In the central nervous system, glycogen-derived bioenergetic resources in astrocytes help promote tissue survival in response to focal neuronal stress. However, our understanding of the extent to which these resources are mobilized and utilized during neurodegeneration, especially in nearby regions that are not actively degenerating, remains incomplete. Here we modeled neurodegeneration in glaucoma, the world’s leading cause of irreversible blindness, and measured how metabolites mobilize through astrocyte gap junctions composed of connexin 43 (Cx43). We elevated intraocular pressure in one eye and determined how astrocyte-derived metabolites in the contralateral optic projection responded. Remarkably, astrocyte networks expand and redistribute metabolites along distances even 10 mm in length, donating resources from the unstressed to the stressed projection in response to intraocular pressure elevation. While resource donation improves axon function and visual acuity in the directly stressed projection, it renders the donating tissue susceptible to bioenergetic, structural, and physiological degradation. Intriguingly, when both projections are stressed in a WT animal, axon function and visual acuity equilibrate between the two projections even when each projection is stressed for a different length of time. This equilibration does not occur when Cx43 is not present. Thus, Cx43-mediated astrocyte metabolic networks serve as an endogenous mechanism used to mitigate bioenergetic stress and distribute the impact of neurodegenerative disease processes. Redistribution ultimately renders the donating optic nerve vulnerable to further metabolic stress, which could explain why local neurodegeneration does not remain confined, but eventually impacts healthy regions of the brain more broadly.

The brain accounts for 30 to 50% of our resting rate of energy consumption, despite comprising only 2% of human body mass (1–3). Neurons rely primarily on glucose as a substrate for ATP production, which depletes quickly during periods of high demand or stress (4–6). In the central nervous system, neuronal energy needs are satisfied through interactions with astrocytic glia, which create and store glycogen as a safeguard against periods of stress (4, 7–11). This metabolic collaboration between neurons and astrocytes is critical to tissue survival during injury, such as in ischemia, and in neurodegenerative disease, such as during Alzheimer’s, Parkinson’s, and Huntington’s diseases, among many others (7, 12, 13). Such conditions tax normal energy utilization through increased oxidative stress and diminished metabolic capacity (7, 14, 15).

Like other age-related neurodegenerations, glaucomatous optic neuropathy (or glaucoma) involves astrocyte remodeling during insult, in this case to the axons of retinal ganglion cell neurons that form the optic projection to the brain (16). These axons include a lengthy unmyelinated segment that traverses the retina that is especially vulnerable to metabolic and oxidative stress in glaucoma (17, 18). This stress arises from sensitivity to intraocular pressure (IOP) conveyed at the optic nerve head (19–21), which underlies glaucoma’s stature as the world’s leading cause of irreversible (i.e., neurological) blindness (22). Early progression involves axonal dysfunction, including degradation of anterograde transport to central projection sites in the brain and accumulation of phosphorylated neurofilaments linked to local depletion of mitochondria (15, 23, 24). These changes map closely to extensive redistribution of astrocyte processes in the optic nerve (23, 25–27). Astrocyte networks share cytoplasmic information via gap junctions comprised primarily of connexin 43 (Cx43) (28, 29), which is up-regulated in the glaucomatous optic nerve head (30). Since Cx43 is permeable to molecules up to 1 to 1.2 kDa (28, 31), we probed to what extent astrocyte gap junction coupling mediates redistribution of much smaller energy substrates in response to neurodegenerative stress in the optic projection. We find that expanding Cx43-mediated networks of astrocytes redistribute metabolites over the exceptional distances between unstressed and stressed optic nerve projections from the two eyes. While preventing this distribution accelerates loss of neuronal function and degradation of visual acuity, redistribution renders donating tissue vulnerable to further metabolic stress. Thus, we posit that astrocyte networks utilize gap junctions through an endogenously protective mechanism that serves as a metabolic buffer against bioenergetic stress to distribute the impact of degenerative processes and preserve neuronal function.

**Significance**

Astrocytes respond to neuronal energy needs by mobilizing metabolic stores. How these resources are mobilized during neurodegeneration, especially in nearby regions that are not actively degenerating, could impact new therapeutic approaches. We modeled neurodegeneration in glaucoma and measured how metabolites mobilize through astrocyte networks. We elevated pressure in one eye and determined how astrocyte-derived metabolites in the opposite eye and nerve responded. Remarkably, astrocyte networks expand and redistribute metabolites along distances nearly 1 cm in length to donate resources from unstressed to stressed tissue. While resource donation improves axon function and visual acuity, it renders the donating tissue susceptible to further stress, which could help explain why local neurodegeneration eventually impacts healthy regions of the brain.
resources occurs between healthy and degenerating tissue, we assessed the function of Cx43 in the optic nerve. We generated a mouse that allowed us to conditionally excise Cx43 from astrocytes under temporal and local control after mice reached 2 mo of age, 1 wk before any IOP elevation. We verified excision of the floxed portion of the Cx43 gene in the retina, optic nerve, and cerebellum of GFAP-Cre-ER$^{12} \times$ Cx43$^{flox/flox}$ mice following tamoxifen gavage (knockout, KO) compared to gavage without tamoxifen (knockout control or KO-Ctrl) (Fig. 1 A and B). Induction successfully reduced label of Cx43 puncta by 98% across the retina, optic nerve, and superior colliculus (SC), the primary retinal projection site in the rodent brain (32, 33), when averaged between independent animals and tissues (4.06 $\times$ 10$^5$ ± 1.28 $\times$ 10$^5$ puncta/mm$^3$ vs. 8.38 $\times$ 10$^3$ ± 6.65 $\times$ 10$^3$; $P < 0.001$) (Fig. 1 C).

