Ask1 and Akt act synergistically to promote ROS-dependent regeneration in Drosophila

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Abstract

How cells communicate to initiate a regenerative response after damage has captivated scientists during the last few decades. It is known that one of the main signals emanating from injured cells is the Reactive Oxygen Species (ROS), which propagate to the surrounding tissue to trigger the replacement of the missing cells. However, the link between ROS production and the activation of regenerative signaling pathways is not yet fully understood. We describe here the non-autonomous ROS sensing mechanism by which living cells launch their regenerative program. To this aim, we used Drosophila imaginal discs as a model system due to its well-characterized regenerative ability after injury or cell death. We genetically-induced cell death and found that the Apoptosis signal-regulating kinase 1 (Ask1) is essential for regenerative growth. Ask1 senses ROS both in dying and living cells, but its activation is selectively attenuated in living cells by Akt1, the core kinase component of the insulin/insulin-like growth pathway. Akt1 phosphorylates Ask1 in a secondary site outside the kinase domain, which attenuates its activity. This modulation of Ask1 activity results in moderate levels of JNK signaling in the living tissue, as well as in activation of p38 signaling, both pathways required to turn on the regenerative response. Our findings demonstrate a non-autonomous activation of a ROS sensing mechanism by Ask1 and Akt1 to replace the missing tissue after damage. Collectively, these results provide the basis for understanding the molecular mechanism of communication between dying and living cells that triggers regeneration.

Author summary

One of the early events that occur after tissue damage consists on the production of Reactive Oxygen Species (ROS), that signal to the surrounding tissue to initiate wound healing and regeneration. Many signaling pathways, such as JNK and p38, respond to oxidative stress and are necessary for regenerative growth. As the link between ROS and...
regenerative signaling is not well understood, we decided to explore the mechanism that underlies this process. To do that, we genetically induced cell death in specific areas of *Drosophila* wing imaginal discs and then studied the mechanism that drives living cells to replace the damage zone until it is completely regenerated. We found that the *Drosophila* Apoptosis signal-regulating kinase 1 (Ask1), a protein that is sensitive to oxidative stress, is a key player in this scenario. This protein acts as an intracellular sensor that upon damage activates the JNK and p38 regenerative signaling pathways. However, high activity of Ask1 can be toxic for the cell. This is controlled by Akt, a kinase downstream the insulin pathway, which attenuates the activity of Ask1 in the living cells that will participate in the regeneration process. In consequence, Ask1 and Akt act synergistically to respond to the stress generated after tissue damage and drive regeneration. Our results provide a first overview within the framework of how insulin signaling inputs could modulate the capacity to overcome tissue damage.

**Introduction**

Organisms are continuously exposed to a wide variety of environmental stressors that can cause deterioration and cell death. Tissues overcome the effect of those stressors by replacing damaged cells to restore homeostasis. Therefore, understanding the early signals that initiate the response to damage is an essential issue in regenerative medicine. Regeneration can be monitored in *Drosophila* imaginal discs, which are well-characterized epithelial sacs capable to regenerate after genetically-induced apoptosis or when parts are physically removed (reviewed in [1]). Compiling evidence supports that reactive oxygen species (ROS) fuel wound healing and oxygen-dependent redox-sensitive signaling processes involved in damage response [2–4]. Actually, genetically-induced apoptosis in the imaginal discs, using the Gal4/UAS system, leads to the production of ROS which propagate to the surrounding neighbors [5–7]. Although oxidative stress has been associated with several pathologies, it has been described that low levels of ROS can be beneficial for signal transduction [8].

The Jun-N Terminal kinase (JNK) and p38 signaling pathways are MAP kinases that respond to many stressors, including ROS, and foster regeneration and cytokine production in *Drosophila* [5,6,16–25,7,26,27,9–15]. Both pathways control numerous cellular processes as diverse as cell proliferation and cell death. For example, ectopic activation of JNK induces apoptosis [28,29], but its inhibition results in lethality [30]. It is though that these disparities could be due to either different levels of activity or different mechanisms of activation.

After genetic ablation in a specific zone of the wing imaginal disc, dying cells produce high levels of ROS, and show high JNK activity [5,7]. High levels of JNK have been associated to dying cells and are involved in a feedback amplification loop to enhance the apoptotic response to stress [31]. However, the living cells nearby the dying zone show low levels of ROS, which are beneficial for the cell as they turn on the activation of p38 and low levels of JNK [5]. Both MAP kinases in the living cells are required for a cytokine-dependent regenerative growth. A key question is how the balance between the beneficial or detrimental effects of ROS is controlled and, in particular, how ROS control JNK and p38 activity. A candidate molecule to perform this function is the MAPKKK Apoptosis signal-regulating kinase 1 (Ask1), which responds to various stresses by phosphorylation of the kinases upstream JNK and p38 [32–34]. Hence, in a reduced environment, thioredoxin (Trx) inhibits Ask1 kinase activity by directly binding to the N-terminal region of Ask1. Upon oxidative stress, the redox-sensitive cysteines of Trx become oxidized, resulting in the dissociation of Trx from Ask1.
Consequently, Ask1 is oligomerized and its threonine-rich kinase domain is phosphorylated, inducing Ask1 activation [35,36]. Mammalian Ask1 is highly sensitive to oxidative stress and contributes substantially to JNK-dependent apoptosis. Nevertheless, recent studies have also revealed other functions of this kinase, including cell differentiation and survival [37]. Ask1-interacting proteins promote conformational changes that lead to the modulation of Ask1 activity and result in various cellular responses. For example, Ask1 is a substrate for phosphorylation by Akt1, a serine/threonine kinase activated by lipid kinase phosphatidylinositol 3-kinase (Pi3K) pathway in response to insulin receptor activation. This phosphorylation is associated to a decrease of Ask1 activity \textit{in vitro} [38]. The Pi3K/Akt pathway, which is conserved between mammals and \textit{Drosophila}, is one of the main effector signals for the regulation of tissue growth [39–41]. In \textit{Drosophila}, loss-of–function mutants of various components of the pathway result in reduced body size or lethality [42]. Conversely, mutants of the phosphatase \textit{PTEN}, an antagonist of Pi3K, result in high activity of Akt and tissue overgrowth [43–45]. Thus, it is conceivable that Pi3K/Akt signaling is involved in regenerative growth, but whether Pi3K/Akt and Ask1 interact for controlling regeneration is unknown and deserves attention.

In this work, we genetically induced cell death in wing imaginal discs to explore the link between ROS production and regeneration. We found that Ask1 acts as a sensor of ROS upstream the JNK and p38 pathways. Moreover, we describe that Akt1 is necessary for modulating Ask1 activity in living cells to trigger regeneration. In addition, our results indicate that oxidative stress generated in the damaged cells signals the neighboring living cells to promote tissue repair.

\section*{Results}

\subsection*{Ask1 is necessary for regenerative growth}

The Gal4/UAS/Gal80\textsubscript{TS} transactivation system has been extensively used to temporarily and spatially activate pro-apoptotic genes such as \textit{reaper (rpr)} [10,11,46]. To study the capacity to regenerate, we induced cell death in the wing disc using the wing-specific \textit{sal\textsuperscript{E/Pv}}-Gal4 strain to activate UAS-rpr, in a temporally controlled manner thanks to the tub-Gal80\textsubscript{TS} thermo sensitive Gal4-repressor (henceforth \textit{sal\textsuperscript{E/Pv}>rpr}). We first kept the embryos at 17\textdegree C until the 8th day (192 hours after egg laying) to activate the \textit{tub-Gal80\textsubscript{TS}} and therefore to prevent \textit{rpr} expression. Subsequently, we moved those larvae to 29\textdegree C for 11 hours, to inhibit \textit{tub-Gal80\textsubscript{TS}}, and activate apoptosis in the \textit{sal\textsuperscript{E/Pv}}-Gal4 zone. Then, larvae were returned to 17\textdegree C to avoid further apoptosis and allow tissue to regenerate. After adults emerged, wings were dissected and regeneration was analyzed. With these experimental conditions, 100\% of the wings of \textit{sal\textsuperscript{E/Pv}>rpr} flies contained the full set of veins and interveins, which demonstrates that the missing parts were regenerated (Fig 1A).

