The nob2 mouse, a null mutation in Cacna1f: Anatomical and functional abnormalities in the outer retina and their consequences on ganglion cell visual responses

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(Received August 23, 2005; Accepted November 9, 2005)

Abstract

Glutamate release from photoreceptor terminals is controlled by voltage-dependent calcium channels (VDCCs). In humans, mutations in the Cacna1f gene, encoding the α1F subunit of VDCCs, underlie the incomplete form of X-linked congenital stationary night blindness (CSNB2). These mutations impair synaptic transmission from rod and cone photoreceptors to bipolar cells. Here, we report anatomical and functional characterizations of the retina in the nob2 mouse, a naturally occurring mutant caused by a null mutation in Cacna1f. Not surprisingly, the b-waves of both the light- and dark-adapted electroretinogram are abnormal in nob2 mice. The outer plexiform layer (OPL) is disorganized, with extension of ectopic neurites through the outer nuclear layer that originate from rod bipolar and horizontal cells, but not from hyperpolarizing bipolar cells. These ectopic neurites continue to express mGluR6, which is frequently associated with profiles that label with the presynaptic marker Ribeye, indicating potential points of ectopic synapse formation. However, the morphology of the presynaptic Ribeye-positive profiles is abnormal. While cone pedicles are present their morphology also appears compromised. Characterizations of visual responses in retinal ganglion cells in vivo, under photopic conditions, demonstrate that ON-center cells have a reduced dynamic range, although their basic center-surround organization is retained; no alteration in the responses of OFF-center cells was evident. These results indicate that nob2 mice are a valuable model in which to explore the pathophysiological mechanisms associated with Cacna1f mutations causing CSNB2, and the subsequent effects on visual information processing. Further, the nob2 mouse represents a model system in which to define the signals that guide synapse formation and/or maintenance in the OPL.

Keywords: Bipolar and horizontal cells, Congenital stationary night blindness, Electroretinogram, ON- and OFF-pathways, Voltage-dependent calcium channel

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Introduction

Rod and cone photoreceptors synapse with second-order retinal neurons in a specialized plexus known as the outer plexiform layer (OPL). In darkness, rods and cones release the neurotransmitter, glutamate, at a continuous rate (Copenhagen & Jahr, 1989; Massey, 1990). Light induces hyperpolarization of photoreceptors, slows calcium influx through voltage-dependent calcium channels (VDCCs), thereby reducing the rate of glutamate release (Schmitz & Witkovsky, 1997). Disruptions in this process reduce the efficacy of synaptic transfer of the light response from photoreceptors to the second-order retinal neurons, the bipolar and horizontal cells. This can be evaluated noninvasively with the electroretinogram (ERG). When only synaptic transmission is compromised, the α-wave component, reflecting activity of rod and cone photoreceptors (Lamb, 1996), is spared, while the β-wave, reflecting predominantly rod or cone depolarizing bipolar cell (DBC) activity (Robson & Frishman, 1995, 1998; Kofuji et al., 2000; Sharma et al., 2005), is reduced. This “negative” ERG pattern has been reported in mice with mutations in genes encoding mGluR6 (Masu et al., 1995), nystatin (Pardue et al., 1998), the G-protein subunit Ga α1 (Dhingra et al., 2000, 2002), the β2 subunit of VDCCs (Ball et al., 2002), the presynaptic cytomatrix protein bassoon (Dick et al., 2003), or calcium-binding protein 4 (Hae-seeler et al., 2004). The wide variety of proteins essential for normal synaptic transmission is suggestive of the intricacy of the synaptic complex linking photoreceptors and DCBs.

In humans, negative ERGs have been observed in two forms of X-linked congenital stationary night blindness (CSNB) (Cacna1f et al., 1986). X-linked congenital stationary night blindness (CSNB) is a recessive, X-linked trait. For the Jackson Laboratory, thenob2 mouse was discovered as part of an ERG-based screen for spontaneous retinal mutants (Chang et al., 2002). All other recombinant inbred strains from the panels (AXB, BXA) had normal α-waves, indicating that the defect in nob2 was the result of a rare spontaneous mutation in the AXB6 line. To determine the chromosomal location of the nob2 gene, affected AXB6 mice were crossed to CAST/EiJ mice. When male nob2 mice were used, all F1 mice were unaffected, consistent with a recessive inheritance pattern. When female nob2 mice were mated to CAST/EiJ males, all F1 female offspring had normal ERGs, while all F1 male offspring were affected, indicating that nob2 was an X-linked trait. For genetic mapping, F1 nob2/CAST/EiJ females were crossed to CAST/EiJ males. The phenotype of a total of 200 F2 male mice was determined by ERG recording and these mice were genotyped using microsatellite markers across the X chromosome.

Materials and methods

Animals

Wild-type (WT) and nob2 mice used in this study were obtained from The Jackson Laboratory (Bar Harbor, ME), either directly or as the progeny from breeders shipped to the institutions involved.

In some experiments, we also studied nob mice, which have a mutation in the nyx gene and are a model of complete CSNB (CSNB1; Pardue et al., 1998; Gregg et al., 2003). nob mice were generated in a breeding colony maintained at the Cleveland VA Medical Center. In all facilities, mice were maintained on a 12-h light/12-h dark cycle and were provided food and water ad libitum. All of the experiments that assessed the visual response properties of RGCs were conducted at the University of Louisville. All procedures involving animals were approved by the individual institute’s Institutional Animal Care and Use Committee and were conducted in accordance with the Institute for Laboratory Animal Research Guide for Care and Use of Laboratory Animals.