In KO-Ctrl mice (without tamoxifen), Cx43 puncta distribute as expected along GFAP-labeled astrocyte processes and not in ganglion cells or their axons; tamoxifen treatment eradicated the majority of puncta (Fig. 1 D–F).

Using these mice, we next measured how conditional loss of Cx43 influences astrocyte glycogen in naïve optic nerve compared to nerves stressed by increasing periods of unilateral elevation in IOP and to their contralateral counterparts (Fig. 2 A). Glycogen within the central nervous system is stored largely within astrocytes and represents the main reserve of metabolites for nervous tissues to utilize during periods of energetic deficit (34, 35). Glycogen stores diminish in the early stages of many

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**Fig. 1.** Cx43 reduction in GFAP-Cre-ER$^{12} \times$ Cx43$^{flox/flox}$ mice. (A) Conditional mutagenesis of Cx43$^{flox/flox}$ showing primers (arrows) for genotyping (blue) and verification of excision (red). (B) PCR products from nervous tissue in Cx43$^{flox/flox}$ with genotype (G) and KO verification following induction. (C) Cx43+ puncta/ mm$^3$ decreased in KO retina, optic nerve, and SC compared to KO-Ctrl ($P < 0.001$; mouse numbers indicated). (D–F) KO reduces Cx43 immunolabel in GFAP-expressing astrocytes compared to KO-Ctrl. CTB labels retinal ganglion cells and their axon projections. (Scale bars, 40 $\mu$m in D and E and 20 $\mu$m in F.)
neurodegenerative diseases and insults that include metabolic stress (12, 36, 37). In the following experiments, we utilized the microbead occlusion model of glaucoma, a highly repeatable, widely utilized tool for IOP elevation (38). In this model, microbeads are injected into the anterior chamber of the eye where they impede aqueous fluid outflow, causing an increase in IOP of about 35%. This increase is similar to that of human patients with elevated IOP (39), causing gradual progression of neurodegeneration over the following 2 to 8 wk in rodents (16, 17).

In WT (C57/BL6) mice (Fig. 2B, Left), 4 d of unilateral IOP elevation significantly diminished glycogen compared to both the optic nerve from naïve mice (−36.9%; P < 0.001) and to the unstressed contralateral nerve from the same animal (−35.5%; P < 0.001). Glycogen did not differ significantly after 1 or 2 wk of elevation compared to 4 d (P ≥ 0.32), although both exposures reduced glycogen compared to naïve mice (−33.1 and −35.1%, respectively; P ≤ 0.001). Surprisingly, following 1 and 2 wk of IOP elevation, glycogen decreased even more dramatically in the unstressed contralateral optic nerve compared to the nerve stressed by IOP (P < 0.001, P = 0.006, respectively). The decrease was also significant compared to naïve mice (−54.3 and −56.4%, respectively; P ≤ 0.001) and to nerves with 4 d of IOP elevation (−53.3 and 55.5%, respectively; P ≤ 0.001). Four weeks of elevated IOP directly diminished optic nerve glycogen even further, compared to 1 wk (−36.0%, P < 0.001) and 2 wk (−39.7%, P < 0.001) of elevation and to naïve mice (−59.6%, P < 0.001). However, at this point glycogen in the contralateral nerve did not differ from that in the IOP-stressed nerve (P = 0.202), suggesting an equilibration.

We made subsequent measurements using IOP elevations of 4 d and 1 wk in KO and KO-Ctrl mice, as these times represent the period during which glycogen in the unstressed contralateral nerve most dramatically changed in the WT. As expected in KO-Ctrl

