γ-Protocadherins regulate neuronal survival but are dispensable for circuit formation in retina

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Twenty-two tandemly arranged protocadherin-γ (Pcdh-γ) genes encode transmembrane proteins with distinct cadherin-related extracellular domains and a common intracellular domain. Genetic studies have implicated Pcdh-γ genes in the regulation of neuronal survival and synapse formation. Because mice lacking the Pcdh-γ cluster die perinatally, we generated conditional mutants to analyze roles of Pcdh-γ genes in the development and function of neural circuits. Retina-specific deletion of Pcdh-γ led to accentuation of naturally occurring death of interneurons and retinal ganglion cells (RGCs) during the first two postnatal weeks. Nonetheless, many neuronal subtypes formed lamina-specific arbors. Blocking apoptosis by deletion of the pro-apoptotic gene Bax showed that even neurons destined to die formed qualitatively and quantitatively appropriate connections. Moreover, electrophysiological analysis indicated that processing of visual information was largely normal in the absence of Pcdh-γ genes. These results suggest that Pcdh-γ genes are dispensable for elaboration of specific connections in retina, but play a primary role in sculpting neuronal populations to appropriate sizes or proportions during the period of naturally occurring cell death.

KEY WORDS: Apoptosis, Interneuron, Laminar specificity, Receptive field, Mouse

INTRODUCTION
The assembly of neurons into complex stereotyped circuits has been hypothesized to require large sets of cell-surface molecules that mediate cell-cell interactions. A group of genes called clustered protocadherins (Pcdhs) has intriguing features that suggest their involvement in these processes. First, they have a remarkable genomic organization in which 58 homologous genes are arrayed in three subclusters (Pcdh-α, -β and -γ) arrayed in tandem on a single chromosome (Kohmura et al., 1998; Obata et al., 1998; Wu and Maniatis, 1999; Wu and Maniatis, 2000). Second, α- and γ-protocadherins arise through the combination of distinct extracellular domains with a common cytoplasmic domain, suggesting a mechanism in which distinct recognition events promote a common cellular response. Third, Pcdhs are members of the cadherin superfamily, other members of which mediate selective intercellular interactions, including synapse formation (Takeichi, 2007). Fourth, Pcdhs are expressed predominantly in the nervous system, with individual family members expressed in combinatorial patterns (Esumi et al., 2005; Frank et al., 2005; Kohmura et al., 1998; Zou et al., 2007). Fifth, Pcdh proteins are associated at least in part with synaptic membranes (Kohmura et al., 1998; Phillips et al., 2003; Wang et al., 2002). Finally, clustered protocadherin orthologs are present in vertebrates but not in invertebrates (Hill et al., 2001; Hirayama and Yagi, 2006; Noonan et al., 2004). Together, these features suggest that Pcdhs might underlie complex patterns of selective neural connectivity in vertebrates.

The first genetic test of this hypothesis led to an unexpected result: targeted mouse mutants lacking all 22 Pcdh-γ genes exhibited massive apoptosis of spinal interneurons during late fetal life and died within hours of birth (Wang et al., 2002). Synapse number was also reduced in mutant spinal cords. This was not a trivial consequence of decreased neuronal number, as synaptic defects and perinatal lethality persisted when apoptosis was blocked (Weiner et al., 2005). Thus, neural connectivity may be defective in the absence of Pcdh-γ genes, and apoptosis may be secondary to circuit defects. However, the associated lethality and the complexity of spinal circuitry have made it difficult to test these possibilities. In addition, it remains unknown whether Pcdh-γ genes are required for neuronal survival and synaptogenesis in other regions of the nervous system.

To address these issues, we generated conditional alleles of the Pcdh-γ cluster, restricting inactivation to defined neuronal populations and bypassing neonatal lethality. Here, we focus on the retina, which has several advantages, including a stereotyped structure, markers for many neuronal and synaptic subtypes, and a clear understanding of the tissue’s function (Masland, 2001; Wässle, 2004). We used Cre recombinase to delete Pcdh-γ genes from retinal neurons and glia, and assessed the consequences for neuronal structure and function. Surprisingly, lamina-specific arbors and complex functional circuits formed in the absence of Pcdh-γ genes, suggesting that these genes play limited roles in synaptic specificity. By contrast, loss of Pcdh-γ genes accentuated naturally occurring death of multiple retinal cell types. These results suggest a primary role for Pcdh-γ genes in neuronal population matching during development.

MATERIALS AND METHODS
Animals/generation of targeted mice
Pcdh-γmut and Pcdh-γdis mutants have been described previously (Wang et al., 2002). Mice in which regulatory elements from the Chx10 gene drive expression of a Cre-GFP fusion protein linked by an internal ribosome entry site (IRES) to placental alkaline phosphatase (Chx10-Cre) (Rowan and Cepko, 2004) were provided by Constance Cepko (Harvard). Mice in which a short enhancer fragment from the Pax6 gene drive expression of a Cre-IRES-GFP cassette (Pax6-Cre) (Marquardt et al., 2001) were provided by Peter Gruss (Göttingen, Germany). Mice in which regulatory elements from the β-actin gene drive expression of Cre (Actin-Cre) (Lewandoski et al., 1997) were provided by Gail Martin (UCSF). Bax mutants (Knudson et al., 1995) and Z/EG reporter mice (Novak et al., 2000) were obtained from Jackson Laboratories.

