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Control of Patterns of Corneal Innervation by Pax6

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Abstract

Purpose—Corneal nerves play essential roles in maintaining the ocular surface through provision of neurotrophic support, but genetic control of corneal innervation is poorly understood. The possibility of a neurotrophic failure in ocular surface disease associated with heterozygosity at the Pax6 locus (aniridia-related keratopathy [ARK]) was investigated.

Methods—Patterns of corneal innervation were studied during development and aging in mice with different Pax6 dosages and in chimeras. Immunohistochemistry and ELISA-based assays were used to determine the molecular basis of defects seen in Pax6 mutants, and wound healing assays were performed.

Results—in adults, the Pax6+/− epithelium was less densely innervated than the wild-type epithelium, and radial projection of epithelial nerves was disrupted. Neurotrophic support of the corneal epithelium appeared normal. Directed nerve projection correlated with patterns of epithelial cell migration in adult wild-types, but innervation defects observed in Pax6+/− mice were not fully corrected in wound healing or chimeric models where directed epithelial migration was restored.

Conclusions—Pax6 dosage nonautonomously controls robust directed radial projection of corneal neurons, and the guidance cues for growth cone guidance are not solely dependent on directed epithelial migration. There is little evidence that ARK represents neurotrophic keratitis.
nerve bundles enter the corneal periphery radially to form the perilimbal ring. Bundles of fibers project and extend through the corneal stroma, subdivide, penetrate the anterior limiting lamina, and innervate the corneal epithelium, forming a basal epithelial nerve plexus. The mechanisms that control corneal epithelial nerve growth are incompletely known, but wound-induced electric currents have been shown to direct nerve regeneration after epithelial wounding.

Corneal nerves maintain a healthy cornea through the release of soluble trophic factors. Substance P (SP) is of particular importance; it acts synergistically with other growth factors, including nerve growth factor (NGF), to accelerate corneal epithelial cell migration and proliferation during healing and regeneration.

Disruption of corneal innervation can cause degenerative conditions ranging from dry eye to neurotrophic keratitis. The symptoms accelerate cell loss leading to corneal epithelial thinning, increased epithelial permeability, and wound healing deficiency. Most neurotrophic keratitis results from herpetic viral infection; however, ocular surgery such as laser-assisted in situ keratomileusis (LASIK) and corneal transplants, even the use of contact lenses, can lead to damage of corneal nerves. We hypothesized that corneal epithelial degeneration in PAX6+/- humans and mice may be in part a type of neurotrophic keratitis.

**Methods**

**Mice**

Pax6+/-Sey-Neu (Pax6+/-) mice were maintained on the CBA/Ca genetic background by heterozygous mating. Pax6 overexpressing transgenics, called PAX77, were maintained on the CBA/Ca genetic background, breeding PAX77 x wild-type CBA/Ca. The H253 (XLacZ) transgenic line, which is mosaic for LacZ expression from an X-linked LacZ reporter, was mated to Pax6+/- as described previously. Experimental Pax6+/+ ↔ Pax6+/-, LacZ+ and control Pax6+/+ ↔ Pax6+/+, LacZ+ chimeras were produced as described previously. All experiments were performed in accordance with the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research.

**Chemicals and Reagents**

NGF, SP, and anti-β-tubulin III were from Sigma-Aldrich (Poole, Dorset, UK). Anti-TrkA was from AbCam (Cambridge, UK), and anti-TrkB and TrkC were from Santa Cruz Biotechnology (Santa Cruz, CA).

**Immunohistochemistry on Whole Mount Corneas**

Embryos or eyes were fixed in 4% wt/vol paraformaldehyde for 2 hours at 4°C and in methanol for 1 hour at −20°C. Dissected corneas were incubated in 1% pepsin in 10 mM HCl, 37°C, 20 minutes followed by 10-minute incubation in 0.1 M sodium borate, pH 8, at room temperature, and 3 x 20 minutes PBS. Corneas were blocked for 30 minutes (0.3% BSA, 4% normal goat serum, 0.1% Triton X-100, PBS) and incubated with anti-β-tubulin III; Tuj 1, 1:1000) overnight at 4°C. After 1 hour in PBS, secondary antibody (Alexa-Fluor 488 donkey anti-rabbit, 1:300, blocking buffer) was added for 3 hours. After a 1-hour wash in PBS, corneas were mounted.