Fig. 2. Glycogen redistribution through astrocyte Cx43. (A) Elevated (Δ, +35%) vs. contralateral (C) IOP in WT (Left) and transgenic (Right) mouse eyes. (B) In WT mice (Left), microbead-elevated (M) IOP reduces glycogen vs. naïve (N) optic nerve (*P ≤ 0.005). Glycogen diminishes with elevated IOP after 4 d but increases at 1 and 2 wk vs. contralateral (C, *P ≤ 0.05). For transgenic mice (Right), elevation decreases glycogen for both KO-Ctrl and KO vs. each respective naïve (−P ≤ 0.02). In KO-Ctrl mice, glycogen decreases after 4 d elevation but increases after 1 wk vs. contralateral nerve (*P ≤ 0.01); for KO, glycogen decreases with both elevations vs. contralateral (*P < 0.001). In KO-Ctrl, elevation increased glycogen compared to contralateral (*P = 0.04), while in KO the opposite occurred (*P < 0.001). (C) Western blots probing pAMPK, AMPK, and GPBB in naïve optic nerves (naïve GPBB Ctrl SEM = 0.149, KO SEM = 0.28; naïve pAMPK/AMPK Ctrl SEM = 0.17, KO SEM = 0.37) and in contralateral (C) vs. 4 d and 1- and 2-wk microbead (M) nerves (Upper). The pAMPK/AMPK ratio (Lower Left) increases in KO-Ctrl nerves (microbead and contralateral) vs. respective naïve (n = 6) for 4 d (n = 5) and 2 wk (n = 5) and in contralateral only at 1-wk (n = 6) microbead elevation (*P ≤ 0.04); ratio in KO nerves (M and C) does not change vs. respective naïve (n = 6, P ≥ 0.20) and is less than ratio in KO-ctrl contralateral for all times (n = 4, n = 7, n = 5; *P ≤ 0.01). Ratio in KO-ctrl contralateral exceeds microbead at 1 wk (*P = 0.05), while microbead exceeds contralateral at 2 wk (*P = 0.02). Fold-change in GPBB (Lower Right) is increased at 4 d in KO-Ctrl contralateral vs. microbead nerves (*P = 0.04) and in KO microbead vs. contralateral and vs. KO-ctrl microbead nerves (*P ≤ 0.03). At 1 wk, GPBB increased vs. respective naïve in KO-Ctrl and KO nerves from both eyes (*P ≤ 0.002) and in KO-Ctrl contralateral vs. microbead and vs. KO contralateral (*P ≤ 0.04); GPBB increased more in KO microbead vs. contralateral (*P = 0.03).
mice, 4 d of unilateral IOP elevation reduced glycogen compared to the unaltered contralateral nerve (−28.41%; P = 0.01); glycogen in each did not differ from their WT counterparts (P = 0.233 and P = 0.466, respectively) (Fig. 2 B, Right). As it did in WT nerves, glycogen diminished following 1 wk of IOP elevation in unaltered contralateral nerve compared to stressed for KO-Ctrl mice (−33.3%; P = 0.012). Following 4 d of IOP elevation, glycogen in KO nerves did not differ from KO-Ctrl (P ≥ 0.08), once again decreasing in the IOP-stressed vs. the contralateral nerve (−30.9%; P < 0.001). However, 1 wk of IOP elevation in KO mice reversed the glycogen pattern observed in KO-Ctrl and WT mice. Although glycogen decreased in both IOP-stressed and contralateral KO nerves at 1 wk compared to 4 d of elevation (−63.4% and −43.1%, respectively; P < 0.001), the unaltered KO contralateral nerve contained significantly more glycogen than the IOP-stressed KO nerve (0.56% ± P = 0.002).

To assess metabolic state in the optic nerve, we measured via Western blots the ratio of pAMPK/AMPK, which increases with metabolic stress (40), and glycogen phosphorylase (GPBB) relative to total protein, which is a marker of glycogen breakdown (35). Only 32% of the mouse optic nerve is occupied by retinal ganglion cell axons, with the remainder representing glial cells (66%) and extracellular space (2%) (41), making the entirety of somatic protein as well as the majority of total protein within these samples glial in origin. In KO-Ctrl mice, nerves from unaltered contralateral eyes as well as nerves stressed by elevated IOP showed increased pAMPK/AMPK compared to naïve KO of Cx43 eradicated this difference (Fig. 2 C, Lower Left). Furthermore, pAMPK/AMPK in KO-ctrl contralateral nerves consistently exceeded that in KO contralateral nerves (P ≤ 0.01), demonstrating the necessity of astrocyte Cx43 for conveying metabolic stress due to elevated IOP. GPBB only differed from naïve at 1-wk IOP elevation, when it significantly increased for both nerve from KO-Ctrl and KO mice alike (Fig. 2 C, Lower Right). This is expected, since glycogen diminished significantly in both nerves for each strain compared to naïve at 1 wk (Fig. 2 B, Right). Importantly, for KO-ctrl nerves, changes in GPBB relative to naïve were opposite for IOP-stressed vs. contralateral: GPBB in IOP-stressed nerves diminished while increasing in the contralateral. This difference was significant at 4-d and 1-wk IOP elevation (P < 0.04). Cx43 KO reversed this trend significantly, so that GPBB within IOP-stressed nerve was diminished that of contralateral (P ≤ 0.04), again consistent with less glycogen in stressed KO nerves (Fig. 2 B, Right). Thus, as with glycogen, astrocyte Cx43 contributes to the activation of metabolic pathways in both optic nerves following IOP elevation in one eye.

Not only does the presence of astrocyte Cx43 allow unilateral IOP elevation to activate AMPK across both visual streams, we also find it influences the distance Cx43-mediated astrocyte networks extend in both eyes. For this experiment, we utilized GFAP-GFP mice, with which we could distinguish astrocyte cell bodies in the retina among both their processes and Müller cell processes. We used these mice for patch-clamp recording and intracellular filling. Compared to naïve retinas (Fig. 3 A), the diffusion of gap junction-permeable dye (neurobiotin-350) injected into an astrocyte following 1 wk of unilateral elevation increased in the contralateral retina as it did in the IOP-stressed retina (Fig. 3 B and C). Diffusion within the retina was abolished by the gap junction blocker carbenoxolone (Fig. 3 D). We confirmed the identity of single astrocytes marked for neurobiotin injection by their lack of action potentials in response to depolarizing voltage steps (Fig. 3 E). Following injection, the most distant biotin-marked astrocyte (defined by a streptavidin signal greater than 20% above background that colocalized with GFAP in a soma) was significantly farther compared to naïve, not only for retina with IOP elevation, but also for the contralateral retina (Fig. 3 F) (P < 0.001). Thus, unilateral IOP stress expands the astrocyte-coupled network in both retinas, allowing astrocytes to communicate quickly over greater distances.