The same experiment was carried out in \textit{Ask1} mutant backgrounds. The \textit{Ask1\textsuperscript{MB06487}} mutant flies are viable in homozygosis but lethal over the deficiency \textit{Df(3R)BSC636}, which suggests that is a hypomorphic allele. The \textit{Ask1\textsuperscript{MI02915}} mutant is lethal in homozygosis as well as over the \textit{Df(3R)BSC636}, which suggests that is an amorphic allele. We found that after genetically-induced cell death in \textit{Ask1\textsuperscript{MB06487}} or \textit{Ask1\textsuperscript{MI02915}} heterozygous background, full regeneration is only achieved in 15\% and 23\% of the wings, respectively (Fig 1A). The phenotypes of the abnormally regenerating wings vary and consisted in lack of some veins or parts of a vein or anomalous vein segments (S1A Fig).

To further address the role of \textit{Ask1} in regeneration, we used a double transactivation system to simultaneously induce cell death in the \textit{sal\textsuperscript{E/Pv}} domain and inhibit \textit{Ask1} with UAS-RNAi (UAS-\textit{Ask1\textsuperscript{RNAi}}) in an adjacent compartment (Fig 1B). RNAi knockdown of \textit{Ask1 (UAS-\textit{Ask1\textsuperscript{RNAi}})} reduces \textit{Ask1} mRNA total levels to approximately 25–30 percent of that observed in controls.
The UAS-Ask1 RNAi transgene was activated in the anterior compartment using the Gal4/UAS system (ci-Gal4>UAS-Ask1RNAi) and cell death was induced in the salE/Pv domain using the Gal80-repressible transactivator system LHG (LexA-Hinge-Gal4 activation domain), a modified form of the lexA lexO system (salE/Pv-LHG>lexO-rpr) (Fig 1B) [5,47]. Both transgenes were activated at the same time, following the same protocol as in the previous experiment. The resulting adult wings (ci>Ask1RNAi salE/Pv>rpr) lacked some veins or interveins and their size was reduced in all cases observed. However, the inhibition of Ask1 in the anterior compartment for the same time but without cell death, did not affect vein pattern nor wing size (ci>Ask1RNAi) (Fig 1B).

Similar results were obtained after driving UAS-Ask1RNAi to a different compartment (dorsal) and inducing cell death in salE/Pv (ap>Ask1RNAi salE/Pv>rpr); in this case, we found that 82% of wings could not regenerate properly. As for the anterior compartment, Ask1RNAi in the dorsal compartment without cell death did not affect vein pattern nor wing size (ap>Ask1RNAi) (Fig 1B).

To discard any toxicity due to the insertion of the transgene, all experiments were carried out in parallel but constantly at 17˚C to maintain tub-Gal80TS activity and block transgene (UAS- or lexO-) expression. In these conditions no defects in the wings were detected (salE/Pv OFF in Fig 1A and 1B).

To evaluate whether these anomalies were due to impairment of proliferation, we analyzed the mitotic index, calculated as the number of cells positive for the phosphorylated form of H3.
Histone 3 (P-H3) in the anterior compartment (ci>), where the Ask1 RNAi transgene was expressed. It is well known that apoptosis in discs induces compensatory proliferation of nearby cells (reviewed in [1]). Accordingly, genetic ablation (salE/Pv>rpr ci>GFP) resulted in an increase of mitosis compared to the neutral (salE/Pv>GFP ci>RFP) or to the Ask1 knockdown (salE/Pv>GFP ci>Ask1 RNAi) conditions. However, combining genetic ablation in the salE/Pv zone with Ask1 knockdown in the nearby anterior compartment, the number of mitosis associated to damage did not increase (salE/Pv>rpr ci>Ask1 RNAi) (Fig 1C). This observation confirms that Ask1 reduction impairs regenerative growth.

### Ask1 activity is higher in apoptotic cells than in regenerating cells

The Drosophila Ask1 locus encodes two protein isoforms of different lengths, Ask1-RB and Ask1-RC, with predicted molecular weights of 136.5 kDa and 155.5 kDa, respectively. Both have a protein kinase-like domain that contains a highly conserved core of threonines conferring functionality to the protein (reviewed in [32]). It has been reported that after Trx release, the Ask1 oligomer undergoes conformational change leading to trans-autophosphorylation of the threonine residue corresponding to Thr838, Thr845 and Thr747 in human, mouse and Drosophila Ask1, respectively [32]. In human cells, phosphorylation of the N-terminus Ser83 by Akt results in attenuation of Ask1 activity in vitro [38]. This is noteworthy, because attenuation of Ask1 could result in moderate levels of activity necessary for turning on JNK and p38 in living cells. Drosophila Ask1 lacks this residue and the Akt consensus motif in the N-terminus (S2 Fig, S1 Appendix, S2 Appendix). However, we found that the longer Ask1-RC isoform contained a conserved domain of unknown function DUF4071 with a highly conserved Ser (human Ser174, Drosophila Ser83), which we hypothesized that could be key in Ask1 regulation (Fig 2A, S2 Fig).

Active Ask1 can be traced with antibodies against the phosphorylated threonine residues of the highly conserved kinase domain (henceforth P-Thr) [48] and the attenuated form with specific antibodies against phosphorylated Ser83 (P-Ser83). We first tested both antibodies in wild-type wing imaginal discs. We detected low levels of P-Thr all over the disc including some transient high activity during mitosis (Fig 2B and S3A Fig), as occurs in dividing mammalian cells [49]. In Ask1 mutant backgrounds, both the general and the mitosis-associated P-Thr levels were reduced but not abolished (S3B Fig), and Ask1 RNAi discs did not show any effect on P-Thr levels in mitosis (Fig 2E). This discrepancy may be due to the hypomorphic condition of the mutant combinations, but also to that the antibody may recognize some unspecific epitope in mitotic cells. We also found basal levels of P-Ser83 in wild-type discs (Fig 2C). Ectopic expression of Ask1 RNAi suppressed the low endogenous P-Ser83 Ask1 levels (S3C Fig).

