

External light activates hair follicle stem cells through eyes via an ipRGC–SCN–sympathetic neural pathway

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Changes in external light patterns can alter cell activities in peripheral tissues through slow entrainment of the central clock in suprachiasmatic nucleus (SCN). It remains unclear whether cells in otherwise photo-insensitive tissues can achieve rapid responses to changes in external light. Here we show that light stimulation of animals' eyes results in rapid activation of hair follicle stem cells with prominent hair regeneration. Mechanistically, light signals are interpreted by M1-type intrinsically photosensitive retinal ganglion cells (ipRGCs), which signal to the SCN via melanopsin. Subsequently, efferent sympathetic nerves are immediately activated. Increased norepinephrine release in skin promotes hedgehog signaling to activate hair follicle stem cells. Thus, external light can directly regulate tissue stem cells via an ipRGC–SCN autonomic nervous system circuit. Since activation of sympathetic nerves is not limited to skin, this circuit can also facilitate rapid adaptive responses to external light in other homeostatic tissues.

melanopsin | circadian rhythm | sympathetic nerve | hair follicle | stem cell

Adult stem cells (SCs) reside in designated niches that tightly regulate them via short-range signals (1). In complex multicellular animals such as mammals, SCs often respond to external cues while insulated from the harsh outer environment (1, 2). For such responses, pathways for detecting and precisely relaying external signals to internal SC niches must exist. Light is an important environmental signal that can modulate cell and tissue activities (1, 2). External light primarily regulates daily tissue rhythmic activities by entraining the central circadian clock located in the suprachiasmatic nucleus (SCN) through the eyes (2–4). The central circadian clock synchronizes the oscillatory activities in peripheral tissues to regulate different physiological functions and help the organism adapt to daily regular environmental changes (2).

Photoentrainment of the central circadian clock is mediated by a group of atypical photoreceptor cells named “intrinsically photosensitive retinal ganglion cells” (ipRGCs) located in the inner layer of the retina which express the photopigment melanopsin (3–7). By integrating the intrinsic melanopsin photodetection pathway and extrinsic input from rod and cone signaling, ipRGCs provide environmental luminance signals for several important non-image-forming functions, such as circadian photoentrainment, pupillary light reflex, alertness, and phototaxis (5, 6, 8–11). Compared with cone opsins and rhodopsin of rods, melanopsin requires higher light intensity for activation due to its lower density on the ipRGC cell surface and has maximum sensitivity in the short-wavelength blue light range (peak absorption ~480 nm) (5, 12, 13). Furthermore, unlike the quick on/off signaling from cone opsins and rhodopsin in response to light, melanopsin signaling in ipRGCs can produce sustained light signals for a long period of

time and has prolonged activation after the light stimulation is turned off (3, 13). In mouse retina, there are at least five subtypes of ipRGCs (M1–M5) with distinct morphological and electrophysiological properties and preferential projections to different brain regions (14–20). Specifically, M1 ipRGCs, which do not express the transcriptional factor Brn3b, innervate the SCN for circadian photoentrainment (9). Most Brn3b-positive and non-M1 ipRGCs innervate other brain regions, such as the olivary pretectal nucleus for pupillary light reflex (9). In addition to modulating animal behavior through the circadian clock, ipRGCs innervate many brain regions in the thalamus and hypothalamus (21). It is unclear whether and how ipRGCs regulate other cellular activities in peripheral tissues.

Skin is a large peripheral organ densely populated by SCs. It contains a huge number of hair follicles (HFs) that undergo

Significance

Intrinsically photosensitive retinal ganglion cells (ipRGCs) exhibit several important functions including the circadian photoentrainment, pupillary light reflex, alertness, and phototaxis. Whether ipRGCs regulate other physiological activities is unknown. We show that external light stimulation can activate hair follicle stem cells through the eyes via an ipRGC–suprachiasmatic nucleus–sympathetic nervous circuit. Immediately after ipRGCs are stimulated by light, the systemic sympathetic activities are activated. In skin, the local release of norepinephrine activates hair follicle stem cells. This neural circuit enables prompt communication between peripheral tissues and the external environment. Due to the systemic activation of sympathetic activities, this circuit can also allow for timely responses to external light in other organs. It also highlights a function of ipRGCs in regulating autonomic nervous activity.

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repetitive cycles of regeneration consisting of growth (anagen), regression (catagen), and rest (telogen) phases (22). Activation of each new regeneration cycle is fueled by HF stem cells (HFSCs) residing in the so-called “bulge” and “secondary hair germ” niche compartments (23, 24). When HFSC activity is suppressed, entrance to anagen is prohibited (25–29). HFSC activity has been shown to be regulated by intrinsic intracellular factors as well as by the signals from the local and systemic environments (1, 26, 30–35). In addition to these internal regulatory factors, HF regeneration can be altered by external cues (1, 36). Therefore, HF provide an opportunity for interrogating the mechanisms by which tissue SCs communicate with the outer environment and how changes in the external environment are perceived and transmitted to an internal SC niche. Using the HF regeneration paradigm, we uncovered a noncircadian neural circuit that activates HFSCs in response to light stimulation through retinal ipRGCs.

Results

Visible Light Stimulation to Eyes Induces Hair Regeneration. In mice, the second telogen in back skin lasts for ~6 wk starting from week 7 of age (22) (Fig. 1A). To test whether ocular light stimulation can influence HF regeneration, 7-wk-old C57BL/6 mice ($n = 10$) were exposed daily to blue light stimulation (463 ± 50 nm, 4 mW/cm^2 or $9.32 \times 10^{15} \text{ photons}\cdot\text{cm}^{-2}\cdot\text{s}^{-1}$; 2 J/cm^2 for

8 min and 20 s) to the eyes under anesthesia for 10 consecutive days at zeitgeber time (ZT)1–2 (1–2 h after light-on during the daily light/dark cycle) (Fig. 1A). During light stimulation, only the animal's head was exposed with the eyes kept open. Activation of a new anagen phase, marked by prominent darkening of the skin due to pigment production in anagen HFSCs (22, 26), was observed between day 5 and day 11 after the initial light stimulation (Fig. 1B and C). Consistently, regions of new hair growth first appeared simultaneously on both sides of the trunk and extended toward the central back (Fig. 1B). Such patterns are distinct from the largely synchronized hair growth on the whole back during the second physiological cycle (22), and also are different from the pattern of hair regrowth induced by direct light irradiation to the dorsal skin that first appeared homogeneously on the back and then extended to the lateral trunk (SI Appendix, Fig. S1D) (37). Skin of unirradiated control mice remained in telogen phase for more than 21 d ($n = 10$) (Fig. 1B and C). We also irradiated mice without anesthesia and found that mice became agitated and tried to avoid intense light, thus reducing the light stimulation to eyes. In this condition, anagen entry could be induced by increasing the daily blue light irradiation (6 J/cm^2 or $9.32 \times 10^{15} \text{ photons}\cdot\text{cm}^{-2}\cdot\text{s}^{-1}$ for 25 min) (SI Appendix, Fig. S1C). To provide a consistent amount of light exposure among animals, we irradiated mice under anesthesia for all the following experiments.