Positional cloning

The nob2 mouse was initially identified in a recombinant inbred strain, AXB6, established between A/J and C57BL/6J strains (AXB6/PgnJ) and maintained at the Jackson Laboratory. nob2 was discovered as part of an ERG-based screen for spontaneous retinal mutants (Chang et al., 2002). All other recombinant inbred strains from the panels (AXB, BXA) had normal α-waves, indicating that the defect in nob2 was the result of a rare spontaneous mutation in the AXB6 line. To determine the chromosomal location of the nob2 gene, affected AXB6 mice were crossed to CAST/EiJ mice. When male nob2 mice were used, all F1 mice were unaffected, consistent with a recessive inheritance pattern. When female nob2 mice were mated to CAST/EiJ males, all F1 female offspring had normal ERGs, while all F1 male offspring were affected, indicating that nob2 was an X-linked trait. For genetic mapping, F1 nob2/CAST/EiJ females were crossed to CAST/EiJ males. The phenotype of a total of 200 F2 male mice was determined by ERG recording and these mice were genotyped using microsatellite markers across the X chromosome.

Polymerase chain reaction (PCR) genotyping

DNA was isolated from tail snips as previously reported (Buffone & Darlington, 1985). For PCR amplification, 25 ng DNA was used in a 10-µl volume containing 50 mM KCl, 10 mM Tris-HCl, pH 8.3, 2.5 mM MgCl2, 0.2 mM oligonucleotides, 200 µM dNTPs, and 0.02 U AmpliTaq DNA polymerase. The reactions were initially denatured for 3 min at 94°C, and then subjected to 40 cycles of 15 s at 94°C, 1 min at 51°C, 1 min at 72°C, and a final 7-min extension at 72°C. PCR products were separated by electrophoresis on 3% MetaPhor (FMC, Rockland, ME) agarose gels, stained with ethidium bromide, and visualized under UV light. Initially a scan of the X chromosome using microsatellite (Mit) DNA markers was carried out on pooled DNA samples (Taylor et al., 1994). After detection of linkage on the X chromosome, microsatellite markers were scored on individual DNA samples. To test the Cacna1f gene as a candidate, four pairs of PCR primers were used to amplify the entire open reading frame of the gene from cDNA. PCR products were sequenced directly using standard fluorescent-based sequencing methods. Total RNA was isolated from retinas of newborn mice using TRIZOL LS Reagent (Invitrogen Life Technologies, Carlsbad, CA) and the SuperScript™ premultiplication system (Invitrogen Life Technologies) was used to synthesize first strand cDNA. The genomic DNA fragment containing the transposon insertion in exon 2 that is illustrated in Fig. 1a and flanking DNA was amplified using the Accuprime DNA polymerase as described by the manufacturer (Invitrogen Life Technologies).
Life Technologies). The PCR fragment containing the transposon insertion was sequenced by fluorescent cycle sequencing.

**Histology**

Mice were killed by CO₂ administration and the eyes were removed and immersion fixed overnight. For light microscopy the eyes were fixed at 4°C in Bouin’s fixative, rinsed in phosphate-buffered saline, dehydrated, and embedded in paraffin. Sections 6-μm thick were cut on a RM 2135 microtome (Leica Microsystems, Wetzlar, Germany), mounted onto slides, rehydrated, and stained using hematoxylin and eosin.

**Preparation of retinal sections for immunohistochemistry**

Mouse eyecups or isolated retinas were prepared as described previously (McCall et al., 2002). Freshly dissected eyecups or retinas were fixed for 5–60 min by immersion in 4% paraformaldehyde in 0.1 M phosphate buffer, pH 7.4 and stored at −80°C until used.

**Immunofluorescent staining**

Sections were brought to room temperature, and incubated in a blocking solution [5% (v/v) normal horse serum, 0.5% (v/v) Triton X-100 in PB] for 30 min. This solution was replaced and sections were incubated with primary antibody diluted in blocking solution for either 1–2 h at room temperature or overnight at 4°C. Primary antibodies and dilutions were as follows: monoclonal anti-αRF (4G2; Morgans et al., 2005, at 1:2); anti-Ctbp2/Ribeye (BD Biosciences Pharmingen, San Diego, CA, at 1:5000); anti-Protein Kinase Cα (PKC; Sigma, St Louis, MO, at 1:5000); anti-calbindin D-28K (Chemicon, Temecula, CA, at 1:1000); anti-Neurokinin-3 receptor (NK3R; Grady et al., 1996, at 1:500); rhodamine-labeled PNA (Vector Labs, Burlingame, CA, at 1 μg/ml); and anti-mGluR6 (NeuroMics, Minneapolis, MN, at 1:1000). After washing in PB three times for 20 min, sections were incubated in fluorescent secondary antibodies at room temperature for 1 h. Secondary antibodies were: 1:500 dilution of CY3 conjugated goat anti-mouse (Jackson ImmunoResearch, West Grove, PA) for αRF; 1:1000 dilution of Alexa 488 conjugated goat anti-mouse (Ribeye); and 1:1000 dilution of Alexa 488 or Alexa 568 goat anti-rabbit (mGluR6, PKC, calbindin, NK3R; Invitrogen Life Technologies, Carlsbad, CA). Slides were washed three times in PB and then coverslipped with Crystal/Mount mounting medium (Biomedica, Foster City, CA). Immunostained sections were imaged on either a Zeiss LSM 510 META or an Olympus FV300 confocal microscope. Images presented are all maximum projections of stacks of confocal sections, adjusted for contrast and brightness with Adobe Photoshop.

**Electroretinography**

After overnight dark adaptation, mice were anesthetized with ketamine (80 mg/kg) and xylazine (16 mg/kg). Eye drops were used to anesthetize the cornea (1% proparacaine HCl) and to dilate the pupil (1% mydriacyl, 2.5% phenylephrine HCl, 1% cyclopentolate HCl). Mice were placed on a temperature-regulated heating pad throughout the recording session.