Eyes that had been XGal-stained were treated similarly, but the secondary antibody was biotin-labeled anti-rabbit (Sigma-Aldrich) diluted 1:300 and added to corneas for 45 minutes at room temperature. Corneas were rinsed four times in TBS for 10 minutes. Avidin/biotin ABC solution (Dako, Ely, Cambridgeshire, UK) was added for 30 minutes, and staining was visualized with diaminobenzidine.
Measurement of Nerve Density and Orientation in Adult Corneas

Adult whole mount corneas were immunostained for β-tubulin III as described. For each cornea, the mean number of nerve fibers transected by 200-μm lines drawn (Photoshop; Adobe, Mountain View, CA) perpendicularly to the radius at the edge of the cornea was measured. This allowed the mean spacing of the neurites to be calculated (t). The mean spacing of fibers transected by 200-μm lines, drawn radially parallel to the neurite projection, was also measured for each cornea (r). For random orientation of nerve projections, r/t would approximate 1, whereas for radially directed nerves, r/t > 1.

Assay of SP Levels

SP levels in Pax6+/+ and Pax6+/− were determined by ELISA immunoassay (Cayman Chemical, Ann Arbor, MI).

Wound Healing

For in vivo wound healing, mice (>8 weeks old, male and female) were anesthetized under veterinary advice. Central circular (1.5-mm diameter) corneal epithelial wounds were made, and the epithelial sheet within the wound was removed with a scalpel blade. Anesthesia was immediately reversed with the use of atipamezole hydrochloride (Antisedan, 0.014 mg/10 g subcutaneously; Pfizer Animal Health, Exton, PA) to facilitate normal blinking and tear production. Twelve hours after wounding, eyes were enucleated, fixed with paraformaldehyde, stained with Hoechst to measure the size of the wound, and immunostained with anti-β-tubulin III as described.

To assay neurotrophic modulation of healing, after wounding the eyes were enucleated and placed into serum-free media with or without SP and NGF, at 37°C, as described previously. 36

Results

Corneal Innervation in Pax6+/+ and Pax6+/− Mice

The pattern of corneal neurons in the basal epithelial nerve plexus of 8- to 12-week-old Pax6+/+ and Pax6+/− littermates was compared (Fig. 1). β-Tubulin III staining in flat mount wild-type corneas showed patterns of centripetal extension of axons from the limbus toward the cornea center terminating in a whorl-like pattern (Fig. 1A), in agreement with previous observations.37 However, in the peripheral corneas of Pax6+/− mice, the nerves were disorganized. Radial orientation was reestablished centrally, but, on reaching the center of the Pax6+/− cornea, the axons become densely knotted within the basal layers of the persistent lens-corneal bridge, with no evidence of swirling (Figs. 1B-D). Centripetal projection and swirling in the PAX77 transgenic line,32 which overexpresses PAX6, was comparable to wild-type (Fig. 1E). Tissue sections confirmed that the immunostaining visualized C-fibers between the basal corneal epithelial cells (Fig. 1F). Thick nerve bundles, not necessarily radially oriented, projected through the underlying corneal stroma, penetrating the basement membrane in the peripheral cornea, from which the basal epithelial fibers directed radially (Fig. 1G). The pattern of corneal nerves in wild-type corneas was reminiscent of the patterns of centripetal epithelial cell migration observed previously (Fig. 1H).35 The swirling patterns were maintained throughout life, although in aging mice 9 to 12 months old, significant loss of nerve terminals from sectors of the peripheral cornea was seen (18 mice; Supplementary Fig. S1, http://www iovs.org/cgi/content/full/50/3/1122/ DC1). The data showed that Pax6+/− corneas do have sensory nerves but that the pattern of innervation is modulated by Pax6 dosage. It was further hypothesized that nerve direction is controlled by epithelial migration.
Immunostaining of corneas from embryonic and neonatal mice showed that the radial patterns of innervation developed between postnatal day (P) 10 and P25 (Supplementary File, http://www.iovs.org/cgi/content/full/50/3/1122/DC1).

**Fiber Density and Orientation**

Fiber density and orientation were measured in Pax6+/+, Pax6+/−, and PAX77+ corneas, as described in Methods. Briefly, the radial (r) and tangential (t) spacing of epithelial nerves within 200 μm of the edge of each cornea were measured, and an r/t ratio was calculated. For randomly oriented fibers, r/t should approximate 1. For fibers with a radially biased orientation, r/t > 1. For Pax6+/− corneas, the tangential spacing between axons was significantly greater than for wild-type, confirming that the heterozygous cornea was less densely innervated (Fig. 2A). Spacing between the nerves in PAX77+ corneas (mean, 6.88 μm) was not significantly different from that in wild-type corneas (mean, 5.67 μm; see Fig. 2A). The mean corneal diameter of PAX77+ adults measures only approximately 1.5 mm, compared with 3.5 mm in wild-type adults,36 indicating that PAX77+ adults have fewer axons entering a significantly smaller cornea.