To directly track the movement of glucose and its metabolites through the astrocyte network between the two eyes after unilateral injury, we elevated IOP in one eye and injected the radioactive glucose analog 18F-FDG (fluorodeoxyglucose) intravitreally into the contralateral eye 1 wk later (Fig. 4 A; IOP in SI Appendix, Fig. S1). Mice without microbead elevation served as control. One hour following 18F-FDG injection, PET and CT scans showed how metabolites originating in the unaltered projection redistributed. In naïve WT mice without IOP elevation (Fig. 4 B), transfer to the contralateral eye and nerve was minimal, only 1.24 ± 0.11% of the injected dose, typical of dissemination in neuronal tissues (42). Remarkably, a 1-wk IOP elevation in WT mice (Fig. 4 C) increased contralateral transfer from the unaltered eye by 324%; 5.26 ± 1.06% of the injected dose (P < 0.001). Thus, IOP elevation has an extensive impact on metabolic state well beyond the directly stressed tissue, causing contralateral tissues to donate bioenergetic resources.

In KO-Ctrl mice, the transfer of 18F-FDG to the IOP-stressed eye and nerve was not significantly different from WT (4.93 ± 1.01%; P = 0.42) (Fig. 4 D). In KO mice, however, transfer of 18F-FDG from the unaltered eye to the IOP-stressed eye and nerve was reduced such that KO-Ctrl transferred 27% more glucose than KO mice (P < 0.001) and did not differ from transfer in naïve KO mice without IOP elevation (1.33 ± 0.17 vs. 1.56 ± 0.13%; P = 0.16) (Fig. 4 E and F). Levels of 18F-FDG contained in the bladder did not differ between any of the groups, indicating that in each a similar amount of metabolite reached the circulatory system (P ≥ 0.18). Additionally, transection of optic nerve that maintained their eye’s vascular input prevented transfer of metabolites to the IOP-stressed contralateral projection (Fig. 4 F). Metabolite transfer for WT sham was significantly elevated above transection (4.89 ± 1.46 vs. 0.78 ± 0.25; P = 0.025), resulting in contralateral 18F-FDG values similar to that of WT microbead alone (Fig. 4 J and K). Both transection and sham conditions resulted in similar bladder 18F-FDG content (P = 0.48) (Fig. 4 L). Together, these results demonstrate that 18F-FDG is redistributed from healthy to stressed tissue through an astrocyte Cx43-mediated network within the optic projection itself.

Optic Nerve Contralateral to Elevated IOP Exhibits Functional Deficits following Metabolic Challenge. Next, we investigated how metabolic transfer between optic projections influences physiological response to induced stress by recording the compound action potential (CAP) from WT and KO optic nerves ex vivo during glucose deprivation and recovery (Fig. 5). We measured the positive voltage response following brief depolarizing voltage pulses, typically in the first 40 to 80 ms following the pulse (Fig. 5 A); the CAP itself was calculated as its integral (4, 8, 37, 43). After an initial period (15 min) of recording baseline CAP to determine each nerve’s maximum, we depleted glucose via subduction with l-glucose, an enantiomer of glucose that is not metabolically utilized by living organisms because it cannot be phosphorylated by hexokinase, the first enzyme in the glycolysis pathway (44). Following glucose depletion for 1 h, we then measured the recovery in v-glucose for the following 30 min (Fig. 5 A, Inset). The CAP for naïve KO optic nerve did not differ from that of naïve WT at any point, including an initial increase upon l-glucose substitution we attribute to glycogen mobilization (P ≥ 0.09) (Fig. 5 B). Following 1 wk of unilaterally elevated IOP (SI Appendix, Fig. S1), the WT CAP did not change compared to naïve (P = 0.10) (Fig. 5 C vs. B). However, the CAP for the unaltered contralateral nerve significantly diminished at 9 of 12 sampled times during glucose depletion (P ≤ 0.04), indicating suppression of the initial rise in response, and recovered only to 45% of
baseline compared to 65% for the IOP-stressed nerve (P ≤ 0.04) (Fig. 5C). Thus, the contralateral, putatively unstressed nerve exhibits a greater deficit in its physiological response to subsequent metabolic depletion than tissue directly stressed by IOP.