ERGs were recorded using a stainless-steel electrode that was coiled in a small loop and made contact with the corneal surface through a thin layer of 0.7% methylcellulose. Needle electrodes placed in the cheek and the tail served as reference and ground leads, respectively. Responses were differentially amplified (0.05–1500 Hz), averaged, and stored using a UTAS E-3000 signal averaging system (LKC Technologies, Gaithersburg, MD). Stimulus flashes were first presented to the dark-adapted eye, and ranged in intensity from −3.6 log cd s/m² to 2.1 log cd s/m². Stimuli were presented in order of increasing intensity. The number of successive responses averaged together decreased from 20 for low-intensity flashes to 2 for the highest intensity stimuli. The duration of the interstimulus interval (ISI) increased from 4 s for low-intensity flashes to 90 s for the highest intensity stimuli. A steady rod-desensitizing adapting field was then presented within the ganzfeld bowl. After allowing a 7-min period of light adaptation, cone ERGs were recorded to flashes superimposed upon an adapting field. Flash intensity ranged from −0.8 log cd s/m² to 1.9 log cd s/m² and responses to 100 flashes presented at 2.1 Hz were averaged at each intensity level.

The major components of the ERG were measured conventionally. The amplitude of the a-wave was measured from the prestimulus baseline to the a-wave trough. The amplitude of the b-wave was measured from the a-wave trough to the peak of the b-wave or, if no a-wave was present, from the prestimulus baseline. The implicit time of the b-wave was measured from the time of flash presentation to the peak of the b-wave. To derive Rmp, representing the maximum a-wave amplitude, the leading edge of the dark-adapted a-wave was fitted with a model of rod phototransduction (Pugh & Lamb, 1993), as described elsewhere (Nusinowitz et al., 2003).

**Extracellular in vivo single-unit retinal ganglion cell recording**

Visual responses were recorded extracellularly from the optic nerve of WT and nob2 mice, *in vivo*, under photopic conditions, using previously published procedures (Sagdullayev & McCall, 2005). Mice were anesthetized with ketamine (127 mg/ml) and xylazine (12 mg/ml), and anesthesia was maintained throughout the course of the experiment with hourly supplemental injections of the original dose diluted to 42%. Mice were placed in a stereotaxic (David Kopf Instruments, Tujunga, CA) and body temperature was maintained at 37°C with a thermostat-controlled heating pad (CWE Instruments, Ardmore, PA). Eyedrops (1% mydriacyl, 2.5% phenylephrine HCl) were used to dilate the right pupil and to paralyze accommodation and a clear, plano, plastic contact lens (Sagdullayev et al., 2004) was placed over each eye to prevent corneal drying. A small craniotomy was performed and the cortex overlying the optic nerve was removed using vacuum suction. Tungsten electrodes (40–70 MΩ impedance) were inserted into the optic nerve and a reference electrode was inserted into the skin on the back of the neck. Action potentials from single RGC axons were isolated, amplified (X3+Cell, slope/height window discriminator/amplifier, FHC, Bowdoinham, ME) and digitized at 21 kHz (Power1401, Cambridge Electronics Design, Cambridge, UK). Ganglion cell responses were played in real time over an audiometer (AM7, Grass Instruments, Quincy, MA) and displayed on both an oscilloscope (Tektronix 2215, Richardson, TX) and a computer monitor, and stored for future analysis using Spike2 software (Cambridge Electronic Design, Cambridge, UK).
Table 1. Stimulus intensities and interstimulus intervals used for RGC intensity–response functions

<table>
<thead>
<tr>
<th>Intestimulus interval (s)</th>
<th>Intensity (cd/m²)</th>
<th>ON-center cells</th>
<th>Intensity (cd/m²)</th>
<th>OFF-center cells</th>
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<td></td>
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<tr>
<td>60</td>
<td>789</td>
<td>0.088</td>
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<td>1860</td>
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The receptive-field center of each isolated RGC was mapped onto a tangent screen positioned between 10–20 cm from the mouse’s eye using a handheld Pantoscope (Keeler Instruments, Broomall, PA). Once the sign and extent of the receptive-field center was determined, a Ganzfeld stimulator (Advanced Illumination, Rochester, VT), which contained 64 light emitting diodes and whose size could be adjusted with an iris diaphragm (Edmund Industrial Optics, Barrington, NJ), replaced the tangent screen and the aperture was adjusted to be slightly smaller than the size of the cell’s receptive-field center. A computer program, Superscope II 2.17.1 (GW Instruments Inc., Somerville, MA) was used to synchronize stimulus presentation and response acquisition (Power1401, Cambridge Electronic Design).

After a 2-min adaptation period to a 30-cd/m² spot, each RGC was stimulated with a spot of optimal size and fixed duration (4 s) whose intensity changed. For ON-center cells, the spot intensity increased in 0.25–0.5 log intensity steps from the adaptation level to 1680 cd/m². For OFF-center cells the intensity decreased similarly from the adaptation level to 0 cd/m². Between the stimuli the screen returned to 30 cd/m². The duration of the ISI needed to maintain responses independent of the preceding stimulus was empirically determined and ranged from 10 s for the smallest intensity steps to 60 s for the largest intensity steps. Table 1 shows the exact intensities and ISIs for both increments and decrements. To account for differences in the total number of spikes that occur between transient and sustained RGC responses, the mean spike rate over the first 0.5 s after stimulus onset was used as a metric of the cell’s response. For each RGC studied, the background firing rate was measured as the mean firing rate over 10 s after a 2-min adaptation period and prior to the presentation of the stimuli. This background rate was subtracted from each of the visually evoked responses. To compare across cells with different peak firing rates, responses were normalized relative to the maximum response for that cell. Responses were plotted against log (normalized intensity) for ON-center cells and –log (normalized intensity) for OFF-center cells, where normalized intensity is the ratio of stimulus intensity to the intensity of the adapting spot (30 cd/m²). A Hill equation was fit to each data set (GraphPad Prism 4.03, San Diego, CA), and the semisaturation constant (log E₅₀) and the Hill exponent (h) parameters were derived.

