Axon degeneration induces glial responses through Draper-TRAF4-JNK signalling

Tsai-Yi Lu1,t, Jennifer M. MacDonald1, Lukas J. Neukomm1, Amy E. Sheehan1, Rachel Bradshaw1, Mary A. Logan1,t & Marc R. Freeman1,t

Draper/Ced-1/MEGF-10 is an engulfment receptor that promotes clearance of cellular debris in C. elegans, Drosophila and mammals. Draper signals through an evolutionarily conserved Src family kinase cascade to drive cytoskeletal rearrangements and target engulfment through Rac1. Glia also alter gene expression patterns in response to axonal injury but pathways mediating these responses are poorly defined. We show Draper is cell autonomously required for glial activation of transcriptional reporters after axonal injury. We identify TNF receptor associated factor 4 (TRAF4) as a novel Draper binding partner that is required for reporter activation and phagocytosis of axonal debris. TRAF4 and misshapen (MSN) act downstream of Draper to activate c-Jun N-terminal kinase (JNK) signalling in glia, resulting in changes in transcriptional reporters that are dependent on Drosophila AP-1 (dAP-1) and STAT92E. Our data argue injury signals received by Draper at the membrane are important regulators of downstream transcriptional responses in reactive glia.
Glial cells are exquisitely sensitive to perturbations of nervous system physiology\(^1\)–\(^7\). In response to neural injury they coordinate a sophisticated cascade of reactive events, including engagement of phagocytic programmes to clear cellular debris, and secretion growth factors to recruit additional immune cells or promote neuronal survival and repair\(^8\). Reactive glial responses are driven in important ways by changes in glial gene expression following trauma. Strong upregulation of expression of GFAP and S100β are among the earliest detectable changes in mammalian neural tissues after brain injury, and similar upregulation occurs in chronic neurodegenerative disease\(^9,\)\(^10\). In peripheral nerves early events include the activation of JNK signalling, which is required for efficient Schwann cell de-differentiation and initiation of regenerative responses\(^1\)\(^1\). Initiating events in glial reactive responses are presumably driven by 'injury signals' released by damaged cells that signal to transmembrane receptors on glia, but whether and how cellular debris activates glial transcriptional responses remains mechanistically unclear.

**Drosophila** glia rapidly respond at the transcriptional level to axotomy, which argues reactive glial responses are an ancient feature of the metazoan nervous system\(^1\)\(^2\). Transcriptional reporters for the sole *Drosophila* STAT molecule, STAT92E, are induced in glial cells within hours after local axons are severed\(^6\). Axotomy also activates the glial JNK signalling cascade and ultimately transcriptional reporters for the *Drosophila* AP-1 complex (dAP-1)\(^7\). Activation of STAT92E-dependent transcriptional responses has been shown to require the engulfment receptor Draper\(^6\), suggesting STAT92E lies genetically downstream of Draper, although how axon degeneration is mechanistically linked to STAT92E- or dAP-1-mediated transcriptional changes is not known.

**Results**

**Axon degeneration activates dAP-1 reporters in glia.** To understand how axonal injury leads to glial transcriptional changes we sought to define the mechanisms regulating activation of the dAP-1 complex in glia after axotomy. We used the dAP-1 transcriptional reporters containing multiple tetradecanoylphorbol acetate response elements (TRE, that is the dAP-1 binding sequence) upstream of either eGFP (TRE-eGFP) or RFP (TRE-RFP)\(^1\)\(^3\). Axon injury was induced by surgical removal of adult *Drosophila* third antennal segments, which severs ~90% of olfactory receptor neurons (ORNs) projecting into the *Drosophila* antennal lobe and axon fragmentation begins within ~12 h by activation of Wallerian degeneration\(^2\). Prior to axonal injury TRE-RFP was not detectable at significant levels in control animals in any part of the adult brain. However, one day after axonal injury (1 dpant, 1 day post antennal ablation), we observed strong activation of TRE-RFP throughout the brain (Supplementary Fig. 1), as previously observed for TRE-eGFP (ref. \(^7\)), along with glial hypertrophy and increased Draper levels in glia adjacent to the injury site. To determine whether activation of TRE-RFP represented a glial response to axon degeneration, we blocked ORN axon degeneration using *highwire* (hiw) mutants\(^1\)\(^4\). Glial cells in *hiw*\(^{1\text{N}}\) mutant animals exhibited no morphological changes after axonal injury; TRE-RFP activation was not observed in any part of the brain, and Draper levels adjacent to the injury site remained similar to the uninjured controls (Fig. 1a). Hiw is not expressed at detectable levels in glia but is prominent in neurons (Fig. 1b), suggesting the lack of glial activation after axotomy in *hiw* mutants resulted from blockade of axon degeneration, rather than a cell autonomous role for *hiw* in glial activation. Consistent with this notion, pan-neuronal expression of *Highwire* in *hiw*\(^{1\text{N}}\) mutant background rescued glial response to axon degeneration (Supplementary Fig. 2b) while glial expression of a *hiw* cDNA in the *hiw*\(^{1\text{N}}\) mutant background did not lead to activation of the TRE-RFP reporter (Supplementary Fig. 2a). Activation of the Wallerian degeneration signalling pathway is therefore essential for brain-wide induction of dAP-1 transcriptional reporters in glia after nerve injury. Furthermore, these data indicate glial cells respond specifically to signals generated by degenerating axons, rather than other alterations in axons caused by axotomy (for example loss of neuronal activity).