This pattern reversed in KO nerves (Fig. 5D). Elevated IOP eradicated the initial rise upon glucose depletion compared to naïve KO (P ≤ 0.04) and IOP-stressed WT nerves (P ≤ 0.05), as in the unstressed contralateral nerve in WT mice. With return to D-glucose, recovery of the CAP for IOP-stressed KO nerves did not differ from either naïve KO or IOP-stressed WT (P ≥ 0.220, P ≥ 0.103, respectively). In contrast to WT, the CAP from the unstressed contralateral nerve in KO mice exceeded that of the IOP-stressed nerve in the initial response to glucose depletion and in the final recovery (both P ≤ 0.04). Remarkably, in KO mice the CAP for the unstressed contralateral nerve did not differ at any time from the naïve response (P ≥ 0.12). As well, the CAP from the unstressed contralateral KO nerve exceeded that of the corresponding WT nerve at each point following glucose depletion (P ≤ 0.03). Thus, metabolic redistribution through astrocyte Cx43 renders the donating tissue more susceptible to subsequent metabolic depletion but preserves physiological responses in the IOP-stressed tissue.

Conditional KO of Astrocyte Cx43 Accelerates Axonopathy and Functional Loss. Similar to other neurodegenerative diseases (45, 46), glaucoma involves deficits in axonal anterograde transport to central brain targets. This represents an early and sensitive degenerative outcome associated with elevated IOP (16). The ratio of hyperphosphorylated vs. phosphorylated intermediate filaments impedes active transport and is another early indicator of axonal pathology in human neurodegeneration (47). For KO-Ctrl mice, Western blots using antibodies against each epitope of intermediate filament (SMI34 and -31, respectively) demonstrate that this ratio increased after 4 d and 1 wk of elevation in the IOP-stressed nerve compared to naïve (Fig. 6A). In contrast, for KO mice at 1 wk the ratio of hyper- to phosphorylated neurofilaments was significantly greater in the microbead nerve compared to the unstressed contralateral nerve (P = 0.003). After 2 wk of unilateral IOP elevation, for KO-Ctrl mice the ratio in the unstressed contralateral nerve exceeded not only naïve, but also the microbead nerve. In contrast, the ratio in the unstressed KO nerve was significantly less than naïve and the IOP-stressed nerve. Therefore, astrocyte Cx43 contributes to bilateral transfer of IOP-related stress even at the level of neurofilament modifications in neuronal axons comprising the optic nerve.
From the preceding data, the contralateral optic nerve appears vulnerable to neurodegenerative stress after donating its energetic reserves. To test directly whether the contralateral nerve would progress through neurodegeneration more quickly if its fellow eye had already experienced a stressor, we tested axonal transport in vivo using three IOP-stressed cohorts of KO and KO-Ctrl mice (Fig. 6B, Top, and SI Appendix, Fig. S2). A cohort with unilateral elevation was killed after 1 wk, while a second cohort was killed following 2 wk. In a third, staggered cohort, following 1 wk of IOP elevation in one eye, we elevated IOP in the contralateral eye and then killed after a further week. We then compared tissues that had experienced either 1 or 2 wk of elevation in this staggered, bilaterally stressed cohort to their corresponding unilateral controls. Both 1- and 2-wk elevations were significant compared to preexperiment IOP and to IOP in naïve eyes ($P < 0.001$), but did not differ in magnitude from one another ($P ≥ 0.24$). Forty-eight hours prior to killing, all eyes received an injection of cholera toxin B (CTB) to assess the retinotopic representation of active axonal anterograde transport to the SC, the primary ganglion cell neuron target in rodents (32, 33).

The unilaterally stressed groups revealed intriguing differences in neuronal degeneration in the presence and absence of astrocyte Cx43. One-week unilateral IOP elevation did not affect the level of active transport compared to the unstressed contralateral SC in KO-Ctrl mice ($P = 0.35$) (Fig. 6B and C, Left). In contrast, the same 1-wk elevation in KO mice reduced transport by 40% ($59.6 ± 6.2%$), significantly less than active transport both in the contralateral unstressed SC ($91.6 ± 6.2%$ intact; $P = 0.002$) and in the corresponding KO-Ctrl SC ($P = 0.03$). For the 2-wk unilateral cohort (Fig. 6B and C, Center), elevated IOP significantly reduced active transport in KO-Ctrl mice compared to the unstressed contralateral SC ($68.8 ± 4.4%$ vs. $92.8 ± 2.1%$; $P = 0.02$). Even so, transport degraded more dramatically in KO SC compared to its contralateral unstressed SC ($48.2 ± 4.9%$ vs. $92.8 ± 1.9%$; $P < 0.001$) and compared to the IOP-stressed SC in KO-Ctrl ($P = 0.01$). These differences in transport following unilateral IOP track well with changes in SMI34/31 ratio in the corresponding optic nerves (Fig. 6A) and demonstrate the impact of astrocyte networks on maintaining axonal function through neurodegenerative stress.

Astrocyte Cx43-mediated gap junctions proved even more important in our staggered, bilaterally stressed cohort. In the
staggered cohort of animals that experienced a second IOP elevation affecting the contralateral, donating tissue, we found that astrocyte Cx43 allows the nervous system to distribute and equalize neurodegenerative outcomes between otherwise disparate regions. For this staggered KO-Ctrl cohort (Fig. 6 B and C, Right), degradation of transport to the SC subjected to 2-wk IOP elevation (M in Fig. 6 B, Top) did not differ significantly from the contralateral SC subjected to microbead elevation 1 wk later (MC in Fig. 6 B, Top) (P = 0.41), despite that eye experiencing only half the IOP elevation. However, the level of intact transport in both SCs significantly diminished compared to their unilateral counterparts, despite similar elevations in IOP (57.5 ± 2.9 and 51.4 ± 2.1, respectively; P ≤ 0.008). Stressing both eyes caused each to be at a significant disadvantage when combating this neurodegenerative stressor, and particularly affected the tissue that had already donated resources to the originally stressed region.