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The Pcdh-γ:\textsuperscript{fusg} targeting vector was modified from the Pcdh-γ:\textsuperscript{fusg} vector shown in Fig. 2B of Wang et al. (Wang et al., 2002) by inserting a loxP sequence into an Nhel site upstream of the final coding exon. The Pcdh-γ:\textsuperscript{fusg} allele was generated by re-targeting the ES cells used to generate Pcdh-γ:\textsuperscript{fusg} with the vector that had been used to generate Pcdh-γ:\textsuperscript{fusg}. This vector inserted a loxP sequence directly upstream of variable exon A1. Homologous recombinants and germ line chimeras were generated by standard methods. Mice were maintained on a C57/B6J background.

Electrophysiology

Dark-adapted retinas were isolated under an infrared microscope into superfused with Ringer’s (Kim et al., 2008). Extracellular action potentials were recorded and single units identified by spike-sorting methods as described previously (Meister et al., 1994). Whole-cell patch clamp recordings were made in the voltage-clamp configuration using a patch pipette filled with an internal solution containing 135 mM K-gluconate, 10 mM HEPES, 5 mM NaCl, 10 mM MgCl₂, 0.5 mM EGTA, 0.5 mM CaCl₂, and 4 mM Na₂ATP (pH 7.3, 280 mosmol/kg H₂O). Membrane potential was clamped to -70 mV and currents were recorded at a holding current of 0 nA. The firing rate analysis was performed using Clampfit (Molecular Devices). RGCs were classified as silent, low-spiking, or high-spiking using an automated classifier that could also be manually adjusted.

RESULTS

Broad expression of gamma protocadherins in retina

We began our study by assessing the distribution of Pcdh-γ genes in retina. We generated transgenic mice in which GFP is fused to the shared C terminus, tagging all 22 Pcdh-γ isoforms (Pcdh-γ:\textsuperscript{fusg}) (Wang et al., 2002). Homozygous Pcdh-γ:\textsuperscript{fusg} mice are viable and fertile, and show none of the defects documented previously in Pcdh-γ mutants (Wang et al., 2002; Weiner et al., 2005). We therefore believe that GFP is a neutral reporter of endogenous Pcdh-γ localization.

The retina consists of three cellular layers separated by two synaptic or ‘plexiform’ layers (Fig. 1A). The outer nuclear layer (ONL), containing photoreceptors; the inner nuclear layer (INL), containing interneurons (horizontal, bipolar and amacrine cells) and Müller glia; and the ganglion cell layer (GCL), containing RGCs and displaced amacrine cells. The outer plexiform layer (OPL) contains synapses of photoreceptors onto horizontal and bipolar cells, and the inner plexiform layer (IPL) contains synapses of bipolar and amacrine cells onto RGCs. As judged by localization of GFP in Pcdh-γ:\textsuperscript{fusg} mice, Pcdh-γ are present in all five retinal layers (Fig. 1A). In the ONL, Pcdh-γ are present in outer segments and around photoreceptor somata (Fig. 1B); in the INL and GCL, Pcdh-γ outline neuronal somata (Fig. 1,C,D). Pcdh-γ levels are highest in the most membrane-rich layers: IPL, OPL and the optic fiber layer that carry RGC axons to the brain (Fig. 1A’,D). In situ hybridization confirmed Pcdh-γ expression by cells in the INL and GCL, though this method did not reliably detect Pcdh-γ RNA in photoreceptors (Fig. 1E).

To determine which retinal cell types express Pcdh-γ genes, we dissociated Pcdh-γ:\textsuperscript{fusg} retinas and immunostained cells with antibodies to cell-type-specific markers (Haverkamp and Wässle, 2000; Wahlin et al., 2004; Zhang et al., 2004). This method circumvented the difficulty of determining which of the cells abutting Pcdh-γ-rich membranes are themselves Pcdh-γ positive. Markers included Brn3a and Thy1 for RGCs, syntaxin 1 for amacrine cells, Chx10 for bipolar cells, calbindin for horizontal cells, recoverin for photoreceptors and glial fibrillary acidic protein, Sox9 and glutamine synthetase for Müller glia. All six cell types were Pcdh-γ positive (Fig. 1F–K; data not shown). Thus, Pcdh-γ are expressed in all cell types of the neural retina.

\begin{equation}
\text{h}(x,t) = \frac{1}{n} \sum_{i=1}^{n} \text{h}(x_{i}, t_{i} + t)
\end{equation}
We asked whether Pcdh-γ genes are present in the retina during early postnatal life, when neural circuits form. At postnatal day (P) 0, the retina contains ganglion cell and neuroblast layers, separated by a nascent IPL. All RGCs have been born by this time, while neurogenesis and migration of newborn interneurons and photoreceptors continue in the neuroblast layer. At this time, Pcdh-γ is present on cells in the neuroblast layer, in the IPL, and on RGC axons (Fig. 2A). At P3, Pcdh-γ is apparent in the layer of horizontal cells that prefigures the OPL (Fig. 2B). By P7, Pcdh-γ appears in the OPL, as it divides the neuroblast layer into INL and ONL (Fig. 2C). By P14, the adult pattern described above is established (Fig. 2D).