Remarkably, upon apoptosis, high levels of P-Thr were localized in dying cells (Fig 2D) and absent or, in some cases, increased weakly above the basal levels in nearby living cells. This increase of P-Thr in apoptotic cells was inhibited after Ask1 RNAi expression (Fig 2E). In contrast, P-Ser83 accumulated in living cells adjoining the apoptotic zone and was completely absent in apoptotic cells. The increase in P-Ser83 varied from discs with strong accumulation near the dying domain to those with an extended increase in the whole wing pouch (2D and S4A Figs). Similar results were obtained after killing cells with a different pro-apoptotic gene (salE/Pv>hid) (S4B Fig). In the presence of apoptosis and blocking Ask1, the P-Ser83 increase in living cells was inhibited (salE/Pv>rpr ci>Ask1 RNAi, Fig 2F). Moreover, P-Ser83 was also found to be elevated at the wound edges after physical injury (S4C Fig). Together, these observations indicate that living cells respond to damage by phosphorylation of Ask1 Ser83.
Fig 2. Ask1 activation in apoptotic cells and in regenerating cells. (A) Structural features of Ask1-RC isoform. The phosphokinase domain (KD) with a highly conserved core of threonines is shown in green. The DUF4071 domain, close to the N-terminal is shown in red. Inactive Ask1 is constituted by association of Ask1 monomers through the coiled coil domain (grey shadow) and the reduced form of thioredoxin (Trx-SH) in the N-terminal domain (left). Upon oxidation, Trx-S-S dissociates, which allows Ask1 autophosphorylation in the phosphokinase domain, leading to an active Ask1. The C-terminal coiled coil domain (dark grey) allows interactions between Ask1 monomers to form oligomers. Active Akt phosphorylates Ask1 on Ser83 residue. (B-C) Characterization of mammalian P-ASK1 antibodies in Drosophila discs. Endogenous levels of P-Thr (B) and P-Ser83 (C) in wild-type (WT) discs. (D) Ask1 activity in saE/Pv>rpvrpr discs. SalE>Pv>rpvrpr disc indicating the area imaged (white square); TP-3 (TO-PRO-3) nuclei; purple area: the salE>Pv>rpvrpr ablated zone. Disc stained with TUNEL assay and Ask1 P-Thr, (n = 21). Disc stained with Ask1 P-Ser83 (n = 35). (E-F) After cell death (purple) and simultaneous inhibition of Ask1 in the anterior compartment (saE>Pv>rpvrpr ci>Ask1WT, P-Thr (E) and P-Ser83 (F). White-dotted line indicates the AP boundary; anterior to the left, posterior to the right. The inset in (E) depicts the Ask1WT area (black) and the salE>Pv>rpvrpr ablated zone (purple). (G) Ectopic expression of the Ask1WT form in the posterior compartment (blue). (H) Ectopic expression of the mutant form Ask1S83A. Scale bars 50μm. (I) Percentage of regenerated wings in controls (rpvrpr, Ask1WT or
In contrast to the high activity of Ask1 in dying cells (high P-Thr), the presence of P-Ser83 in living cells could be indicative of tolerable levels of Ask1 achieved by attenuation of P-Thr activity. To test this hypothesis, we mutated Ask1 at serine 83 to alanine and cloned it into a UAS vector (UAS-Ask1S83A). In parallel, we also cloned a wild-type form of Ask1 (UAS-Ask1WT). We ectopically expressed Ask1WT in the posterior compartment of the wing disc (hh-Gal4 UAS-Ask1WT), which resulted in an increase of P-Ser83, in addition to low levels of P-Thr (Fig 2G). Interestingly, the ectopic expression of Ask1S83A in the posterior compartment (hh-Gal4 UAS-Ask1S83A), did not show a rise in P-Ser83, but strong activation of P-Thr (Fig 2H). The high levels of P-Thr were not homogeneously distributed, varied from disc to disc, and were always found within the posterior compartment (hh>), where the transgene was activated. This observation concurs with the P-Ser83 residue as responsible for the attenuation of P-Thr.

We next analyzed whether the Ser83 residue was key for the damage response. Using the double transactivation system, we found that the expression of UAS-Ask1WT or UAS-Ask1S83A transgenes in the anterior compartment without cell death did not cause any defects in wing morphology, size or vein and intervein patterning (salE/Pv>rpr ci>Ask1WT or Ask1S83A in Fig 2I). After genetic ablation, expression of UAS-Ask1WT resulted in full regeneration in 89% of wings (salE/Pv>rpr ci>Ask1WT). In contrast, expression of UAS-Ask1S83A led to a fall to only 29% of individuals being capable of regenerate and the rest showed strong effects on wing morphology, with altered pattern of veins and interveins, as well as the appearance of notches, which together are indicative of disrupted regeneration (salE/Pv>rpr ci>Ask1S83A in Fig 2I). The suppression of the ability to regenerate by UAS-Ask1S83A, which may act as a dominant negative allele, is likely due to the lack of non-autonomous P-Ser83 increase in the living cells near the damaged zone.

**Pi3K/Akt signaling attenuates Ask1 activity in living cells**

Next, we decided to identify the upstream signal responsible for Ser83 phosphorylation. In mammalian cells, attenuating phosphorylation of Ask1 is driven by the serine-threonine Akt kinase, the core kinase of the insulin pathway [38]. We wondered whether the *Drosophila* Akt1 as well as its upstream Pi3K92E kinase (the *Drosophila* Pi3K kinase also known as dp110), were required for Ser83 phosphorylation.

We first found that the active phosphorylated form of Akt (P-Akt) was increased in the living tissue and decreased or was absent in the dying zone (Fig 3A–3D). To test whether the Akt1 phosphorylated the endogenous Ask1 Ser83 in vivo, we specifically inhibited this kinase with a UAS-AktRNAi in a stripe of cells at the center of the disc, the ptc domain (ptc>AktRNAi), and found that P-Ser83 was reduced (Fig 3E and 3F). In addition, ectopic activation of Akt1 (UAS-myr-Akt1S), a constitutively activated membrane-anchored form of Akt1 [50], resulted in an increase in the levels of P-Thr (Fig 3G and 3H).

We next studied the role of P-Akt on Ask1 in the context of regeneration. In the presence of cell death and blocking Pi3K92E, with the dominant negative form dp110DN in the adjacent compartment, we observed a reduction in the accumulation of P-Akt and P-Ser83 induced after genetic ablation (salE/Pv>rpr hh> dp110DN in Fig 3I–3K). Moreover, in the overlapping zone of dying cells and dp110DN, the accumulation of P-Thr was unaffected (Fig 3L and S5 Fig). We also scored the effects on wing regeneration after blocking Akt and Pi3K92E kinases.
Fig 3. Phosphorylation of *Drosophila* Ask1 Ser83 depends on the Pi3K/Akt1 pathway. (A–C) P-Akt staining in wild-type discs (A), and after *ptc>*rpr (B), and *sal>*rpr induction (C). Apoptotic zone (pyknotic nuclei) in purple. (D) Mean pixel intensity for the wild type and *ptc>*rpr taken from regions of interest in the A and P compartments as shown in the images. (E,F) RNA interference of Akt1 in the *ptc* stripe of cells (*ptc>*Akt1<sup>DN</sup>) and (G,H) ectopic expression of a constitutively active form of Akt1 (*ptc>*myr-Akt1S), stained with P-Akt and Ask1 P-Ser83. (I) Design of the experiments in J,K and L with *sal>*rpr induction (purple) and blocking Pi3K (*ih>*dp110<sup>DN</sup>) in the posterior compartment (black). Staining with anti-P-Akt (I), with anti-Ask1 P-Ser83 (K) with Ask1 P-Thr (L). White-dotted line indicates anterior-posterior boundary. TP-3; TO-PRO-3 nuclear staining. Scale bars 50 μm. (M) Percentage of fully regenerated wings after genetic ablation and Pi3K pathway inhibition alone or in combination with *Ask1* mutant backgrounds (genotypes indicated). Right: *sal>*rpr ON, examples of wings with full regeneration (controls *sal>*rpr or ci>*dp110<sup>DN</sup> alone) and anomalous regeneration (*sal>*rpr with the genotypes indicated); *sal>*rpr OFF wings: examples of control wings in the absence of rpr-ablation (maintained at 17˚C). Error bars indicate standard error of sample proportion. ***P<0.001.

