Diabetic retinopathy (DR) is the fifth leading cause of blindness worldwide and the third in terms of retinal disorders (~170 million, World Health Organization). It constitutes the leading cause of vision loss for working-age persons in developed countries. DR has been viewed principally as a microvascular disease, and hyperglycemia induces the disruption of cell junctions within the retinal endothelium, thickening of basement membranes and pericyte loss. Vessels become leaky and acellular, circulation is compromised leading to the formation of ischemic zones, and exudation from the capillaries contributes to macular edema. However, many studies have indicated that abnormal visual function, such as decreased contrast sensitivity or color vision defects, actually precede clinically detectable signs of retinopathy, suggesting that vascular abnormalities are not the first events to occur in DR.

Numerous rodent models of DR exist, with varying degrees of similarity to the human disease. No single animal model recapitulates all the neural and vascular complications associated with DR, especially the late-stage proliferative form, PDR. Animal models can be classified as drug-induced, as with the destruction of pancreatic beta cells by streptozotocin (SZT); genetic, as with the widely used Ins2Δmmt mouse strain; and spontaneously occurring diabetes. This last category includes the sand rat Psammomys obesus, long known to develop metabolic stress syndrome in captivity. In contrast to animals maintained on a natural plant-rich diet, when reared on a high-calorie regimen many animals exhibit hallmark features of type 2 diabetes. The similarity between metabolic, physiological, and endocrine changes in this species and those occurring in human type 2 diabetes makes it a highly relevant animal model with which to understand the pathogenesis of this condition.

Aside from a single study on tyrosine hydroxylase levels during diabetes, retinal modifications occurring in this species are unknown. We reasoned that, given the other numerous similarities between type 2 diabetes in Psammomys and humans, the rodents may also develop DR and serve as a novel model for unraveling the complex pathogenesis in this blinding disease. The extensive morphologic and immunochemical studies presented here support this hypothesis.

**Materials and Methods**

**Experimental Animals**

All animal experimentation adhered to the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research and abided by national and local laws of ethical conduct. Young adult Psammomys were captured in southern Tunisia using baited traps, transferred to animal facilities, and maintained under cyclic lighting conditions (12 hours white light [~300 lux]/12 hours dark) with free access to food and water. Data presented in this study came from two independent field excursions, the first concerning 15 animals and the second concerning 26 animals. Male animals were separated into two groups. The low calorie diet control group (n = 5 from the first capture, n = 8 from the second capture; n = 13 total) was raised on a natural diet (ND) and was fed only halophilic plants, rich in water and mineral salts (0.4 kcal/g wet weight). The other animals (n = 10 from the first capture, n = 16 from
the second capture; \( n = 26 \) total) received a standard laboratory rat chow feed (3.3 kcal/g; Purina). Both groups were followed up for 4 months (first capture) or 7 months (second capture), with measurements of their body weight, plasma glucose (One Touch Ultra Blood Glucose Monitoring System 20247; LifeScan, Inc., Milpitas, CA), and hematocrit every 5 days. Based on weight and glucose measurements (see Results), a total of 15 animals from the hypercaloric groups (\( n = 5 \) from the first capture, \( n = 10 \) from the second capture) were considered diabetic (blood glucose levels \( > 200 \) mg/dL after 6 weeks of diet) and termed high-protein diet diabetic (HDD). The remaining animals from each of the hypercaloric groups (\( n = 5 \) from the first capture, \( n = 6 \) from the second capture; \( n = 11 \) total) were classified as high-protein diet nondiabetic (HDDN). At the end of the experimental periods (4 and 7 months, respectively, for the two series), animals were euthanized by CO\(_2\) inhalation and decapitation. Eyes were either fixed in 4% paraformaldehyde in PBS (0.01 M, pH 7.4) for 12 hours at 4°C or the retinas and vitreous bodies were dissected free under a binocular microscope and snap frozen in liquid nitrogen. Samples were processed for sectioning on a cryostat (CM 3050 S; Leica, Wetzlar, Germany) using standard protocols. For flat-mounted preparations, after fixation as described above, the eyeballs were opened at the limbus, and the cornea and lens were discarded. The retinas were carefully separated from the posterior eyecup composed of the retinal pigment epithelium (RPE), choroid, and sclera, and both tissues were washed in PBS. Radial incisions were made in the retinas and eyecups to permit flattening.

### Immunohistochemistry

Sections were processed for immunohistochemistry using conventional procedures. Primary antibodies are listed in Table 1 and were diluted in saturation buffer (PBS containing 0.1% bovine serum albumin and 0.1% Tween 20) and left overnight at 4°C. Sections were washed and then incubated with the appropriate secondary antibodies, either goat anti-rabbit or anti-mouse IgG-conjugated antibodies coupled with Alexa 594 or Alexa 488 (Molecular Probes Ltd., Eugene, OR) diluted 1:400. Cell nuclei were stained with 4,6-diamino-phenyl-