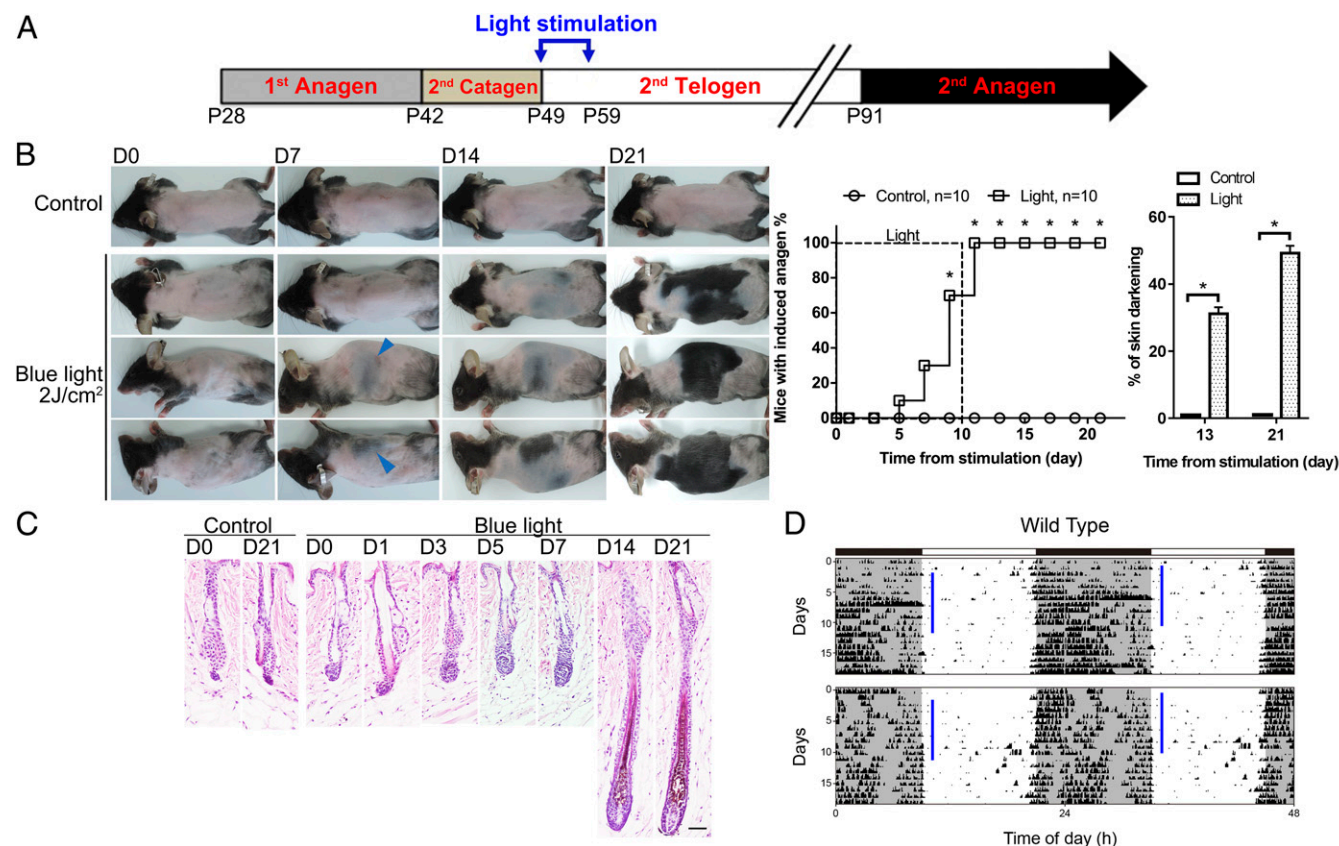


Fig. 1. Optic blue light stimulation promotes anagen entry without changing the ongoing circadian rhythm. (A) Postnatal mouse hair cycle and light stimulation. Daily light stimulation to the eyes was initiated on postnatal day 49 for 10 consecutive days. (B) Effect of daily optic blue light irradiation (463 ± 50 nm, 2 J/cm^2) on anagen entry. Successful induction of anagen was defined as more than 10% of the dorsal skin turning from pink to gray/black or with hair emergence. Anagen was first induced on sides of the trunk (blue arrowheads) and extended toward the central back. No hair regrowth was observed in unirradiated control mice. Dorsal skin with anagen entry (percentage of skin darkening) was significantly higher in the irradiated group on both day 13 and day 21. $*P < 0.05$ ($n = 10$). (C) Histology. Under blue light stimulation, premature anagen entry was induced, and HFSCs started to enlarge and elongate from day 3. In unirradiated control mice, HFSCs remained in telogen. (Scale bar, $50 \mu\text{m}$.) (D) Effect of daily optic blue light stimulation on the ongoing circadian rhythm. The ongoing circadian rhythm was not changed by daily blue light stimulation. Similar results were obtained in five independent experiments. Vertical blue bars indicated the time of light irradiation each day.