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Regeneration was impaired when the dominant negative form of Pi3K<sup>92E</sup> was expressed in the anterior compartment (*sal<sup>E/Pv</sup>*rpr ci>*dp110<sup>DN</sup>) as well as in the Akt1<sup>1</sup> heterozygous background (*sal<sup>E/Pv</sup>*rpr Akt1<sup>1/+</sup>), an allele that encodes a catalytically inactive protein [51] (Fig 3M). Moreover, regeneration was severely affected in double heterozygous flies containing Akt1<sup>1</sup> and *Ask1<sup>MB06487</sup>* or *Ask1<sup>MI02915</sup>* alleles (Fig 3M). Neither the dominant negative form of Pi3K<sup>92E</sup> nor the allelic combination Akt1<sup>1</sup> and *Ask1<sup>MB06487</sup>* or *Ask1<sup>MI02915</sup>* affected wing development in the absence of cell death (*sal<sup>E/Pv</sup>* OFF wings in Fig 3M). These results further support the notion that Pi3K<sup>92E</sup>/Akt1 and Ask1 genetically interact and that their epistatic interaction is key to drive regeneration.

**ROS promote Ask1 activation**

Next, we modulated the ROS levels in order to determine if Ask1 can sense oxidative stress after damage. We fed *sal<sup>E/Pv</sup>*rpr larvae with food supplemented with N-acetyl cysteine (NAC), a potent non-enzymatic scavenger that decreases ROS production, and examined Ask1 phosphorylation (Fig 4A). After induction of cell death, we found a significant decrease in both P-Thr and P-Ser83 in discs from NAC-fed larvae (Fig 4B and 4C; S3 Appendix). In addition, we fed larvae with H<sub>2</sub>O<sub>2</sub>-supplemented food and observed a significant increase in both P-Thr and P-Ser83 levels (Fig 4D–4F). This oxidative stress-induced increase was blocked in the hypomorphic *Ask1<sup>MB06487</sup>* homozygous mutant (Fig 4E and 4F; S3 Appendix). It is known that in mammalian cells Ask1 can also be activated by endoplasmic reticulum (ER) stress [52]. Feeding larvae with tunicamycin, an inhibitor of N-glycosylation in the ER that induces ER stress, led to an increase in Ask1 activation, which was also blocked in *Ask1<sup>MB06487</sup>* mutant discs (Fig 4E and 4F; S3 Appendix).

Furthermore, we examined whether Akt1 activation in the living cells is targeted non-autonomously by ROS produced by the dying cells. To examine this hypothesis, we enzymatically blocked ROS production using ectopic expression of the ROS scavengers *Super-oxide dismutase 1* and *Catalase* (*Sod1:Cat*) in the rpr ablated region (*sal<sup>E/Pv</sup>*rpr, *Sod1:Cat*) and monitored the pixel intensities of two adjacent zones in each disc, the anterior (A) and posterior (P) to the *sal<sup>E/Pv</sup>* (Fig 4G). The activation of the *Sod1:Cat* transgene concomitantly with *sal<sup>E/Pv</sup>*rpr did not inhibit apoptosis and P-Thr was detected in the dying cells (S6 Fig). However, in these conditions, we found that P-Akt levels in neighboring cells were reduced in both A and P zones when *Sod1:Cat* was co-expressed with rpr. Accordingly, we found that the increase of P-Ser83 in neighboring cells was blocked when *Sod1:Cat* was co-expressed with rpr (Fig 4G; S3 Appendix). Together, these results demonstrate that the oxidative stress generated from the dying cells targets Akt1 and Ask1 in the neighboring undamaged tissue.
Fig 4. Ask1 is activated by ROS. (A) Design of experiments after feeding larvae with the NAC antioxidant. (B) Examples of Ask1 P-Thr after salE/Pv>rpr induction in standard or NAC-supplemented food. Mean pixel intensities of P-Thr fluorescent labeling in the apoptotic zone of salE/Pv>rpr discs from larvae fed with a standard or NAC-supplemented diet. (C) Examples of Ask1 P-Ser83 after salE/Pv>rpr induction with standard or NAC-supplemented food. White dotted line indicates the salE/Pv>rpr dying domain. Mean pixel intensities of Ask1 P-Ser83 fluorescent labeling in the neighboring cells near the apoptotic zone of salE/Pv>rpr discs from larvae fed with a standard or NAC-supplemented diet. (D) Design of experiments after enhancing oxidative stress. (E) Left: P-Thr mean pixel intensity of wild-type wing discs from larvae fed with standard, H2O2- and tunicamycin-supplemented food. Right: P-Thr mean pixel intensity of Ask1>hox mutant discs from larvae fed with standard, H2O2- and tunicamycin-supplemented food. (F) Left: P-Ser83 mean pixel intensity of wild-type wing discs from larvae fed with standard, H2O2- and tunicamycin-supplemented food. Right: P-Ser83 mean pixel intensity of Ask1>hox mutant discs from larvae fed with standard, H2O2- and tunicamycin-supplemented food. (G) P-Akt (top) and P-Ser (bottom) mean pixel intensity from discs with rpr activation (salE/Pv>rpr, GFP; high oxidative stress) and from discs with simultaneous Sod1 Cat and rpr activation (salE/Pv>rpr, Sod1 Cat; low oxidative stress). Purple area shows where transgenes were activated. A and P are the zones where pixel intensity was measured. S.f.: standard food. *P<0.05. **P<0.01, ***P<0.001. Error bars indicate standard deviation. TP-3: TO-PRO-3 nuclear staining. Scale bars: 50μm.

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Ask1 activates p38 and JNK in regeneration

We have shown that the Ask1 P-Ser83 attenuating signal is essential for regeneration, and that this signal depends on ROS and Akt. We next hypothesized that this P-Ser83-mediated attenuation does not silence Ask1, but instead maintains the low levels of Ask1 activity necessary for regeneration. Therefore, we analyzed the expression of its known targets, the stress-activated MAP kinases JNK and p38 [48,53], both required for ROS-dependent regeneration [5,6]. We induced cell death in the salE/Pv domain and simultaneously blocked Ask1 in the anterior compartment, and used the posterior as an internal control for the same disc (ci>Ask1 RNAi salE/Pv>rpr). We found most phosphorylated p38 (P-p38), as an indicator of p38 activity, in the posterior compartment (Fig 5A and 5B). The effects of Ask1 on JNK activity were monitored by matrix metalloproteinase 1 (Mmp1) expression as it is one of the bona fide transcriptionally regulated read-outs of the JNK pathway [54]. After salE/Pv>rpr cell death, Mmp1 was found in both compartments, but in ci>Ask1 RNAi salE/Pv>rpr discs Mmp1 mostly accumulated in the posterior (Fig 5C and 5D). The effects of Ask1 interference on JNK pathway were strengthened after staining with P-JNK antibody. P-JNK was found accumulated around the dead zone in salE/Pv>rpr discs. However, when Ask1 was knockdown in the anterior compartment (ci>Ask1 RNAi salE/Pv>rpr discs), P-JNK was reduced in this compartment and accumulated in the posterior (S7A and S7B Fig). To confirm that Ask1 is responsible for JNK and p38 activation, we tested whether Ask1 was required for their activation after physical damage. We made two incisions in hh>GFP,Ask1 RNAi discs, one in the UAS-Ask1 RNAi compartment and another into the control compartment. We found that P-p38 localized at the wound edges in the control compartment, whereas the levels of P-p38 were reduced at the wound edges of the Ask1 RNAi compartment (Fig 5E and 5G). Likewise, cut Ask1 RNAi discs resulted in less Mmp1 than in the control compartment (Fig 5F and 5H). In addition, we tested the puckered (puc) reporter of the JNK pathway, which normally accumulates in wounds, and found to drop in heterozygous Ask1 discs after a physical injury (S7C and S7D Fig). Together, these results indicate that Ask1 is necessary for the activation of p38 and JNK during regeneration.