#### Table 1. Antibodies Used in Immunohistochemistry and Western Blot Analysis

<table>
<thead>
<tr>
<th>Antibody/Antigen</th>
<th>Type</th>
<th>Retinal Staining</th>
<th>Dilution</th>
<th>Manufacturer/Supplier</th>
</tr>
</thead>
<tbody>
<tr>
<td>VEGF</td>
<td>Monoclonal (mouse)</td>
<td>—</td>
<td>WB 0.1 µg/mL</td>
<td>Santa Cruz Biotechnology, Santa Cruz, California</td>
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<tr>
<td>PEDF</td>
<td>Polyclonal (rabbit)</td>
<td>—</td>
<td>WB 0.1 µg/mL</td>
<td>Kind gift of Joyce Tombran-Tink, Pennsylvania State College of Medicine, Hershey, PA</td>
</tr>
<tr>
<td>α-Smooth muscle actin</td>
<td>Monoclonal (mouse)</td>
<td>Pericytes (blood vessels)</td>
<td>IHC 1 µg/mL</td>
<td>Sigma Aldrich Life Sciences, Cergy Pontoise, France (A7607)</td>
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<tr>
<td>GFAP (whole protein from cow)</td>
<td>Polyclonal (rabbit)</td>
<td>Astrocytes/Müller glia</td>
<td>IHC 2 µg/mL</td>
<td>DAKO A/S, Glostrup, Denmark (Z 0334)</td>
</tr>
<tr>
<td>Glutamine synthetase (mouse peptide)</td>
<td>Monoclonal (mouse)</td>
<td>Glia</td>
<td>WB 0.2 µg/mL</td>
<td>Millipore (mAb 502)</td>
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<tr>
<td>GLAST-1 (rat C-terminal peptide)</td>
<td>Polyclonal (rabbit)</td>
<td>Glia</td>
<td>WB 0.2 µg/mL</td>
<td>Alpha Diagnostic International (GLAST 11-S)</td>
</tr>
<tr>
<td>Neurofilament (200 kDa) pPKC ζ (human peptide)</td>
<td>Monoclonal (mouse)</td>
<td>RGC, HC</td>
<td>IHC 1 µg/mL</td>
<td>Sigma (N-20)</td>
</tr>
<tr>
<td>PKCa</td>
<td>Monoclonal (mouse)</td>
<td>Rod and cone ON BC</td>
<td>IHC 5 µg/mL</td>
<td>Santa Cruz Biotechnology, sc-12894-R (Thr 410)</td>
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<tr>
<td>Parvalbumin</td>
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<td>Rod and cone ON BC</td>
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<td>Synaptophysin (rat protein)</td>
<td>Monoclonal (mouse)</td>
<td>HC</td>
<td>IHC 1 µg/mL</td>
<td>Sigma P3088</td>
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<tr>
<td>Rhodopsin (Rho4D2)</td>
<td>Monoclonal (mouse)</td>
<td>ROS +++, and RCB+</td>
<td>IHC 1 µg/mL</td>
<td>Santa Cruz Biotechnology, sc-12737 (SVP38)</td>
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<tr>
<td>MW cone opsin (mouse peptide)</td>
<td>Polyclonal (rabbit)</td>
<td>COS (MW cones)</td>
<td>WB 0.1 µg/mL</td>
<td>Kind gift of Robert Molday, University of British Columbia, Vancouver, BC, Canada</td>
</tr>
<tr>
<td>SW cone opsin (mouse peptide)</td>
<td>Polyclonal (rabbit)</td>
<td>COS (SW cones)</td>
<td>IHC 1 µg/mL</td>
<td>Kind gift of Robert Molday, University of British Columbia, Vancouver, BC, Canada</td>
</tr>
<tr>
<td>Peripherin (rat protein)</td>
<td>Monoclonal (mouse)</td>
<td>OS, CB</td>
<td>IHC 1 µg/mL</td>
<td>Kind gift of Robert Molday, National Institutes of Health, Bethesda, Maryland</td>
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<tr>
<td>Arrestin (cow protein)</td>
<td>Polyclonal (rabbit)</td>
<td>OS</td>
<td>IHC 1 µg/mL</td>
<td>Kind gift of Alexander Dizhoor, Northwestern University, Chicago, Illinois</td>
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<tr>
<td>Recoverin</td>
<td>Polyclonal (rabbit)</td>
<td>OS, CB, BC</td>
<td>IHC 1 µg/mL</td>
<td>Kind gift of Cheryl Craft, Doheny Eye Institute, University of Southern California, Los Angeles, California</td>
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<tr>
<td>Ga t1 (rod transducin)</td>
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<tr>
<td>Ga t2 (cone transducin)</td>
<td>Polyclonal (rabbit)</td>
<td>COS, CCB</td>
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The table shows antibody/antigen type, cellular localization within the retina (if used in IHC), working dilution for IHC and WB, and commercial origin or reference. WB, Western blot; IHC, immunohistochemistry; BC, bipolar cells; (C/R)OS, (cone/rod) outer segments; (C/R)CB, (cone/rod) cell bodies; HC, horizontal cells; RGC, retinal ganglion cells.
indolamine (DAPI; Sigma Aldrich Life Sciences, Cergy Pontoise, France) diluted 1:500. Incubation was performed at room temperature for 2 hours, and sections were washed thoroughly in PBS and coverslipped. Retinal and posterior eyecup wholemount immunohistochemistry was performed as described previously. Some tissues were incubated in lectin from Bandeiraea simplicifolia directly conjugated to tetramethylrhodamine isothiocyanate (25 μg/mL; Sigma Aldrich Life Sciences), or to phalloidin- tetramethylrhodamine isothiocyanate (100 μg/mL; Invitrogen-Molecular Probes, Eugene, OR) or a confocal microscope (LSM510 version 2.5; Carl Zeiss Meditec, Jena, Germany). Photographs were taken with identical exposure times between control and treated specimens. Any graphics editing of images (Photoshop; Adobe, San Jose, CA) was performed using identical parameters between the two sets. Tissues from a minimum of three animals per group (ND, HDnD, and HDD) were examined for each condition.