**Glia activate dAP-1 reporters after injury through Draper.** The engulfment receptor Draper activates glial phagocytic function during Wallerian degeneration\(^2\). We found *draper*\(^{45}\) null mutant animals also exhibited a striking loss of TRE-eGFP induction 1 day after axon injury compared to controls (Fig. 1c,d), indicating that Draper is also required for dAP-1 reporter activation in response to axonal degeneration. ORN axon terminals only project to the antennal lobe. The lack of TRE-eGFP activation in *draper* mutants observed throughout the adult brain implied that Draper was required for TRE-eGFP activation even in glia at sites distant from degenerating axons in the antennal lobe. To characterize the autonomy of Draper function in activation of dAP-1 transcriptional reporters in brain glia, we generated *draper*\(^{45}\) mutants clones using the mosaic analysis with a repressible cell marker (MARCM) approach\(^1\)\(^5\), severed antennal ORN axons, and assayed TRE-RFP activation 1 day after axotomy. Based on morphological analysis of clones we determined that activation of TRE-RFP occurred in ensheathing (Fig. 1e) and cortex glia (Fig. 1f), but not in astrocytes (Supplementary Fig. 3), consistent with a previous study demonstrating Draper is expressed solely in these glial subtypes in the *Drosophila* adult brain\(^1\)\(^6\). While control clones in each subtype of glial cell showed strong activation of TRE-RFP in response to axonal injury, *draper*\(^{45}\) mutant clones did not activate the TRE-RFP reporter (Fig. 1c,f). This was even true when *draper*\(^{45}\) mutant clones were made in cortex glia, which are not in direct contact with severed axons (for example Fig. 1f). This was surprising in light of the fact that Draper/Ced-1 is considered a recognition receptor for 'eat me' cues on the surface of engulfment targets such as axonal debris\(^1\), which we presumed would require cell–cell contact. This observation suggests that Draper may be activated by a soluble ligand released by severed ORNs, or alternatively that Draper-dependent glial-glial signalling may occur in cortex glia upstream of dAP-1 reporter activation, perhaps similar to MEGF10 signalling in starburst amacrine cells during tiling where MEGF10 appears to act genetically as both a receptor and ligand\(^1\)\(^7\).

**TRAF4 signals downstream of Draper after axonal injury.** Based on the potent regulation of TRE-RFP by Draper, we sought to determine how Draper activation might lead to its activation through DAP-1. We screened a large collection of RNAi lines for novel signalling molecules and assayed for roles in glial clearance of degenerating axons. Interestingly, we found that pan-glial expression of UAS-traf4IR (ref. \(^1\)\(^8\) altered strength of glial clearance of axonal debris, compared to controls (Fig. 2a,b, Supplementary Fig. 4c,d). TRAF4 is known to act upstream in activation of the JNK signalling pathway through association with the MAP kinase kinase kinase (MAP4K) *missihapen* (MSN) (ref. \(^1\)\(^9\), which then phosphorylates the downstream MAP3K Slipper and ultimately drives DAP-1 activation of transcriptional targets\(^2\)\(^0\). A role for TRAF4 in glial engulfment was intriguing given that we previously implicated a number of downstream components of the JNK signalling...
pathway in glial engulfment of axonal debris including dAP-1 (ref. 7), but how this pathway might be activated remained unclear.

Pan glial expression of a dominant negative version of MSN \((\text{msn}^{\text{DN}})\) strongly suppressed glial engulfment of axonal debris 5 days after axotomy (Fig. 2c,d). Consistent with a role for TRAF4 and MSN in activation of the JNK signalling pathway we found that glial expression of \(\text{traf}^{\text{IR}}\) (Fig. 2e,f) or \(\text{msn}^{\text{DN}}\) (Fig. 2g,h,k) suppressed activation of the TRE-eGFP reporter. Moreover, we observed that depletion of TRAF4 or expression of MSN\(^{DN}\) suppressed injury-induced increases in Draper that are normally observed 1 day after axonal injury,
and which we previously reported for downstream components of the JNK signalling pathways. We further noted that in **tra**^IR^ animals Draper expression was reduced ~50% even before axon injury (Fig. 2i)). This phenotype is similar to glial stat92eRNAi animals, where Draper levels are reduced in uninjured animals, and engulfment defects can be rescued by overexpression of a *draper* cDNA. In contrast, we found defects in glial responses to axonal degeneration in the **tra**^IR^ background did not result from insufficient Draper levels: re-expression of a functional Draper-1 transcript (Supplementary Fig. 4a,b) in **tra**^IR^ animals failed to rescue axonal debris clearance defects (Supplementary Fig. 4c,d). Together these data support a model whereby TRAF4 signals downstream of Draper to promote target engulfment and glial transcriptional reporter activation mediated by STAT92E and dAP-1.