This effect was absent for KO mice in the staggered cohort, where transport degradation in SC from neither the initial 2-wk elevation (M) nor the staggered contralateral 1 wk (MC) differed from deficits in the corresponding unilateral IOP elevation SC (P ≥ 0.11). In addition, unlike KO-Ctrl SC, transport deficits for 2-wk elevation were significantly worse than the contralateral 1 wk (43.7 ± 5.2% vs. 61.3 ± 3.5% intact; P = 0.04). These findings suggest that resource redistribution through astrocyte Cx43 renders neurons within the donating tissue more susceptible to neurodegenerative damage during subsequent stressors. Additionally, by eliminating astrocyte Cx43, functional outcomes between the two projections were no longer equilibrated.

Finally, to determine whether Cx43 KO influences visual function after IOP elevation, we examined contrast spatial acuity (Fig. 7). Acuity in KO-Ctrl eyes that received the initial microbead-induced IOP elevation (M) diminished 10% by day 6 compared to baseline (0.45 ± 0.01 vs. 0.51 ± 0.01 cycles/degree; P < 0.001), with a total decline of 20% by 2 wk (0.41 ± 0.01 cycles/degree; P < 0.001). Importantly, acuity in the KO-Ctrl contralateral eye that received subsequent IOP elevation for 1 wk (MC) declined by a similar amount from its baseline (0.46 ± 0.01 cycles/degree; P < 0.001). Thus, although isolating each visual stream from the broader astrocyte network transiently protected the contralateral nerve from subsequent damage, both the initially stressed and contralateral KO visual streams eventually demonstrated visual acuity either equivalent to or even worse than both control conditions.

Discussion

Astrocytes are the most numerous cell type in the central nervous system, forming interconnected networks mediated by gap junctions primarily composed of Cx43. Astrocyte gap junctions have been implicated in a variety of neurodegenerative conditions (28, 30), but the beneficial aspects of gap junction alterations through disease progression are less well understood. As neurons signal and utilize their metabolites under healthy conditions, local astrocyte gap junctions allow mobilization of stored energetic reserves (31, 48). To determine the mechanism by which these energetic reserves are mobilized during disease, we utilized conditional mutagenesis of floxed Cx43 under a GFAP promoter (GFAP-Cre-ER<sup>+</sup> × Cx43<sup>fl/fl</sup>). We found astrocyte gap junction-mediated networks influence metabolite distribution more globally, between distant regions of the optic projection during stress induced by elevated IOP.

Sensitivity to IOP is a defining feature of axonal degeneration in the optic nerve during glaucoma (17), which also involves enhanced neuronal excitation early in progression (49). Our fundamental finding is that in response to IOP elevation in one eye, astrocyte gap junctions redistribute metabolic resources from the other optic projection, across the large distance separating the two eyes (Fig. 4). While beneficial to the stressed optic projection, redistribution renders contralateral donating tissue
Fig. 6. Cx43 KO accelerates axonopathy in the optic projection. (A, Upper) Western blots probing hyperphosphorylated (SMI34) and phosphorylated (SMI31) intermediate filaments in KO and KO-Ctrl optic nerves following unilateral IOP elevation. (Lower) For KO-Ctrl, the ratio of SMI34/31 increased significantly over genotype naïve (#; Ctrl SEM ± 0.09, KO SEM ± 0.22) in both microbead and contralateral nerves after 4 d of unilateral elevation (n = 5), in microbead at 1 wk (n = 6), and in the contralateral nerve at 2 wk (n = 5). At 2 wk, KO-Ctrl microbead was less than its contralateral and the KO microbead (*). For KO microbead, SMI34/31 exceeded its genotype naïve at 4 d (n = 5), 1 and 2 wk (n = 8, 5). In contrast to KO-Ctrl, at 2 wk KO microbead exceeded naïve while contralateral was less (#). *P ≤ 0.02; **P ≤ 0.006. (B, Top) Contralateral (C) eye is naïve for unilateral 1- and 2-wk IOP elevations via microbead (M) injection, in staggered cohort (Right), contralateral received microbead injection (MC) 1 wk later. Both eyes underwent OMR assessment. All eyes received CTB prior to killing. IOPs in SI Appendix, Fig. S2. (Bottom) Coronal section (Middle) through SC demonstrating intact (solid white line) vs. degraded (dashed) anterograde transport. Reconstructed SC retinotopic maps range from 100% (red) to 0% (blue) transport. M: medial; R: rostral. (Scale bars, 500 μm.) (C) Intact transport (percent map with ≥70% CTB) for KO-Ctrl vs. KO in 1-wk (Left), 2-wk (Center), and staggered (Right) cohorts. One-week elevation in KO mice reduced transport vs. the contralateral SC (*P = 0.002) and vs. the corresponding KO-Ctrl SC (*P = 0.03). Two weeks reduced transport in KO-Ctrl (*P = 0.020) and KO (*P < 0.001) vs. unstressed contralateral, while KO deficits exceeded KO-Ctrl (*P = 0.01). In the staggered cohort, transport reduced after 1-wk (MC) and 2-wk (M) elevations in KO-Ctrl vs. unilateral 1- and 2-wk (*P ≤ 0.008) and in 2- (M) vs. 1-wk (MC) KO SC (*P = 0.04).
pattern reverses; after 1 wk of elevation, glycogen in the ipsilateral to IOP elevation. In GFAP-driven Cx43 KO mice, this
in size compared to naïve (Fig. 3).

