Corneal epithelial regeneration initiates a temporary subbasal nerve plexus composed of fine nerve fibers after photorefractive keratectomy (PRK). First, branches of the subepithelial nerve plexus at the margin of the injury site regenerate and infiltrate the tear film. These nerves develop a biphasic pattern: during the first 24 hours, a transient subbasal plexus forms, but by 3 months postoperatively, nerve density is almost restored. Electrical nerve impulse activity of single sensory terminals is recorded. At 3 months, nerve density is normal, but cold thermoreceptors are still scarce. The primate model offers an opportunity to study the regeneration of corneal sensory nerves postoperatively. The functional recovery is faster in corneas than in rodents. The postoperative level of cold thermoreceptors is significantly lower compared to that of normal control eyes. The corneal regeneration pattern of each fiber type may be associated with the variable regeneration pattern of each fiber type.
important role in the modulation of basal tearing and blinking rates associated with the homeostasis of eye surface wetness.24,25 Surgical injury acutely affects the different classes of sensory axons innervating the ocular surface.26 However, the degree of residual functional impairment in mecano-nociceptor, polymodal nociceptor, and cold thermoreceptor fibers following PRK, leading eventually to dysesthesias and reflex autonomic disturbances in operated eyes, is still poorly known.

In the present work, we analyzed the changes in spontaneous and stimulus-evoked nerve impulse activity experienced by corneal sensory terminals, immediately and at different times up to 30 days after PRK, while observing the parallel evolution of corneal nerve architecture within and around the wounded area. We discovered that cold-sensitive endings regenerate early albeit a fraction of them, located both in the injured and in the intact peripheral cornea, exhibit an abnormal impulse activity attributable to the incomplete branching of regenerating cold thermoreceptor nerve axons. Polymodal nociceptors and mecano-nociceptors required longer times to recover the normal responsiveness to their natural stimuli.

METHODS

Animals

Male, 3- to 6-month-old C57Bl/6 mice were purchased from Charles River Laboratory (L’Arbresle Cedex, France). Animals were handled and housed according to the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research and the applicable guidelines of the European Union (2010/63/EU) and the Spanish Government (RD 53/2013). The ethics committee of the University of Oviedo approved all procedures.

Corneal Nerve Injury Through Photorefractive Keratectomy (PRK)

Surgical corneal injury was performed by using a VISX Star S2 excimer laser (VISX, Inc., Santa Clara, CA, USA) as previously described.29–32 Right eyes were subjected to PRK surgery with a 1.5-mm 2 circular area, 45 μm of the anterior stroma (representing approximately 20% of the total stromal thickness), removing all subbasal nerve fibers and intraepithelial terminals of the treated area.

Before surgery, mice were deeply anesthetized by intraperitoneal injection of a mixture of ketamine hydrochloride (80 mg/kg) and xylazine hydrochloride (5 mg/kg), and 0.13 mm 2. This circular area was divided in three concentric circles (Supplementary Fig. S1), with the first and biggest circle corresponding to the border of the injury in PRK-ablated corneas. For comparison, in uninjured corneas the central corneal area was also divided into three similar concentric circles of 25 mm 2 (parallel to the limbus; see Supplementary Fig. S1). Whole-mounted corneas were incubated in blocking serum solution for 24 hours at RT against rabbit anti–neuronal class III β-Tubulin (1:250; Cell Signaling Technology, Boston, MA, USA) in blocking serum solution. After three rinses with 0.2% BSA (washing solution), 0.2% goat serum, 0.2% sodium azide, and PBS-Triton, corneas were incubated in blocking serum solution for 24 hours at RT with secondary antibody against anti-rabbit IgG Alexa Fluor 594 (1:500; Molecular Probes, Eugene, OR, USA). Afterwards, corneas were rinsed three times with washing solution, followed by incubation for 10 minutes at RT with 4,6-diamidino-2-phenylindole (DAPI, 2 μg/mL; Molecular Probes). Finally, corneas were mounted in slides with fluorescent mounting medium (DAKO, Glostrup, Denmark).