Based on our observation that TRAF4 and MSN regulated injury-induced changes in Draper levels and transcriptional reporters in glia, we sought to define signalling interactions between TRAF4 and MSN, the JNK signalling pathway, and STAT92E. STAT92E activity after axotomy was assayed using the degradable GFP reporter 10xSTAT92E-dGFP (ref. 22), which is transiently activated in glia 1 day after axotomy. In control uninjured animals, GFP expression was absent in brains, but strongly activated in glia surrounding the antennal lobe 1 day after antennal ORN axotomy (Fig. 2i). Glial expression of **tra**^IR^ suppressed this increase of GFP expression after axon injury (Fig. 2l,m), indicating that TRAF4 is upstream of the injury-induced activation of 10xSTAT92E-dGFP. Similarly, blockade of the JNK signalling pathway with basketRNAi (*Drosophila* JNK), or overexpression of the JNK inhibitor Puckered eliminated the injury-induced upregulation of 10xSTAT92E-dGFP (Fig. 2l,m).

Finally, knockdown of Kayak, a component of the dAP-1 transcriptional complex, also led to a significant decrease in 10xSTAT92E-dGFP activation (Fig. 2l,m). Thus STAT92E is genetically downstream of dAP-1 in the context of 10xSTAT92E-dGFP reporter activation. Taken together, these genetic data argue that glia activate these transcriptional reporters after axonal injury through a TRAF4→JNK pathway→dAP-1→STAT92E signalling cascade that lies downstream of Draper.

**TRAF4 binds the Draper intracellular domain.** The mechanism by which Draper activates JNK signalling and dAP-1-mediated transcriptional reporter activation has not been defined. The only molecule known to physically associate with Draper to promote downstream signalling events is the Src family kinase Shark, which binds an ITAM in the Draper intracellular domain after phosphorylation by Src42a (ref. 4). We found that depletion of Shark from all glia by pan-glial expression of UAS-sharkRNAi (using the repo-Gal4 driver) blocked activation of TRE-eGFP 1 day after axotomy (Supplementary Fig. 5a,b), and similar RNAi-mediated knockdown of racl or src42a also significantly decreased injury-induced activation of TRE-eGFP (Supplementary Fig. 5c,d). We therefore conclude that dAP-1 transcriptional reporters are genetically downstream of Draper/Shark signalling interactions.

Given that TRAF4 is an adaptor protein, we speculated that TRAF4 might interact with Draper and thereby directly couple Draper to activation of JNK kinase signalling. TRAF4/MSN have well-defined roles in the activation of Slipper (ref. 20), and in turn the *Drosophila* JNK signalling pathway. We therefore performed co-immunoprecipitation (co-IP) experiments using *Drosophila* S2 cell lysates with a Myc-tagged TRAF4 (TRAF4-myc) and HA-tagged Draper I (Draper-I-1A). We found that pull down of Draper-I-1A led to the co-precipitation of TRAF4-myc, indicating that TRAF4 can physically associate with the Draper signalling complex (Fig. 3a). To identify the region through which TRAF4 associates with Draper we assayed the ability of epitope-tagged versions of the N-terminal TRAF4, which is enriched with zinc-finger domains (ZF, aa 1-927) or the C-terminal TRAF domain (TD, aa 928-1458) to associate with Draper. We found that the N-terminal TRAF4 domain co-immunoprecipitated with Draper I-1A, while the C-terminal TRAF domain, which is known to bind MSN (ref. 19), was not detected in the same assay (Fig. 3b,c). We therefore conclude that Draper is bound by the N-terminal domain of TRAF.