more susceptible to further stressors (Figs. 3, 6, and 7). Bilateral
glial activation after unilateral optic nerve stress has been ob-
erved previously (50), in microglia (51) as well as astrocytes (52).
One study (50) confirmed bilateral axonal loss after uni-
ilateral optic nerve crush, finding similarities in the contralateral
nerve to an incomplete crush and postulating an adaptive func-
tion wherein the SC equilibrated inputs from both eyes. We too
found evidence that when the donating tissue is exposed to el-
evated IOP after it has donated its energetic reserves, functional
outcomes are equilibrated between the two optic projections.
Equilibration no longer occurred following astrocyte Cx43 KO,
implying that astrocyte connectivity is necessary for this process.
There is evidence in human glaucoma progression as well for
extensive cross-talk between the two optic projections that pro-
duces a complete bilateral visual field by fusing the fragmented,
yet complementary representations from the two eyes (53).

While Cx43 may be retained in vascular endothelial cells, in
retinal Müller glia, or in astrocytes due to variability in Cre-ER\textsuperscript{T2}
induction, our transgenic experiment yielded significant reduc-
tion of astrocyte Cx43 in the retina and optic projection (Fig. 1),
which otherwise is elevated in glaucoma (30). Optic nerve stress
from elevated IOP in glaucoma involves diminished ATP and
other metabolic challenges that contribute to early axon dys-
fuction, including degradation of anterograde transport to the
brain (15–18). We found IOP elevation for only 1 wk influences
protein expression in the unstressed, contralateral optic nerve,
increasing stress-response proteins, glycogen catabolism, and
hyperphosphorylation of neurofilaments (Figs. 2 and 6). This
induced contralateral response travels as far as the opposite eye,
where the coupled retinal astrocyte network more than doubles
in size compared to naïve (Fig. 3).

In WT and KO-Ctrl tissues, we see that glycogen stores in the
contralateral optic nerve diminish more than their counterparts
ipsilateral to IOP elevation. In GFAP-driven Cx43 KO mice, this
pattern reverses; after 1 wk of elevation, glycogen in the
ipsilateral nerve is significantly diminished compared to that in
the contralateral nerve (Fig. 2B). This reversal was significant,
but not complete; the contralateral tissue still showed a signifi-
cant loss of glycogen when compared to naïve tissues. This may
be from the nature of a tamoxifen-induced KO, which is unlikely
to have induced excision in every GFAP-expressing astrocyte.
Additionally, GFAP expression and transcription varies greatly
across astrocytes (54), so by its nature our KO would not reach
every astrocyte in the brain. Incomplete reversal may also be due
to resource donation through other connexins astrocytes may
express (29). Each of these factors could contribute to the
amount of resource donation and glycogen loss we still detect
in the contralateral tissue. That these findings were still of such
magnitude speaks to the remarkable capacity of astrocyte Cx43-
mediated networks.

Intriguingly, the pAMPK/AMPK ratio in our GFAP-driven
Cx43 KO mice did not increase (Fig. 2C), even in the nerve
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Fig. 7. Cx43 KO causes rapid deterioration of visual function. Spatial acuity
(cycles/degree) determined from OMR for KO-Ctrl (n = 9) and KO (n = 10)
mice in staggered cohort following initial microbead injection (M, arrow)
and subsequent injection in contralateral eye (MC, arrow); IOP in SI Ap-
pendix, Fig. 52. Acuity did not differ for the first four measurements in any
group (P ≥ 0.11). Acuity in eye with first elevation (M) for KO-Ctrl and KO
diminished below baseline by day 6 (P < 0.001); decline in KO acuity was
worse than KO-Ctrl (P < 0.001). For KO-Ctrl, acuity in the contralateral eye
(MC) diminished below baseline by day 9 and continued to decline until the
final measurement (P < 0.001), where it reached acuity in the M eye (P = 0.33).
For MC KO eyes, acuity remained above M eye throughout (P < 0.001)
and did not decline from baseline until the final measurement (P < 0.001).

Metabolite redistribution (as measured by 18F-FDG PET imaging)
from unstressed to stressed optic projections after elevated
IOP requires Cx43-mediated astrocyte coupling, since condi-
tional KO inhibits redistribution (Fig. 4G). This coupling
likely allows transfer via the optic chiasm, where the predomi-
nantly contralateral projections from the two eyes cross (41).
When the experiment is repeated but the contralateral optic
nerve is transected, metabolites no longer transfer between the
two projections, but still circulate through the bloodstream to the
same extent as the sham condition (Fig. 4 K and L). Similar
metabolic transfer occurs in corpus callosum, with astrocytes and
oligodendrocytes coupling in response to glucose deprivation
to maintain action potential generation (56, 57).