**Western Blot Analysis**

Western blot analysis was performed on proteins extracted from entire vitreous and retinas (n = 3 per group), as described. After incubation on ice for 30 minutes, proteins were suspended in Laemmli buffer and separated by SDS-PAGE (4%–10%; 40–50 μg protein/lane; total protein concentrations were initially determined using the Bradford colorimetric assay) and then transferred onto a polyvinylidene fluoride membrane. Membranes were incubated with primary antibodies (Table 1) overnight at 4°C under agitation, washed, and incubated with corresponding secondary antibodies coupled to horseradish peroxidase (1:10,000; Jackson ImmunoResearch Laboratories, West Grove, PA). Immunoreactive bands were detected by chemiluminescence (Immobilon Western; Millipore, Bedford, MA). Relative abundance was quantified by scanning densitometry (ImageJ software, developed by Wayne Rasband, National Institutes of Health, Bethesda, MD; available at http://rsb.info.nih.gov/ij/index.html), and expressed relative to tubulin immunoreactive bands in the same blots (detected after membrane stripping and reprobing with monoclonal antitubulin antibody, diluted 1:10,000 [Jackson ImmunoResearch Laboratories]). Experiments were performed on extracts from a minimum of three retinas from separate animals for each group.

**Measurement of Retinal Vasculature**

The entire vasculature, from the optic nerve head toward the periphery, was visualized on retinal flatmounts under a fluorescence microscope (Optiphot 2; Nikon). The entire field was mapped using rectangular grids (810.87 μm²), and the surface was divided into nine concentric zones (see Results; Fig. 3E). The retinal vascular caliber was measured along the entire length of each major vessel, at 10 regularly spaced points (~5 μm) for each rectangle, for three separate retinas from ND, HDnD, and HDD animals. We further measured the number of branch points in the most peripheral zone (lying beyond zone 9) for three retinas from each ND and HDD group, averaged per square millimeter.

**Trypsin Digestion**

To isolate the retinal vascular bed from fixed retinas, we followed published procedures. Quantification of acellular cordlike capillaries was performed using the criteria described by these authors, under 40× microscope objectives, using a 100 × 100 μm grid superimposed over 10 random fields of HD and HDD preparations. Measurement of major vessel thickness was performed as described, beginning at the widest (central) end of isolated vessels.

**Cell Counting**

The density of cell nuclei and cell layer thickness was determined for each condition (three fields of each of three animals per group) by light microscope examination of retinal sections. Images were obtained within the posterior retina, within 1 mm of the optic nerve head. Cell layer thickness and number of nuclei per layer were calculated within a standard rectangle placed across captured images, equivalent to an area measuring 0.07 mm².

**Statistical Analysis**

Results are expressed as mean ± SD. Comparisons of the two groups (cell counting and Western blot analyses) were performed using factorial analysis of variance (ANOVA) followed by post hoc Tukey HSD test. P < 0.05 was considered statistically significant. Calculations were performed using statistical analysis software (Statistica 8.0; StatSoft, Maisons-Alfort, France). For vessel parameter analysis, we used two-way ANOVA for independent and repeated measures, followed by post hoc Tukey HSD test.

**RESULTS**

**Development of Obesity and Hyperglycemia**

Figure 1 depicts results from the second capture, involving 24 animals. Whereas body mass and blood glucose levels re-
mained stable over time in captivity in plant-fed animals (ND; \( n = 8 \)), many animals raised on standard rat chow (total \( n = 16 \)) became obese and developed hyperglycemia (HDD; \( n = 10 \)). Six animals on high-calorie diets did not show increased blood glucose levels (except slight increases during the final month of captivity, which were not significantly different from the ND values), although they showed intermediate values of weight gain (HdND; Fig. 1). Values for body weight (Supplementary Table S1, http://www.iovs.orglookup/suppl/doi:10.1167/iovs.11-8423/-/DCSupplemental) and blood glucose (Supplementary Table S2, http://www.iovs.orglookup/suppl/doi:10.1167/iovs.11-8423/-/DCSupplemental) for each animal as a function of time in captivity are given in Supplementary Data, http://www.iovs.orglookup/suppl/doi:10.1167/iovs.11-8423/-/DCSupplemental. Although ND animals exhibited a stable weight of \( \sim 115 \) g throughout the experimental period, HDD animals gained 10 to 20 g per month to reach \( \sim 250 \) g at 7 months. HdND animals gained \( \sim 10 \) g per month to reach \( \sim 180 \) g in the same period. ND animals had \( \sim 80 \) mg/dL (5 mM) blood glucose levels throughout the time in captivity, whereas HDD animals showed permanently increasing concentrations with monthly increments of \( \sim 100 \) mg/dL to reach 400 mg/dL (25 mM) by 7 months (Fig. 1B). Hematocrit values for the three groups did not vary significantly over the course of captivity or between groups (mean \( \pm 4\% \) for both; data not shown). Results from the first capture were largely similar, with ND, HdND, and HDD groups (\( n = 5 \) in each group) maintained for 4 months (Supplementary Fig. S1, Table S1, http://www.iovs.orglookup/suppl/doi:10.1167/iovs.11-8423/-/DCSupplemental). Although captivity was shorter, the HDD group increased body weight by approximately 80% and blood glucose by \( \sim 550\% \) compared with starting values (ND group = 10% weight gain, 0% glucose increase; HdND group = 40% weight gain, 0% glucose increase over same period).