Results

nob2 is caused by a mutation in the Cacna1f gene

The nob2 mutation was mapped between the microsatellite markers, DXMit123 and DXMit124, a region that includes Cacna1f. Sequence analysis revealed an insertion of a transposable element (Mus musculus transposon ETn) in exon 2 of the Cacna1f gene (Fig. 1a). This insertion results in an out-of-frame insertion, which is predicted to produce a stop codon after synthesis of only 32 amino acids. In view of these results, nob2 mice have been renamed Cacna1f⁻nop2. In this manuscript, however, affected mice will continue to be referred to simply as nob2, regardless of their gender (i.e., Cacna1f⁻nop2/Cacna1f⁻nop2 or Cacna1f⁻nop2/Y). To examine whether the α₁β subunit protein was expressed in the nob2 retina, we reacted retinal cross sections with an antibody against the C-terminal 177 amino acids of rat α₁β, which recognizes α₁β in the OPL and inner plexiform layer (IPL) of WT mice (Morgans et al., 2005). As can be seen in Fig. 1b, there is no expression of α₁β in either synaptic layer in the nob2 retina, which confirms that this spontaneous mutation results in a null allele. As can be seen by comparing the hematoxylin and eosin-stained transverse retinal sections from WT and nob2 (Fig. 1c), adult nob2 mice exhibit a disorganized IPL. In contrast, the lamination pattern of the rest of the nob2 retina appears normal, despite the absence of expression of the α₁β subunit in the IPL.

ERGs are abnormal in nob2 mice

To evaluate overall retinal function, ERGs were recorded to full-field flashes using stimulus conditions that isolated rod- or cone-driven activity (Pardue et al., 1998). The b-wave component, reflecting activity of rod DBCs (Kofuji et al., 2000), was reduced in dark-adapted ERGs from nob2 mice (red traces, Fig. 2a) compared to those obtained from B6AF1/J WT controls (black traces, Fig. 2a) throughout the stimulus range. The b-wave was not abolished, however, as nob2 responses were consistently more positive than those obtained from all mice (blue traces, Fig. 2a) that completely lack a dark-adapted b-wave (Pardue et al., 1998). The nob2 responses also included high-frequency oscillatory potentials superimposed upon the b-wave. While diminished in comparison to WT, the presence of oscillatory potentials in nob2 ERGs provides additional evidence that light-evoked activity reaches the inner retina (Wachtmeister, 1998; McCullough et al., 2002). The maximum amplitude of the initial cornea-negative a-wave, reflecting the light-induced closure of sodium channels along the rod outer segment (Lamb, 1996), was similar in all mice tested (Figs. 2a & 2b), which is consistent with the absence of photoreceptor degeneration in these mice (Fig. 1c; Pardue et al., 1998). Between −1.2 and 0.0 log cd s/m², however, a-waves were actually larger in nob2 and nob mice than in B6Af1/J WT mice. This apparent enhancement occurs because normally the b-wave component truncates the a-wave (Hood & Birch, 1992). In nob2 mice, the implicit time of the dark-adapted ERG b-wave was delayed (Fig. 2c).

CSNB in humans is a nonprogressive disorder (Greenstein et al., 1988). Thus, we examined whether the dark-adapted ERG abnormalities of nob2 mice also were stable and recorded ERGs at 3, 7, and 15 months of age. To compensate for the reduction in overall amplitude that occurs in older WT mice (Li et al., 2001), values are expressed as a percentage of the mean for age-matched WT mice recorded under the same conditions. At each age tested, the maximum a-wave amplitude, represented by the Rm₁₀₀ values, from nob2 mice were not different from those of WT animals (Fig. 2d). In addition, the relative amplitude of the nob2 b-wave was diminished to the same extent at all ages examined (Fig. 2d).

The nob2 mutation also affects the cone pathway. Light-adapted ERGs from nob2 mice (red tracings in Fig. 3a) were consistently reduced in amplitude and had slower kinetics than
those of B6AF1/J WT controls (black traces in Fig. 3a). In comparison to nob responses (blue traces in Fig. 3a), however, nob2 responses were more positive and had faster kinetics. Although cone ERGs of nob2 and nob mice had distinct waveforms, their overall peak-to-trough amplitudes were similar (Fig. 3b). The implicit time of the cone ERG b-wave was delayed in both nob2 and nob mice, although the magnitude of this delay was less pronounced in nob2 responses (Fig. 3c). These data predict that deficits should be expected within the ON-pathway, mediated by either rods or cones. Unfortunately, the rodent ERG cannot easily

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**Fig. 1.** (a) Diagram of Cacna1f, indicating the location of the intragenic insertion into exon 2. Exon 2 nucleotides are shown as capitals. The 6-bp duplication at the insertion site is underlined. (b) Immunofluorescent staining for α1F is localized to the OPL and the IPL of the WT retina but is absent in the nob2 retina. (c) Comparison of retinal anatomy in adult WT and nob2 mice. In the nob2 retina, the OPL is disorganized but other retinal layers appear normal. GCL, ganglion cell layer; IPL, inner plexiform layer INL, inner nuclear layer; OPL, outer plexiform layer; and ONL, outer nuclear layer. Scale bar indicates 20 μm (b) or 40 μm (c).
be used to predict whether this mutation affects the OFF-pathway (Sharma et al., 2005).

Anatomical abnormalities in nob2 mice

Since the ERG b-wave abnormalities of nob2 mice are likely to reflect a synaptic defect between photoreceptors and DBCs, we examined the morphology of dendrites of these postsynaptic cells. On the postsynaptic side, DBC and horizontal cell dendrites form an invagination in each rod spherule or multiple invaginations in each cone pedicle. These invaginations are associated with presynaptic structures referred to as synaptic ribbons, which are thought to facilitate the constant release of glutamate that occurs in the dark (von Gersdorff, 2001). Because of the disorganization of the nob2 OPL that we observed at the light-microscope level, we also examined the localization of specific elements in the presynaptic terminals using immunohistochemistry and confocal microscopy. To examine the morphology of the entire rod bipolar cell, we
reacted retinal sections with antibodies to PKCa (Fig. 4a). Similar to previous reports (GREFERATH ET AL., 1990; HAVERKAMP & WÄSSELE, 2000), dendrites of WT rod bipolar cells terminate in short tufts uniformly in the OPL. In comparison, in nob2 retina the rod bipolar cell dendrites extend beyond the OPL into the outer nuclear layer (ONL) and, in many cases reach the retinal pigment epithelium (Fig. 4b). To examine the structure of horizontal cells we reacted retinal sections with an antibody to calbindin (PEICHL & GONZALEZ-SORIANO, 1994; HAVERKAMP & WÄSSELE, 2000). In WT retinas, calbindin-positive horizontal cell processes define a dense layer within the OPL (Fig 4c). In the nob2 retina, the horizontal cell dendrites, like those of bipolar cells, extend across the ONL to the retinal pigment epithelium (Fig. 4d). Careful examination of confocal images indicated that every rod bipolar and horizontal cell showed these ectopic dendrites. By comparison the inner retina of the nob2 mouse appears qualitatively normal. The cell bodies of the rod bipolar and calbindin-positive amacrine cells are appropriately located in the inner nuclear layer (INL) of the nob2 retina and their axon terminals and processes laminate normally within the IPL. To assess the status of a subset of cone hyperpolarizing bipolar cells (HBCs) in the nob2 retina, we used an antibody against NK3R (HIRANO & BRECHA, 2002; HAVERKAMP ET AL., 2003). The dendritic arbors of this class of HBCs in the nob2 mice are indistinguishable from WT (Figs. 4e & 4f). Thus, it appears that the dendrites of the NK3R-positive HBCs are unaffected by the lack the α1F in photoreceptors.