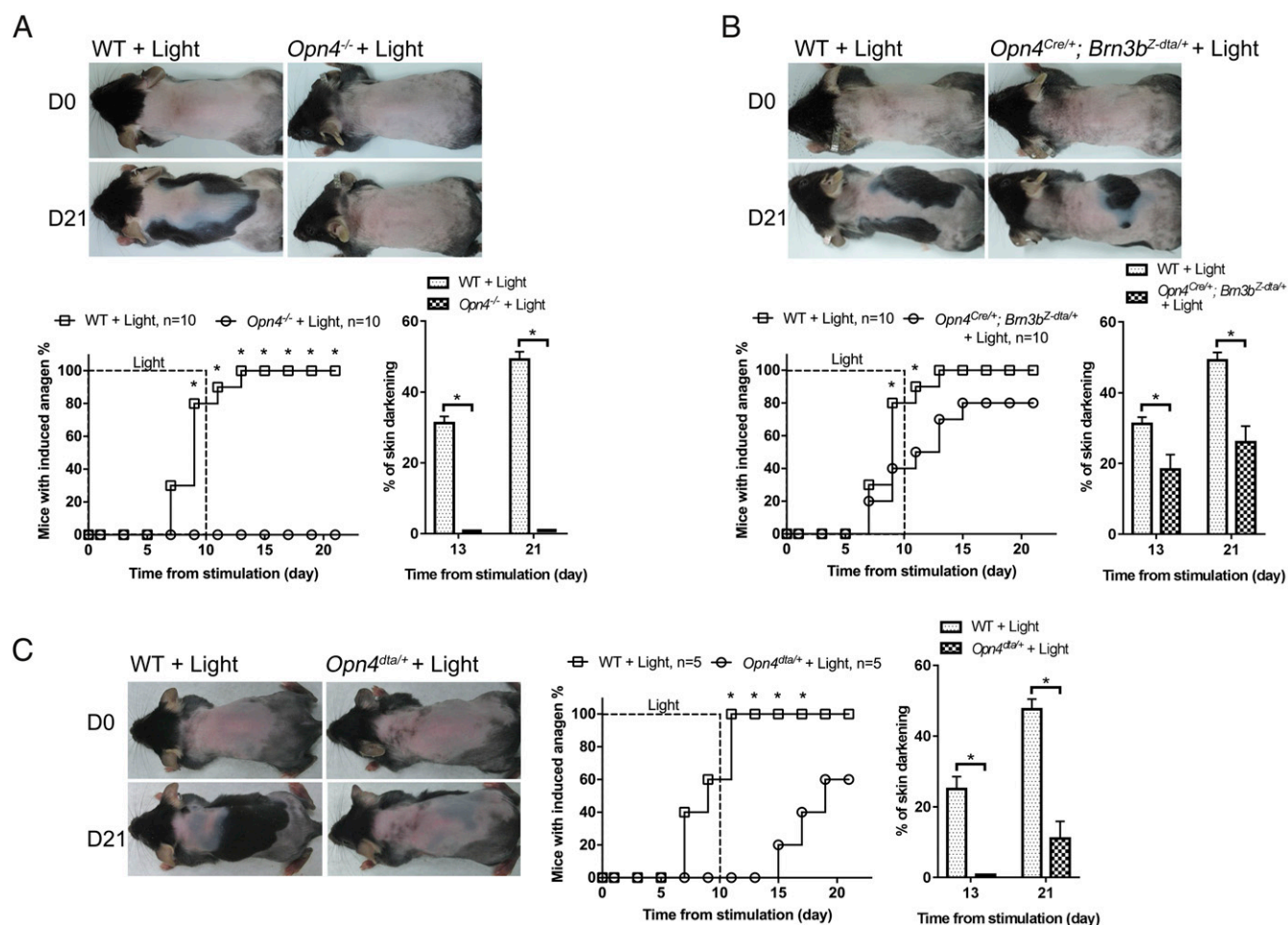


Fig. 3. Light triggers anagen entry through melanopsin and the ipRGC-to-SCN projection. (A) Effect of *Opn4* deletion. Light-induced anagen entry was completely abolished in *Opn4*^{-/-} mice. **P* < 0.05 (*n* = 10). (B) Effect of depleting *Brn3b*⁺ ipRGCs. Anagen could still be induced in *Opn4*^{Cre/+}; *Brn3b*^{Z-dta/+} mice, but light-induced anagen entry was slightly delayed and reduced in *Opn4*^{Cre/+}; *Brn3b*^{Z-dta/+} mice compared with wild-type mice. **P* < 0.05 (*n* = 10). (C) Effect of reduced neuronal input from ipRGCs. Reduced ipRGC numbers in *Opn4*^{dta/+} mice delayed and reduced light-induced anagen entry. **P* < 0.05 (*n* = 5).

sympathetic nervous system (41, 42). Whether light stimulation to ipRGCs can immediately alter the activity of the systemic endocrine or the sympathetic nervous system is unknown. We examined whether signaling relay to HF is mediated by systemic circulating factors. We generated parabiotic mice and illuminated the eyes of only one of the two paired animals. We confirmed the successful establishment of parabiosis by i.v. injecting a fluorescent dye in one mouse and detecting the fluorescence in the paired mouse before light irradiation. Accelerated hair growth was induced only in light-treated (*n* = 5) mice, but not in conjoint nontreated mice (*n* = 5) (Fig. 4A). It was shown that circulatory prolactin and melatonin might regulate HF growth (43–46). We found that, compared with control mice, only melatonin was slightly reduced 50 min after light exposure when mice were irradiated at ZT1–2 (SI Appendix, Fig. S6A). The results suggest that circulatory factors might not play a major role in light-induced HF regeneration.

Autonomic nerves regulate many cellular activities in peripheral tissues (47–49). Consistent with previous reports (50), we found that sympathetic nerves not only innervate arrector pili muscles of the HF (SI Appendix, Fig. S6C) but also project to the HFSC niche (Fig. 4B and SI Appendix, Fig. S6B). To determine whether sympathetic nerves are activated through the melanopsin photodetection system, we measured sympathetic activities in light-treated wild-type and *Opn4*^{-/-} mice. Ocular

light stimulation increased the heart rate, perspiration, and the renal sympathetic activity in wild-type mice but not in *Opn4*^{-/-} mutants (Fig. 5A–C). The increased heart rate and sympathetic activity continued for more than 40 min after light stimulation. With respect to HF, light-induced hair growth was blocked when cutaneous sympathetic innervation was disrupted by 6-hydroxydopamine (6-OHDA) treatment (Fig. 4B and C and SI Appendix, Fig. S6D). These results suggest that light stimulation to the eyes immediately activates sympathetic activities in multiple organs through a melanopsin signaling pathway from ipRGCs and promotes HF regeneration through cutaneous sympathetic nerves.

To further confirm the involvement of the cutaneous sympathetic nervous system, we quantified changes in cutaneous norepinephrine, the neurotransmitter released by peripheral sympathetic nerve endings. Five minutes after ocular illumination, norepinephrine levels increased by about 10-fold in the skin of wild-type mice but not in the skin of *Opn4*^{-/-} mice (*n* = 3) (Fig. 5D). Furthermore, consistent with previous reports (50, 51), we found that HFSCs express β 2-adrenergic receptors in all the HFSCs examined (*n* = 30) (Fig. 5E). We then treated skin topically with propranolol, a β -adrenergic antagonist, and found that light-induced HF regeneration was suppressed (*n* = 10) (Fig. 5F and SI Appendix, Fig. S6E). Together, these results reveal that

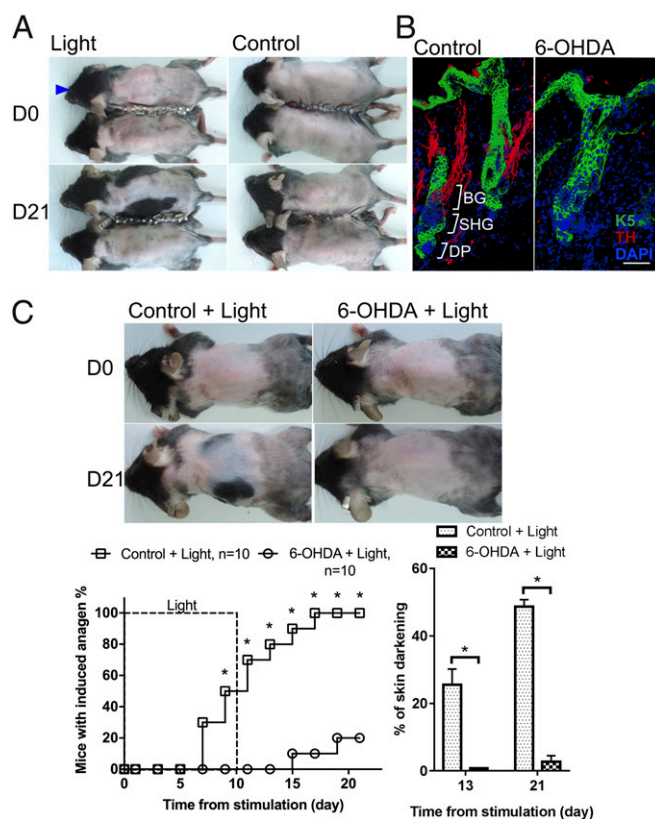


Fig. 4. Sympathetic nerves innervating HF are required for light-induced anagen entry. (A) Effect of parabiosis. Light stimulation induced anagen only in the irradiated mouse (blue arrowhead) but not in the paired non-irradiated mouse. Similar results were obtained in five independent experiments. (B) Compressed immunostaining images of thick (34- μ m) sections. (Left) In telogen, sympathetic nerves were in close proximity to bulge (BG) and secondary hair germ (SHG) areas before looping around arrector pili muscles. (Right) 6-OHDA treatment disrupted sympathetic innervation in all HF examined ($n = 30$). DP, dermal papilla; K5, keratin 5; TH, tyrosine hydroxylase. (Scale bar, 50 μ m.) (C) Effect of pharmacological sympathectomy with 6-OHDA. 6-OHDA treatment inhibited light-triggered anagen entry. * $P < 0.05$ ($n = 10$).