Image Acquisition

Fluorescence images were obtained with a Leica TCS SP8 confocal microscope (Leica Microsystems). For whole-mounted corneas, a mosaic of images was obtained at magnification of ×200 by using tile scan utility of LASX and confocal z stacks, spaced 3.5 μm in the Z-axis.

Morphologic Data Analysis

Images were analyzed by using also the image analysis software Fiji. Whole-mounted corneas were divided in five areas of study (Supplementary Fig. S1): four peripheral zones, which were defined each as two 0.25 mm 2 square regions (a 500 × 500-μm box) whose peripheral borders were approximately 200 μm away from the limbus border and were separated approximately 500 μm from each other; and a fifth area, defined as the injured area in the center of the cornea, which was determined by the absence of DAPI staining in injured corneas (mean injured area, 1.46 ± 0.13 mm 2). This circular area was divided in three concentric circles (Supplementary Fig. S1), with the first and biggest circle corresponding to the border of the injury in PRK-ablated corneas. For comparison, in uninjured corneas the central corneal area was also divided into three similar concentric circular areas of 1.5, 1.0, and 0.5 mm 2.

Subbasal nerves were quantified in the periphery by counting the number of nerves intersecting a line drawn inside the square area (parallel to the limbus; see Supplementary Fig. S1). In the central cornea of both injured and control eyes, subbasal nerves were quantified as the number of nerves intersecting the circumferences defining each of the concentri...
Morphofunctional Changes in Corneal Nerves Following PRK

The eye was then placed in a recording chamber and was superfused with the same physiological saline solution. Temperature during the experiment was controlled continuously and was continuously superfused with the same physiological saline solution. Temperature during the experiment was controlled with a homemade Peltier device.

Extracellular electrical activity of single sensory nerve endings on the corneal surface was recorded with a borosilicate glass microelectrode with tip diameter of approximately 50 μm that was filled with saline solution. An Ag/AgCl wire located in the bath serves as indifferent electrode. With the aid of a micromanipulator, the recording pipette was placed on the corneal surface with slight suction at different points both in the injured area and in the periphery (mean of 15 seals per cornea, 10 in injured area and 5 in periphery), searching for nerve terminal impulse (NTI) activity. NTIs were amplified with an AC amplifier (Neurolog NL104; Digitimer, Welwyn, UK) and stored at 10 kHz into a computer, using a CED micro 1401 interface and Spike 2 software (both from Cambridge Electronic Design, Cambridge, UK).

Only recordings containing NTIs originating from a well-defined single nerve terminal were analyzed. In these recordings NTIs were clearly distinguished from noise (~10 μV peak-to-peak) and had similar amplitudes and waveforms, indicating that they originated from the same sensory nerve ending. To minimize deterioration of the preparation with time, the total duration of the experiment was limited to a maximum of 120 minutes per eye (30 minutes for preparation; 90 minutes for recording). Percentage of successful attempts refers to the number of seals made on the cornea in which nerve terminal activity was recorded.

### Experimental Protocol

The recording pipette was placed at regularly aligned points on the corneal surface, separated by an approximate distance of 0.2 mm. First, one half of the cornea was explored starting in the center and descending to the periphery; then, the eye was rotated and the opposite side of the cornea was explored.33 After 90 minutes of recording attempts, the eye was fixed and prepared for morphologic studies.

Cold stimulation was performed by decreasing the background temperature of the perfusing solution (from ~34°C to ~14°C or ~5°C, in cold thermoreceptors or polymodal nociceptors and mecano-nociceptors, respectively), generating a cooling ramp at mean cooling rate of 0.6°C·s⁻¹, when the peak temperature of ~14°C or ~5°C was attained, warming was applied at a similar speed to return to the basal temperature. Following a resting period of 120 seconds, mechanical stimulation was performed by moving forward the pipette with a controlled displacement of the micromanipulator. Pressure with the pipette was maintained for 2 seconds. Two minutes later, a heating ramp (from 34°C to ~5°C at 0.5°C·s⁻¹, ~30 seconds’ duration) was applied, returning to the initial control value of 34°C when the peak temperature was reached.