**Vitreal Protein Analysis in *P. obesus***

Total protein concentrations in HDD vitreous were approximately 2.2-fold greater than those of ND animals, with mean values of 0.32 and 0.71 \( \mu \)g/\( \mu \)L in ND and HDD groups, respectively (Table 2). Quantitative Western blots of vascular endothelial growth factor (VEGF) and pigment epithelium derived growth factor (PEDF) from each group showed highly significant statistical differences in both cases, with VEGF values \( \sim 8\)-fold higher, and PEDF values \( \sim 100\)-fold lower, in HDD animals (Fig. 2). VEGF/PEDF ratios calculated from these data showed very large differences (up to 340-fold) between the two groups (Table 2). VEGF and PEDF concentrations were not studied in HdND animals.

**Table 2. Vitreal Protein, Proangiogenic, and Antiangiogenic Growth Factor Levels in ND and HDD *P. obesus***

<table>
<thead>
<tr>
<th></th>
<th>ND (( n = 8 ))</th>
<th>HDD (( n = 10 ))</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vitreous protein levels, ( \mu )g/( \mu )L</td>
<td>0.32 \pm 0.01</td>
<td>0.71 \pm 0.03*</td>
</tr>
<tr>
<td>Mean VEGF/PEDF ratio</td>
<td>0.239 \pm 0.007</td>
<td>71.364 \pm 7.686*</td>
</tr>
</tbody>
</table>

Values are mean \( \pm \) SD. Measurement of total soluble protein content of vitreous isolated from ND and HDD animals showed highly significant differences, with 2.2-fold greater levels in diabetic samples. As calculated from Western blots using anti-VEGF and anti-PEDF, ratios of the two proteins were calculated from ND (\( n = 8 \)) and HDD (\( n = 10 \)) individual vitreal samples.

* \( P < 0.001 \).

Examination of isolectric-labeled blood vessels in flat-mounted whole retinas showed a clear demarcation into large retinal vessels (arteries and veins) and retinal capillaries (Fig. 5). Although blood vessels in the ND group showed a regularly branching uniform aspect (Fig. 3A), HDD retinas showed a variety of abnormal features: vessels displayed an irregular width with alternating constrictions and dilations (Fig. 3B) as well as focal vascular sprouts with blind ends (Fig. 3C) and balloon-like swellings resembling microaneurysms (Fig. 3D). Quantitative measures of major blood vessel diameter as a function of distance from the optic nerve head (Fig. 3E) showed that mean values for HDD were always higher than for ND retinas and were statistically significantly different in the midperiphery (Fig. 3F). Mean values for HdND blood vessels were intermediate but were not statistically significantly different from ND values (Fig. 3F). Quantification of branch points in the far peripheral capillaries also showed highly significant differences between ND (119.7 \( \pm \) 19.2/mm\(^2\)) and HDD (225.5 \( \pm \) 22.8/mm\(^2\); 88% increase; \( P < 0.001 \)). Similar analyses of HdND retinas revealed intermediate values, which, although higher than plant-fed animals, did not attain statistical significance (190.7 \( \pm \) 29.9/mm\(^2\); \( P = 0.059 \)). Analysis of the retinal vasculature after trypsin digestion revealed a regular network of uniform capillaries in ND tissue (Fig. 3G) but the presence of thin cord-like acellular strands in HDD samples (Fig. 3H). Similar measures of major blood vessel diameter as performed in intact flat-mounted retinas also showed differences in ND and HDD thickness in isolated vascular net preparations (Fig. 3I). Furthermore, counting of acellular strands in these trypsin digests confirmed the large differences between ND and HDD samples (Fig. 3J). Trypsin digest experiments were not performed for HdND retinas.

Staining of ND retinal flat-mounts with isoelectric (to outline vessels) and smooth muscle actin antibody (to label pericytes) revealed that ND blood vessels were decorated densely with pericytes (Figs. 4A, 4B). In contrast, although HDD retinas showed extensive isolectric staining of vessels (Fig. 4C), there was almost complete absence of pericytes (Fig. 4D). At the level of the blood-retina barrier within the posterior pole, phalloidin staining of Factin bundles within flat-mounted ND eye cups revealed a highly regular honeycomb-like aspect of the RPE monolayer, formed of tightly linked hexagonal cells compared with the irregularity of the HDD retinas, which showed extensive isolectric staining of vessels. The far peripheral capillaries also showed highly significant differences between ND and HDD retinas (Fig. 3E).
ple by tight junctions (Fig. 4E). On the other hand, phalloidin staining of HDD eyecups showed weakened junction, as evidenced by splitting between neighboring cells (Fig. 4F). ND eyecups also displayed strict localization of the tight junction marker protein kinase C (PKCζ) to cell borders (Fig. 4G), whereas HDD eyecups showed generalized staining across the cell indicative of delocalization (Fig. 4H); the same was true for another tight junction marker, occludin (data not shown). Additional examples of pericyte loss in diabetic retinas are shown in Supplementary Fig. S1, http://www.iovs.org/lookup/suppl/doi:10.1167/iovs.11-8423/-/DCSupplemental. Pericyte and RPE barrier immunostaining were not performed for HDnD retinas.