melanopsin-dependent local norepinephrine release in skin promotes HF regeneration.

Ocular Light Irradiation Activates HFSCs by Enhancing Hedgehog Signaling. Since new anagen entry is driven by HFSC activation (23, 24), we examined how light affects HFSC quiescence. BrdU pulse-labeling showed that HFSCs started to proliferate between day 3 and 5 after initial light stimulation (Fig. 6A). Gene ontology analysis of whole-skin transcriptomes 1 d after the initiation of light exposure revealed that ocular light stimulation activated multiple signaling pathways (Fig. 6B). Hedgehog signaling and hedgehog-associated basal cell carcinoma signaling were among the top five most significantly changed ontologies (Fig. 6B).

During the physiological telogen-to-anagen transition, Wnt signaling, required for anagen entry, is first activated in the HFSC compartment (24, 25, 52). In telogen, there is localized hedgehog signaling activity in the upper bulge and the secondary hair germ due to hedgehog ligand production from sensory nerves surrounding the upper bulge region and from dermal papilla cells below the hair germ (53, 54). After HFSCs enter anagen with prominent HFSC proliferation, hedgehog signaling is highly activated in the entire HFSC population due to the enhanced production of hedgehog ligands from early transit-amplifying cells (55, 56). Although not required for the initia-

tion of physiological anagen, the enhanced activation of hedgehog signaling early after anagen onset is essential for further growth and maturation of the lower segment of anagen HFSCs (55–57). We sorted HFSCs for further analysis 1 d after the initiation of light stimulation when HFSCs were still quiescent (Fig. 6A). We found that *Gli1* and *Gli2*, downstream targets of hedgehog signaling, were up-regulated in the HFSC compartment, including bulge SCs and hair germ SCs (Fig. 6C). These results showed that ocular light stimulation enhanced hedgehog signaling in HFSCs before they were activated to initiate anagen. To further test whether activation of hedgehog signaling is required for light-triggered anagen initiation, we topically administered cyclopamine, a Smoothened inhibitor, and GANT61, a *Gli1/2* inhibitor. Topical treatment with cyclopamine ($n = 10$) or GANT61 ($n = 5$) inhibited light-induced HFSC activation and hair cycle entry (Fig. 6D and E and *SI Appendix*, Fig. S7A and B). These results suggest that hedgehog signaling is activated and required for ocular light stimulation-induced HFSC activation and anagen initiation.

Discussion

In summary, we discovered a neural circuit initiating from M1-type ipRGCs in the retina that regulates HFSC activation in response to external photic cues (Fig. 7). Since ipRGCs also contribute to the photoentrainment of the central circadian clock in the SCN (3, 4), our results demonstrate a function of ipRGCs in the immediate conveyance of the external light message to internal peripheral tissues.

In *Opn4*^{−/−} mice and mice with crushed optic nerves, the light-induced anagen entry was completely abolished. In the mouse eyes, melanopsin is expressed only in ipRGCs and iris muscles (5, 58). Our results suggest the essential role of retinal ipRGCs in mediating the effect. The optic nerve crushing could potentially damage trigeminal nerves, and it was shown that optic nerve crushing could not completely block the activation of melanopsin-expressing trigeminal neurons (59). Our study showed that optic nerve crushing completely suppressed light-induced hair regeneration. Although it is unlikely that melanopsin-expressing trigeminal neurons are the primary photon detector for light-induced hair regeneration, we cannot completely rule out the potential contribution of this non-retinal pathway. Because light is able to promote hair regeneration in *Opn4*^{Cre/+}; *Bmn3b*^{Z-*tda*/+} mice in which only M1 ipRGCs (~200 cells per eye) (*SI Appendix*, Fig. S5A) that selectively innervate the SCN are preserved, our data suggest that M1 ipRGCs and the SCN are part of the neural circuitry that can transmit light information to alter peripheral SC activity. This mechanism is mediated through the retinohypothalamic tract that signals to the SCN to activate the sympathetic nervous system (Fig. 7). The stimulated sympathetic activity here is distinct from its daily oscillation controlled by the central circadian clock (49). Compared with the circadian clock that optimizes cellular activities in anticipation of daily rhythmic environmental changes, the circuit described here allows immediate cellular responses to external environmental cues. Since projection of ipRGCs through the retinohypothalamic tract to SCN is also essential for photoentrainment of the central circadian clock (3), our results also show the multifunctionality of the ipRGC-SCN pathway both in regulating long-term daily oscillatory activities and in relaying immediate photic information to peripheral tissues. Therefore, in addition to its known functions, we demonstrate another important role of ipRGCs in regulating physiological functions in peripheral tissues through the sympathetic nervous system. Since photoentrainment of the central circadian clock and direct optic stimulation of sympathetic activities both require the projection of ipRGCs to the SCN (60, 61), it is an intriguing question whether distinct ipRGCs or the same ipRGCs innervating SCN are employed for these two functions.

Due to the sophisticated network of both sensory and sympathetic nerves around HFSCs and their dynamic remodeling