### Analysis of NTI Activity

Success percentage indicates the percentage of successful attempts at recording a nerve terminal activity in relation to the total number of attempts. Background activity is defined as the mean basal ongoing frequency in impulses per second (imp·s⁻¹) at the basal temperature (33.9°C ± 0.07°C) measured during the 60-second period that preceded the onset of a stimulus.

The following parameters of the NTI activity were analyzed in the different terminal types.

#### Cold Thermoreceptor Terminals

**Cooling Threshold.** Temperature (°C) during a cooling ramp at which NTI frequency increased to a value that was the mean NTI frequency measured during the 10-second period preceding the onset of a cooling ramp plus three times its standard deviation.

**Cooling Response.** Expressed as the mean in NTI frequency between the cooling threshold and the peak response to the cooling ramp.

#### Polymodal and Mechano-nociceptor Terminals

**Response to Cooling.** The total number of spikes during the 45 seconds following the onset of the cooling ramp was compared with the total number of spikes during 45 seconds immediately before the cooling ramp.

**Response to Heating.** The total number of spikes during the 30 seconds following the onset of the heating ramp was compared with the total number of spikes during 30 seconds immediately before the heating ramp.

**Response to Mechanical Force.** The total number of NTIs during the 10 seconds following the onset of mechanical stimulation was compared with the total number of NTIs during the 10-second period immediately before applying the mechanical stimulus.

### Statistical Analysis

Statistical comparisons were performed using Microsoft Excel 2010 (Microsoft Corporation, Redmond, CA, USA), Origin 8 (OriginLab Corporation, Northampton, MA, USA), and InStat 3 (GraphPad Software, Inc., La Jolla, CA, USA). Data are expressed as mean ± SD. For electrophysiology data unpaired Student’s t-test was used. The morphologic measurements were compared by using 1-way ANOVA with Tukey-Kramer multiple comparisons post hoc test. Significance threshold was set at P < 0.05.

### Results

Sixty-five mice were used in this study. PRK was performed in a total of 54 mice. Immediately after laser ablation, fluorescein staining in seven mice delimited in the cornea a circular area of epithelial damage of 2.2 ± 0.1 mm². Twenty-four hours later, the size of the injured area in the same animals had decreased to 0.46 ± 0.22 mm². The injury was not visible after 2 days in four of the seven mice and was very small in the remaining three (0.05 ± 0.08 mm²). On the third day, no fluorescein staining was apparent in any of the eyes. PRK caused initially a
marked opacity that reversed progressively with time. Still, 28% of the corneas continued to show light opacity spots 30 days after surgery.

From the 54 operated animals, 26 corneas (at T0, n = 5; T3, n = 4; T7, n = 6; T15, n = 5; T30, n = 6) and six corneas of intact mice were studied morphologically (Fig. 1A). Electrophysiological recordings were performed in 43 eyes at various times after PRK (T0, n = 5; T3, n = 5; T7, n = 15; T15, n = 10; T30, n = 8) and in five control eyes where the cornea was intact. The corneas of 15 eyes previously used for electrophysiological recordings were also studied morphologically. In total, electrical activity was recorded from 117 corneal nerve terminals: 66 with receptive fields located within the central area, in the region where the wound was performed, and 51 terminals located in the surrounding peripheral area.

In all recording experiments, the search for nerve terminal impulse activity started by placing the recording electrode on the corneal surface at different pre-established points defined both in the central area where the injury was made and in the intact periphery. From previous studies in intact mice corneas,35 we defined the fraction of the nerve terminals of the intact cornea responding only to mechanical force as mechano-nociceptors and those activated by heat and mechanical force as polymodal nociceptors. Finally, cold thermoreceptors, initially defined by their responsiveness to cooling, were subclassified by the value of their spontaneous firing at 34°C and the temperature threshold required to increase their firing rate with a cooling ramp35 as high-background (≥1.5 NTI s⁻¹), low-threshold (≥30.5°C) cold thermoreceptors (HB-LT), or low-background (<1.5 NTI s⁻¹) high-threshold (<30.5°C) cold thermoreceptors (LB-HT). Altogether, in the central cornea 54.1% of the terminals were identified as cold thermoreceptors, 26.7% as polymodal nociceptors, and 10% as mechano-nociceptors. In the peripheral area, 46.7% were cold thermoreceptors; 26.7%, polymodal nociceptors; and 10%, mechano-nociceptors. The remaining units recorded at the central and peripheral cornea could not be accurately classified.