Discussion

We have shown here that Ask1 acts as a sensor of ROS after damage, and that synergizes with Pi3K/Akt1 to phosphorylate the Ask1 Ser83, a key phosphorylation event to initiate regeneration. In addition, we have demonstrated that the activation of the Pi3K/Akt1 and...
Ask1 in undamaged cells is originated non-autonomously by ROS produced by the damaged tissue.

An essential question to understand regeneration is how damaged cells communicate to the nearby living tissue to initiate regenerative growth. One of the most classical and determining studies on epithelial regeneration was the discovery that massive cell death caused by ionizing radiation in *Drosophila* larvae resulted in proliferation of the unaffected neighbors to compensate the lost [55]. Since then, compensatory proliferation, a cellular response linked to
regeneration, has been considered as a result of signals released from apoptotic cells (reviewed in [56]). We propose here that the oxidative stress generated by damaged or apoptotic cells signals undamaged tissue to initiate the Ask1/Akt1 machinery that will culminate with repair and compensatory proliferation.

ROS acting in signaling after wounding has been subject of extensive research in various organisms and tissues [5,6,64–67,26,57–63]. Here, we have uncovered two scenarios of Ask1 activity: high activity in apoptotic cells, with high levels of P-Thr; and low activity in undamaged cells, where P-Ser83 prevails over P-Thr. This fits with the high levels of ROS and JNK in dying cells and the low levels of ROS, low JNK and P-p38 in living cells reported previously [5]. Together, our observations address the question of how JNK signaling selectively fosters apoptosis or proliferation, and how this signal is related to high or low ROS levels. The same applies to p38, which is only activated in unharmed cells when neighbors enter in apoptosis or are damaged [5,53].

Our study demonstrates that Ask1 operates in an Akt1-dependent manner in living cells and in an Akt1-independent manner in dying cells. It is conceivable that Akt1-mediated attenuation results in either low or transient levels of Ask1 activity necessary to stimulate P-p38 and low JNK levels for regeneration. In contrast, apoptotic cells induce high or sustained levels of JNK because high levels of ROS result in high Ask1 P-Thr in the absence of Akt1 attenuation.

Based on the results presented here and previous observations, we propose a model in which oxidative stress and the absence or presence of Pi3K/Akt1 survival signals are integrated in Ask1 to control the selective activity of JNK and p38, and in turn regenerative growth (Fig 5I). In this model, only dying cells produce high ROS levels and high JNK activity [5,7]. In addition, other mechanisms can exacerbate apoptosis through the lethal levels of JNK, and therefore increase ROS production. For example JNK, as well as p53, functions in a positive feedback after apoptosis as it transcriptionally activates the pro-apoptotic genes hid and rpr, thus amplifying apoptosis [31,68]. It is therefore likely that the high ROS that result in high Ask1 activation, inabilities the dying cells to respond to PI3K/Akt1, preventing the Ask1 attenuation and resulting in deleterious JNK activity. We also have demonstrated that P-Akt1 is absent, or low, in dying cells, which supports a lack of PI3K/Akt1-dependent Ask1 attenuated activity in dying cells. These cells will enter into a committed-to-die status with no return to survival. In contrast, the living neighbor cells, albeit adjacent to dying cells, show tolerable levels of ROS [5,7], and activate moderate levels of Ask1 through the Akt1-dependent phosphorylation of Ask1 P-Ser83. In these cells, Ask1 will turn on the activation of tolerable levels of JNK and p38, leading to a committed-to-regenerate status. Eventually, the JNK and p38 pathways will promote the transcription of the unpaired cytokines necessary for survival and regeneration [5,15,69–72]. In addition to cytokines, Wnt/Wg signaling pathway acts as a mitogenic signal involved in wing disc regeneration that responds to JNK [11,18,69,73] as well as the inhibition of the Hippo pathway, which results in increased Yki activity [74–76].

Several phosphatases are sensitive to ROS, and could be also integrated into the model presented here. This is the case of the JNK phosphatases, which are critical molecular targets of ROS. Oxidative inhibition of JNK phosphatases results in sustained JNK activation in order to promote caspase 3 cleavage and eventually cell death [77]. Therefore, the early burst of ROS in dying cells could activate lethal levels of JNK activity through inhibition of the Drosophila JNK phosphatase puckered concomitantly with the high levels of Ask1, and together boost the committed-to-die status. Additionally, the PTEN phosphatase, which regulates many cellular processes through direct antagonism of PI3K signaling, is also ROS sensitive. Increasing ROS can enhance insulin signaling attributable to the oxidative inhibition of PTEN [78]. PTEN inactivation enhances the activation of AKT signaling, which in turn promotes the expression of cell survival genes [78]. Thus, PTEN could be involved in the living cells to release the Akt1 activity
that is necessary to reach the tolerable levels of Ask1/JNK/p38. Therefore, the low oxidative stress that reaches the living cells can trigger not only Trx2 dissociation from Ask1, but also the PTEN-oxidation dependent Pi3K/Akt1 activity required for attenuated Ask1 necessary for survival and proliferation.

Our model concerns to an inherent response in the imaginal disc epithelium. However, other signals could enhance or stabilize the ROS-dependent regenerative response. For example, ROS produced after wounding are necessary for recruitment and activation of the Drosophila macrophage-like immune cells, which will secrete the TNF ortholog Eiger. Eiger activates JNK signaling in the epithelial cells to contribute to the regenerative response [6].

Because the moderate activity of Ask1 for regeneration is achieved in an Akt1-dependent manner, and because Akt1 functions downstream of the insulin signaling, it seems conceivable an implication of nutrition in imaginal disc regeneration. The production and secretion of insulin and AKH, the fly analog of mammalian glucagon, vary depending of the food content. Thus, a challenge for the future will be to analyze how these hormones can modulate the regenerative response to damage. For example, the Drosophila TNF Eiger is produced by the fat body when larvae are exposed to low protein diet [79]. As Eiger activates JNK non-autonomously in many tissues, it could have a central role in imaginal disc regeneration, depending on food availability. The relation between nutrition and metabolism, as well as their involvement in regeneration deserves further analysis in order to achieve an integrative view of regeneration.

Materials and methods

Drosophila strains

The Drosophila melanogaster strains, salE/Pv-LHG and lexO-rpr, are previously described [5]; lexA-GFP (gift from K. Basler). UAS-H2B-RFP (from J. Knoblich), Akt1 (gift from H. Stocker), sal-Gal4, salE/Pv-Gal4 (gift from J.F. de Celis), ci-Gal4 (from R. Holmgren) and en-Gal4 (gift from G. Morata). The following strains were provided by the Bloomington Drosophila Stock Center: tubGal80TS (RRID:BDSC_7017), ptc-Gal4 (RRID:BDSC_2017), Ask1MB06487 (RRID:BDSC_36048) Ask1Mn02915 (RRID:BDSC_36163), UAS-GFP (RRID:BDSC_4776), UAS-dp110DN (RRID:BDSC_25918), Df(3R)BSC636 (RRID:BDSC_25726), UAS-Ask1RNAi (RRID:BDSC_35331), UAS-Cat.A (RRID:BDSC_24621), UAS-Sod.A (Sod1) (RRID:BDSC_24754), UAS-rpr on the X chromosome (RRID:BDSC_5823), UAS-rpr on the third chromosome (RRID:BDSC_50791). These strains are described in FlyBase: hh-Gal4, ap-Gal4, UAS-hid. The UAS-AktRNAi (2902) strain was obtained from the Vienna Drosophila Resource Center (VDRC). We used the UAS-myr-Akt1.S [50]. Canton S and w118 were used as controls. A full list of genotypes is provided at the end of this section.