To identify the region of Draper-I to which TRAF4 bound, we next made serial deletions in the intracellular domain (Fig. 3d), and performed co-IP with full-length TRAF4-myc. We found that aa 826-894 were essential for Draper-I-TRAF4 interactions (Fig. 3e, Δ826-894), while all other regions were dispensable. We note this includes the evolutionarily conserved NPXY motif that is likely bound by the PTB domain binding protein dCED-6 (ref. 23), which is consistent with the observation that dCED-6 is not required for activation of 10xSTAT92E-dGFP (ref. 6) or JNK signalling (Supplementary Fig. 6) in glia after axotomy. Further analysis of this subdomain revealed that aa 826-854 and 855-594 were both able to bind TRAF4 (Fig. 3d,f), suggesting there may be multiple TRAF4 binding sites in the Draper intracellular domain. The NPXY motif is not absolutely required for TRAF4 binding, as TRAF4 associated strongly with the Δ855-894 construct lacking this domain (Fig. 3f). Finally, given that both Src42A and Shark are required for activation of TRE-eGFP in response to injury, we assayed whether TRAF4 and Shark binding were reciprocally interdependent. Phosphorylation of Y949 in ITAM, which is
mediated by Src42A and is bound by Shark (ref. 4), was also dispensable for TRAF4 to associate with Draper (Fig. 3g). Reciprocally, elimination of the TRAF4 interaction domain in Draper I (Δ826-894) did not alter Shark association with Draper (Supplementary Fig. 7). Taken together these data indicate that TRAF4 physically associates with the Draper intracellular domain in the region of the NPXY motif, and this interaction occurs in a Shark-independent fashion. These data argue Draper activates the JNK signalling pathway and dAP-1- and STAT92E-dependent transcriptional reporters in...
response to axon degeneration by direct association with TRAF4 and activation through MSN.

Discussion

Activation of JNK/AP-1 and STAT transcriptional activity during reactive glial responses is an evolutionarily conserved feature of Drosophila and mammalian glia\(^24,25\). However, transmembrane receptors on glia that receive extracellular injury signals and in turn activate AP-1 or STAT-dependent transcriptional changes have not been identified. Our work supports a model whereby the engulfment receptor Draper is activated in response to axonal injury, which in turn promotes JNK signalling through TRAF4 and MSN to alter glial transcription via dAP-1 and STAT92E. Our reliance on well-characterized transcriptional reporters for dAP-1 and STAT92E supports this conclusion, but we note that we have not formally shown transcriptional activation of a downstream target gene, primarily because none have been identified.

Axon degeneration is required to activate glial cells after injury. Blockade of axon degeneration using hiw mutations fully suppresses glial responses to axon injury, including morphological changes and activation of transcriptional reporters, indicating that axonal self-destruction leads to the production of cues that activate glial responses through Draper. Interestingly, based on the activation of glial transcriptional reporters (that is TRE-RFP) even at a distance from the injury site (Fig. 1f), it appears that glia can be activated by degenerating axons even when not in direct contact with severed axons. This observation suggests that axonal ‘injury signals’ that stimulate Draper signalling may be soluble, or, alternatively that glial-glial signalling may propagate through the glial network in a Draper-dependent manner. Our identification of Draper/TRAF4 interactions as a mechanism for JNK activation raises the intriguing possibility that MEGF10 might play a conserved role in mammalian reactive astrocytes to activate AP-1 or STAT signalling, or other contexts. Indeed, MEGF10 and several other engulfment genes are expressed in astrocytes and MEGF10 is required for developmental synapse pruning\(^26\).

We propose a simple model where axon self-destruction leads to injury signals that activate Draper, which in turn drives two primary downstream signalling events (Fig. 3h). In the first, Draper promotes cytoskeletal rearrangements essential for phagocytic activity through Src42a, Shark and Traf4 in direct association with Draper, and downstream GEF complexes such as Crk-II/Mbc/dCed-12 and DRK/DOS/SOS (refs 5,27). These association with Draper, and downstream GEF complexes such as non-phagocytic signalling roles including activation of autophagy in Drosophila salivary glands\(^28\) and the tiling of starburst amacrine cells ref. 17. Exploring the roles for TRAF4, dAP-1 and STAT92E could provide further important molecular insight into Draper/MEGF10 regulation of transcription changes in these physiological contexts.

Methods

Fly strains. Drosophila melanogaster strains used in this study include: OR56a-mCD8::GFP/CyO (ref. 29), repo-Gal4/TM3 (ref. 30), nSyb-Gal4, 5xUAS-mCD8:GFP (ref. 31), UAS-traf4IR (gift from T. Xu, Yale U)\(^3\), UAS-drpr(24127)-HA, TRE-eGFP-16 (ref. 13), TRE-RFP (II)\(^8\), UAS-tub-Gal80ts, FRT2A, FRT82B (III), repo-Gal4, UAS-mCD8::GFP/TMV (ref. 35), UAS-gfp-hiw (ref. 31), UAS-drpr-GFP; repo-Gal4, UAS-msnDN:UAS-mCD8::GFP/TMV (ref. 27), UAS-Rac1\(^\Delta\)N(TF) (ref. 36) and UAS-shark\(^\Delta\)Namb (ref. 4).