**Nerve Terminal Impulse Activity Was Absent From the PRK-Injured Area at Early Stages of Corneal Nerve Regeneration**

At day 0, immediately after laser ablation, the corneal epithelium had disappeared and the normal innervation of the central cornea was virtually absent. Only midstromal nerve trunks, often terminating abruptly, could be identified in the stroma, while subepithelial and subbasal plexuses were lost (Figs. 1B, 2B, 3A), and no nerve penetration points were identified in the wounded area (Table). In accordance with this morphologic observation, electrical activity was undetectable within the lesioned area in the animals studied immediately after PRK (Fig. 3A).

In the peripheral cornea surrounding the wounded area, the density of nerve fibers measured after surgery and the number of nerve penetration points in the basol lamina per mm² was slightly lower in day 0 corneas than in the same area of the corneas of control mice (Fig. 3B; Table). At this time, the incidence of NTI activity in the peripheral area surrounding the wound was similar in operated and intact corneas (32.4% ± 9.3%, n = 5 at T0 versus 39.1% ± 7.7%, n = 5 in intact corneas; Fig. 3B). At day 3, regenerating individual rectilinear axons, sprouting from some of the cut subbasal nerves, started to enter the injured region (Figs. 1C, 2C), although most appeared abruptly interrupted at the wound border. The overall density of these regenerating fibers within the damaged area was only 14.3% ± 3.6% of the density values at the central cornea of intact mice (P < 0.001; Fig. 3A, Table). At this time, NTI activity was still undetectable in the wounded area (Fig. 3A).

In the peripheral cornea surrounding the wound, the density of nerve fibers at T3 was significantly lower than in control corneas (Fig. 3B; Table). Likewise, the probability of finding an active nerve terminal at day 3 in the noninjured peripheral cornea was also lower (13.9% ± 7.9%, n = 5; Fig. 3B). Moreover, cold thermosensory terminals exhibiting abnormal low background activity combined with a low cooling threshold, never observed in uninjured corneas, were recorded at the periphery for the first time (Fig. 4).

At day 7 an incipient process of remodeling of the stromal nerves below the wounded area was evident, leading to new basal lamina nerve penetration points that reached near-normal numbers (Table). Entering subepithelial axons were short, tortuous, and ramified (Figs. 1D, 2D). However, at T7 most of the axons present in the ablated area originated from peripheral stumps of subbasal leashes that were growing centrally, but the nerve density remained significantly lower (P < 0.001) than in control mice, with nerve density values 51.7% ± 4.9% of control values (Table). The first nerve terminal electrical activity within the injured corneal area was recorded at this time point and only in 7 of the 13 explored corneas, with the probability of finding an active unit being significantly lower than for the control value (Fig. 3A). Ten active terminals were clearly identified as cold-sensitive. One active terminal with low background activity (0.03 imp s⁻¹) responded only to mechanical stimulus (from 0 imp/10 s to 71 imp/10 s). In addition, there were three sites with low-amplitude NTIs that could not be identified functionally.

In the peripheral cornea, the mean nerve density increased slightly at T7. In addition, the percentage of points where NTI activity was recorded over the total number of attempts increased to 23.1% ± 6.5% (Fig. 3B). Six of ten terminals were classified as cold thermoreceptors, while the rest were low-amplitude, functionally unidentifiable terminals.

**One Month After PRK, a Subset of the Corneal Endings Exhibited Abnormal Activity Despite the Apparent Morphologic Recovery of the Innervation**

At day 15 after injury, the overall density of nerve fibers in the injured area became greater than in the periphery (Table). This was due to a higher number of stromal nerve penetrations (Table) and the incorporation of nerve fibers spreading from the periphery of the wound, forming tortuous and ramified arborizations (Figs. 1E, 2E), although nerve density was still lower (58.9% ± 5.2%, P < 0.05) than in control eyes. At this time, active terminals were found in 6 of 10 corneas, with a higher probability of finding active recording points than at T7 (Fig. 3A). Six of the identified terminals were characterized as cold-sensitive and one as a mechano-nociceptor. In six instances, units with spontaneous activity were observed that had a low amplitude, which decreased further with cooling, preventing reliable identification of their function.