The r/t value for Pax6+/− nerves (1.43 ± 0.087 [mean ± SEM], n = 9) was significantly less than that for Pax6+/+ or PAX77+ (3.23 ± 0.265, n = 11; 2.72 ± 0.280, n = 10, respectively; t-test, P < 0.005). Hence, radial orientation of Pax6+/− nerves was less robust, and more random, than for Pax6+/+ nerves. PAX77+ corneal nerve orientation was normal (Fig. 2B).

**Nerve Projection Disruption by a Lens-Corneal Bridge**

The lens-corneal bridge in mice is a result of the failure of the invaginating lens placode to separate from the overlying facial epithelium.38 We investigated whether the tangle of nerves at the center of the Pax6+/− cornea was a direct consequence of Pax6 mutation on projection or a secondary consequence of the presence of the lens-corneal bridge. The Pax6 null mutation was bred onto a mixed CBA/Ca × C57BL/6 background, which, though it shares all the other characteristics as the CBA/Ca Pax6+/− mouse, lacks the lens-corneal bridge.39 On this background, r/t ratios (calculated as before) were 3.70 ± 0.22 (n = 12) for wild-type corneas and 2.55 ± 0.22 (n = 8) for Pax6+/− (t-test, P = 0.002), confirming that the less radial orientation of heterozygous nerves is robust across genetic backgrounds. Lens-corneal bridges were not present, and no neural knots were observed; rather, the nerves swirled as in wild-type corneas (n = 6; Fig. 3). It was concluded that neural knotting was a secondary consequence of the lens-corneal bridge that might have disrupted nerve guidance by acting as an attractant.

**Nerve Fiber Guidance**

Swirling of nerves at the center of the wild-type cornea and disrupted radial projection in the Pax6+/− mouse recapitulated the patterns of corneal epithelial cell migration described previously in LacZ mosaics (XLacZ) and chimeras.34,35 We tested whether growth cone projection is dependent on directed epithelial cell migration and predicted that projection and swirling would correlate with patterns of epithelial X-Gal staining in XLacZ mosaic reporters. The nerves of Pax6+/+ and Pax6+/− XLacZ-mosaic littermates were immunostained against β-tubulin III (Fig. 4). In wild-type 15-week-old corneas, nerve projections coincided with radial stripes, and there was a 1:1 correlation between the direction of epithelial swirling (clockwise or counterclockwise) and that of the nerves (Fig. 4A; n = 17 [11 clockwise, 6 counterclockwise]). Pax6+/− XLacZ+/− eyes displayed disorganized clonal patterns of X-gal–stained cells, giving a patchwork appearance (Fig. 4B; n = 8). However, the nerves toward the center of the cornea projected radially and were able to cross clonal patches of epithelial cells. Our previous observations were that limbal stem cells and radial epithelial migration become active at 5 to 6 weeks, and in 3-week-old wild-
type mice radial patterns of epithelial stripes are not yet clearly defined. However, nerves were able to project radially and to cross patches of randomly orientated blue cells in \(Pax6^{+/+}\) eyes in a 3-week-old mosaic (Fig. 4C). The data suggested that corneal nerve projection does not require directed epithelial cell migration.

