57 mm Hg, respectively. The mean IOPs of both groups are shown in Figure 2B with error bars; the difference between the two groups was statistically significant \((P < 0.01)\).

Optic Nerve Atrophy and Impaired Retinal Development in ODAG Tg

To investigate neuronal abnormalities in the ODAG Tg, the optic nerve (ON) was examined. Macroscopically, the ONs of the ODAG Tg were very thin and small compared with those of the WT (Fig. 3A, top). Histologic examination showed that the ONs of ODAG Tg were hypoplastic and contained fewer retinal nerve fibers than those of the WT (Fig. 3A, bottom). In addition, optic discs of the ODAG Tg were poorly formed and difficult to detect (data not shown).

To investigate intraocular changes, we subjected the eye specimens of the ODAG Tg and WT to HE staining. Under low magnification, the ODAG Tg eye specimen exhibited thin retinal layers and extended cornea (Fig. 3B, top, black and white triangles). Examination with high magnification revealed that, although the basic structures were retained in the retina of the ODAG Tg, all the cell layers were poorly developed, irregularly arranged, and contained fewer cells than those of the WT (Fig. 3B, middle). In some of the ODAG Tg samples, only a single intraretinal nuclear layer was formed, instead of separate INL and ONL (Supplementary Fig. S1, http://www.iovs.org/cgi/content/full/50/1/242/DC1).

We then examined the anterior chamber angle to see whether the high IOPs in the ODAG Tg are caused by a closed angle. The results showed that the angle of the anterior chamber was open in the ODAG Tg as well as the WT (Fig. 3B, bottom, arrows), indicating that the ocular hypertension in ODAG Tg was not attributable to angle closure.
Expression of Transgenic ODAG in the Presumptive INL and ONL in the Developing Retina

To investigate the mechanism underlying the impaired retinal development observed in the ODAG Tg, we attempted to detect transgene-expressing cells in the retina. Since IRES/EGFP was attached to the ODAG coding region in the transgene (Fig. 1A), EGFP should have coexpressed with transgenic ODAG. To examine whether EGFP expression could be detected in the retina, we observed whole retinas of WT and ODAG Tg under a fluorescence microscope at P6, when the Crx promoter drives high transgene expression. Green fluorescence was clearly detected in the ODAG Tg retinas (Fig. 4A, right bottom), whereas no signal was observed in the WT retinas (Fig. 4A, left bottom), indicating that transgenic ODAG-expressing cells could be identified as EGFP-positive cells at this stage. To identify which cells in the retinal layers expressed EGFP, we observed retinal sections of the WT and ODAG Tg eyes under a confocal microscope. Among three retinal layers at P6 (presumptive ONL, presumptive INL, and GCL) in the ODAG Tg specimens, strong signal was detected in the presumptive ONL and weak signal was observed in the presumptive INL, whereas no positive signal was found in the GCL (Fig. 4B, bottom right), which contrasted with the WT retina, which exhibited only autofluorescence (Fig. 4B, bottom left). These results indicate that transgenic ODAG was highly expressed in the presumptive ONL, weakly expressed in the presumptive INL, but not expressed in GCL in the developing retina.