**Synaptic localization of Pcdh-γ proteins in the retina**

To evaluate the subcellular localization of Pcdh-γ proteins, we focused on the OPL, because its synapses are larger than those in the IPL. We labeled photoreceptor terminals with antibodies to bassoon, which is present in both rod terminals (spheres) and cone terminals (pedicles), and labeled spheres and pedicles selectively with anti-PSD-95 and peanut agglutinin, respectively (Blanks et al., 1987; Koulou et al., 1998; tom Dieck and Brandstätter, 2006). Pcdh-γ was associated with both spheres and pedicles (Fig. 3A-C). We labeled bipolar cell dendrites with antibodies to protein kinase Ca and neurokinin receptor 3, which mark rod and cone bipolars, respectively, and to G protein γ13 (Gγ13), which is present in subsets of both rod and cone bipolars (Haverkamp et al., 2003; Huang et al., 2003). Pcdh-γ was associated with both rod and cone bipolar dendrites (Fig. 3D; data not shown). Thus, Pcdh-γ proteins were present in pre- and postsynaptic compartments of rod and cone synapses. By contrast, although horizontal cell processes labeled with antisera to calbindin (Sharma et al., 2003) were Pcdh-γ-positive, little Pcdh-γ was present in their synaptic varicosities (Fig. 3E).

Pcdh-γ was also present throughout the IPL. GFP-positive puncta often overlapped with bassoon-positive presynaptic ribbons in bipolar cells, glutamate decarboxylase- and GlyT1-positive terminals of inhibitory amacrines, and PSD-95-positive postsynaptic membranes of excitatory synapses (Fig. 3F; data not shown). Taken together, these results suggest that Pcdh-γ is present at many synapses in the retina, although they are not confined to synapses.

**Inactivation of Pcdh-γ null in the retina leads to neuronal and synaptic loss**

Pcdh-γ-null and hypomorphic mice die shortly after birth (Wang et al., 2002; Weiner et al., 2005). We examined retinas of Pcdh-γ-null mutants at late embryonic stages [embryonic day (E) 17-18; birth is at E19] but found no obvious defects in retinal structure (see below). However, as the development of retinal circuitry occurs largely during postnatal life, roles of Pcdh-γ in circuit formation and function could not be studied in these mutants. We therefore...
generated two conditional inactivation alleles to bypass neonatal lethality (Fig. 4A). In *Pcdh-γ*<sup>fdel</sup>, loxP sites flank the entire *Pcdh-γ* locus, such that Cre-mediated recombination generates a null allele. In *Pcdh-γ*<sup>fcon3</sup>, the C-terminal exon shared by all isoforms is flanked by loxP sites, such that Cre truncates all *Pcdh-γ* genes in both alleles, GFP is fused to this C-terminal exon, allowing us to use loss of GFP as an indicator of Cre-mediated *Pcdh-γ* excision. The *Pcdh-γ*-GFP fusion protein was identical to that in the *Pcdh-γ*<sup>fusg</sup> allele described above.

In initial tests, we excised the floxed segments of the *Pcdh-γ*<sup>fdel</sup> and *Pcdh-γ*<sup>fcon3</sup> alleles in the germline by mating them to transgenic mice in which Cre is expressed ubiquitously (*Actin-Cre*) (Lewandoski et al., 1997). Homozygotes generated from these animals died at birth and exhibited spinal cord phenotypes similar to those described previously in null mutants (Wang et al., 2002), indicating that recombination inactivates the *Pcdh-γ* gene (data not shown). Western blotting reported by Prasad et al. (Prasad et al., 2008) failed to detect *Pcdh-γ* protein in *Actin-Cre;Pcdh-γ*<sup>fcon3</sup>*fcon3* mice, indicating that this allele is effectively a protein null. The truncation in *Pcdh-γ*<sup>fcon3</sup> is more extensive than the hypomorphic allele described previously (*Pcdh-γ*<sup>tr</sup>), in which *Pcdh-γ* levels were decreased several-fold (Weiner et al., 2005). We speculate that increased truncation of *Pcdh-γ*<sup>fcon3</sup> and lack of a polyadenylation signal led to greater destabilization of *Pcdh-γ* protein and mRNA, respectively.