To better define alterations in the relationship between photoreceptors and second-order neurons, we examined the expression pattern of one postsynaptic (mGluR6) and two presynaptic markers [Ribeye and peanut agglutinin (PNA)] in transverse retinal sections. Ribeye (also referred to as Ctbp) is a component of the presynaptic ribbon complex and is present in both rod and cone

Fig. 3. Light-adapted electroretinograms. (a) Comparison of ERGs recorded from nob2 mice (red tracings) with those obtained from WT (black tracings) or nob (blue tracings) mice. Numbers indicate flash intensity in log cd s/m². Amplitude (b) and implicit time (c) of the cone ERG b-wave for nob2 (red circles), nob (blue circles) and WT (black circles) mice. Data points in (b & c) indicate the mean ± S.E.M. response for four mice tested at 6 weeks of age.
punctate staining both in the residual OPL and throughout the ONL, where the bipolar and horizontal cells extend their ectopic processes. Cone terminals, as shown by PNA labeling, are restricted to the OPL in WT retina (Fig. 5a) and also to the residual OPL of the nob2 retina (Fig. 5b). In the nob2 retina, however, the pattern of PNA staining is disorganized (Fig. 5b), an effect that may be secondary to the gross disorganization of the OPL.

Figs. 5c–5f compare the expression patterns for Ribeye and PNA with that of the postsynaptic marker, mGluR6. In WT mice (Fig. 5c), mGluR6 puncta (red) align with the horseshoe-shaped structures defined by Ribeye (green) expression. In the nob2 retina (Fig. 5d), however, the morphology of the Ribeye-positive profiles is altered and the overall density of structures stained for Ribeye and mGluR6 are reduced. While there remain cases in which a close apposition of staining between Ribeye and mGluR6 in nob2 mice is maintained, this association is reduced compared to WT retina. It remains to be determined whether these associations represent ectopic synapses, as has been observed in bassoon knockout (KO) mice (Dick et al., 2003).

To determine if the dendrites of the cone DBCs colocalize with cone terminal profiles, we double labeled sections with an antibody to mGluR6 and with PNA. Fig. 5e shows a representative section through the OPL of a WT mouse retina. Both mGluR6 (red) and PNA (green) colocalize at the cone terminals (yellow); the additional punctate staining of mGluR6 represents rod bipolar cell dendritic tips. In the section from the nob2 mouse, while cone terminals appear less organized, mGluR6-positive profiles continue to colocalize with cone terminals (Fig. 5f, yellow). Thus, both the ERG and morphological analyses point to a major disruption in the signaling patterns subserved by DBCs. In contrast, our morphological data suggest little impact on the cone-to-HBC pathway.

**ON-center RGCs visual responses are altered in nob2 mice**

Abnormalities in both rod- and cone-mediated vision have been documented in patients with CSNB1 and CSNB2 (Miyake et al., 1986), although quantitative characterizations of either central or peripheral vision have yet to be published. Our ERG results indicate that ON-pathway signaling through both rod and cone pathways should be compromised, while OFF cone pathway signaling might be spared. To compare the outcome of this potential alteration in signaling through the independent ON and OFF cone pathways, we examined basic receptive-field structure and visual response properties of RGCs in nob2 mice under light-adapted conditions. Responses of 24 RGCs from nob2 mice and 27 RGCs from WT mice were characterized by their responses to spots of light. Regardless of genotype, all RGCs had a discrete receptive-field center that could be localized to the ON-center or OFF-center. Some RGCs were characterized by their responses to spots of light. Regardless of genotype, all RGCs had a discrete receptive-field center that could be localized and mapped subjectively with hand-mapping techniques and there were no gross abnormalities evident between WT and nob2 cells or when ON- and OFF-center cells were compared separately.

We determined both the spontaneous activity to a photopic adapting stimulus (30 cd/m²) and the evoked activity to stimulus increments or decrements from that adaptation level for the same ON- and OFF-center RGCs in both WT and nob2 mice. When we compared the spontaneous activity for ON-center cells, we found a significant difference between nob2 and WT mice. The spontaneous activity of ON-center cells in nob2 mice was less than half of that of WT mice.