**General Structural Modifications in* P. obesus**

The general retinal aspect of the three groups showed clear differences. ND retinas exhibited regular layers with a thin inner limiting basement membrane (Fig. 5A, 5B), whereas sections of HDD retina exhibited a wider nerve fiber layer with a membranous aspect (Fig. 5C) and displayed thin irregular nuclear layers (Fig. 5D). This was also observed in HDnD retinas (data not shown). Quantitative measures showed that the widths of the outer segment (OS) layer and that of the outer nuclear layer (ONL), inner nuclear layer (INL), and inner plexiform layer (IPL) were significantly reduced in HDnD and HDD retinas (by up to 40%, 36%, 38%, and 22% respectively) (Fig. 5E). Similar reductions were noted when cell density was counted within the ONL (up to 26% less), INL (up to 30% less), and ganglion cell layer (GCL) (up to 44% less) (Fig. 5F).

In support of these histologic observations, plexiform layers were clearly visualized by synaptophysin immunostaining of synaptic vesicles. Immunolabel of the two synaptic zones was different between ND and HDD groups. In ND retinas, the synaptic layers were regular and linear (Fig. 5G), whereas in HDD tissue, the OPL was irregular and the IPL was markedly thinner (Fig. 5H). However, and in contrast to other proteins, synaptophysin levels were significantly increased in diabetic retinas (Fig. 5I). Synaptophysin immunoreactivity was not analyzed in HDnD retinas.

**Retinal Glial Modifications in* P. obesus**

Visualization of glial fibrillar acidic protein (GFAP) immunostaining at the vitreal surface of flat-mounted ND retinas showed filamentous staining, particularly marked around blood vessels (Fig. 6A). However, GFAP staining intensity appeared higher in HDnD retinas and much higher in HDD retinas, both along and between blood vessels (Figs. 6B, 6C). Examination of frozen sections from each group showed that GFAP expression was confined to the nerve fiber layer but with more intense staining because of the hypertrophy in HDD (data not shown). Western blot analysis of protein extracts from each group showed a small but significant increase in intensity ending (C). Using a morphometric approach in which major blood vessels were traced from the center to the periphery and were divided into nine equivalent zones in which repeated measures of vessel width were made (E), a significant increase in the midperiphery of HDD vessels was observed (F, filled circles) (*P < 0.05, ANOVA). HDnD vessels (filled squares) showed diameters intermediate between those of ND (filled diamonds) and HDD tissues, but these were not statistically significant. Trypsin-digested preparations of ND (G) and HDD (H) tissue showed the presence of cord-like acellular strands in the disintegrated tissue (white arrows). Quantitative analysis of vessel caliber in major blood vessels in trypsin digests also showed a significant increase in HDD tissue (**P < 0.001). Quantification of acellular cords in trypsin digests showed them to be restricted to HDD tissue (**P < 0.001). Scale bars: 100 μm (A, B, E, G, H); 25 μm (C, D).
in the HDnD group compared with the ND group and increases of approximately twofold in HDD compared with ND (Fig. 6D).

In stark contrast to GFAP immunoreactivity, staining for glutamine synthetase (GS) showed an inverse pattern. In ND retinas, labeling was intense across the tissue, outlining the cell bodies in the three cellular layers and filling the plexiform layers (Fig. 6E). HDnD retinas were similar but appeared less intense (Fig. 6F), whereas staining was greatly reduced in sections of HDD retina, particularly in the outer half of the tissue, with radial staining seen in individual Müller glia (Fig. 6G). Again, these qualitative differences were confirmed by Western blot analysis and were seen to be highly significantly different between ND and HDD samples (Fig. 6H). Glutamate-aspartate transporter (GLAST) immunoreactivity was also intense in ND retinas, extending from the Müller glial apical microvilli through the nerve fiber layer (Fig. 6I). HDnD retinas displayed similar patterns, although there appeared to be less staining in the INL (Fig. 6J), whereas intensity was much reduced in HDD retinas (Fig. 6K). Western blot analysis showed the difference was statistically significant between ND and HDD retinas (Fig. 6L).

Retinal Neuronal Modifications in Diabetic P. obesus

Retinal ganglion cells (RGCs) were visualized by staining against the 200-kDa subunit of neurofilament, in both retinal wholemounts and frozen sections. As viewed in retinal wholemounts, axonal morphology differed between control and diabetic groups. Axons were smooth and organized into smooth bundles in ND tissue (Fig. 7A) but exhibited numerous varicosities and were often seen as individual fibers in HDD tissue (Fig. 7B). Frozen sections of retinas from ND and HDD animals showed a fairly similar pattern, with strong immunostaining along the nerve fiber layer visible as discrete axon bundles in ND (Fig. 7C) but more diffuse in HDD (Fig. 7D). HDnD RGCs resembled those seen in ND retinas (data not shown). Bipolar cells were stained with antibodies against PKCα and PKCζ (both restricted to this cell type within the neural retina). In ND tissue, PKCα immunoreactivity was present in cell bodies, axons, and synapses of this population (Fig. 7E). The diabetic state led to a marked reduction in staining intensity so that only background staining was seen in HDD retinas (Fig. 7F). Quantification of PKCα by quantitative Western blot analysis confirmed a significant reduction in HDD retinas (Fig. 7G). Very similar observations were made for PKCζ, with immunostaining visible throughout the bipolar cells in ND (Fig. 7H) and HDnD (albeit at reduced intensity) retinas (Fig. 7I) but was reduced to faint synaptic staining in HDD tissue (Fig. 7J). These qualitative results were confirmed by quantitative Western blot analysis, with HDnD samples showing significant reductions and HDD tissue showing highly significant reductions (Fig. 7K). Horizontal cell staining with parvalbumin antibody was not visibly different between the groups (Figs. 7L, 7M).