It remained possible that cell migration and axon guidance were responding to the same extrinsic guidance cues and that the epithelial Pax6 dosage was controlling these cues. Therefore, we produced chimeric mice to investigate whether disrupted neural patterning in the \(Pax6^{+/−}\) mouse was a nonautonomous effect of reduced Pax6 dosage in the corneal epithelium (i.e., whether the genotype of the epithelium, not of the neurons, was modulating patterns of neural projection in this system). Control \(Pax6^{+/+}\), LacZ\(↔\)Pax6\(^{+/+}\), LacZ\(^+\) and experimental \(Pax6^{+/+}\), LacZ\(↔\)Pax6\(^{+/−}\), LacZ\(^+\) chimeras were produced as described previously.34 In these chimeras, radial patterns of blue and white sectors were observed in the corneal epithelium that resulted from inward immigration of LacZ\(^+\) and LacZ\(^−\) patches of limbal-derived epithelial cells. In the control chimeras blue and white epithelial cells were Pax6\(^{+/+}\), but in experimental chimeras the white cells were Pax6\(^{+/+}\) and the blue cells were Pax6\(^{+/−}\). The lenses in our \(Pax6^{+/+}\) ↔ \(Pax6^{+/−}\) chimeras were composed entirely of wild-type cells.40 Collinson et al.34 showed that centripetal migration of \(Pax6^{+/−}\) corneal epithelial cells was restored in \(Pax6^{+/+}\) ↔ \(Pax6^{+/−}\) chimeras and that hence epithelial cell migration was, at least in part, controlled nonautonomously by cues extrinsic to the migrating cells. Nerve fibers in white, \(Pax6^{+/+}\) patches of cells were on average denser and more radially oriented that those in \(Pax6^{+/−}\) blue patches (Fig. 4D). Control r/t values for epithelial nerves projecting under LacZ\(^+\) Pax6\(^{+/+}\) and LacZ\(^−\) Pax6\(^{+/−}\) corneal epithelia were 3.76 ± 0.32 (n = 9) and 3.60 ± 0.43 (n = 9), respectively, showing that the presence or absence of LacZ does not itself affect directed migration (t-test; \(P = 0.77\)). In contrast, r/t for neurites projecting under LacZ\(^−\) Pax6\(^{+/−}\) corneal epithelial stripes was 1.88 ± 0.13 (n = 9), significantly less than LacZ\(^+\) Pax6\(^{+/+}\) controls (t-test; \(P = 0.003\)). Robust radial projection of epithelial neurons was, therefore, shown to be a nonautonomous function of the genotype (\(Pax6^{+/+}\) or \(Pax6^{+/−}\)) of the corneal epithelium, and a peripheral nerve projection defect was retained even when epithelial migration was corrected.

Epithelial cells have been shown to support corneal nerves.41-44 These data suggested that epithelial cells showed clone-specific variation in their ability to provide neurotrophic support, either in an experimental chimeric model or (in the case of aging, Supplementary Fig. S1) because of possible stochastic variation in gene expression in clones of limbal cells from which the epithelium is derived.45 The neuropeptide SP, released by sensory neurons, has been shown to upregulate the expression of NGF by skin keratinocytes, which may in turn feed back to promote nerve survival.46 Neurotrophic support of the cornea was investigated further.

### Neurotrophic Support of the Cornea

We quantified the SP levels in \(Pax6^{+/+}\) and \(Pax6^{+/−}\) corneas (Fig. 5A) but found no significant difference (219.3 ± 16.95 pg/cornea and 179.3 ± 28.86 pg/cornea, respectively; t-test, \(P > 0.05\)). Although the trend was downward in \(Pax6^{+/−}\) corneas, the diameter of the \(Pax6^{+/−}\) cornea was approximately 90% that of wild-type on the CBA/Ca background.40 On a per unit area basis, it can be calculated there is no deficiency of SP in the \(Pax6^{+/−}\) cornea: 179.3/(0.9)^2 = 221.4. By Western blot analysis, no difference was detected in levels of expression of NK1, the SP receptor. NGF receptors TrkA, TrkB, and TrkC were also assayed by Western blot, and all were expressed at normal levels in the \(Pax6^{+/−}\) cornea (Fig. 5B).

The effect of SP and NGF on wound closure in whole eye culture was investigated. \(Pax6^{+/+}\) corneas healed slightly faster in the presence of a combination of SP and NGF (controls...
Nerve Regeneration after Wounding

In vivo epithelial wounding was performed to determine whether Pax6+/− corneal nerves respond normally by projecting to the wound edge in the wake of epithelial cell migration. In Pax6+/+ and Pax6+/− wounds after 12 hours of healing, nerve sprouting was observed with regenerating nerves projecting usually toward the wound edge (Fig. 6). An estimate of whether the Pax6+/+ and Pax6+/− nerves were equally robustly directed after wounding was obtained by measuring the tortuosity of 100 to 200 μm of each visible nerve closest to the wound edges. The straight-line distance between the growth cone (the point nearest to the wound edge) and a proximal branch point (100 to 200 μm further from the wound edge) was measured. Next the actual distance traced along the track of the nerve was measured, and this value was divided by the straight line distance to give a value of tortuosity. The tortuosity value of Pax6+/− nerve sprouts (1.16 ± 0.02; n = 5 mice, 65 nerves) was slightly but significantly greater than that of wild-type littermates (1.08 ± 0.01; n = 6 mice, 66 nerves; t-test, P = 0.039). Cell migration after wounding was much greater (20–50 μm/h) than that observed in the uninjured corneal epithelium (20–30 μm/d). Our data therefore suggested that defective Pax6+/− nerve projection was not corrected by the increased speed of epithelial migration.