**Fig. 4.** Bipolar and horizontal cells extend dendritic processes into the ONL in the nob2 but not WT retina. Retinal sections from WT (a, c, & e) or nob2 (b, d, & f) retinas immunolabeled with antibodies against the rod bipolar cell marker, PKCa (a & b), the horizontal cell marker, calbindin (c & d), or the HBC marker NK3R (e & f). INL, inner nuclear layer; OPL, outer plexiform layer; and ONL, outer nuclear layer. DIC images were obtained from a WT retina. Scale bar indicates 20 μm. Mice were studied at 5 months of age.
that in WT mice (14.8 ± 1.5 spikes/s vs. 31.6 ± 5.8 spikes/s; Student’s t-test; P = 0.017). In contrast, spontaneous rates for OFF-center cells were similar between the two groups (nob2 vs. WT, 9.8 ± 1.6 spikes/s vs. 6.0 ± 1.7 spikes/s; Student’s t-test; P = 0.13). These results suggest that steady state excitation through the ON-pathway is decreased in nob2 mice compared to WT, whereas the cone-driven OFF-pathway remains unaffected. This is consistent with the reduced amplitude of the cone ERG b-wave throughout the range of intensities tested. To examine whether the range of intensities over which RGC responses could be evoked was also altered, we measured the responses of ON-center cells to optimally sized spots over a range of incremental stimuli increasing from the adapting level (30 cd/m²) up to a maximum of 1680 cd/m². Similarly, responses of OFF-center cells were measured to decrements from the adapting level. Responses from a representative nob2 ON-center RGC are shown in Fig. 6a. The normalized responses of the cells were plotted as a function of intensity to produce intensity–response functions. Mean response functions for all nob2 (n = 13) and WT (n = 16) ON-center cells are compared in Fig. 6b. There is a significant decrease in the range of intensities over which ON-center RGCs in nob2 mice can encode intensity changes. When a Hill equation was fit to the individual data sets, the intensity needed to reach a half-maximum response (log EI₅₀) was almost three times higher in WT than in nob2 cells (log EI₅₀ =

Fig. 5. Synaptic abnormalities in nob2 retina. (a–f) micrographs of vertical sections through WT (a, c, & e) or nob2 (b, d, & f) retinas immunolabeled with PNA (red)/Ribeye (green; [a & b]), mGluR6 (red)/Ribeye (green; [c & d]) or mGluR6 (red)/PNA (green; [e & f]). Scale bar indicates 10 μm (a, b) or 5 μm (c–f). Mice were studied at 5 months of age.
0.86 ± 0.14 vs. 0.42 ± 0.03; ANOVA; $P = 0.004$). In addition, nob2 ON-center cell responses saturated well before our highest intensity spot, an effect that we rarely observed in WT ON-center cells. These results indicate that the dynamic range and thus the ability of nob2 ON-center cells to encode increments in intensity is dramatically reduced and predict that their increment thresholds would be significantly altered. In contrast, the intensity–response functions of OFF-center cells were similar in WT and nob2 animals (log $E_{100} = 0.47 ± 0.04$ vs. 0.39 ± 0.03; ANOVA; $P > 0.05$; Fig. 6c). This differential affect on the ON-pathway of the nob2 retina is consistent with the alterations seen in the ERG $b$-wave and in the propensity for second-order retinal neurons to develop ectopic neurites.

**Discussion**

We describe anatomical and functional changes in the retina of a mouse with a spontaneous mutation in the *Cacna1f* gene, which encodes the $\alpha_{1F}$ subunit (Cav1.4) of the VDCC. We show that the mutation is caused by the insertion of a transposon that results in a null allele and the absence of expression of this VDCC subunit protein in the retina. Similar to its manifestations in the human disease, ERG $b$-waves are significantly reduced under both dark- and light-adapted conditions, while $a$-waves are unaffected. This indicates that the primary defect in both mouse and human resides in altered synaptic transmission from both rods and cones to DBCs. In addition to this defective signal transmission, the dendrites of the rod DBCs and horizontal cells in nob2 mouse retinas extend ectopic processes throughout the outer retina, which leads to the collapse of the OPL. In contrast, at least one class of HBCs are unaffected. In spite of these significant changes in the structure and function of the outer retina and an absence of expression of the $\alpha_{1F}$ subunit in the inner retina, visual responses and receptive fields of RGCs in the light-adapted nob2 retinas exhibit relatively minor changes and those are limited to the ON-center RGCs.

**The role of the $\alpha_{1F}$ subunit of the VDCCs in retinal synaptic transmission**

Expression studies in heterologous systems show that $\alpha_{1F}$ combines with $\beta$ and $\alpha_{2}/\delta$ subunits to form functional L-type VDCCs (Koschak et al., 2003; Baumann et al., 2004; McRory et al., 2004; Hoda et al., 2005; Morgans et al., 2005). In the outer retina, the $\alpha_{1F}$ subunit of the VDCC is expressed in the OPL, and it has been localized specifically to both rod and cone photoreceptor terminals (Morgans et al., 2005). These data, and the fact that the $\alpha_{1F}$ subunit is critical to producing a VDCC on photoreceptor terminals that does not inactivate during sustained depolarization (Taylor & Morgans, 1998), indicate that this subunit may be the primary VDCC $\alpha_{1}$ subunit involved in mediating tonic glutamate release from rods and cones. The results of the current study are concordant with this hypothesis. Under dark-adapted conditions, nob2 mice exhibit a marked reduction in the ERG $b$-wave, while the $a$-wave component is spared. This indicates that in the absence of the $\alpha_{1F}$ subunit, photoreceptors are able to hyperpolarize in response to light but that there is a significant loss of transmission.

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Fig. 6. Ganglion cell responses in nob2 mice. Representative raw traces from an ON-center RGC in a nob2 mouse are shown in (a) and were obtained by recording from the optic nerve in vivo. The cell’s response to the 30 cd/m$^2$ adapting background is shown at the top and the responses to increasing intensity spot stimuli from 90 cd/m$^2$ to 1860 cd/m$^2$ are shown from top to bottom. A profile of the stimulus is shown at the bottom of the column. Intensity–response functions are shown in (b) for ON- and (c) for OFF-center RGCs for nob2 and WT animals. Responses are plotted as the mean spike rate (spikes/s) determined for the first 0.5 s of the stimulus, expressed as a percentage of the maximum response obtained for that cell. Responses are plotted against the log(normalized intensity) for ON-center cells and $-\log$ (normalized intensity) for OFF-center cells, as defined in the Methods. Data points indicate mean ± S.E.M. for 16 WT ON-center cells, 13 nob2 ON-center cells, 11 WT OFF-center cells, or 11 nob2 OFF-center cells. Curves represent the Hill equation fitted to the averaged data set, to derive the semisaturation constant and slope. Mice were studied at 5 months of age.
from photoreceptors to second-order neurons. The role of the $\alpha_{1F}$ VDCC subunit in the inner retina is less clear, although its expression is widespread. As in the outer retina, it would be expected to mediate glutamate release from the bipolar cells. The continued presence of high-frequency oscillatory potentials in the ERG indicates that function in the inner retina of nob2 mice may be less compromised than in the outer retina in the absence of $\alpha_{1F}$. Our characterization of visual responses and receptive fields in RGCs is consistent with this observation and could indicate the use of, or compensation by, another $\alpha_{1}$ subunit in the inner retina.