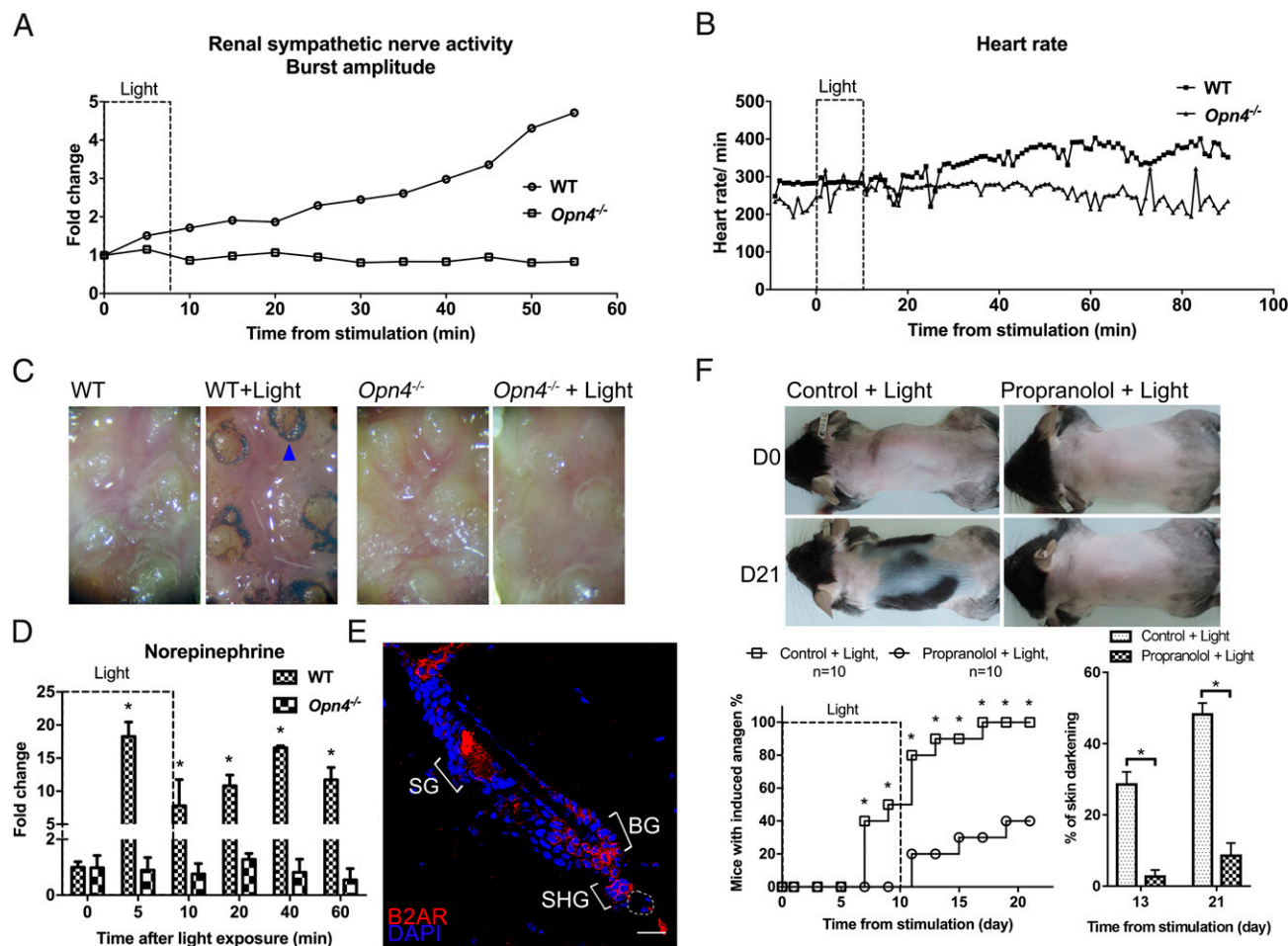


Fig. 5. Optic light stimulation activates sympathetic nerves and promotes anagen entry by enhancing norepinephrine release in skin. (A) Effect of light stimulation on renal sympathetic nerve activity. Sympathetic activity was increased by optic blue light stimulation in wild-type mice but not in *Opn4*^{-/-} mice. Similar results were obtained in three independent experiments. (B) Effect of light stimulation on heart rates. The heart rate was increased by optic blue light stimulation in wild-type mice but not in *Opn4*^{-/-} mice. Similar results were obtained in three independent experiments. (C) Effect of light stimulation on foot-pad sweating. Optic blue light stimulation induced sweating (blue arrowhead) in wild-type mice but not in *Opn4*^{-/-} mice. Similar results were obtained in three independent experiments. (D) Effect of light on cutaneous norepinephrine release. Norepinephrine was highly increased after light irradiation in wild-type mice but not in *Opn4*^{-/-} mice. **P* < 0.05 compared with time 0 (*n* = 3). (E) Immunostaining. Enriched expression of β -2 adrenergic receptor (B2AR) in the HFSC compartment. A similar pattern of expression was observed in all HFSCs examined (*n* = 30). BG, bulge; SG, sebaceous gland; SHG, secondary hair germ. Dashed line represents dermal papilla. (Scale bar, 50 μ m.) (F) Effect of β -adrenergic antagonist. Topical propranolol inhibited light-induced anagen entry. **P* < 0.05 (*n* = 10).

around hair cycles, piloneural interactions have been suggested to be involved in regulating the physiology of HFSCs (50, 62, 63). However, whether the piloneural network can directly activate HFSCs for hair regeneration was unknown. Sensory nerves have been shown to densely innervate the upper bulge region of HFSCs and to regulate a small population of SCs in the upper bulge area by providing the ligand of sonic hedgehog (53, 62). The focally intensified hedgehog signaling in the upper bulge region defines a population of SCs that are able to become interfollicular epidermal cells upon skin wounding but is not sufficient to activate the general HFSC population to initiate a new anagen (53). Sympathetic nerves also loop around the HFSC niche when they travel from the deep dermal nervous plexus to innervate the arrector pili muscle for the piloerection function (50). Sympathetic nerves were suggested to promote HF growth after a new anagen is already initiated (50). In humans, β -blockers can induce hair loss, and defective sympathetic regulation is also associated with dysregulated hair growth (64, 65). It remained unclear whether sympathetic nerves can directly activate HFSCs. Our results show that activation of sympathetic nerves can acti-

vate HFSCs by enhancing hedgehog signaling to promote anagen entry. This also supports the findings that enhancing hedgehog signaling by forced dermal or epidermal expression of sonic hedgehog ligands or by treatment with hedgehog agonists can promote the telogen-to-anagen transition (55, 57, 66). In our other testing, we irradiated the mice at ZT5–8 and found that hair regeneration could also be induced, but the hair regeneration was a bit slower than during ZT1–2. It has been shown that intrafollicular clock activity regulates the priming state of HFSCs as well as the activity of other follicular cells and that autonomic nerves can regulate local tissue clock (67–70); the responsiveness of HFSCs to light irradiation might vary as the circadian time of light irradiation is changed. Whether sympathetic nerves regulate hedgehog signaling through modulating the local clock is to be determined. Since intense white light stimulation to eyes not only activates sympathetic nerves but also suppresses the parasympathetic activity (71, 72), it is possible that blue light here might also suppress parasympathetic activities in skin to regulate HFSC activity. In addition to neural control, HF growth is also regulated by neurohormones, such as prolactin and melatonin,