Both the speed at which metabolites migrate and the relative
deficit in glycogen within the unstressed compared to stressed
nerve imply an active process. A relative metabolic “sink” in the
stressed tissue could explain the eventual equilibrium between
the two nerves we see after 4 but not 1 or 2 wk of elevation.
Perhaps through a heterotypic connexin configuration (28, 58)
astrocytes can dynamically regulate the flow of ions and mole-
cules through their network. Another possibility involves the
charge of the metabolites specifically traced here, which all de-
rive from glucose, the brain’s main source of energy (3). Glucose
and the metabolites derived from it are polar molecules (59),
subject to interaction with other charged entities, such as calcium
ions that could influence their flow.

Our results show that nerves contralateral to short-term (1 wk)
unilateral IOP elevation exposed to a further metabolic stressor
ex vivo exhibit much weaker compound action potential than
their directly stressed counterparts (Fig. 5). In the KO this pat-
tern reverses, with the directly stressed nerve demonstrating a
weaker signal than its contralateral counterpart. This finding
suggests that astrocyte-mediated transfer of metabolites between
projections depletes the donor tissue and renders it susceptible
to further metabolic stressors. Yet, astrocyte Cx43 KO reduced
physiological measures of visual function in the directly stressed
tissue extensively, including both active anterograde transport to
the SC (Fig. 6) and visual acuity (Fig. 7), indicating the systemic
importance of Cx43 gap junction-mediated mobilization of
metabolic resources. Even the donating KO-Ctrl tissue, which
indeed was more susceptible to neurodegenerative stress initially

\[ P < 0.001 \]

\[ P < 0.001 \]

\[ P < 0.001 \]
following IOP elevation, still demonstrated more intact visual function than its KO counterpart after 6 d of elevation (Fig. 7). These results demonstrate that metabolic components of the Cx43 astrocyte network are critical for maintaining function bilaterally early in neurodegenerative disease (Fig. 8). By comparison, pharmacological blockade of neuronal gap junctions generally or genetic knockdown of connexin 36 (Cx36), which links retinal neurons of specific types to one another, significantly slowed retinal and nerve degeneration in the same model of glaucoma (60). Thus, retinal and nerve gap junctions serve multiple purposes, depending on whether coupling is glial (Cx43) or neuronal (Cx36).

Astrocyte Cx43 serves a number of important purposes in addition to metabolic redistribution worthy of consideration. Unpaired connexins have the capacity to act as hemichannels, opening to the extracellular space rather than to an individual cell (61). These hemichannels release gliotransmitters (such as glutamate) (62), facilitate purinergic signaling (63), and release molecules such as nicotinamide adenine dinucleotide and α-serine into the extracellular milieu (64). Paired gap junctions formed by Cx43 can transmit other molecules and ions as well, provided the molecules are generally below 1 to 1.2 kDa (28), including the well-documented example of calcium signaling (65). Each of these functions would also be disturbed in our KO and could have contributed to our results.

Evolving evidence suggests that astrocyte reactivity in response to disease-relevant stress, although most often considered pathogenic, also includes protective outcomes (66). Our results demonstrate that local neurodegenerative stress induces a much broader astrocyte response capable of mediating metabolic redistribution across bilateral regions of the central nervous system. That the donating, contralateral optic nerve itself becomes functionally more intact visual function than its KO counterpart after 6 d of elevation (Fig. 7). These results demonstrate that metabolic components of the Cx43 astrocyte network are critical for maintaining function bilaterally early in neurodegenerative disease (Fig. 8). By comparison, pharmacological blockade of neuronal gap junctions generally or genetic knockdown of connexin 36 (Cx36), which links retinal neurons of specific types to one another, significantly slowed retinal and nerve degeneration in the same model of glaucoma (60). Thus, retinal and nerve gap junctions serve multiple purposes, depending on whether coupling is glial (Cx43) or neuronal (Cx36).

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We utilized a Siemens Inveon PET scanner (Siemens Preclinical), a NanoSPECT/CT ( Bioscan), and the radioactive glucose analog 18F-FDG. Mice were briefly anesthetized with 2.5% isoflurane and then increased 25% to ensure stimulus strength was supramaximal. All animals were then returned to their cages and fed ad libitum for 60 min. An analog 18F-FDG. Ani-

mals were then returned to their cages and fed ad libitum for 60 min. An-
imals were again anesthetized with 2% isoflurane and imaged for 40 min in the Inveon microPET in static mode. PET and CT images were normalized to the injected radioactive dose of 18F-FDG. Regions-of-interest (ROIs) were drawn as 7 x 14 x 7-mm cubes around the entire optical projection contra-
larateral to 18F-FDG injection, and mean radiotracer concentrations within the top 90% of voxels within ROIs were measured in units of percent injected dose per unit volume (%ID/g).

OMR Testing. Visual acuity was measured as described previously (49 (OptoMotry; CerebralMechanics Inc.).

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