To selectively inactivate *Pcdh-γ* in retina, we crossed *Pcdh-γ*<sup>fdel</sup> and *Pcdh-γ*<sup>fcon3</sup> mutants with mice in which a GFP-Cre recombinase fusion protein is expressed under the control of regulatory elements from the *Chx10* gene (*Chx10-Cre*) (Rowan and Cepko, 2004). These elements drive expression of GFP-Cre transiently in embryonic retinal progenitors and postnatally in bipolar cells. To assay recombination in retinas of *Chx10-Cre* mice, we crossed them to a reporter line in which β-galactosidase (*lacZ*) and GFP label non-recombined and recombined cells, respectively (*Z/EG*) (Novak et al., 2000). Recombination was extensive (>90%) in the INL and ONL, but occasional columns of cells were spared (see Fig. S1A in the supplementary material). By contrast, approximately half of the cells in the GCL were GFP-negative and *lacZ* positive (see Fig. S1B in the supplementary material). This pattern may reflect the fact that many RGCs are born by E12 (Farah and Easter, 2005), before Cre accumulates in progenitors. We then used loss of GFP to assay *Chx10-Cre*-mediated loss of *Pcdh-γ*-GFP from the *Pcdh-γ* alleles. This method did not allow us to assess excision in bipolar cells, in which GFP was expressed from the *Chx10* transgene (see Materials and methods). Nonetheless, *Chx10-Cre* excised *Pcdh-γ*<sup>fdel</sup>, *Pcdh-γ*<sup>fcon3</sup> and *Z/EG* genes in similar patterns and to a similar extent (see Fig. S1C-E in the supplementary material). The efficacious excision of the *Pcdh-γ*<sup>fdel</sup> allele is surprising given the length of the floxed segment.

*Chx10-Cre;Pcdh-γ*<sup>fdel</sup> mice are healthy and outwardly normal. We first examined these mutants at P18, by which time the retinal architecture is well developed. Labeling of nuclear layers with DAPI and plexiform layers with antibodies to the synaptic vesicle protein synaptophysin revealed that mutant retinas were properly laminated (Fig. 4B-G). However, mutant retinas were ~25% thinner than those of wild-type mice or heterozygote littermates. The difference resulted from a selective reduction of ~40% in the thickness of the INL and the IPL (Fig. 4H). Thus, *Pcdh-γ* are required for
development or maintenance of retinal interneurons and the layer in which they form synapses. The INL and IPL were thinned to the same extent in Chx10-Cre;Pcdh-γfcon3/fdel mice and Chx10-Cre;Pcdh-γfdel/fdel mice (Fig. 4C,D,F,G), consistent with the idea that Pcdh-γfcon3 is functionally a null. In subsequent studies, we used the two alleles interchangeably, but most of the results reported here are from Pcdh-γfdel mice.

The mosaicism described above for Chx10-Cre retinas made labels with DAPI or Pax6 immunohistochemistry to identify α-amacrines; RGC, Brn3a+ retinal ganglion cells; MG, Sox9+ Müller Glia. Other abbreviations as in Fig. 1. Error bars indicate s.e.m. ***P<0.001, Mann-Whitney non-parametric test. Scale bars: 50 μm.

Increased postnatal apoptosis in the absence of Pcdh-γ genes

We next asked when retinal defects arise in Pcdh-γ mutants, and whether they are progressive. We detected no differences in laminar arrangement or thickness between mutant and control retinas perinatally (E17.5-P3) (Fig. 5A,B,G,H and data not shown). By P7, however, shortly after the ONL and INL form, mutant retinas were thinner than those of controls (Fig. 5C,D). We used Chx10-Cre: Pcdh-γfcon3 mice for quantification of these defects. Both layers were ~40% thinner in mutants than in controls by P14, then changed little over the following months (Fig. 5E-I). Thus, the difference between mutant and control retinas appears during the first postnatal week, is maximal by the end of the second postnatal week, and neither abates nor worsens substantially thereafter.

A process of naturally occurring programmed cell death eliminates many retinal neurons during the first two postnatal weeks (Pequignot et al., 2003; Young, 1984). Apoptosis followed a similar time course in Pcdh-γ-deficient retinas, but levels were significantly higher in mutants than in controls (Fig. 5J-L). Although increased apoptosis was seen in both neuroblast and ganglion cell layers at P10, it was confined to the INL at P7 (Fig. 5M). This pattern is consistent with the finding that naturally occurring cell death in the GCL is complete several days before that in the INL (Farah and Easter, 2005; Pequignot et al., 2003; Young, 1984). These results suggest that Pcdh-γ genes regulate neuronal survival during the period of naturally occurring programmed cell death.