Genetic ablation and dual Gal4/LexA transactivation system

Cell death was genetically induced as previously described [5]. We used the salE/Pv-Gal4 as a driver, which consists of the spalt wing enhancer with expression confined to the wing to score adult wing parameters. The UAS lines used to promote cell death were UAS-rpr or UAS-hid, two pro-apoptotic genes, controlled by the thermo-sensitive Gal4 repressor tubGal80TS. We also used the salE/Pv-LHG and LexO-rpr strains [5] for genetic ablation, utilizing the same design as for Gal4/UAS.

Embryos were kept at 17°C until the 8th day/192 hours after egg laying to prevent rpr expression. They were subsequently moved to 29°C for 11 hours and then back to 17°C to allow tissue to regenerate. Two types of controls were always treated in parallel; individuals
without rpr expression (UAS-GFP, moved to 29°C for 11 hours) and individuals kept continuously at 17°C to avoid any transgene activation.

In most of the experiments shown here, we induced cell death for 11h. However the induction of cell death for testing the effects of signals emerging from the dying zone was longer. This is the case of the P-Akt experiment in Fig 3B and 3C and for the genetic scavenging of ROS (Sod1:Cat) (Fig 4G and S6 Fig), in which sal>rpr was activated for 24h.

In dual transactivation experiments, we used salE/Pv-LHG LexO-rpr to ablate the salE/Pv domain (abridged as salE/Pv>rpr), whereas Gal4 was used to express different transgenes under the control of Gal4 drivers (ci-Gal4 for anterior compartment; ap-Gal4 for dorsal compartment and hh-Gal4 for posterior compartment).

**Test for regenerated adult wings and statistics**

To test the capacity to regenerate in different genetic backgrounds, we used adult wings emerged from salE/Pv>rpr individuals in which patterning and size defects can be scored easily. Flies were fixed in glycerol: ethanol (1:2) for 24 hours. Wings were dissected in water and then washed with ethanol. Subsequently, they were mounted in lactic acid:ethanol (6:5) and analyzed and imaged under a microscope.

The percentage of regenerated wings refers to fully regenerated (for genetic ablation genotypes) or normally developed wings (for testing transgenes) and was calculated according to the number of wings with a complete set of veins and interveins, as markers of normal patterning. For each sample, we scored the percentage of individuals belonging to the “regenerated wings” class. We calculated the standard error of the sample proportion based on a binomial distribution (regenerated complete wing or not) SE = √(p(1-p)/n), where p is the proportion of successes in the population.

**Computational characterization of the fly Ser83 domain**

Serine/threonine putative phosphorylation sites were determined by scanning the orthologous human/fly sequences with the RxRxx[ST] domain motif for the Akt kinase (purple block on S2 Fig) and the degenerated pattern xxRxRxx[ST] (green blocks on S2 Fig). Strikingly the human Ser83 site was not found on *Drosophila melanogaster*; which was confirmed by using GeneWise (version wise2.4.1) [80] to map the M3K5_HUMAN protein sequence (Q99683), downloaded from UniProt [81], over a window of 10kbp centered at the Ask1 transcription start site; this sequence segment was retrieved from FlyBase FB2017_06 genome version ([82], chr3R:19,875,000–19,885,000[+]). The GeneWise output is provided as S1 Appendix, the software tool was not able to find a match on the fly genome for the initial 126 residues from the human amino term. In order to assess the relevance of the fly Ser83 two homology-based approaches were implemented. First, a search on PFAM database [83] returned the domain of unknown function DUF4071 (PF13281). The first amino acid positions for this domain are highly conserved across a set of diverse homologs, including fly Ask1-PC and human MAP3K5 but also paralogs sharing the domain. The consensus sequence for DUF4071 has been included on the right alignment block from S2 Fig for illustrative purpose. The second approach was considering a precomputed set of homolog sequences downloaded from EggNOG database (version 4.5) [84]. A cluster of homologous sequences already included fly Ask1 (KOG4279), providing a total of 338 protein sequences for 164 species; a total of 49 sequences were selected from that set, fly Ask1-PC isoform among them, that were also reliably annotated as orthologs of human MAP3K5. Fly Ask1-PB isoform protein sequence was retrieved from FlyBase and appended to the selected MAP3K5 orthologs. Then, the 50 protein sequences were aligned by MAFFT (version 7.271) [85] using the following
parameters: maxiterate = 1000, localpair, op = 10. The complete alignment is available from S2 Appendix, from which two conserved blocks centered on the human and fly Ser83 domains were trimmed and processed with TeXshade [86] to produce the S2 Fig. Note that on S2 Fig the domain highlighted in cyan, which includes the fly equivalent column highlighted in red, is highly conserved in a wide range of taxa, from sponges to humans. Other domains are conserved only within chordata or even tetrapoda species. Such conservation signature makes Drosophila Ser83 a good candidate for functional and mutational studies.

**Generation of UAS-Ask1\(^{WT}\) and UAS-Ask1\(^{S83A}\) flies**

pUASt-attb\_Ask1\(^{WT}\) was constructed by cutting Ask1 cDNA EcoRI /BamHI from DGRC clone FI02066 and cloning it into pUASt-attb.

pUASt-attb\_Ask1\(^{S83A}\). Two serines in the 75 and 83 residues in the DUF4071 domain showed putative non-canonical AKT phosphorylation sites (ILTQRPRLSHYGYVRESF). Both residues are spatially exposed similarly to human Ser83 of ASK1, which makes them accessible to kinases. pUASt-attb\_Ask1 S83A was constructed by mutating serine 83 to alanine by PCR using oligos Ask1Mut-Fwd and Ask1S83A-Rev for partial PCR1 and Ask1S83A-Fwd and Ask1Mut-Rev for partial PCR2. Complete PCR was performed using the two partial PCRs as templates with oligos Ask1Mut-Fwd and Ask1Mut-Rev. The complete PCR was then cut with EcoRI /PflMI and cloned into FI02066-cut EcoRI/PflMI. The mutation introduced a new StuI site that was used to check the mutated clones. Mutated Ask1S83A cDNA was then cut from EcoRI/BamHI and cloned into pUASt-attb. Both clones were injected by standard procedures in line zh-86Fb-attP and transgenic lines were selected.

Ask1Mut-Fwd: AAT ACA AGA AGA GAA CTC TGA ATA CGG AAT
Ask1Mut-rev: CGG CGG TGT GGT TTT GTG CAC AAA CCG ATC
Ask1S83A-Fwd: CGT TAG GGA GGC CTT CGG GAT GAA GGA GA
Ask1S83A-Rev: CGG CGG TGT GGT TTT GTG CAC AAA CCG ATC

**Immunostaining**

Immunostaining was performed using standard protocols. The primary antibodies used in this study were the polyclonal Ask1 P-Ser83 (Santa Cruz Biotechnology sc-101633 1:100, which recognizes the conserved region surrounding human P-Ser83) and Ask1 P-Thr (Santa Cruz Biotechnology sc-109911 1:100, which labels the preserved region nearby mouse P-Thr845). We selected the anti-Ask1 P-Ser83 because of the following in vivo analysis: (1) its localization dropped under a UAS-Ask1\(^{RNAi}\) (S3C Fig); (2) Its localization increased after ectopic activation of Ask1\(^{WT}\) (Fig 2G); (3) Mutation in the Drosophila Ser83 residue (UAS-Ask1\(^{S83A}\)), inhibited its localization (Fig 2H); (4) Akt activation or repression, resulted in increased or decreased localization of P-Ser83 (Fig 3E–3H). Other antibodies used were: Ptc (DSHB, 1:100), P-p38 (Cell Signalling 1:50), Mmp1 (cocktail of three antibodies: DSHB 3A6B4, 5H7B11, 3B8D12 1:100), P-Akt (S473 Cell Signalling 1:100) and P-Histone-H3 (Millipore, 1:1000).