Retinal Photoreceptor Modifications in Diabetic P. obesus

There were notable differences in the responses of the two photoreceptor populations (rods and cones) to the diabetic-like state. Expression of rhodopsin did not differ in intensity or distribution between the three groups (Figs. 8A–C). However, cone mid-wavelength opsin (MOp) staining was detectably reduced in HDD compared with ND and HDD retinas (Figs. 8D–F). This was also the case for cone short-wavelength opsin (SOp) (Figs. 8G, 8H). These qualitative observations were all confirmed by Western blot analysis. Rhodopsin immunostaining was equal between the groups (Fig. 8I), whereas MOp and SOp were reduced to approximately 50% the levels in HDD.
General structural modifications in *P. obesus* retina. General histologic aspect of ND (A) and HDD (C) retinas, with corresponding DAPI-stained nuclear structures (B, ND; D, HDD). The reduced width of HDD retinas is apparent, together with the less orderly organization of the cellular layers. HDD retinas showed a membranous appearance of the nerve fiber layer in many sections (C, asterisk). Quantitative measures of retinal layer thickness (E) and cell density (F) in ND (black columns), HDnD (gray columns), and HDD (white columns) animals. There were significant decreases in layer thicknesses (OS, ONL, INL, IPL) and cell numbers (ONL, INL, GCL) in HDD compared with ND animals (P < 0.05; ANOVA). HDnD measures were also significantly different from ND values for ONL, INL, and IPL for layer width and for INL and GCL for cell density (P < 0.05; ANOVA). Synaptophysin (SVP38) immunostaining was intense in both plexiform layers in ND (G) and HDD *P. obesus* (H). Western blot analysis and quantification of synaptophysin expression showed it was significantly increased in HDD retinas (I) (*P < 0.05, ANOVA). OS, outer segments. Scale bar, 50 μm (A–D, G, H).

**FIGURE 5.** General structural modifications in *P. obesus* retina. General histologic aspect of ND (A) and HDD (C) retinas, with corresponding DAPI-stained nuclear structures (B, ND; D, HDD). The reduced width of HDD retinas is apparent, together with the less orderly organization of the cellular layers. HDD retinas showed a membranous appearance of the nerve fiber layer in many sections (C, asterisk). Quantitative measures of retinal layer thickness (E) and cell density (F) in ND (black columns), HDnD (gray columns), and HDD (white columns) animals. There were significant decreases in layer thicknesses (OS, ONL, INL, IPL) and cell numbers (ONL, INL, GCL) in HDD compared with ND animals (P < 0.05; ANOVA). HDnD measures were also significantly different from ND values for ONL, INL, and IPL for layer width and for INL and GCL for cell density (P < 0.05; ANOVA). Synaptophysin (SVP38) immunostaining was intense in both plexiform layers in ND (G) and HDD *P. obesus* (H). Western blot analysis and quantification of synaptophysin expression showed it was significantly increased in HDD retinas (I) (*P < 0.05, ANOVA). OS, outer segments. Scale bar, 50 μm (A–D, G, H).

**DISCUSSION**

Diet-induced metabolic stress syndrome in the sand rat *P. obesus* leads to a number of pathologic alterations in their eyes. These modifications include a range of abnormal features associated with control of the vasculature (within the vessels themselves: vasodilation, increased tortuosity, vascular sprouts, occluded vessels and small swellings; increased total protein and radically opposing concentrations of VEGF and PEDF within the vitreous, splitting of the RPE monolayer, and delocalization of tight junction proteins forming the blood-retina barrier). There was an extensive series of remodeling events and phenotypic changes in retinal glia and neurons. To the extent to which data are available for human DR, the two conditions are remarkably similar.

There is extensive literature on the use of *P. obesus* as an experimental model for investigating the development and effects of metabolic syndrome leading to obesity, dyslipidemia, hypertension, and type 2 diabetes.18 Studies indicate that overly rich diets may lead to the development of hyperinsulinemia through ineffective extraction of excess insulin by liver receptors. Hyperinsulinemia, in turn, causes a reversible reduction in receptor kinase activity, leading to insulin resistance.19 Furthermore, marked insulin depletion is observed in diabetic islets, probably because of a hyperglycemia-driven increase in secretory demand that cannot be compensated.20 The similarities among metabolic, physiological, and endocrine changes in this species compared with those in human type 2 diabetes make it a highly relevant animal model with which to understand the pathogenesis of this condition.12,13,18 In the present study, >50% of the animals maintained on hypercaloric diets spontaneously developed characteristics of diabetes (obesity and elevated blood sugar levels), and all exhibited DR. The intermediate group was not examined for every parameter, but it displayed a variable phenotype ranging from no difference to large differences compared with the plant-fed group. This group experienced significant weight gains without elevated blood glucose levels and also displayed moderate retinal symptoms, as indicated by reduced width and cell numbers, in-
creased expression of GFAP, and decreased expression of PKC. However, most of these changes were smaller than those seen in HDD animals and were not significantly different for GS or mid- or long-wavelength cone opsin. This intermediate group may represent latent animals who responded to dietary imbalances with delayed-onset or refractory diabetes compared with the HDD group. Similar percentages of diabetes-susceptible (40%) and diabetes-resistant animals have been reported in previous studies on Psammomys,21 and analysis of the two populations has permitted identification of a major susceptibility gene.22