Cell autonomy of Pcdh-γ-dependent cell survival in retina

Are Pcdh-γ genes required for cell survival in cells that express them, in neighboring cells, or in both? As a first step to distinguish between these possibilities, we capitalized on the recombination pattern of Pax6α-Cre transgenic mice. As noted above, Cre is expressed in all progenitors in peripheral retina as well as in a subset of amacrine cells marked by the GFP transgene in the GCL is complete several days before that in the INL (Farah and Easter, 2005; Pequignot et al., 2003; Young, 1984). These results suggest that Pcdh-γ genes regulate neuronal survival during the period of naturally occurring programmed cell death.
neighboring amacrines are Pcdh-γ deficient in the central region, this result does not demonstrate cell autonomy sensu strictu, but does indicate that loss of Pcdh-γ genes from a single cell type impairs survival, even when the majority of its synaptic inputs (bipolar cells) and targets (RGCs) are wild type. This result also rules out the possibility that neuronal apoptosis in the absence of Pcdh-γ is secondary to a defect in surrounding glial cells. Furthermore, loss of Pax6α-positive amacrines is equivalent in peripheral and central retina (Fig. 6F), indicating that Pcdh-γ-negative cells are not protected from apoptosis when surrounded by Pcdh-γ-positive cells.

We also asked whether loss of Pcdh-γs from Pax6α-amacrines was detrimental to survival of neighboring cells (Fig. 6F). Loss of Pcdh-γs from the Pax6α-positive amacrines had no detectable effect on the survival of bipolar cells, horizontal cells or Müller glia. Likewise, survival of a distinct, intermingled subpopulation of amacrines – the cholinergic starburst cells – was unaffected in central retina. By contrast, we detected a small (~15%) but significant loss of Brn3a-positive RGCs from central retina of Pax6α-Cre;Pcdhγfcon3/fcon3 mice. We do not know whether this loss reflects absence of Pcdh-γs per se, or death of amacrines, which regulate at least some aspects of RGC development (Goldberg et al., 2002).

### Pcdh-γ genes are dispensable for laminar targeting of retinal neurons

Although the width of the IPL is dramatically reduced in Pcdh-γ-deficient retina, it nonetheless contains synapses, as judged by the presence of pre- and postsynaptic markers such as synaptophysin and bassoon (Fig. 4E-G; Fig. 8A,B). Are they appropriate synapses? The retina is well-suited to test specificity, because discrete subsets of bipolar and amacrine cells arborize and synapse in just one or a few of at least 10 closely spaced parallel sublaminae within the IPL (Pang et al., 2002; Roska and Werblin, 2001; Wässle, 2004).

We used markers of 10 lamina-specified amacrine and bipolar subtypes to assess lamina-specific arborization and connectivity in the absence of Pcdh-γs. We follow a scheme in which five sublaminae of equal width are numbered, from the border of the INL (S1) to the border of the ganglion cell layer (Ghosh et al., 2004).
in mutant retinas at P18, although disruptions or gaps were
In all 10 cases, processes were arrayed in appropriate sublaminae
(anti-NK3R); and rod bipolars (anti-protein kinase C
Pax6
Pcdh-
positive amacrine cells (asterisks) and Pcdh-
GFP and Brn3a antibodies. Anti-GFP labels both Pax6
positive amacrines and RGCs; OFF bipolar cells (anti-
hydroxylase); type AII amacrines (anti-disabled); calbindin-
glutamatergic amacrines (anti-vGlut3); GABAergic amacrines
starburst amacrines, labeled by choline acetyltransferase;
2004; Yamagata and Sanes, 2008). Populations examined were
γ
protocadherins in retina
α
Cre;Pcdh-
egFP proteins are visible in the
α
γ
- Cre;Pcdh-
egFP proteins (arrows). In
γ
single mutants (Fig. 8M,N; data not shown).
Deletion of Bax also resulted in expansion of the IPL. The IPL in
Chx10-Cre;Pcdh-γ
fcon3/+/fcon3;Bax–/– mice was thicker than that in
Pcdh-γ mutants and indistinguishable from that in Pcdh-γ-positive
Bax–/– mutants (Fig. 8A-D). The density of synapses in the IPL and
OPL, as judged by staining for PNA or bassoon, did not differ
significantly between Pcdh-γ-positive and Pcdh-γ-negative
Bax–/– retinas (Fig. 8E-L; see Fig. S2 in the supplementary material). Thus,
loss of Pcdh-γ had little effect on synapse number in the IPL when
apoptosis was prevented. This result is consistent with the idea that
much of the synapse loss in the IPL of Pcdh-γ-deficient retina is a consequence of decreased neuron number.
To assess the laminar targeting of interneurons that would have
died in the presence of Bax, we stained Chx10-Cre;Pcdh-γ
fcon3/fcon3;Bax–/– retinae with the panel of markers listed above. In all cases,
targeting of processes to appropriate laminae was as specific in
double mutants as in Pcdh-γ-single mutants (Fig. 8M,N; data not
shown). Moreover, thinning and disruptions of layers observed in
Pcdh-γ single mutants were rescued in Pcdh-γ-positive
Bax–/– double mutants (compare Fig. 7B,D with Fig. 8M,N). We therefore
conclude that the gaps observed in Pcdh-γ deficient retina are secondary to the loss of cells rather than a manifestation of improper laminar targeting.