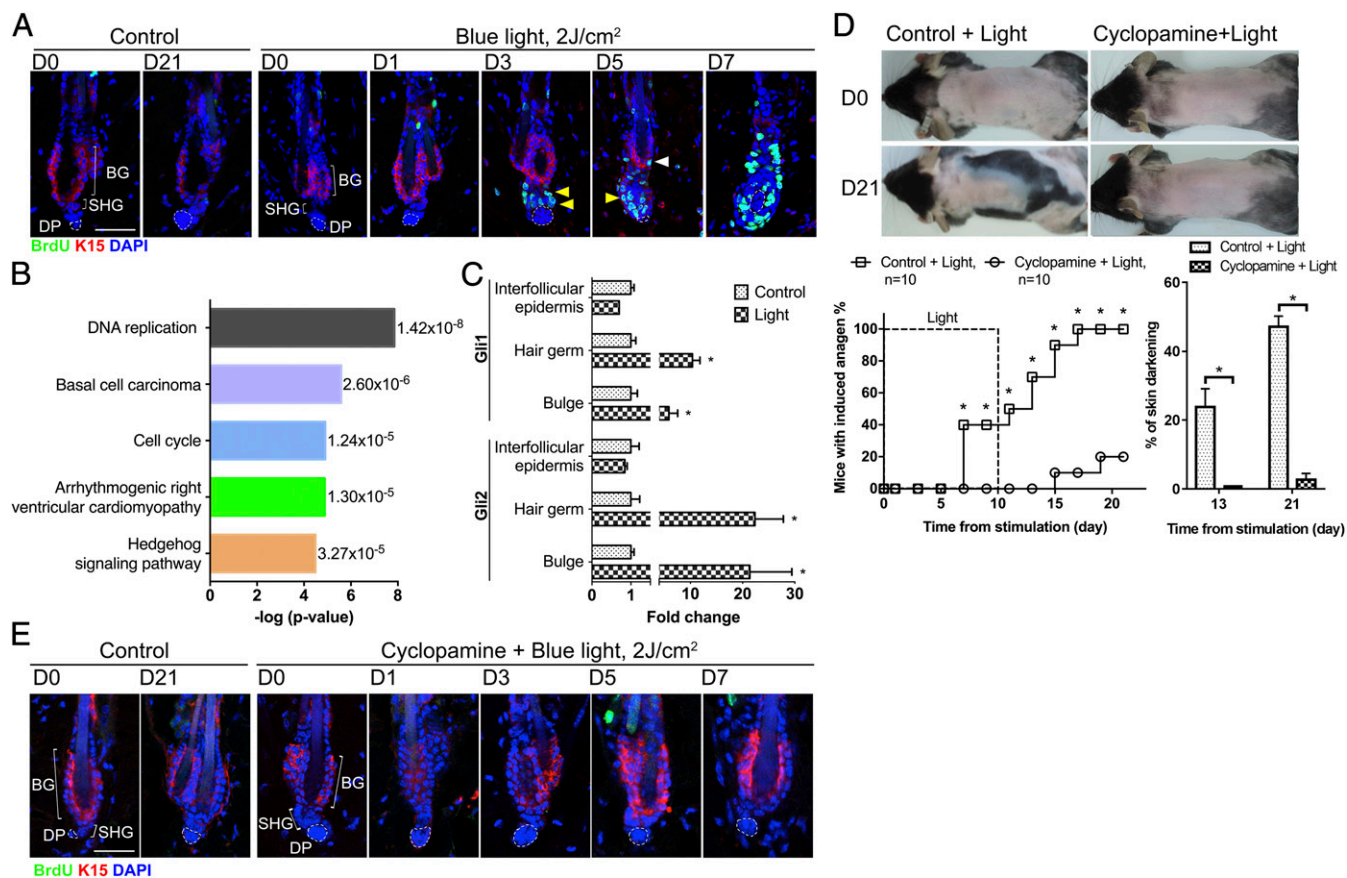


Fig. 6. Light enhances hedgehog signaling in HFSCs to promote anagen entry. (A) Pulse BrdU labeling under daily 2 J/cm² blue light stimulation. Cell proliferation of HFSCs was first observed in secondary hair germs on day 3 (yellow arrowheads) and then in the bulge area on day 5 (white arrowheads). In the nonirradiated control group, no cell proliferation was observed in the HFSC compartment. Similar results were observed in all HFSCs examined (*n* = 30). (Scale bar, 50 μm.) BG, bulge; DP, dermal papilla; SHG, secondary hair germ. (B) Analysis of gene expression 1 d after initiation of light stimulation by RNA-sequencing. The top five signaling pathways enhanced by light stimulation are shown here. *P* values are shown on the right. (C) Gene expression in HFSCs determined by quantitative PCR 1 d after initiation of light stimulation. Expression of *Gli1* and *Gli2* was up-regulated in bulge and secondary hair germ SCs after light irradiation. In the inter-follicular epidermis, expression of *Gli1* and *Gli2* was not significantly increased. **P* < 0.05, control vs. light-treated group (*n* = 3). (D and E) Effect of hedgehog pathway inhibitor. (D) Topical cyclopamine inhibited light-triggered anagen entry. **P* < 0.05 (*n* = 10). (E) Pulse BrdU labeling under cyclopamine treatment. Cyclopamine inhibited light-triggered HFSC activation. Similar results were observed in all HFSCs examined (*n* = 30). (Scale bar, 50 μm.)

and these neurohormones can also be produced intrafollicularly (43–46). Although we did not detect extensive changes in plasma prolactin and melatonin after blue light irradiation, the intrafollicular melatonin might be changed due to the local release of noradrenaline (45, 46).

HFSCs respond to both short-range and long-range signals from the surrounding tissues and systemic environment, including HF dermal papilla cells, fibroblasts, adipose tissue, macrophages, resident regulatory T cells, neighboring HFs, and systemic hormones (1, 26, 30–35). The response of HFSCs to external light not only adds another layer to HF regeneration mechanisms but also raises a broader question with respect to the definition of SC niches. Conventionally, the SC niches have been defined as a constellation of neighboring cells and local extracellular factors that constitute the microenvironment in proximity to SCs (73). Here we show that the external environment can also regulate the internal SCs through an ipRGC–SCN–sympathetic nervous pathway. Hence, the external environment should also be considered part of the SC niche components, and internal SCs can be linked to the external component of the SC niche by the neural circuit we show here. We can consider the SC niche at three levels, i.e., the local SC microenvironment as the “microniche,” the systemic environment (such as circulatory hormones) inside the body as the

“macroniche,” and the external environment as the “meganiche.” The ipRGC–SCN–sympathetic nervous pathway enables quick regulation of tissue SCs by the meganiche through modulating the microniche.

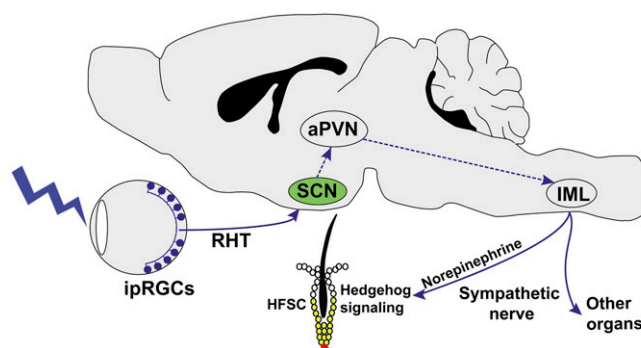


Fig. 7. Depiction of the neural pathway for light-stimulated sympathetic activity and HFSC activation. aPVN, autonomic neurons of the paraventricular nucleus; HFSC, hair follicle stem cell; IML, intermediolateral cell column; RHT, retinohypothalamic tract. Dashed lines indicate proposed neural relays for this pathway.