Figure 2 | TRAF4 and MSN are upstream of JNK- and STAT92E-mediated responses to axotomy. (a) RNAi-mediated depletion of TRAF4 in glia (repo > traf4\(^\Delta\)R) OR85e-mCD8::GFP/UAS-traf4IR; repo-Gal4/+ resulted in a delay in axonal debris clearance 5 days after maxillary malp ablation (5dpam) compared to controls (OR85e-mCD8::GFP; +/ +; repo-Gal4/+). OR85e\(^a\) maxillary nerves were labelled with mCD8::GFP. Scale bar for all = 10 µm. (b) Quantification from (a). n = 10. Unpaired t-test, 2-tailed (for all). Expression of dominant-negative MSN (msn\(^\Delta\)N) suppressed glial clearance of axonal debris. Control: OR85e-mCD8::GFP, tub-Gal80\(^+/+\); repo-Gal4/+ +. Repo > msn\(^\Delta\)N; OR85e-mCD8::GFP, tub-Gal80\(^+/+\); UAS-msn\(^\Delta\)N, repo-Gal4/+ +. Representative images were shown (control: n = 10; repo > msn\(^\Delta\)N; n = 9). (d) Quantification from (c). (e) TRE reporter activity was suppressed in TRAF4 RNAi animals (repo > traf4\(^\Delta\)R) 1 day after antennal ablation. Control: TRE-eGFP-16+/ +; repo-Gal4/+ +. Repo > traf4\(^\Delta\)R; TRE-eGFP-16/UAS-traf4IR; repo-Gal4/+ +. (f) Quantification of the eGFP immunoreactivity in (e). n = 25 each. (g) TRE reporter activity was inhibited 1 day after antennal ablation by glial expression of Msn\(^\Delta\)N. Control: TRE-eGFP-16, tub-Gal80\(^+/+\); repo-Gal4/+ +. Repo > msn\(^\Delta\)N; TRE-eGFP-16, tub-Gal80\(^+/+\); UAS-msn\(^\Delta\)N; repo-Gal4/+ +. (h) Quantification from (g). n = 15 each. (i) Draper expression was reduced on western blots when TRAF4 was knocked down in glia from adult dissected central brain. α-tubulin was used as loading control. Representative images were shown from three repeats. Control: repo-Gal4/+ +. Repo > traf4\(^\Delta\)R; UAS-traf4IR/+; repo-Gal4/+ +. Q) Quantification of (i). n = 3. Unpaired t-test, 2-tailed. (k) Higher magnification images of dashed squares in (g) showed the lack of Draper increase 1 day after antennal ablation in msn\(^\Delta\)N animals. (l) STAT92E-mediated dGFP (degradable GFP) expression was induced in glia 1 day after antennal ablation in control animals (10xSTAT92E-2dGFP; +/ +; repo-Gal4/+ +), but blocked in animals where JNK signalling cascade was interrupted by glial expression of traf4\(^\Delta\)R (10xSTAT92E-2dGFP/UAS-traf4IR; repo-Gal4/+ +), basket\(^\Delta\)N (10xSTAT92E-2dGFP/UAS-basket\(^\Delta\)N\(^\Delta\)3473; repo-Gal4/+ +) and puckered\(^\Delta\)WT (10xSTAT92E-2dGFP; +/ +; repo-Gal4/UAS-puckered), or kayak\(^\Delta\)N (repo > kayak\(^\Delta\)N; repo-Gal4/UAS-kayak\(^\Delta\)N\(^\Delta\)620)), or kayak\(^\Delta\)N (repo > kayak\(^\Delta\)N; repo-Gal4/UAS-kayak\(^\Delta\)N\(^\Delta\)620)). (m) Quantification of the increase of dGFP immunoreactivity in (l), Repo > traf4\(^\Delta\)R; n = 15 each. Repo > basket\(^\Delta\)N, repo > puckered\(^\Delta\)WT, and repo > kayak\(^\Delta\)N; n = 25 each. Unpaired t-test for each group, 2-tailed.
Figure 3 | TRAF4 physically associates with Draper independently of Shark. (a) Western blots of co-immunoprecipitations from Drosophila S2 cells showed TRAF4-myc associated with Draper I-HA. All the blots shown are representative images after three repeats. (b) Schematic drawing of Drosophila TRAF4. ZF: zinc-finger enriched domain. TD: TRAF domain. (c) Draper I-HA interacted with the ZF region of TRAF4 but not the TRAF domain. (d) Schematic drawing of deletions in the Draper I intracellular for different constructs. The extracellular domain (//) was not drawn to scale. All the Draper I constructs were C-terminally tagged with HA. (e) The deletion of a.a. 826-894 decreased the association between Draper I and TRAF4. V: vector alone. (f) Draper I Δ826-854 and Δ855-894, bound TRAF4-myc at similar levels, suggesting these two regions are equally important in mediating the interaction between TRAF4 and Draper I. (g) The mutation Y949F in Draper, which is required for Shark to bind Draper I, did not affect the interaction between Draper I and TRAF4, indicating that the interaction between TRAF4 and Draper I does not require Shark association. (h) Proposed model of Draper-mediated gial transcriptional responses to axon degeneration. Draper is cell-autonomously required for glia to respond to axon degeneration and genetically upstream of JNK signalling cascade activation. The two binding partners of Draper, TRAF4 and Shark, are also upstream of JNK signalling cascade, likely through the activation of Slpr. We propose activation of JNK signalling cascade induces dAP-1-mediated gene transcription, which is also upstream of STAT92E-dependent transcriptional activation. The activation of STAT92E leads to the upregulation of Draper and facilitation of gial engulfment of axonal debris.