A major question concerns the potential advantages and disadvantages of this new animal model for DR. According to the Animal Models of Diabetic Complications Consortium (AMDCC), useful models should exhibit capillary and neuronal loss and capillary obliteration (features of early-stage nonproliferative diabetic retinopathy [NPDR]), retinal edema, and pre-retinal neovascularization (associated with PDR). Many experimental models have been developed and characterized for their pertinence to the human disease, but most replicate only aspects of NPDR. Some do progress to PDR, which is responsible for severe vision loss but at advanced age (e.g., spontaneously diabetic Torii rats >50 weeks of age).23 Diet-induced DR in Psammomys satisfies most of the criteria cited by the AMDCC. It is typified by a range of vascular abnormalities and widespread neuronal loss, and the presence of acellular capill-

**Figure 6.** Glial alterations in retinas of *P. obesus*. Immunohistochemical and Western blot analyses of glial fibrillar acidic protein (GFAP) (A–D), glutamine synthetase (GS) (E–H), and GLAST (I–L). GFAP-containing astrocytes were distributed across the vitreal face of flat-mounted ND retinas, especially concentrated along major blood vessels (A). Similar views of HDnD retinas showed intense staining around blood vessels and higher amounts between vessels (B). Observation of the vitreal face of HDD retinas showed very high numbers of GFAP-immunopositive astrocytes across the surface (C). Increased GFAP expression in HDnD and HDD animals was confirmed by densitometric scanning of Western blots, expressed in arbitrary units (a.u.) normalized to tubulin (*n* = 3 per retina, shown above each corresponding column) (D). Compared with ND samples (black columns), there were large increases in HDnD (gray columns; **P < 0.01, ANOVA) and HDD (white columns. ***P < 0.001, ANOVA). GS immunoreactivity was robust across the entire retina in ND (E) and HDnD (F) animals but was greatly reduced in HDD animals (G). Western blot analysis and densitometric scans confirmed the decreased expression for GS (H). Triplicate Western blots for GS and tubulin are shown above each column. GLAST immunostaining was also strong in ND (I) and HDnD (J) retinas and decreased in HDD retinas (K). As for GS, quantitative Western blots confirmed the decrease (L), and triplicate Western blots for GLAST and tubulin are shown above the columns (**P < 0.001, ANOVA). MV, Müller glial microvilli; OS, outer segments; Tub, tubulin. Scale bars: 20 μm (A–C); 50 μm (E–G, I–K).
laries indicates the formation of obliterated nonfunctional microvessels, as seen in humans. Psammomys showed dramatic pericyte loss in diabetic retinas, with almost complete disappearance from vessel surfaces. Pericyte dropout represents the earliest detectable vascular change in animal models of DR and is a hallmark feature of human DR. In the present study, we did not specifically examine preretinal hemorrhage and neovascularization, but two observations argue that it develops in our model: increased vitreal protein concentrations indicate leakage from new blood vessels, and the huge imbalance in proangiogenic and antiangiogenic factors measured in the vitreous of diabetic animals is strongly correlated with the development of PDR in humans.

We especially focused on nonvascular aspects of DR because these often precede blood vessel remodeling and may be causal events in functional decline. Such alterations are difficult to investigate in animal models, but they are likely to play a role in the pathogenesis of diabetic retinopathy. For example, increased VEGF/PEDF ratios were directly correlated with the degree and progression of retinal neovascularization in another pathologic mouse model. We did not investigate retinal edema in the present study because eyes were dissected and fixed before examination. To the best of our knowledge, only one animal model has so far been described that exhibits PDR: the Akimba mouse, a genetic model of type 1 DR (double transgenic Ins2αkita/VEGF mice). It should be emphasized that unlike the Akimba mouse, the current model is generated solely by mild dietary overload, with no genetic overexpression of VEGF, and constitutes a faithful model of metabolic syndrome/type 2 diabetes. Furthermore, the observed changes occurred within 7 months of hypercaloric diet, considerably faster than the passage to PDR in other animal models.