**Role of the $\alpha_{1F}$ subunit of the VDCC in synaptic development or maintenance in the outer retina**

In WT retina, central DBC dendrites and flanking horizontal cell processes invaginate into the spherule of the rod photoreceptor. In cone pedicles, multiple invaginations form, each containing a similar triadic configuration. In contrast, HBCs make flat synapses with cone terminals outside of the invaginations (Haverkamp et al., 2003). In the mouse retina, the development of the invaginating synapses is established by P14 (Blanks et al., 1974). The early stages of synapse development appear to be genetically determined, although full maturation is thought to depend upon activity-dependent processes that occur after eye opening (Mumm et al., 2005). In the adult nob2 retina, every rod bipolar and horizontal cell extends ectopic dendrites across a disrupted OPL into the ONL, and these processes sometimes reach the retinal pigment epithelium. In addition, the morphology of the Ribeye-positive arcuate forms, which characterize the ribbon in the presynaptic process (tom Dieck et al., 2005), is abnormal. Thus, it is highly unlikely that rod spherules contain conventional invaginating synapses, although electron microscopy will be required to definitively resolve this issue. Certainly, the apposition of these Ribeye-positive profiles with mGluR6-positive puncta in the ONL indicates the possibility of ectopic synapses. Whether these alterations are the result of abnormal synapse formation or failure of synapse maintenance also remains to be determined. By contrast, the mGluR6-positive puncta that reflect DBC synapses onto cone pedicles appear to be less affected. It is possible that another VDCC $\alpha_{1}$ subunit may be expressed and support at least partial synaptic formation with the cone pedicle. This idea is supported by reports of $\alpha_{1D}$ expression in cone terminals (Wilkinson & Barnes, 1996; Taylor & Morgans, 1998; Morgans, 1999; Morgans et al., 2001, 2005) and is consistent with the greater ERG b-wave deficit in mice lacking the VDCC $\beta_{2}$ subunit (Ball et al., 2002), the only VDCC $\beta$ subunit known to be expressed in the OPL (Ball et al., 2004).

Two other mutant mice have been reported that alter neurotransmitter release from photoreceptors and exhibit abnormal bipolar cell dendritic morphology and reduced ERG b-waves. These mice involve the $\beta_{2}$ subunit of the VDCC (Ball et al., 2002; Gregg et al., 2002) or calcium-binding protein 4 (Haeseleer et al., 2004). Similar abnormalities also have been noted in mice lacking expression of bassoon (Dick et al., 2003) as well as in mice that lack functional rod or cone photoreceptors (Claes et al., 2004). In comparison, mice with mutations in the postsynaptic proteins mGluR6, nycatalopin or G_{60}, do not share these morphological abnormalities (Pardue et al., 2001; Ball et al., 2003; Masu et al., 1995; Tagawa et al., 1999; Dhingra et al., 2000, 2002).

One common feature of all presynaptic mutants is that they have decreased ERG b-waves most likely because of a reduction in glutamate release. Some, including nob2 mice or mice lacking the VDCC $\beta_{2}$ subunit (Ball et al., 2002), clearly also have a decrease in calcium entry. In these mutants the abnormal calcium signaling could involve processes other than neurotransmitter release (Ghosh & Greenberg, 1995). For example, calcium influx can activate intracellular pathways, which can act locally at the synapse or mediate changes in nuclear gene transcription (Rosen et al., 1994; Ghosh & Greenberg, 1995; Curtis & Finkbeiner, 1999), processes that have the potential to alter cell surface molecules that may be critical to normal synapse formation. *In vitro* studies have presented a possible link between VDCCs and photoreceptor development in which VDCC blockade inhibits process outgrowth, varicosity formation, and vesicle protein synthesis in developing rods (Zhang & Townes-Anderson, 2002; Nachman-Clewner et al., 1999). These results indicate that the presence and/or activity of VDCCs is required for normal development and synapse formation and/or maintenance in the outer retina.

It is also possible that glutamate release is a key component guiding bipolar and horizontal cell neurites to the photoreceptor active zones. Neurotransmitters have been proposed to act as an activity-dependent signal during synaptic development and neuronal plasticity in the retina (Redburn & Rowe-Rendleman, 1996). In particular, glutamate has been shown to play a role in synaptic motility (Engert & Bonhoeffer, 1999; Maletic-Savatic et al., 1999; Wong et al., 2000; Wong & Wong, 2001) and in the growth and maintenance of dendritic spines (Passafaro et al., 2003). Glutamate may also cell-autonomously affect presynaptic development using local feedback loops through presynaptic a-amino-3-hydroxy-5-methylisoxazole-4-propionic acid (AMPA)/kainate-type receptors (Chang & De Camilli, 2001). This local feedback loop could affect the growth and/or maintenance of the photoreceptor terminal through neuronal activity, although the presence of such presynaptic glutamate receptors in adult mice is debated (see review by Yang, 2004) and their expression during development is unknown. Further studies are required to determine the contributions of abnormal glutamate release, intracellular calcium signaling, or absent VDCCs to the nob2 phenotype.