Identification of Rab6-GAP and Rab6 as Recombinant ODAG-Binding Proteins

We finally investigated molecules involved in the ODAG-mediated signaling pathway, by isolating protein(s) that forms a complex with ODAG by a GST pull-down assay in combination with proteomics. We generated a GST-fused ODAG protein (GST-ODAG) and confirmed its purity by SDS-PAGE (Fig. 5A, left lane). GST alone was used as a negative control (Fig. 5A, right lane). After these proteins were incubated with whole-retinal lysates of WT and ODAG Tg, we performed a GST pull-down assay, and the bound proteins were analyzed by SDS-PAGE. We observed a band corresponding to Rab6-GAP in the ODAG Tg sample but not in the WT sample (Fig. 5B, left lane). This result suggests that Rab6-GAP is a binding partner of ODAG in the retina.
cell extracts of R28 rat retinal precursor cells, binding proteins with GST-ODAG or GST alone were resolved by two-dimensional gel electrophoresis and the protein maps were compared. We used R28 cells for this assay, since it has been used for many ophthalmic examinations and was found to express ODAG at a high level (data not shown). In four independent experiments, protein spots reproducibly detected only in GST-ODAG pull-down samples (Fig. 5B, arrow and dotted circles) were further analyzed by mass spectrometry. These spots were identified as rat Rab6-GTPase-activating protein (Rab6-GAP, NCBI ID: XP222799.2, pI: 5.1, 94.38 kDa) with high probability (P = 9.7e−0.01; estimated z = 0.66; sequence coverage, 13%). To investigate the possibility that GST-ODAG may also bind to Rab6, the substrate of Rab6-GAP, we performed a GST pull-down/Western blot analysis. The lysates of R28 cells were incubated with GST-ODAG or GST alone, and the bound proteins were analyzed by Western blot with an antibody against Rab6. The results show that Rab6 was included in the binding proteins of GST-ODAG (Fig. 5C), indicating that not only Rab6-GAP but also Rab6 was an ODAG-associated protein.

**DISCUSSION**

Glaucoma is one of the most commonly observed disorders in the ophthalmic field, and elevated IOP is the major risk and causative factor for glaucomatous optic neuropathy. Attempts have been made to develop animal models that replicate the clinical features of the disease and several mouse models with elevated IOP have been reported. DBA/2 is a model of secondary open-angle glaucoma and its substrain DBA/2Nia is used as a secondary closed-angle glaucoma model. These models provide insights into the molecular mechanisms underlying disease development and also help in designing novel therapies for the disease.

In this study, we generated transgenic mice overexpressing ODAG in the eye and showed that the ODAG Tg exhibited gradual eye dilation with ocular hypertension, optic nerve atrophy, and impaired retinal development. The macroscopically apparent aspect was gradual eye protrusion with high ocular pressure, which resembled that of human congenital glaucoma. However, in contrast to the fact that high IOP in...
human congenital glaucoma is caused by a closed angle, pathologic analysis using HE-stained specimens and physiological examination using a gonioscope disclosed that the angle was open in ODAG Tg (Fig. 3B and not shown). The latter two abnormalities can be secondarily caused by consistent ocular hypertension. However, this was not the case in ODAG Tg, since these abnormalities already existed at 3 weeks of age, when corneal protrusion or high ocular pressure was not present (Supplementary Fig. S2, http://www.iovs.org/cgi/content/full/50/1/242/DC1).

It should be clarified that these abnormalities were primarily caused by ODAG overexpression or a secondary effect. To detect transgenic ODAG-expressing cells, we attached IRES/EGFP to ODAG coding region as a reporter gene (Fig. 1A), which would enable us to identify transgenic ODAG-expressing cells by green fluorescence. Fluorescence analysis of the retina demonstrated that EGFP expression was high in the presumptive ONL, weak in the presumptive INL, and not detectable in the GCL (Fig. 4B), which is in good accordance with the results of previous reports of the mouse Crx promoter.5,6 Since most of the layers in adult retina (OS, ONL, OPL, INL, and IPL) originated from the presumptive ONL and INL and their fibers, the impaired development of these layers would be the direct effect caused by deregulated ODAG expression. On the other hand, since ODAG was not expressed in the GCL, the reason that the GCL was also poorly developed remains unknown. Considering that ganglion cell development occurs by physically and functionally interacting with other types of cells existing in ONL and INL, such as photoreceptor cells, horizontal cells, bipolar cells, and amacrine cells,20 it would be likely that deregulated expression of ODAG in the presumptive ONL and presumptive INL secondarily affected ganglion cells growth and consequently induced poor GCL development. In addition, since the optic nerve is the axonal neurite elongated from the GCL, the optic nerve atrophy observed in ODAG Tg may also be a secondary effect of overexpressed ODAG.