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**Figure 7.** Neuronal modifications in *P. obesus* retina. (A) Flat-mounted ND retina immunostained with neurofilament (NF200 kDa) antibody showing smooth axon bundles (arrow). (B) On the contrary, axons in HDD retinas revealed moderate defasciculation and numerous varicosities (arrow). A similar result was seen in frozen cross-sections, with large neurofilament-immunopositive bundles seen along the nerve fiber layer in ND samples (C) but thinner variable staining in HDD animals (D). In ND retinas, PKCa antibody showed clear staining in cell bodies within the inner nuclear layer (INL), as well as their axons and synaptic endings in the inner plexiform layer (IPL), corresponding to rod and cone ON bipolar cells (E). HDD retinas showed almost complete disappearance of PKCa staining (F), which was confirmed by quantitative Western blot analysis as described earlier (triplicate lanes for ND and HDD retinas, top) (G). PKCζ immunostaining revealed a pattern similar to that of PKCa in ND retinas (H) and a less intense signal in HDnD retinas (I), whereas in HDD retinas residual staining was visible in the synapses (J). Reduced staining was again confirmed by Western blot analysis (K). Quantitative densitometric measurement of band densities compared to tubulin (Tub) confirmed the decreases in HDnD (*P < 0.01, ANOVA) and HDD (*P < 0.001, ANOVA) samples. Triplicate blots of PKCa and tubulin are shown above each corresponding column: ND (black), HDnD (gray), HDD (white). Histologic sections of *P. obesus* retina immunostained with parvalbumin antibody were very similar in ND (L) and HDD (M) animals. OS, outer segments. Scale bars: 20 μm (A, B); 50 μm (C–F, H–J); 80 μm (L, M).
to study in a methodical manner in humans. We paid great attention to cone photoreceptor changes because deficits in color vision and contrast sensitivity are among the earliest indications of DR in humans, occurring long before clinically detectable vascular defects.5–7 We saw no significant changes in rhodopsin levels between control and diabetic animals. In contrast, both short-wavelength and mid-/long-wavelength cones were strongly affected, with significant reductions in both populations. Importantly, cone opsin expression was not altered in the obese nondiabetic group, indicating it is causally linked to the hyperglycemic state. Perturbations in the blue-yellow axis have been particularly implicated in DR,7 and blue cone dropout has been observed in postmortem human DR.32 Recent studies33 in a zebrafish model of type I NPDR revealed that cones were highly sensitive to hyperglycemia, independent of vascular changes. Differential susceptibility of rods and cones in hyperglycemic conditions might be attributed to glucose use; cone physiology is known to have a higher dependence on correct glucose balance.54 With respect to permitting detailed examination of the pathogenesis involved in cone degeneration, P. obesus offers the major advantage of a high cone percentage (~40%)35 compared with conventional laboratory rats and mice with <5% cones.36,37 Although there are clear differences in diurnal rodent and human retina (e.g., rodents do not have a macula and lack a distinct long-wavelength cone), their rod/cone ratio and axonal density in the optic nerve resemble human central retina (Matter JM, personal communication, 2011). Their retinas have a structural organization that facilitates quantitative measurement of rod and cone density and survival,16,38 and visual function can be readily followed by standard electroretinography.38 In the latter, diurnal rodents show robust photopic responses to high-frequency flicker electroretinography, similar to that of humans but not seen in mice and rats.38

It is likely that glial malfunction is a major factor in DR. Retinal Müller glial cells are clearly implicated in its pathogenesis,39 perturbation of their function impinges directly on neuronal survival,40 and they are implicated in insulin signaling and glucose metabolism in the retina.40 In the present study, retinal glia displayed marked downregulation of two key proteins involved in glutamate detoxification, GLAST and glutamine synthetase (GS). Although GS showed no differences between normal and diabetic postmortem human retinas,41 numerous studies in rodent models show clear reductions of both.42,43 Furthermore, elevated vitreal glutamate concentrations are found in both animal44 and human27 DR, indicating a failure of glutamate metabolism. The large upregulation of the glial stress marker GFAP is commonly seen in different models.

FIGURE 8. Rod and cone photoreceptor opsins in P. obesus retina. Rhodopsin (Rho) immunolabeling showed strong expression in rod outer segments (OS, green) in ND (A), HDnD (B), and HDD (C) animals. Staining of cone mid wavelength opsin (MOp) was localized to numerous cone outer segments (OS, red) in ND (D) and HDnD (E) animals, whereas there were many less immunolabeled cells in HDD tissue (F). This was also the case for cone short wavelength (SOp) immunostaining in OS of ND (G) and HDD animals (H). Western blot analysis and quantification of Rho, MOp, and SOp expression. (I) Rho and tubulin (Tub) immunoreactivity showed no difference in intensity between samples. Triplicate samples are shown above corresponding columns: ND (black), HDnD (gray), HDD (white). (J) MOp immunoreactivity showed a roughly 50% decrease in HDD retinas when normalized to Tub (***P < 0.001, ANOVA). (K) Similarly, SOp immunoreactivity was also reduced by 50% in HDD retinas (***P < 0.001, ANOVA). Scale bar, 80 μm.
of retinal degeneration. Some reports show increased expression of GFAP in human DR, and all describe animal models of DR that demonstrate an upregulation of GFAP.

We observed neuronal loss in all layers, as occurs during DR. However, because these decreases were primarily also seen in obese nondiabetic animals, they may not be linked to elevated glucose levels but to some other aspect of metabolic syndrome. Ganglion cells are the best-studied neuronal type in diabetes, and several reports show cell loss and axonal damage in both human and animal models of DR. The ganglion cell axonal swellings seen in the present study have been observed in human DR, and the Ins2Akita transgenic mouse model of DR and are considered to indicate axonal degeneration. To the best of our knowledge, this is the first report of diabetes-induced decreases in PKC expression in bipolar cells and may be related to perturbed inner retinal function seen in DR, such as alterations in oscillatory potentials.

In conclusion, *P. obesus* represents a potentially valuable novel animal model of human type 2 DR. Not only is disease induction (enriched diet) very similar to a major cause of human type 2 diabetes, but the wide spectrum of diabetic modifications is not seen in other laboratory models and arises relatively rapidly. Alterations in cone-mediated color vision constitute one of the earliest characteristics of human DR and may represent a “biomarker” for early diagnosis. The animal presents a major advantage in terms of analyzing the pathogenic mechanisms underlying structural and functional cone degeneration. We are undertaking functional studies of retinal visual responses to diabetes and mapping the progression of retinal changes in respect to the duration of hyperglycemia. The model will furthermore permit testing of potential therapeutic strategies such as insulin treatment, dietary control, and pharmacologic intervention.

**Acknowledgments**

The authors thank Patricia Tiazzzi (INSERM U872, Paris, France) for helpful discussion, Jorgé Mendoza (Département de Neurobiologie des Rythmes, CNRS UPR 3212, INCI, Strasbourg, France) for assistance with statistical analysis, and Claudine Boissier for expert technical assistance.

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