In the peripheral cornea, nerve density continued to increase at T15 in comparison with earlier stages of the study, and the probability of finding NTIs also increased (Fig. 3B; Table), reaching values near those in the periphery of intact corneas. Four active terminals responded clearly to cold stimuli, while four others presented a low amplitude that prevented a reliable functional identification. Another terminal showed the characteristics of a mechano receptor (background activity of 0.05 imp s⁻¹, and 0 imp/10 s before and 14 imp/10 s after mechanical stimulation). One polymodal nociceptor
FIGURE 1. Subbasal and stromal corneal nerves at different times after PRK. Whole-mount corneas showing at low magnification (×200) the entire corneal innervation by stromal and subbasal corneal nerves, stained against β-Tubulin III (red). (A) An intact cornea with the area in which the PRK wound is performed has been marked with a circle. (B–F) Corneas excised and stained at different times after PRK (T0, T3, T7, T15, and T30, respectively). The circle indicates in each case the extension of the initial PRK-injured area. Scale bars: 500 μm.
FIGURE 2. Subbasal and stromal corneal nerves at different times after PRK. z-stack images of whole-mount corneas stained against β-Tubulin III (red). Images from an intact cornea (A) and PRK-operated corneas at different times after PRK ((B–F): T0, T3, T7, T15, and T30, respectively) are shown. Asterisks mark the penetration points. No subbasal nerves were observed immediately after PRK (B). Three days after ablation, some sprouting subbasal nerves started entering the injured area (arrowheads, C). Scale bars: 100 μm.
FIGURE 3. Density of subbasal nerve fibers (left axis, black bars) and percentage of successful attempts at recording a terminal (right axis, gray bars) in control eyes and at the different time points after PRK, in the injured area (A) and the peripheral, uninjured area (B). Nerve density is expressed as percentage of the mean number of axons/mm² measured in control corneas in the central (A) or the peripheral (B) cornea. Successful recording is expressed as percentage of electrophysiological recording attempts where an active corneal unit is found. Statistical analysis compared differences with control group values: *P value < 0.05, **P value < 0.01, ***P value < 0.001.

TABLE. Subbasal Nerve and Epithelial Penetration Point Density Values in the Injured and Peripheral Areas at Different Time Points Post PRK and in Intact Corneas (Control)

<table>
<thead>
<tr>
<th>Experimental Groups</th>
<th>Nerve Density, Axons/mm²</th>
<th>Penetration Point Density, Points/mm²</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Injured Area</td>
<td>Periphery Area</td>
</tr>
<tr>
<td>Control, n = 6</td>
<td>574.5 ± 66.5</td>
<td>233.1 ± 6.3</td>
</tr>
<tr>
<td>T0, n = 5</td>
<td>0.0 ± 0.0*</td>
<td>200.0 ± 7.2†</td>
</tr>
<tr>
<td>T3, n = 4</td>
<td>42.9 ± 11.0*</td>
<td>166.1 ± 8.2*</td>
</tr>
<tr>
<td>T7, n = 6</td>
<td>143.0 ± 16.3*</td>
<td>185.4 ± 5.9*</td>
</tr>
<tr>
<td>T15, n = 5</td>
<td>266.5 ± 12.5‡</td>
<td>190.0 ± 6.2*</td>
</tr>
<tr>
<td>T30, n = 6</td>
<td>351.6 ± 36.1</td>
<td>208.5 ± 5.4</td>
</tr>
</tbody>
</table>

Corneas were processed at 0, 3, 7, 15, and 30 days after surgery (T0, T3, T7, T15, and T30, respectively).

* P value < 0.001.
† P value < 0.01.
‡ P value < 0.05.
terminal that responded to mechanical stimulation and heat was also found.