Functional visual circuits form in the absence of
Pcdh-γ
To test whether circuits that form in Pcdh-γ-deficient retina are
functional, we recorded light responses from RGCs. These cells
integrate signals from amacrine and bipolar interneurons and send
the resulting spike trains to the brain. RGCs differ in their
responses to visual stimuli, depending on the synaptic inputs they
receive. Thus, ON RGCs, which respond primarily to light onset,
receive synapses from ON bipolar cells in the inner half of the IPL
(near the GCL); OFF RGCs receive synapses from OFF
bipolars in the outer IPL; and ON-OFF RGCs receive both types
of synapses. Further specializations, such as responses that are
transient, sustained or selective for moving objects, result from
innervation by specific subsets of bipolar and amacrine cells
(Masland, 2001; Wässle, 2004). Accordingly, the presence of
diverse, specific responses from RGCs is a sensitive indicator of

2004; Yamagata and Sanes, 2008). Populations examined were
starburst amacrines, labeled by choline acetyltransferase;
glutamatergic amacrines (anti-vGlut3); GABAergic amacrines
(anti-GADB5/67); dopaminergic amacrines (anti-tyrosine
hydroxylase); type AII amacrines (anti-disabled); calbindin-
positive amacrines and RGCs; OFF bipolar cells (anti-
synaptotagmin 2); ON bipolar cells (anti-GY13); OFF bipolars
(anti-NK3R); and rod bipolars (anti-protein kinase Cα) (Fig. 7E).
In all 10 cases, processes were arrayed in appropriate sublaminae
in mutant retinas at P18, although disruptions or gaps were

sometimes present (Fig. 7A-D; data not shown). We also observed
proper laminar targeting of amacrine subsets at P7, and of bipolar
subsets at P14, in each case soon after these synapses formed in
controls (data not shown).

Synapses and arbors of Pcdh-γ-deficient neurons
destined to die
Results illustrated in Fig. 7 are consistent with the idea that Pcdh-γs
are dispensable for formation of neural circuits in retina. Alternatively, however, some INL interneurons might fail to target
appropriate sublaminae and then die. In this case, neuronal apoptosis in Pcdh-γ mutants might be secondary to circuit defects. To test this
possibility, we blocked apoptosis in Pcdh-γ
fcon3/fcon3 mice by
deleting the pro-apoptotic gene, Bax. Naturally occurring death in
many regions of the nervous system, including retina, is dramatically
reduced in Bax–/– mice (Mosinger Ogilvie et al., 1998; Pequignot et
al., 2003; White et al., 1998), and deletion of Bax preserves spinal
interneurons that would otherwise die in Pcdh-γ null spinal cord
(Weiner et al., 2005). Likewise, Bax deletion rescued neurons destined
to die in Pcdh-γ mutant retina: the thickness of the INL and
GCL and the number of Chx10-positive bipolar cells were indistinguishable in
Chx10-Cre;Pcdh-γ
fcon3/fcon3;Bax–/– and Chx10-
Cre;Pcdh-γ
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(Masland, 2001; Wässle, 2004). Accordingly, the presence of
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Fig. 6. Cell autonomous and non-autonomous components of
Pcdh-γ-dependent cell survival. (A) Schematic of the recombination pattern in Pax6α-Cre retina. In Pax6α-Cre;Pcdh-γ
fcon3/fcon3 mutants, all cells are Pcdh-γ-negative in peripheral retina, but only Pax6α-positive (+) amacrine cells are Pcdh-γ-negative in the central sector.
(B-E) Immunolabeling of central and peripheral regions of Pax6α-
Cre;Pcdh-γ
fcon3/fcon3 and Pax6α-Cre;Pcdh-γ
fcon3/fcon3 mutant retinas with GFP and Brn3a antibodies. Anti-GFP labels both Pax6α-Cre-ires-GFP positive amacrine cells (asterisks) and Pcdh-γ-GFP proteins (arrows). In uncombined portions of the central Pax6α-Cre;Pcdh-γ
fcon3/fcon3 and Pax6α-Cre;Pcdh-γ
fcon3/fcon3 retinas, Pcdh-γ-GFP proteins are visible in the outer segments, OPL, and retinal axons (arrows). In peripheral regions, Pcdh-γ-GFPs are absent. In both regions, mutant retinas have reduced numbers of Pax6α-positive amacrine cells; Brn3a-positive RGCs are dramatically decreased in the mutant peripheral sector while slightly decreased in the central sector. (F) Quantification of retinal cell types in central regions of Pax6α-Cre;Pcdh-γ
fcon3/fcon3 retinas, expressed as a percentage of cells in Pax6α-Cre;Pcdh-γ
fcon3/fcon3 litters. Six to eight animals per genotype were analyzed, *P<0.05, **P<0.01; ***P<0.0001, by Student’s t-test or Mann-Whitney test. Scale bar: 50 μm.
precisely patterned synaptic connectivity. We therefore monitored action potentials simultaneously from large populations of RGCs in control and Pcdh-γ mutant retinas, using a multi-electrode array (Meister et al., 1994). Results from Chx10-Cre;Pcdh-γ<sup>fcon3/fcon3</sup> retinas and peripheral regions of Pax6α-Cre;Pcdh-γ<sup>fcon3/fcon3</sup> retinas were similar, so they are combined here. We did not use Pcdh-γ:Bax double mutants for this study, because visual responses are compromised in Bax single mutants (Pequignot et al., 2003).