For animals, especially nocturnal mice, intense ambient light can be a danger signal that warrants immediate reaction. In physiological conditions, mice would not receive prolonged intense light stimulation to the eyes during the daytime unless they are threatened, such as being chased by predators or forced to move due to harsh environmental changes. Without plucking, the old mouse pelage hair within an HF is normally not shed as the new hair forms (74). Initiation of a new round of hair growth by light stimulation may make a denser hair coat for additional protection from the harsh environment or timely replenish hair accidentally lost in a stressed condition. Here we demonstrated the indirect stimulating effect of light on hair regeneration through melanopsin in ipRGCs. We have shown that direct irradiation of skin by visible light can also promote the telogen-to-anagen transition by enhancing epithelial-mesenchymal interactions and that red light is more potent than green/blue light (37). Direct irradiation to HFs also promoted hair growth in vitro (75). Hence, visible light of different spectra possesses differential direct and indirect stimulatory cues to HFSCs. In the wild, the murine body is covered by a dense hair coat, and indirect stimulation by light through the eyes would be a more prominent mechanism in the regulation of HFSCs. Although melanopsin (*Opn4*) is not expressed in HFs, rhodopsin (*Opn2*) and panopsin (or encephalopsin) (*Opn3*) are present in HFs (75). The stimulating effect of direct light irradiation to the skin may be mediated in part through nonvisual effects of these opsins (75).

Our result is consistent with previous reports showing that intense white light can alter the autonomic nervous activities in rodents (41, 71). We show that, compared with green light of a longer wavelength, blue light is more efficient in activating the ipRGC-SCN-sympathetic nervous pathway due to melanopsin's preferential absorption of light in the blue light spectrum (76). In addition to HFSCs, SCs in other tissues are also regulated by autonomic nerves (47, 49, 77, 78). The activity of tissue SCs exhibits circadian changes corresponding to the daily oscillation of autonomic nervous activities governed by the central circadian clock (49). The ability to modulate SC activities beyond its circadian changes might be applied for clinical purposes (79). Since the light-activated sympathetic activity is not limited to skin (Fig. 5A–C) (41, 71, 72), the ipRGC-SCN-sympathetic circuit might have a role in coordinating systemic responses and hence have a broader effect on the behavior of SCs and physiological activities in other organs.

Materials and Methods

Animals. The animal experiments were approved by Animal Care and Use Committee of National Taiwan University. Female mice were used in this study. Animals were fed ad libitum and housed in animal facilities with 12-h light/12-h dark cycles. C57BL/6 mice were purchased from the Taiwan National Laboratory Animal Center. The transgenic mice *Opn4*^{−/−}, *Gnat1*^{−/−}, *Gnat2*^{cpfl3/cpfl3}, *Opn4*^{Cret/+}, *Brn3b*^{Z-dta/+}, and *Opn4*^{dta/+} were generated with a mixed background (C57BL/6; 129SvJ) and were backcrossed to C57BL/6 mice; they have been described in our prior work (9, 80). They were kindly provided by Samer Hattar (Johns Hopkins University, Baltimore). For experimental procedures, mice were anesthetized with a mixture of zolazepam (Zoletil veterinary anesthesia; Virbac Laboratories) and xylazine (Rompun; Bayer) (4:1, vol/vol) unless stated otherwise.

Light Irradiation. A tailor-made light box with light-emitting diodes of adjustable light intensity on the ceiling (Lustrous Technology Ltd.) was employed (37). Light-emitting diodes of two different wavelengths were used: blue light (463 ± 50 nm, 4 mW/cm² or 9.32 × 10¹⁵ photons·cm^{−2}·s^{−1}) and green light (522 ± 50 nm, 3 mW/cm² or 7.88 × 10¹⁵ photons·cm^{−2}·s^{−1}). Light intensity was measured by a digital light meter (DLM 530; Tecpel) at the level of the mouse eyes, and the intensity of light of each wavelength was kept constant in our experiments. The dose of light exposure was adjusted by varying the time mice spent in the light box. The temperature fluctuation within the light box was less than 1 °C during light irradiation. The dorsal skin of 7-wk-old female mice whose dorsal hair cycle was in the second telogen phase was carefully shaved first with an electric clipper (Oster Golden A5 Single Speed Animal Grooming

Clipper with Oster Cryogen-X #50 Pet Clipper Blades) and then with a razor (Braun MobileShave M-60) without damaging the skin. The mice were shielded by aluminum foil from the neck to the tail to avoid light irradiation, and only the heads were exposed. The mice were in a prone position with eyes facing up toward the light-emitting diodes. Under anesthesia, their eyes opened spontaneously. The mice were exposed daily to blue light irradiation (4 mW/cm² or 9.32 × 10¹⁵ photons·cm^{−2}·s^{−1}; 8 min 20 s/d) or to green light (3 mW/cm² or 7.88 × 10¹⁵ photons·cm^{−2}·s^{−1}; 11 min 7 s/d) at 2 J/cm² at ZT1–2 (1–2 h after lights on during the daily light/dark cycle) for 10 consecutive days. To determine the dose effects, irradiation was reduced to 0.5 J·cm^{−2}·d^{−1} with the same light intensity. For control, mice were given the same treatment without the light on. To compare hair regrowth patterns, mice were irradiated by blue light (2 J·cm^{−2}·d^{−1}) to the dorsal skin only at ZT1–2 for 10 consecutive days, as we previously described (37). Mice were irradiated under anesthesia in our light stimulation experiments unless stated otherwise. We also irradiated the mice with blue light (6 J/cm² or 9.32 × 10¹⁵ photons·cm^{−2}·s^{−1} for 25 min) in the unanesthetized condition.

Anagen initiation was quantified by the change in skin color as we previously described (26, 37). Anagen was determined by the skin color changing from pink to gray/black (22, 26). The skin color of the mice was examined visually and photographed to determine the area of anagen entry. The proportion of the dorsal pigmented region over the entire dorsum was calculated by the software Image J (NIH; <https://rsb.info.nih.gov/ij/>) as we previously described (37). Successful induction of anagen entry was defined as more than 10% of the dorsal skin turning from pink to gray/black or with hair emergence from the skin.

Wheel-Running Activity. The mice were placed individually in cages with a running wheel under 12-h light/12-h dark cycles. Their activity was recorded, and the circadian period was calculated based on the ClockLab algorithm (Actimetrics). *Opn4*^{dta/+} mice were given light stimulation at CT1 (circadian time) for 10 d.

Histology and Immunofluorescence. To collect skin specimens for histological analysis and immunostaining, mice were killed at different time points before and after the start of light irradiation. To gauge cell proliferation, BrdU pulse labeling was performed by i.p. injection of 10 mg/kg BrdU (Sigma-Aldrich) 1 h before mice were killed. The skin samples were fixed in 4% paraformaldehyde (PFA) (AMRESCO) in PBS AMRESCO overnight, serially dehydrated with alcohol, and embedded in paraffin. Paraffin sections (5 μm in thickness) were stained with H&E or were prepared for immunostaining following routine procedures. Antibodies and additional details are described in *SI Appendix, SI Materials and Methods*.