Figure 3 | TRAF4 physically associates with Draper independently of Shark. (a) Western blots of co-immunoprecipitations from Drosophila S2 cells showed TRAF4-myc associated with Draper I-HA. All the blots shown are representative images after three repeats. (b) Schematic drawing of Drosophila TRAF4. ZF: zinc-finger enriched domain. TD: TRAF domain. (c) Draper I-HA interacted with the ZF region of TRAF4 but not the TRAF domain. (d) Schematic drawing of deletions in the Draper I intracellular for different constructs. The extracellular domain (//) was not drawn to scale. All the Draper I constructs were C-terminally tagged with HA. (e) The deletion of a.a. 826-894 decreased the association between Draper I and TRAF4. V: vector alone. (f) Draper I Δ826-854 and Δ855-894, bound TRAF4-myc at similar levels, suggesting these two regions are equally important in mediating the interaction between TRAF4 and Draper I. (g) The mutation Y949F in Draper, which is required for Shark to bind Draper I, did not affect the interaction between Draper I and TRAF4, indicating that the interaction between TRAF4 and Draper I does not require Shark association. (h) Proposed model of Draper-mediated gial transcriptional responses to axon degeneration. Draper is cell-autonomously required for glia to respond to axon degeneration and genetically upstream of JNK signalling cascade activation. The two binding partners of Draper, TRAF4 and Shark, are also upstream of JNK signalling cascade, likely through the activation of Slpr. We propose activation of JNK signalling cascade induces dAP-1-mediated gene transcription, which is also upstream of STAT92E-dependent transcriptional activation. The activation of STAT92E leads to the upregulation of Draper and facilitation of gial engulfment of axonal debris.
Antibodies. For primary antibodies used in this study: Mouse anti-GFP clone 3E6 ( Molecular Probes A-11201 2.000 for IFA; 1.2000 for WB), rabbit anti-Draper ( ref. 37) (1.500 for IFA and 1.000 for WB after pre- absorption with yw embryos overnight), mouse anti-t-tubulin (DM1A) (Sigma T9026, 1.000), rat anti-m-Cherry (Invitrogen M12117, 1.000), rat anti-HA (3F10) (Roche 1.867-423, 1.2000 for WB), mouse anti-myc (9E10) (Millipore 05-419, 1.5000), mouse anti-HiW (6H4) (Developmental Studies Hybrido Bank, 1.000) and mouse anti-Repo (Developmental Studies Hybrido Bank, 1.5).FITC anti-mouse IgG (714-095-150), Cy3 anti-rabbit IgG (715-165-150), Cy3-anti-rat IgG (712-165-150) and Cy5 anti-rabbit IgG antibodies (715-095-150), Cy3 anti-rabbit IgG (712-1165-150), Cy3 anti-mouse IgG (715-165-150) and Cy5 anti-rabbit IgG antibodies (715-165-150) were from Jackson ImmunoResearch Laboratories (all 1.000), as well as horseradish peroxidase-conjugated anti-rabbit antibodies (712-035-150, 1.6000). Both horseradish peroxidase-conjugated anti-mouse (ab6808) and anti-rabbit IgG (ab6271-1) were from Abcam (1.6000).

Injury protocol and adult brain dissection. Drosophila cultures were kept at 25 °C except for the experiments involving UAS-mst9 and UAS-dra2.8 to 10 days due to the lethality they caused when overexpressed using repo-Gal4. Temperature-sensitive Gal80 (Gal80p)38 was used to inhibit the activity of Gal4 at 18 °C before fly eclosion and then deactivated at 29 °C for at least 5 days before injury. Flies continued to grow at 29 °C until the day of dissection. Maximal pupal and the third antennal segments were used as a pair of fine forceps for microdissection. For immunocytochemistry, heads were first fixed and permeabilized in 4% formaldehyde/0.1% Triton X-100/PBS for 20 min at room temperature (R.T.). After washed with PBST (0.1% Triton X-100/PBS) for 5 times, brains were dissected in PBST and post-fixed with 4% formaldehyde/0.1% Triton X-100/PBS for 15 min. Fixed brains were then washed for five times with PBST, blocked in 0.1% Triton X-100/5% BSA/PBS for 30 min at R.T., and incubated with primary antibodies in PBST at 4 °C overnight or two nights for anti-Draper antibody. The next day, the brains were washed five times with PBST and then incubated with secondary antibodies in PBST for 2 h at R.T. or 4 h for Draper staining. After washed with PBST for five times, the brains were then mounted with Vectashield Mounting Medium (Vector Labs) and stored at 4 °C before imaging within 2 weeks. For western blot analysis, heads from adult flies aged at 25 °C for at least 5 days after eclosion were isolated and immediately dissected in chilled PBS. Only the central brains (without the optic lobes) were collected for western blot analysis. Equal number of males and females were used for each experiment. Dissected brains were spun down at the highest speed for 1 min and then lysed in 1X Laemmli lysis buffer (2 m, 25 °C for 1 min) and boiled at 100 °C for 5 min. After centrifugation, the supernatants were collected for western blot analysis. The molecular weight of proteins was determined using SDS-PAGE analysis and western blotting. Images have been cropped for presentation in Fig. 2i. Full-size images are presented in Supplementary Fig. 8.