RGCs in Pcdh-γ mutant and control retinas showed a similar variety of responses to small flashing spots, including sustained and transient ON, OFF and ON-OFF responses (Fig. 9A).

Fig. 7. Sublamina-specific targeting of amacrine and bipolar processes in the IPL in the absence of Pcdh-γ genes. (A,B) ChAT- (red) and vGluT3-positive amacrine subsets (green) in Chx10-Cre;Pcdh-γ<sup>con3/</sup>(control) and Chx10-Cre;Pcdh-γ<sup>fcon3/fcon3</sup> retinas. ChAT-positive processes ramify in sublaminae (S) 2 and 4, and vGluT3-positive processes ramify in S3. (C,D) Synaptotagmin 2 (SytII)-positive OFF bipolar processes (red) and G<sub>y13</sub>-positive ON bipolar processes (green) ramify in the outer and inner regions of the IPL, respectively. In all cases, laminar specificity is retained in mutants, but marker-laminae are reduced and disrupted. (E) Sketch of IPL sublaminae stained by the markers used in this study. Scale bars: 50 μm.

Fig. 8. Laminar specificity and synapse formation by Pcdh-γ-deficient neurons rescued from apoptosis. (A-D) Sections from retinas mutant for Bax (Chx10-Cre;Pcdh-γ<sup>fcon3/fcon3</sup> Bax<sup>+/–</sup>), Pcdh-γ (Chx10-Cre;Pcdh-γ<sup>fcon3/fcon3</sup> Bax<sup>+/–</sup>), both (Chx10-Cre;Pcdh-γ<sup>fcon3/fcon3</sup> Bax<sup>+/–</sup>) or neither (Chx10-Cre;Pcdh-γ<sup>fcon3/fcon3</sup> Bax<sup>+/–</sup>). Sections were stained with anti-bassoon (red) and Po-pro1 (blue). Thickness of IPL and INL are similar in Bax mutants and Bax, Pcdh-γ double mutants; both are thicker than those in Pcdh-γ mutants. (E-L) High power images of OPL (E,G,I,K) and IPL (F,H,J,L) from retinas in A-D. Density of synaptic puncta is similar in Bax mutants and Bax, Pcdh-γ double mutants; both are thicker than those in Pcdh-γ mutants. (M,N) Chx10-Cre;Pcdh-γ<sup>fcon3/fcon3</sup> Bax<sup>+/–</sup> mutants immunostained for ChAT (red) and vGluT3 (green) (M) and synaptotagmin 2 (SytII, red), and G<sub>y13</sub> (green) (N). All processes make lamina-specific arbors (compare with Fig. 7) and disruptions seen in Pcdh-γ single mutants are absent in double mutants. Scale bars: 100 μm in A-D; 10 μm in E-L; 50 μM in M,N.
Proportions of ON- and OFF-dominated responses were identical in mutants and controls (Fig. 9C). Mutant RGCs responded to very dim flashes, which excite only rods, and also to bright flashes, which predominantly excite cones (data not shown), indicating that both rod- and cone-activated pathways were functional. We also probed the retina with moving bars and gratings to elicit direction-selective responses, which are known to depend on specific patterns of connectivity in the IPL (Masland, 2001). Both ON and ON-OFF direction-selective cells were encountered in mutant retinas (Fig. 9B; data not shown). Some control and mutant RGCs had receptive field surrounds, where light has the opposite effect of the center (see Fig. S3 in the supplementary material), indicating that lateral inhibitory connections are functional in these circuits.

To survey response properties quantitatively, we stimulated the retina with randomly flickering bars and applied a reverse correlation method (Chichilnisky, 2001; Meister et al., 1994). This measures spatio-temporal receptive fields, revealing how RGCs respond to light intensity at different points on the retina and at different times in the past (see Fig. S3 in the supplementary material). The size of the receptive field center varied greatly among RGCs, but the distribution was similar in control and mutant retinas (Fig. 9D). However, the time course of the light response was significantly slower in mutant retina (Fig. 9E). Moreover, mutant RGCs fired at much lower rates in response to flicker stimuli (Fig. 9F). In principle, this could result from an elevated response threshold; alternatively, the gain of the response might be lower once the threshold is crossed. Based on fitting with a linear-nonlinear model (Chichilnisky, 2001), we found that the threshold is unaltered, but the gain is reduced in mutants (Fig. 9G-H).