Discussion

Control of Nerve Projection in the Corneal Epithelium

Radial patterns of centripetal corneal nerve projection and swirling were observed previously in a transgenic reporter mouse and by neurotubulin staining in rats.37,49 In vivo confocal microscopy of human corneas has revealed similar, if less regular, projections and swirls.50,51 In the adult, there is continuous centripetal corneal epithelial cell migration, and we initially assumed that corneal nerves follow a so-called path of least resistance, swept along by the guided epithelial migration, as tentatively suggested previously.50 The 1:1 correlation between direction of epithelial and nerve swirling at the center of the cornea is consistent with that hypothesis. However, other aspects of our data appear to preclude the possibility that nerves are directed solely by epithelial migration, as follows: the evident development of radial projection of axons in young mice before migration of epithelial cells; the radial projection of Pax6+/− nerves centrally in the cornea across randomly arranged clonal patches of epithelial cells; the retention of a quantitative guidance defect in wound healing with greatly increased rates of epithelial migration; the less robust radial projection of axons through Pax6+/− sectors of chimeric Pax6+/− ↔ Pax6+/− chimeras in which patterns of normal radial epithelial cell migration have been restored. The data are more consistent with a hypothesis that epithelial cells and subepithelial nerves respond to the same external guidance cues on slightly different timescales. These cues may be electrical.

Electrical cues (ion flow) may control epithelial migration and growth cone guidance in the cornea.16,52,53 Vertebrate-stratified epithelia, including the corneal epithelium, exhibit a net influx of sodium and potassium ions through cellular transport, combined with a net efflux of chloride ions such that basal epithelial cells are at a positive potential difference (approximately 25 mV) to the environment. Wounding and abrasion short-circuits this potential difference, leading to a standing voltage gradient from the intact epithelium into the wound site (the cathode) that directs epithelial cells and corneal nerves to migrate to fill the wound.16,52 There is usually no overt wound during normal epithelial homeostasis, but
persistent minor abrasion and cell loss could create a gradient of cumulative subacute wound load from the limbus to the center of the cornea, maintaining the electrical signal for cells and nerves to move centrally. We have recently shown that the barrier function of the Pax6+/− corneal epithelium is disrupted 54; therefore, it seems unlikely that the mutants can maintain the epithelial potential difference. This may underlie the disrupted long distance cell migration and axon guidance in Pax6+/− corneal epithelia.

Why the Swirl?

Vortex formation has been observed and modeled mathematically in a cellular system (the chick epiblast), and though the formation of vortices may be initiated randomly, once formed they are mathematically impossible to eliminate.55 The variation in orientation (clockwise or counterclockwise) in the vortex patterns observed in our system suggest that they are a stochastic consequence of minor targeting errors in the epithelial or growth cone centripetal movement. However, it is also possible that physical or other forces influence the formation of vortices, such as shearing forces exerted during blinking.50 The vortex patterns we observed in wild-type corneal epithelia were consistent with a model whereby two discrete force vectors act on nerves and epithelial cells, a radial force directed centripetally and a tangential force directed perpendicularly to the direction of migration. Exposure to magnetic fields modulates directed cell growth in corneal epithelial cells, and a standing electric current that pushes epithelial cells toward the center of the cornea could elicit a concomitant magnetic field that may direct them sideways, explaining vortical patterns of cell migration.56,57 It is possible that similar forces act on wild-type corneal epithelial nerves.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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References