At day 30, the overall density of subbasal nerve fibers in the injured area remained significantly lower (75.7% ± 4.1% of control corneas, \( P < 0.05 \)) than in the central region of intact corneas (Figs. 1E 2F 3A; Table). Most of the nerve fibers started to organize, forming a vortex in the apex of the cornea. Nonetheless, a low number of axons still showed an erratic distribution. For the first time in the study, NTI activity was present in the injured area of all corneas tested (\( n = 8 \)). The probability that the recording at defined points contained an active terminal was 47.5% ± 7% (\( n = 8 \)), still below the values found in the central cornea of control eyes (65% ± 10.5%, \( n = 5 \)) (Fig. 3A). Of the 27 active terminals found in operated corneas at T30, 5 were defined as polymodal nociceptors and 7 as cold thermoreceptors. At the remaining 15 recording sites the NTIs had a low amplitude and could not be accurately categorized.

In the corneal periphery surrounding the wounded area, the mean nerve density at T30 was similar to the same area in the intact corneas. At this time the probability of finding sites with nerve activity was 32.1% ± 10.1% (Fig. 3B; Table). Of the eight units recorded, two were characterized as cold thermoreceptors, one as polymodal, and one as a mechano-nociceptor. The remaining four units had low-amplitude NTIs, preventing reliable functional identification.

**A Fraction of Cold Thermoreceptor Endings Remain Functionally Altered 30 Days After PRK**

Collectively, at all times after injury, the firing of high-background cold thermoreceptor terminals found in the injured area was abnormal. HB-LT cold thermoreceptor terminals found in the injured area had on the average lower background activity and cooling response values than those of the same area seen in the healthy cornea (background activity: 3.4 ± 0.5 versus 6.1 ± 0.7 imp s\(^{-1}\), \( P < 0.05 \); cooling response: 21.5 ± 4.2 versus 34.2 ± 2.3 imp s\(^{-1}\), \( P < 0.01 \); injured, \( n = 20 \) versus healthy, \( n = 10 \)). LB-HT terminals also responded less to cooling (3.3 ± 0.9 versus 6.1 ± 0.7 imp s\(^{-1}\), \( P < 0.05 \); injured, \( n = 6 \) versus healthy, \( n = 6 \)). Moreover, in the injured and also in the surrounding undamaged area, we frequently observed terminals exhibiting low background activity (<1.5 imp s\(^{-1}\) at 34°C), but combined with an abnormally low cold threshold (LB-LT). These units were observed first in the uninjured area at T3 and in the injured and uninjured area since T7. Conversely, other terminals had a high background frequency at 34°C (>1.5 imp s\(^{-1}\)) but a cooling threshold below 30.5°C (HB-HT). Figure 4 shows an example of the firing pattern of these abnormal cold thermoreceptor terminal types. The overall incidence of abnormal units decreased during the healing period, from 37.5% 1 week after injury to approximately 22.2% 1 month later.
Conversely, polymodal terminals responded clearly to heating ramp, the 45-second cooling ramp to 16.4 °C before cooling (mean, 25.6 ± 0.4 NTIs; n = 5). As occurs with polymodal nociceptors of the intact cornea, this background activity decreased during a cooling ramp so that the total number of impulses fired during the 45-second period before cooling (mean, 25 ± 11.8 NTIs; n = 5) decreased during the 45-second cooling ramp to 16.4 ± 5.4 NTIs (n = 5). Conversely, polymodal terminals responded clearly to heating (7.3 ± 1.9 imp/30 s before and 37.8 ± 10.7 imp/30 s during a heating ramp, n = 5) and clearly increased their firing rate in response to mechanical stimulation (2.8 ± 0.4 NTIs/10 s before and 22 ± 13.8 imp/10 s during mechanical stimulation).

In the periphery, polymodal nociceptor activity was also practically absent until T7, with only one terminal responding to heat and to mechanical stimulation at T3. Terminals presenting the characteristics of polymodal nociceptors were identified at T15 and T30; altogether, their incidence was lower, around 10% in lesioned versus 27% in intact corneas.