The anti-P-JNK used was the Anti-ACTIVE\(^+\) JNK pAb, Rabbit, (V7931, Promega, 1:100). The images were taken using the Thermal LUT from the 3D Surface Plot of FIJI, with maximum values of 50% and minimum 24% to minimize the background. Calculation of pixel intensities was obtained from raw images.

Fluorescently labeled secondary antibodies were from ThermoFisher Scientific. Discs were mounted in SlowFade or ProLong (ThermoFisher Scientific) supplemented with 1 µM TO-PRO-3 (TP-3) or YO-PRO-1 (YP1) (Life Technologies) to label nuclei. For apoptotic cell detection, we used the TUNEL assay. We employed the fluorescently labeled Alexa Fluor®
The number of mitotic cells per mm² (Fig 1C) was calculated after counting the number of P-Histone-H3 (P-H3) positive cells in the anterior compartment of the wing pouch and hinge, excluding the notum, for all the genotypes shown. Note that two genotypes in Fig 1C show the anterior compartment labeled with the transgene ci-Gal4 UAS-GFP (ci> GFP), whereas in the two ci>Ask1RNAi experiments the anterior compartment was labeled with anti-Ci. In these experiments shown in Fig 1C, the induction of cell death or activation of transgenes was done for 16h at 29°C. The number of mitosis after analyzing the stacks of confocal images was calculated using Fiji software.

**Imaginal disc culture and physical injury**

Wing discs were dissected from third instar larvae in Schneider’s insect medium (Sigma-Aldrich), and a small fragment was removed with tungsten needles. To visualize Mmp1 staining, the discs were cultured in Schneider’s insect medium supplemented with 2% heat-activated fetal calf serum, 2.5% fly extract and 5 μg/ml insulin, for 5 hours at 25°C. For P-p38, discs were injured in Schneider’s insect medium and immediately fixed and stained. Ex vivo images were taken using a Leica SPE confocal microscope and processed with Fiji software. Ex vivo discs were also monitored live after a cut, using puc-Gal4 UAS-GFP as a JNK reporter, using the same medium. For nuclei visualization in ex vivo culture, we used NucRed Live647 (Life Technologies; 1 drop per slide) added 20 minutes before imaging.

**ROS scavenging**

To prevent ROS production in salE/Pv>rpr discs we pursued two different protocols. First, we inhibited ROS chemically (Fig 4A–4F). To do this, standard fly food was supplemented with the antioxidant N-acetyl cysteine (NAC 100 μg/ml; Sigma-Aldrich). NAC treatment was dispensed on the 7th day of development at 17°C. On the 8th day, experimental larvae were moved to 29°C for 11 hours to promote cell death, whereas controls were transferred to a vial with standard food and moved to 29°C for the same time period. Afterward, the larvae were move back to 17°C to allow tissue recovery. Second, we decreased ROS production genetically by the ectopic expression of the Sod1 and Catalase enzymes using a recombinant fly UAS-Sod1/UAS-Cat (Sod1:Cat) in the salE/Pv>rpr domain (Fig 4G and S6 Fig). For those experiments, Sod1:Cat and rpr were activated for 24 hours at 29°C.

**Oxidative stress induction**

Third instar larvae were transferred to vials with 5mL of special medium containing 1.3% UltraPure™ LMP agarose (Invitrogen), 5% sucrose (Fluka) and the desired concentration of 0.1% H₂O₂ (Merck) or 1ng/μl tunicamycin (Sigma-Aldrich). To avoid loss of oxidative capacity, these substances were added to the media at a temperature below 45°C. The larvae were fed for 2 hours prior to dissection and fixation of the discs. Controls without H₂O₂ or tunicamycin were always handled in parallel.

**Quantitative real-time PCR analysis**

Control (hh>RFP) and experimental (hh>Ask1RNAi) conditions were induced for 16h at 29°C (192h after egg laying) using the Gal4/tub-Gal80TS system. Wing imaginal discs (n = 40) were dissected after induction in three biological replicates for each condition. RNA was extracted with the Zymo Research ZR RNA MicroPerp (R1060/R1061) and RNA Clean and
Concentrator (R1015/R1016) kits following standard protocols. RT-PCR was performed using SYBR green Master Mix (Roche). Specific primers for Ask1 and Rps18 are detailed in S1B Fig. ΔΔCt method was used to normalize the data. Ask1RNAi and control samples were normalized against the housekeeping gene Rps18. Average standard error of the mean (SEM) of the three biological replicates was computed for each one based on three technical replicates by ΔΔCt method.

Genotypes

Fig 1.

(A) +/- wUAS-rpr/+; saEPv-Gal4/++; tubGal80TS/+  
Ask1MB0291S/+ wUAS-rpr/+; saEPv-Gal4/++; tubGal80TS/Ask1MB0291S  
Ask1MB06487/+ wUAS-rpr/+; saEPv-Gal4/++; tubGal80TS/Ask1MB06487  
(B) saEPv-rpr -> w; ci-Gal4/LexO-rpr; saEPv-LHG:tubGal80TS/UAS-GFP  
Ask1RNAi w; ci-Gal4/LexO-GFP; saEPv-LHG:tubGal80TS/UAS-Ask1RNAi  
Ask1RNAi w; ci-Gal4/LexO-rpr; saEPv-LHG:tubGal80TS/UAS-Ask1RNAi

Fig 2.

(D) saEPv-rpr -> wUAS-rpr/+; saEPv-Gal4/+; tubGal80TS/+  
(E,F) saEPv-rpr ci Ask1RNAi w; ci-Gal4/LexO-rpr; saEPv-LHG:tubGal80TS/UAS-Ask1RNAi

Fig 3.

A) Canton S

(B) ptc->rpr -> wUAS-rpr/+; ptc-Gal4; tubGal80TS/+  
(C) sal->rpr -> wUAS-rpr/+; sal-Gal4/+; tubGal80TS/+  
(D,E) ptc->AktRNAi w; ptc-Gal4; tubGal80TS/+; UAS-AktRNAi/+  
(F,G) ptc->myrAkt -> w; ptc-Gal4; tubGal80TS/+; UAS-myrAkt/+  
(H-K) sal->rpr hh dp110DN w; LexO-rpr/UAS-dp110DN; salEPv-LHG:tubGal80TS/+ tubGal4

(L) ci dp110DN w; ci-Gal4; UAS-dp110DN; salEPv-LHG:tubGal80TS/+  
+/+ w; LexO-rpr/+; salEPv-LHG:tubGal80TS/+ (control for LexO-rpr in the second chromosome)
Fig 4.

(B,C) Std food and NAC → wUAS-rpr/+; sal<sup>E/Pv</sup>-Gal4/+; tubGal80<sup>TS</sup>+/Akt<sup>1</sup>/Ask1<sup>MI02915</sup>

(E,F) CTRL → w<sup>118</sup>; +; +

Ask1<sup>1</sup>/Ask1<sup>MB06487</sup>/Ask1<sup>MB06487</sup>

(G) sa>rp, GFP → w; sal<sup>E/Pv</sup>-Gal4/UAS-GFP; tubGal80<sup>TS</sup>/UAS-rpr

sa>rpr, Sod1:Cat → w; sal<sup>E/Pv</sup>-Gal4/UAS-Sod1:UAS-Cat; tubGal80<sup>TS</sup>/UAS-rpr

Fig 5.