Parabiosis, Nerve Crushing, Pharmacological Sympathectomy. Parabiosis was performed in 6-wk-old mice to connect them along the adjacent flanks as described (81). Their skin was connected with sutures or surgical autoclips (9-mm; Clay Adams). To confirm the successful establishment of parabiosis, one of the paired C57BL/6 mice was i.v. injected with 100 μL of fluorescent vascular imaging reagent [20 μL of Qtracker 655 (Thermo Fisher) dissolved in PBS]. Fluorescent images from the shaved dorsal side of the mice were taken by an in vivo imaging system (IVIS; Xenogen; Perkin-Elmer) at 30 and 90 min after injection. Successful parabiosis was determined by the progressive increase of fluorescence across the dorsal skin in the other of the paired mice. Only mice with confirmed successful parabiosis were used for the experiments.

To disrupt the optic nerves, optic nerves were crushed as previously described at the age of 6 wk (82, 83). The pupil light reflex was determined before and after nerve crushing, as we previously described (9). The inhibition of light reflex indicated the success of optic nerve crushing. Anterograde tracer fluorescent cholera toxin B (recombinant cholera toxin subunit B, Alexa Fluor 488 conjugate; Thermo Fisher) was also injected intravitreally to label the ganglion cells and the axons to confirm the success of optic nerve crushing (84).

Sympathectomy by s.c. injection of 10 mg/kg 6-OHDA (Tocris 2547) was performed 3 d before light stimulation (85). The 6-OHDA was divided into four equal doses in saline, which were injected into each of the four quadrants of the back. For controls, vehicle alone was used. To determine the completeness of pharmacological sympathetic denervation by 6-OHDA, the innervation of sympathetic nerves to HFs was compared in serial sections in 25–35 HFs.

Topical Drug Treatment. Propranolol (10 mg/kg body weight; Sigma P0884), cyclopamine (2 mg/kg body weight; ApexBio A8340), or GANT61 (50 mg/kg body weight; ApexBio) was dissolved in hand cream (Neutrogena Norwegian Formula Concentrated Hand Cream) at a concentration of 10 mg drug/1 g

cream. The propranolol cream, cyclopamine cream, or GANT61 cream was applied to the shaved back of mice twice a day (50 mg cream per mouse) for 10 d beginning on the first day of light irradiation. For controls, vehicle alone was used.

Heart Rate Recording. During the experiment, body temperature was maintained at 37 °C using a thermostatically regulated heating platform. Under anesthesia, heart rate and blood pressure were measured using the tail-cuff method (BP-2000 blood pressure analysis system; Visitech Systems, Inc.). The measurement was recorded at baseline for 10 min, and then the mouse eyes were exposed to blue light (2 J/cm², 8 min 20 s of light exposure). The recordings then lasted for 1.5 h. The control group received the same treatment without exposure to blue light.

Starch/Iodine Sweat Test. The starch/iodine method was conducted as previously described to detect sweating (86). Briefly, iodine/alcohol (2 g iodine/100 mL ethanol) was painted on the plantar surface of the rear paw. After the iodine/alcohol had dried, the rear paw surfaces were covered in a starch-oil suspension (100 g starch powder/100 mL castor oil). The mice were then irradiated with blue light (2 J/cm², 8 min 20 s of light exposure). After light exposure, sweating was determined by the change in skin color from brown to dark blue/black. The control group received the same treatment without exposure to blue light.

Recordings and Analysis of Renal Sympathetic Nerve Activity. To monitor sympathetic nerve activity, surgery was performed to expose the renal sympathetic nerve, and the renal sympathetic activity was recorded as previously described (87). Additional details can be found in *SI Appendix, SI Materials and Methods*.

Quantification of Cutaneous Norepinephrine Level and Plasma Prolactin and Melatonin Levels. To measure the changes of skin norepinephrine level, full-thickness mouse skin was obtained before and after light irradiation at different time points. The skin specimen was homogenized, and the norepinephrine concentration was determined by a norepinephrine ELISA kit (KA1891; Abnova) according to the manufacturer's instructions. The plasma levels of prolactin and melatonin were determined by ELISA kits for prolactin and melatonin (E-EL-M0788 and E-EL-M0083; Elabscience) according to the manufacturer's instructions. The lowest detection limits were 0.16 ng/mL for prolactin and 7.81 pg/mL for melatonin.

RNA-Sequencing and Analysis. Two hours after blue light stimulation on day 2, mouse skin was collected for RNA extraction using TRIzol reagent (Invitrogen) and the RNeasy Mini Kit (Qiagen). The cDNA was synthesized with SuperScript II reverse transcriptase (Invitrogen), and the resulting short DNA fragments were purified by the QIAquick PCR Purification Kit (Qiagen). Further sequencing and analysis are described in *SI Appendix, SI Materials and Methods*.

FACS. Bulge stem cells, hair germ stem cells, and interfollicular keratinocytes were isolated from the dorsal skin through FACS 2 h after blue light stimulation on day 2. Cells were prepared for FACS as described (88). Single-cell suspensions were then incubated with antibodies for 30 min at 4 °C. The following antibodies were employed: CD34-eFluor 660 (1:100; eBioscience); Sca1-PE-cyanine7 (1:100; eBioscience); α 6-integrin-phycoerythrin (1:500; eBioscience); and CD200-FITC (1:50; Bio-Rad). Cells were sorted by a FACSAria III cell sorter (BD Biosciences). Cells were sorted as follows: bulge stem cells were Sca1⁺/ α 6-integrin⁺/CD200⁺/CD34⁺; hair germ stem cells were Sca1⁺/ α 6-integrin⁺/CD200⁺/CD34⁺; and interfollicular keratinocytes were Sca1⁺/ α 6-integrin⁺/CD200⁺/CD34⁺.

Real-Time PCR. Total RNA was extracted from sorted cells by TRIzol reagent (Invitrogen). cDNAs were synthesized with a RevertAid H Minus First Strand cDNA Synthesis Kit (Thermo Scientific). Quantitative real-time PCR was performed as described (89). The sequences of the primers of mouse genes were as follows: *Gapdh* (GU214026.1, 5'-CGTAGACAAAATGGTGAAGTCCG-3' and 5'-AAGCAGTTGGTGCAGGATG-3'); *Gli1* (NM_010296.2, 5'-ATCACCTGTGGGATGCTGGAT-3' and 5'-GGCGTGAATAGGACTTCCGACAG-3'); and *Gli2* (NM_001081125.1, 5'-GTTCGAAGGCCTACTCTCGCTG-3 and 5'-CTTGAGCAGTGGAGCACGGACAT-3').

Statistics. χ^2 test was employed for the comparison of premature anagen entry in different groups using GraphPad Prism version 5.00 (GraphPad Software). In most light-irradiation experiments, each group contained 10 mice; in some experiments, each group contained five to seven mice. Differences were considered statistically significant when the *P* value was <0.05. In the analysis of gene expression, data were expressed as the mean $\pm$ SEM. A two-tailed Student's *t* test was performed for comparisons among groups using GraphPad Prism software. A difference was considered significant when the *P* value was <0.05. Data were plotted using GraphPad Prism. At least two independent experiments were performed for comparisons.

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