**DISCUSSION**

**Protocadherins and neural specificity**

Interest in the clustered protocadherins has centered on the tantalizing idea that their molecular diversity may underlie the extraordinary synaptic specificity of the brain (Benson et al., 2001; Hamada and Yagi, 2001; Hirayama and Yagi, 2006; Kohmura et al., 1998; Morishita and Yagi, 2007; Serafini, 1999; Shapiro and Colman, 1999; Washbourne et al., 2004; Wu and Maniatis, 1999; Yagi and Takeichi, 2000). Several observations that led to this notion are summarized in the Introduction. Moreover, Hasegawa et al. (Hasegawa et al., 2008) recently showed that olfactory axons bearing a single type of odorant receptor fail to coalesce properly onto glomeruli in olfactory bulbs of mice lacking Pcdh-α genes.
These considerations, coupled with the finding that most retinal cells express Pcdh-γ genes, led us to expect that retinal circuitry might be grossly defective in their absence. Surprisingly, it was not. Synaptic specializations were present in the OPL and IPL of Pcdh-γ mutants, and the light-responsiveness of RGCs indicates that synapses in both laminae were functional. Moreover, synapses in the IPL were sublaminae specific as judged by distribution of arbors. This distribution provides a stringent test of targeting, in that 10 or more IPL sublaminae are separated by only a few tens of micrometers (Roska and Werblin, 2001; Wässle, 2004).

The loss of neurons in Pcdh-γ-deficient retinas potentially complicates this interpretation: neurons making improper arbors or connections could be selectively eliminated, so only neurons that wired up properly would be retained. The ability to block apoptosis in Pcdh-γ mutant retinas by deleting the Bax gene allowed us to test this possibility. Lamina-specific targeting was if, anything, more precise in the absence of Bax than in its presence, in that disruptions and irregularities seen in Pcdh-γ+ retinas were absent in double mutants. Therefore, IPL disruptions observed in Pcdh-γ−/− retinas presumably reflected loss of cells rather than mistargeting of neurites. Moreover, with apoptosis prevented by Bax deletion, loss of Pcdh had no detectable effect on the number of synapses in either the IPL or the OPL.

The ability of mutant retinas to process visual information was also remarkably preserved. RGCs exhibited a wide range of complex responses, and their receptive field sizes were normal. Because the spatial extent of RGC receptive field centers are largely determined by their dendritic fields, which collect input from bipolar cells (Wässle, 2004), this result suggests that mutant RGC arbors are normal in size. As bipolar cells provide the main excitation to RGCs, their decreased number could account for the lower firing rate and normal in size. As bipolar cells provide the main excitation to RGCs, their receptive field sizes were normal. Because the electrophysiological evidence for maintained synaptic function, it seems unlikely that any synaptic defect is sufficient in magnitude to explain the massive apoptosis we observe. Likewise, synaptic patterns in the IPL of Pcdh-γ mutants are at least as well preserved in the absence of Bax as in its presence, ruling out the possibility that apoptosis reflects selective elimination of inappropriate synapses.

Thus, synapses can be lost in the absence of neuronal loss in the spinal cord, and neurons can be lost in the absence of major synaptic defects in retina. These results suggest that Pcdh-γ regulates neuronal survival and synaptic maturation by distinct mechanisms, and that effects on these two processes differ in severity among brain regions. The combinatorial diversity provided by the Pcdh-γ genes may therefore be useful for selectively controlling the size of diverse neuronal populations.

Synaptic circuitry and neuronal survival

Patterns of apoptosis in Pcdh-γ-deficient retina are similar to those in spinal cord (Wang et al., 2002; Weiner et al., 2005) (see also Prasad et al., 2008) in several respects. First, approximately half of the interneurons in each region are lost in the absence of Pcdh-γ genes. Second, some interneuron subtypes and primary sensory neurons (dorsal root ganglion cells and photoreceptors) are spared in both regions, even though they express Pcdh-γ genes. Third, the loss of neurons in Pcdh-γ mutants occurs during the period of naturally occurring cell death. One apparent difference is that the output neurons of the spinal cord, motoneurons, are spared in Pcdh-γ-deficient mice, whereas those of retina, RGCs, are affected. However, at least some apoptosis of RGCs is cell-nonautonomous, reflecting either loss of Pcdh-γ from presynaptic cells or loss of input cells themselves. It is possible that in the mutants analyzed to date, motoneurons retain a larger fraction of their inputs than do RGCs, and that this contributes to their survival.

Retina and spinal cord phenotypes are also similar in that loss of Pcdh-γ genes leads to decreased numbers of synapses in both tissues. In spinal cord, synapse loss does not result simply from neuron loss, as shown by analysis of Pcdh-γ-deficient mice in which apoptosis was blocked: neuronal number was normal in these animals, but synapse number was still reduced (Weiner et al., 2005). This result is consistent with the idea that failure of synapse formation or function impairs neuronal survival (see also Prasad et al., 2008). In fact, complete blockade of synaptic function in embryonic brain leads to increased apoptosis (Verhage et al., 2000). By contrast, deletion of Pcdh-γ in a Bax−− background does not decrease synapse density in retina. In addition, given the electrophysiological evidence for maintained synaptic function, it seems unlikely that any synaptic defect is sufficient in magnitude to explain the massive apoptosis we observe. Moreover, the ability to block apoptosis in Pcdh-γ−/− retinas by deleting the Bax gene allowed us to test this possibility.


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Supplementary Fig. 2
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