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Centripetal projection of β-tubulin–expressing nerve fibers in Pax6<sup>+/+</sup>, Pax6<sup>+/−</sup>, and Pax7<sup>+/−</sup> mice. Anti–β-tubulin III immunofluorescence on whole mount corneas. (A) Centripetal projection of nerves from the limbal region to the central cornea ending in a whorl-like pattern in Pax6<sup>+/+</sup> corneas. (B-D) Pax6<sup>+/−</sup> corneas fibers were disorganized with radial projections disrupted in the peripheral area. Although radial projection was reestablished centrally (B), spiraling was absent and nerves became densely knotted on entering the lens-corneal bridge (C, D). Images of same field in superficial epithelial (C) and deeper stromal (D) planes of focus. (E) Normal nerve projection and central swirling in the corneal epithelium of the Pax6-overexpressing mouse of the PAX7<sup>+/−</sup> line. (F) Immunostaining of fibers was observed in basal corneal epithelium layers with projections upward to superficial cells. (G) Image of peripheral wild-type corneal epithelium showing radial projection (toward top left) of epithelial nerve fibers from underlying thick stromal bundles that are not themselves radially oriented. (H) Central swirl of XGal staining in corneal epithelium of LacZ<sup>+</sup> ↔ LacZ<sup>−</sup> chimera, produced by centripetal migration epithelial cells. Part H reprinted with permission from Collinson JM, Morris L, Reid AI, et al. Clonal analysis of patterns of growth, stem cell activity, and cell movement during the development and maintenance of the murine corneal epithelium. Dev Dyn. 2002;224:432–440. © 2002 Wiley-Liss, Inc. Scale bars (A, B, G) 100 μm; (F) 20 μm; (H) 50 μm.
Figure 2. Nerve density and orientation. Nerve fiber density and orientation were measured in Pax6\(^{+/+}\) (darkest shading), Pax6\(^{+/-}\), and PAX7\(^{+}\) (lightest shading) corneas. For each cornea the mean number of epithelial fibers was transected by 200-\(\mu\)m lines, drawn perpendicularly (tangential) to the radius at the edge of the cornea measured (t) giving the mean spacing of the neurites (t; A). Mean spacing of neurites transected by 200-\(\mu\)m lines, drawn radially parallel to nerve projection, was also measured for each cornea (r). For randomly oriented nerves, r/t should approximate 1. For nerves with a radially biased orientation, r/t > 1 (B).
Figure 3.
Corneal nerve projection is disrupted by the lens corneal bridge. Although radial projection is not as robust as wild-type in Pax6+/− cells on the mixed genetic background, normal central swirling is restored in the absence of a lens-corneal bridge.
Figure 4.
Comparison of nerve projection and epithelial cell migration. Whole mount β-tubulin immunostaining of X-Gal–stained mosaic and chimeric corneas. (A) Central cornea of 15-week-old Pax6<sup>+/+</sup> X<sup>LacZ</sup> mosaic. LacZ<sup>+</sup> epithelial cells are blue, epithelial nerve fibers are brown (β-tubulin immunostaining). Projections of the nerves lie parallel to the tracks of epithelial cell migration revealed by X-Gal staining, and there is anticlockwise swirling of both in parallel (top left). (B) Radial projection (to the left) of corneal epithelial nerve fibers across patch boundaries in a 15-week-old Pax6<sup>−/−</sup> X<sup>LacZ</sup> mosaic. (C) Epithelial nerves in a 3-week-old Pax6<sup>+/+</sup> X<sup>LacZ</sup> mosaic. Radial orientation of neurite projection is emerging.
(though not yet as regular as adults), crossing randomly oriented LacZ+ epithelial patches. 

(D) X-Gal–stained corneal epithelium of a 15-week-old Pax6+/+, LacZ− ↔ Pax6+/−, LacZ+ chimera showing centripetal migration of axons. The blue stripe of epithelium is Pax6+/−, the white is Pax6+/+. Neurites were less dense and more randomly orientated in Pax6+/− blue stripes than in Pax6+/+ white stripes. (E) Histogram showing r/t values for control Pax6+/+ LacZ− nerves (white bar) compared with Pax6+/+ LacZ+ and Pax6+/− LacZ+ (blue bars). Scale bars, 50 μm.
Figure 5. Neurotrophic support of the $\text{Pax}_6^{+/−}$ cornea. (A) Substance P concentrations from corneal extracts derived from $\text{Pax}_6^{+/+}$ and $\text{Pax}_6^{+/−}$ mice, expressed as picogram per cornea. (B) Western blot analysis of NK1, TrkA, TrkB, and TrkC levels in $\text{Pax}_6^{+/+}$ and $\text{Pax}_6^{+/−}$ corneas. $\beta$-Actin was assayed as a loading control.
Figure 6.
Nerve regeneration after wounding in vivo. (A) Representative images of a Pax6+/+ eye stained with β-tubulin III, 12 hours after wounding, showing radial projection of regenerating nerves into the wound site (right). (B) Representative image of a β-tubulin III–stained Pax6+/− eye showing less robust radial nerve fiber projection toward wound (right). Scale bar, 50 μm.