Confocal microscopy and image analysis. Confocal microscopy settings were always kept constant throughout the same set of experiments. Zeiss LSM5 Pascal confocal microscopy was used for all axonal debris clearance, TRE-eGFP reporter assay (except for T4A19-1r37), and STAT92E reporter assay. Zeiss Plan-Apochromat × 63/1.4NA oil objective lens. Glial MARCM clones, glial activation in hiw14n animals and TRE-eGFP reporter activity in mst9n animals were analysed using Zeiss Axio Imager.M2 spinning-disk confocal microscope with Zeiss C-Apochromat × 40/1.2NA water objective lens. For quantification of the fluorescence intensity, the centre single x-section was identified from each image stack of the relevant glomeruli or central adult brain, and pixel intensity was measured using ImageJ (National Institute of Health).

DNA cloning and Draper I intracellular domain deletions. To clone TRAF4 cDNA, forward primer containing KpnI and reverse primer containing SacII restriction site (all the primer sequences are listed in Supplementary Table 1) were used to identify TRAF4 cDNA from BDPG (Berkeley Drosophila Genome Project) cDNA clone LD20987. An approximate 1.6 kb cDNA was amplified by Phusion PCR Purification (QIAGEN) and then digested by KpnI and SacII. Digested DNA fragment was ligated to pUAST-CT-myc vector that has 6x myc sequence (MEQKLISEEDLNE) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs). The N-terminal (ZF, a.a. 1-309) and the C-terminal TRAF domain (MEQKLISEED) at the C-terminus, using T4 DNA ligase (New England BioLabs).
11. Arthur-Farraj, P. J. et al. c-Jun reprograms Schwann cells of injured nerves to generate a repair cell essential for regeneration. *Neuron* 75, 633–647 (2012).

12. Logan, M. A. et al. Negative regulation of glial engulfment activity by Draper terminates glial responses to axon injury. *Nat. Neurosci.* 15, 722–730 (2012).

13. Chatterjee, N. & Bohmann, D. A Versatile Fc31 based reporter system for measuring AP-1 and Nrf2 signaling in Drosophila and in tissue culture. *PLoS ONE* 7, e34063 (2012).

14. Xiong, X. et al. The Highwire ubiquitin ligase promotes axonal degeneration by tuning levels of Nmnat protein. *PLoS Biol.* 10, e1001440 (2012).

15. Lee, T. & Luo, L. Mosaic analysis with a repressible cell marker for studies of gene function in neuronal morphogenesis. *Neuron* 22, 451–461 (1999).

16. Doherty, J., Logan, M. A., Tajedemir, O. E. & Freeman, M. R. Ensheathing gli function as phagocytes in the adult *Drosophila* brain. *J. Neurosci.* 29, 4768–4781 (2009).

17. Kay, J. N., Chu, M. W. & Sanes, J. R. MEGF10 and MEGF11 mediate homotypic interactions required for mosaic spacing of retinal neurons. *Curr. Biol.* 9, 101–104 (1999).

18. Igaki, T., Pagliarini, R. A. & Xu, T. Loss of cell polarity drives tumor growth and invasion through JNK activation in *Drosophila*. *Curr. Biol.* 16, 1139–1146 (2006).

19. Liu, H., Su, Y.-C., Becker, E., Treisman, J. & Skolnik, E. Y. A nuclear migration in the developing eye. *Mech. Dev.* 122, 97–108 (2005).

20. Bach, E. A. et al. GFP reporters detect the activation of the Drosophila JAK/STAT pathway in vivo. *Gene Expr. Patterns* 7, 323–331 (2007).

21. Houalla, T., Hiernaux, D., Ruan, W., Suter, B. & Rao, Y. The Ste20-like kinase misshapen functions together with Bicaudal-D and dynein in driving nuclear migration in the developing *Drosophila* eye. *J. Neurosci.* 23, 11772–11779 (2002).

22. Gadea, A., Schinelli, S. & Gallo, V. Endothelin-1 regulates astrocyte proliferation and reactive gliosis via a JNK/c-Jun signaling pathway. *J. Neurosci.* 28, 2394–2408 (2008).