The mechanism underlying gradual eye protrusion with high ocular pressure is to be clarified. As shown in Figure 3B, since the anterior chamber angle was open in ODAG Tg as well as in WT, increased IOP was not attributable to a mechanical blockage of the aqueous humor outflow, thus indicating another mechanism. One possibility is that Crx promoter-mediated ODAG expression may induce dysregulated aqueous humor production. Crx promoter has been shown to drive transgene expression in photoreceptor cells.5,6 In the embryonic stage, since photoreceptor and ciliary body cells originate from the same source, neural ectoderm, it would be possible that the Crx promoter induces ODAG overexpression in the ciliary body as well as photoreceptor cells. Since the ciliary body produces aqueous humor, it is possible that overexpressed ODAG in the ciliary body impairs regulation of aqueous humor production, with excessive aqueous humor causing ocular hypertension.

To investigate molecular mechanism(s) regulating the ODAG-mediated signaling pathway, we performed a GST-ODAG pull-down assay with mass spectrometry in retinal precursor cells and identified Rab6 and Rab6GAP as recombinant ODAG-binding proteins (Fig. 5). Rab6 belongs to the group of small GTPase family proteins that trigger various cellular processes, including cell proliferation, differentiation, intracellular trafficking, membrane ruffling, and assembly of actin stress fibers.21-25 Rab6 is localized in the Golgi complex and plays an important role in intra-Golgi transport.21,22 In one study of the frog eye, Rab6 was reported to be a part of the sorting machinery of photoreceptors that delivers newly synthesized rhodopsin from the trans-Golgi to the site of the new rod outer segment.24,25 and a subsequent study demonstrated that defects in rhodopsin transportation in Drosophila induces retinal degeneration.26 These results indicate that Rab6 plays an essential role in photoreceptor and retinal development through rhodopsin transportation.

Of interest, a recent study demonstrated that another small GTPase, Rab8, participates in vesicular transport27 and is involved in the pathogenesis of human glaucoma.28 Optineurin, whose mutations are found in a subset of patients with glaucoma,29 is physically associated and functionally interacted with Rab8. On apoptotic stimuli, optineurin moves from the Golgi complex to the nucleus, and this subcellular translocation is dependent on the GTPase activity of Rab8. In addition, Rab6 and Rab8 were reported to function sequentially in rhodopsin transport and synergistically participate in rod outer segment disc morphogenesis.24 These results suggest that aberrant expression of ODAG might impair Rab6/Rab6-GAP function, which eventually perturbs ocular homeostasis and consequently induces various abnormalities observed in ODAG Tg. Further studies are necessary to clarify the mechanism(s) of whether and how overexpressed ODAG affects functional properties of small GTPases, including Rab6.

The primary structure of mouse ODAG showed a zinc-finger domain at the N terminus, suggesting that ODAG may function as a transcription factor.4 To address this possibility, we generated ODAG-EGFP fusion protein and investigated its subcellular localization. We found that ODAG-EGFP was mainly localized in the nucleus and partly in the cytoplasm (Supplementary Fig. S3, http://www.iovs.org/cgi/content/full/50/1/242/DC1), suggesting that ODAG may play a role as a nuclear protein as well as a Rab6/Rab6-GAP-interacting molecule, and possibly transport signals from the cytoplasm to the nucleus. Human counterpart of ODAG, which has a 92% amino acid identity with mouse ODAG, was reported and deposited as “GATA zinc finger domain containing 1 (GATAD1)”.30 Therefore, it would be intriguing to examine whether human ODAG/GATAD1 may be involved in human disease with ocular hypertension.

In this study, aberrant expression of ODAG in the eye induced ocular hypertension, optic nerve atrophy, and impaired retinal development, and Rab6-GAP and its substrate, Rab6, were found to be ODAG-associated proteins. These results indicate that physiologically controlled ODAG expression is essential for normal ocular development, possibly through a Rab6/Rab6-GAP-mediated signaling pathway. Our findings provide new insight into the mechanisms regulating ocular homeostasis and ODAG Tg would be a novel animal model for diseases that are caused by ocular hypertension.

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