**Discussion**

The present work confirms the occurrence of extensive morphologic and functional damage to corneal nerves after application of the photoablation procedure used in refractive surgery, and uncovers the marked differences in time course required by corneal sensory nerve fibers of different modality (polymodal nociceptors, mechano-nociceptors, and cold thermoreceptors) to regenerate, and to recover, albeit partially, their normal responsiveness to natural stimuli. Our study additionally revealed that the functional disturbances caused by injury affect also nerve terminals outside the directly wounded area, especially in cold thermoreceptors, possibly reflecting damage by PRK to some terminal branches of parent axons that entered the wound area but that predominantly had their sensory nerve terminals in the noninjured parts of the cornea.

PRK is a procedure that completely removes the epithelium and approximately 20 μm of the anterior stroma of treated corneas, thus affecting the small and medium-sized nerve bundles running in the anterior stromal plexus, particularly its more anterior, dense part, the subepithelial plexus. The laser beam directly destroys the more superficial nerve branches in the treated area, while the distal segments of axons running in a large extent outside the lesion, but affected by the laser beam, degenerate rapidly. Our work confirms previous studies reporting that within hours after PRK, nerves are totally absent from the injured area and those in the surrounding periphery are reduced in number.
In human patients, nerve regeneration is slow. Subbasal nerve fiber bundles visualized with tandem scanning confocal microscopy are present only in 17% of the corneas 1 month after PRK, and the density of these nerve fibers is 98% less than preoperatively. After approximately 3 months, no branched nerve fibers can be seen in the center of the ablation zone, mean subbasal nerve density remains reduced by 59% at 1 year, when compared with preoperative values. By 2 years, subbasal nerve density after PRK is similar but morphologic alterations are still present even 5 years after PRK. Despite the timescale differences in regeneration speed between mice and humans, our study showed that recovery of nerve fiber density in the cornea of mice following PRK is also slow, and 1 month later has not fully reestablished the original architecture of corneal innervation.

Maintenance and remodeling of mature axons depend under normal circumstances on extrinsic signals from the environment, which activate their translational machinery, inducing the synthesis of proteins involved in the dynamic regulation of the cytoskeleton. Nerve injury locally generates a cascade of retrograde signaling events that ultimately activate the transcription of genes required for axon survival and a cascade of retrograde signaling events that ultimately activate the transcription of genes required for axon survival and abnormal firing observed in cold thermoreceptor terminals. Extrinsic signals acting on growth cones of the damaged axons shape the outgrowth pattern regulating the local synthesis of positive and negative regulators of cytoskeletal dynamics, thereby defining the trajectory and length of the growing fibers. In addition to chemotropic cues, mechanical forces exerted by the extruding basal cells during epithelial cell proliferation accompanying corneal wound healing may also influence axonal growth. These forces are transmitted to the subbasal axons and to the reassembling epithelial basement membrane, likely affecting adhesion between epithelial cells, basement membrane, and subbasal axons and thereby, the navigation of regenerating axons. Considering the numerous positive and negative regulators influencing the final trajectory and elongation of axons after injury, the slow and incomplete recovery of the prelesion innervation architecture after PRK is not surprising.

Our study revealed that the appearance of sprouting axons within the injured area does not imply an immediate and parallel recovery of their function. In fact, we observed that corneal sensory terminals of different modality required variable times to recuperate responsiveness to their natural stimuli. Cold thermoreceptor terminals began to respond 1 month after surgery as occurs with mechano-nociceptors, while stimuli. Cold thermoreceptor terminals began to respond 1 month after surgery in 5% to 25% of patients, while nearly 50% report feelings of “dryness.” Corneal cold thermoreceptor fibers are implicated in the regulation of basal tearing rate and have been proposed also as the source of the unpleasant dryness sensation associated with dry eye disease. The differences in time course and characteristics of disturbances in corneal nerve activity observed in mice after PRK confirm the important role played by aberrant impulse activity in axotomized nerves for the appearance of dysesthesias following PRK and lend support to the proposal that, among the different functional subclasses of corneal nerves, cold thermoreceptors are major players in the development of phantom unpleasant dryness sensation after photorefractive surgery procedures.

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