23. Chung, W.-S. *et al.* Astrocytes mediate synapse elimination through MEGF10 and MERTK pathways. *Nature* 504, 394–400 (2013).

24. Oliva, A. A., Kang, Y., Sanchez-Molano, J., Furonos, C. & Atkins, C. M. STAT3 signaling after traumatic brain injury. *J. Neurochem.* 120, 710–720 (2012).

25. Lu, T.-Y., Doherty, J. & Freeman, M. R. DRK/DOS/SOS converge with Crk/Mbc/dCed-12 to activate Rac1 during glial engulfment of axonal debris. *Proc. Natl Acad. Sci. USA* 111, 12544–12549 (2014).

26. Sepp, K. J., Schulte, J. & Auld, V. J. Peripheral glia direct axon guidance across the CNS/PNS transition zone. *Dev. Biol.* 238, 47–63 (2001).

27. Nagarkar-Jaiswal, S. *et al.* A library of MiMICs allows tagging of genes and reversible, spatial and temporal knockdown of proteins in *Drosophila*. *Elife* 4, 2743–2758 (2015).

28. McPhee, C. K., Logan, M. A., Freeman, M. R. & Baehrecke, E. H. Activation of autophagy during cell death requires the engulfment receptor Draper. *Nature* 465, 1093–1096 (2010).

29. Couto, A., Alenius, M. & Dickson, B. J. Molecular, anatomical, and functional organization of the *Drosophila* olfactory system. *Curr. Biol.* 15, 1535–1547 (2005).

30. Sepp, K. J., Schulte, J. & Auld, V. J. Peripheral glia direct axon guidance across the CNS/PNS transition zone. *Dev. Biol.* 238, 47–63 (2001).

31. Nagarkar-Jaiswal, S. *et al.* A library of MiMICs allows tagging of genes and reversible, spatial and temporal knockdown of proteins in *Drosophila*. *Elife* 4, 2743–2758 (2015).

32. Mathew, S. J., Rembold, M. & Leptin, M. Role for Traf4 in polarizing adherens junctions as a prerequisite for efficient cell shape changes. *Mol. Cell Biol.* 31, 4978–4993 (2011).

33. Neumann, L. J., Burdett, T. C., Gonzalez, M. A., Züchner, S. & Freeman, M. R. Rapid in vivo forward genetic approach for identifying axon death genes in *Drosophila*. *Prot. Natl Acad. Sci. USA* 111, 9965–9970 (2014).

34. Stork, T., Sheehan, A., Taydemir-Yilmaz, O. E. & Freeman, M. R. Neuron-glia interactions through the heatless FGF receptor signaling pathway mediate morphogenesis of *Drosophila* astrocytes. *Neuron* 83, 388–403 (2014).

35. Wu, C., Wairkar, Y. P., Collins, C. A. & DiAntonio, A. Highwire function at the *Drosophila* neuromuscular junction: spatial, structural, and temporal requirements. *J. Neurosci.* 25, 9557–9566 (2005).

36. Luo, L., Liao, Y. J., Jan, L. Y. & Jan, Y. N. Distinct morphogenetic functions of similar small GTPases: *Drosophila* Drac1 is involved in axonal outgrowth and myoblast fusion. *Genes Dev.* 8, 1787–1802 (1994).

37. Freeman, M. R., Delrow, J., Kim, J., Johnson, E. & Doe, C. Q. Unwrapping glial biology: Gcm target genes regulating glial development, diversification, and function. *Neuron* 38, 567–580 (2003).

38. Guigue, S. E., Mao, Z. & Davis, R. L. Spatial-temporal gene expression targeting with the TARGET and gene-switch systems in *Drosophila*. *Sci. STKE* 2004, p6 (2004).

Acknowledgements
We thank Tian Xu (Yale U) and Maria Leptin (Cologne U) for providing fly strains, and Jaeda Coutinho-Budd for critical reading of the manuscript. This work was supported by NIH RO1 NS053538 to M.R.F., and during the period of this study M.R.F. was an Investigator with the Howard Hughes Medical Institute.

Author contributions
T.-Y.L., J.M.M. and M.R.F. designed research. All the experiments were performed by T.-Y.L., except the re-expression of Highwire in the neurons of Highwire null animals (performed by L.J.N.). A.E.S. cloned full-length TRAF4 cDNA. R.B. assisted with cell culture and co-IP. M.A.L. provided transgenic flies.

Additional information
Supplementary Information accompanies this paper at http://www.nature.com/naturecommunications

Competing financial interests: The authors declare no competing financial interests.

Reprints and permission information is available online at http://npg.nature.com/reprintsandpermissions/

How to cite this article: Lu, T.-Y. *et al.* Axon degeneration induces glial responses through Draper-TRAF4-JNK signalling. *Nat. Commun.* 8, 14355 doi: 10.1038/ncomms14355 (2017).

Publisher’s note: Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/

© The Author(s) 2017