**Ganglion cell response properties in nob2 mice**

Our ERG and anatomical characterizations of the nob2 mouse retina indicate that the impact of this mutation on the rod and cone pathways should differ. By definition, patients with CSNB are night blind, but those with CSNB2 also show deficits at photopic levels (Miyake et al., 1986). In characterizing RGC visual responses, we focused on a comparison of signaling through the ON and OFF cone pathways. Our results indicate that center-surround organization is unaffected in nob2 mice and that their receptive-field organization is similar to WT cells. When we compared the spontaneous activity and responses to changes in intensity, we found that nob2 OFF-center cells were unaffected. In contrast, ON-center cells in nob2 mice showed both lower spontaneous activity and a change in the gain of their response to incremental light stimuli that significantly narrowed their dynamic range. This deficit in the ON-center pathway is consistent with the reduction in the mouse cone ERG b-wave that is known to have a major contribution from cone DBCs (Sharma et al., 2005). In addition, our results may indicate that the invaginating synapses characteristic of the ON-pathway are more dependent on the presence of the $\alpha_{1F}$ subunit, while the flat synapse with HBCs may normally rely on another mechanism or substitute another $\alpha_{1}$ subunit in the absence of $\alpha_{1F}$. While the primary defect in visual processing must occur at the first synapse in the nob2 retina, it should be noted that...
there also could be a secondary effect of the absence of $\alpha_{1F}$ subunit expression in the inner retina. Several possible $\alpha_{1F}$ containing VDCC-dependent synapses in either the inner or outer retina could be involved in the changes that we report. Further dissection and characterization of light responses will be required to determine the sites and underlying mechanisms of action.

Comparable studies have been made of RGCs in the nob mouse (Gregg et al., 2003b, 2005). While nob RGCs could be characterized as responding to the onset of either a bright or dark stimulus, their receptive fields show no evidence of center-surround organization (Gregg et al., 2005).

nob2 mice provide a model for human disease

CSNB2 is a human retinal disorder that also is caused by mutations within the Cacna1f gene (Bech-Hansen et al. 1998; Strom et al. 1998; Zeitz et al., 2005). While most CSNB disorders are the result of null mutations, some may actually increase calcium entry but reduce overall function by shifting the operating range (Hemera-Wahanui et al., 2005; Hoda et al., 2005). The ERG is commonly used to confirm a diagnosis of CSNB (Fishman et al., 2001). If the nob2 mouse is an accurate model for CSNB2, then the ERG features of the mouse should match those of human patients. The results reported here indicate that they do. Under dark-adapted conditions, both nob2 mice and CSNB2 patients exhibit a marked reduction in the ERG b-wave, while the a-wave component is spared. In both cases, the ERG results are stable across age, consistent with a nonprogressive disorder. While diminished, both the b-wave and high-frequency oscillatory potentials are retained in nob2 mice and CSNB2 patients, indicating the retention of some inner retinal activity in the absence of $\alpha_{1F}$. While there are important differences in the origins of the primate and mouse cone ERG (Sieving et al., 1994; Sharma et al., 2005), these responses also indicate the presence of inner retinal activity in the absence of $\alpha_{1F}$. A recent publication has described a KO mouse for $\alpha_{1F}$ (Mansergh et al., 2005). Like the nob2 mouse, $\alpha_{1F}$-KO mice do not express the $\alpha_{1F}$ subunit protein in the OPL, have a thinner than normal OPL, and develop ectopic neurites from DBCs and horizontal cells. However, the functional deficits reported in $\alpha_{1F}$-KO mice are more severe than those we report here for nob2 mice. For example, cone-driven RGC responses are preserved innob2 mice, while in the $\alpha_{1F}$-KO mice under similar stimulation conditions, light-evoked responses were not observed in the superior colliculus, which is a central target of RGCs. In addition, in $\alpha_{1F}$-KO mice, cone ERGs were absent and under dark-adapted conditions the $\alpha_{1F}$-KO a-wave was reduced and oscillatory potentials were absent. It is not clear if these functional abnormalities noted in the $\alpha_{1F}$-KO mouse are more severe than those we report here for nob2 mice. For example, cone-driven RGC responses are preserved innob2 mice, while in the $\alpha_{1F}$-KO mice under similar stimulation conditions, light-evoked responses were not observed in the superior colliculus, which is a central target of RGCs. In addition, in $\alpha_{1F}$-KO mice, cone ERGs were absent and under dark-adapted conditions the $\alpha_{1F}$-KO a-wave was reduced and oscillatory potentials were absent. It is not clear if these functional abnormalities noted in the $\alpha_{1F}$-KO mouse are more severe than those we report here for nob2 mice.

In contrast, the phenotype of the nob2 mouse is more consistent with the phenotype seen in CSNB2 patients. The changes observed in thenob2 RGC function predict alterations in the peripheral vision of CSNB2 patients since the mouse retina has been considered in many important respects to resemble the primate retina outside of the fovea (Jeon et al., 1998). Currently, little is known about more complex visual phenotypes of CSNB2 patients, but our results would predict a deficit in vision mediated by the ON-center pathway under light-adapted conditions. In particular, our results would predict that CSNB2 patients would show significant deficits in increment, but not decrement thresholds and discriminations.

Our results also predict that CSNB2 patients will have a reduced OPL thickness, as well as ectopic bipolar and horizontal cell dendrites. Confirmation of these anatomical abnormalities will require examination of donor eyes from patients with CSNB2, although advances in optical coherence tomography may soon allow the status of the OPL to be visualized in the living human eye (Wollstein et al., 2005). Human studies aside, further analysis of nob2 mice are likely to contribute to our understanding of the pathophysiology of CSNB2 and synaptogenesis in the OPL of the retina.

Acknowledgments

We are grateful to Allison Crosby and Susan Eichholtz for excellent technical assistance. This work was supported by National Institutes of Health Grants R01EY04067 (N.C.B.), RR001183 (M.T.D.), R01EY012354 (R.G.G.), R01EY007758 (J.R.H.), R01EY04710 (M.A.M.), R01EY014700 (C.W.M), R01EY14465 (N.S.P.), R24EY15638 (N.S.P.), and awards from the Medical Research Service, Department of Veterans Affairs to N.C. Brecha (Merit Award; Senior Research Career Scientist) and to N.S. Peachey (Merit Award; Research Career Scientist).

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nob2 mouse model of CSNB2


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