(A,C) sa>rp → w; LexO-rpr/+; sa<sup>E/Pv</sup>-LHG:tubGal80<sup>TS</sup>+/+

(B,D) sa>rp ci>Ask1<sup>RNAi</sup> → w; ci-Gal4/LexO-rpr; sa<sup>E/Pv</sup>-LHG:tubGal80<sup>TS</sup>/UAS-Ask1<sup>RNAi</sup>

(E-H) en>Ask1<sup>RNAi</sup>, GFP → w; en-Gal4/UAS-GFP; UAS-Ask1<sup>RNAi</sup>/+  

S1 Fig.

(A) sa<sup>E/Pv</sup>-rpr Akt<sup>1</sup>/Ask1<sup>MI02915</sup>/+ → wUAS-rpr/+; sa<sup>E/Pv</sup>-Gal4/+; tubGal80<sup>TS</sup>/Ask1<sup>1</sup>/Ask1<sup>MB06487</sup>

sa<sup>E/Pv</sup>-rpr Akt<sup>1</sup>/Ask1<sup>MB06487</sup>/+ → wUAS-rpr/+; sa<sup>E/Pv</sup>-Gal4/+; tubGal80<sup>TS</sup>/Ask1<sup>MB06487</sup>/Ask1<sup>MB06487</sup>/Ask1<sup>MB06487</sup>

sa<sup>E/Pv</sup>-rpr ci>Ask1<sup>RNAi</sup> → w; ci-Gal4/LexO-rpr; sa<sup>E/Pv</sup>-LHG:tubGal80<sup>TS</sup>/UAS-Ask1<sup>RNAi</sup>

(B) Rt-q-PCR

Controls: w; tubGal80<sup>TS</sup>/+; hh-Gal4/UAS-RFP

UAS-Ask1<sup>RNAi</sup> w; tubGal80<sup>8</sup>/+; hh-Gal4/UAS-Ask1<sup>RNAi</sup>

S3 Fig.

(A) w<sup>118</sup>; +; +

(B) Ask1<sup>1</sup>/Ask1<sup>+</sup> → w<sup>118</sup>; +; +

Ask1<sup>MB06487</sup>/Ask1<sup>MB06487</sup> → w<sup>118</sup>; +; Ask1<sup>MB06487</sup>/Ask1<sup>MB06487</sup>

Ask1<sup>MB06487</sup>/Def(3R)BSC636 → w<sup>118</sup>; +; Ask1<sup>MB06487</sup>/Def(3R)BSC636

(C) hh>Ask1<sup>RNAi</sup>, GFP → w; UAS-GFP/+; hh-Gal4/UAS-Ask1<sup>RNAi</sup>

S4 Fig.

(A) wt and ptc>rpr → wUAS-rpr/+; ptc-Gal4: tubGal80<sup>TS</sup>/+  

(B) sa<sup>E/Pv</sup>-hid → w; sa<sup>E/Pv</sup>-Gal4/+; tubGal80<sup>TS</sup>/UAS-hi

(C) wt and physically injured → Canton S

S5 Fig.

(A-D) sa<sup>E/Pv</sup>-rp ci>dpi110<sup>DN</sup> → w; ci-Gal4/UAS-dpi110<sup>DN</sup>; sal<sup>E/Pv</sup>-LHG: tubGal80<sup>TS</sup>/LexO-rpr

S6 Fig.

w; sa<sup>E/Pv</sup>-Gal4/UAS-Sod1:UAS-Cat; tubGal80<sup>TS</sup>/UAS-rpr

S7 Fig.

(A) wt → Canton S  

sa<sup>E/Pv</sup>-rpr → w; LexO-rpr/+; sa<sup>E/Pv</sup>-LHG: tubGal80<sup>9</sup>

sa<sup>E/Pv</sup>-rpr, ci>Ask1<sup>RNAi</sup> → w; ci-Gal4/LexO-rpr; Sa<sup>E/Pv</sup>-LHG: tubGal80<sup>9</sup>/UAS-Ask1<sup>RNAi</sup>

(C) w; UAS-GFP/+; puc-Gal4/+  

(D) w; UAS-GFP/+; puc-Gal4/Ask1<sup>MB06487</sup>

Supporting information

S1 Fig. Supporting information for Fig 1. (A) Spectrum of wings with anomalous regeneration, ranging from complete regeneration (sa<sup>E/Pv</sup>-rpr in wild type, top) to a severe effect found after Ask1<sup>RNAi</sup> (sa<sup>E/Pv</sup>-rpr in mutant conditions, bottom). It includes a range of defects found after inducing cell death in Ask1 heterozygous background and after UAS-Ask1<sup>RNAi</sup>. Arrowheads indicate some defects. (B) Real time quantitative PCR of the UAS-Ask1<sup>RNAi</sup>. The table indicates the primers used. The RT-qPCR analysis is shown for Ask1 expression in wing
discs of controls (hh>RFP) and Ask1RNAi (hh>UAS-Ask1RNAi). qPCR results are presented as relative quantity of mRNA between control and Ask1RNAi. Error bars represent the standard error of the mean from three biological replicates (n = 40 discs, each replicate). (TIF)

S2 Fig. Alignment of Ask1 D. melanogaster orthologs focusing on human and fly Ser83 residues. Two blocks of conserved columns pinpointing the aforementioned residues were selected from the whole-length alignment of 50 MAP3K5/Ask1 protein orthologs provided on the S2 Appendix. From top to bottom: sequence logos displaying information content based on amino acid frequencies; an estimate of hydrophobicity of the amino acid residues; the aligned sequences split into chordate and invertebrate sequences, above or below the two Drosophila melanogaster Ask1 isoforms respectively; finally, the similarity is depicted as a color gradient ranging from low (blue) to high (red). Bottom reference coordinates for the aligned residues are based on human reference sequence, which is listed on the first row. Serines, threonines, and arginines were labeled in red, green and blue when appearing outside the domain blocks, but shown in white and boldface within those blocks. Numbers on the left side of the alignment blocks correspond to the relative position of the first amino acid within those blocks for each species sequences. Left alignment block contains a domain in purple that matches the consensus RxRxx[ST] pattern, although conservation is limited to birds and mammals (dark versus light shades of violet). On the right, the domain highlighted in cyan is conserved across all the diverse taxa, from human (top) to sponges (bottom), and it contains a serine (red background) that corresponds to the fly Ser83. On both panels, another domain matching a degenerated xxRxx[ST] pattern that is only conserved within vertebrate sequences is shown as a green block. Right alignment also includes at the bottom the consensus sequence of a PFAM domain of unknown function (DUF4071) that starts two residues before the Ser83 conserved block; this block within the domain shows high conservation across the different protein families considered when building the corresponding model. (TIF)

S3 Fig. Supporting information for Fig 2. (A) Mitotic (P-H3 positive) and Ask1 P-Thr. Lower right images: Zoom of mitotic cells from the white square. TP-3: TO-PRO-3 nuclear staining. (B) The basal levels of P-Thr in wild-type (WT) discs (mitotic and interphase cells) decrease in the Ask1MB06487 homozygous mutant background and are strongly reduced in the Ask1MB06487/Ask1MB06487 mutant background over the deficiency Def(3R)50636. Quantification of mean pixel intensity for P-Thr in mitotic and for P-Thr in non-mitotic (basal levels) cells are shown in separate graphics. P-Thr mean Pixel intensity in mitotic cells of the wild-type discs (+/+), 113.92±20.60, S.D. (n = 10); of the Ask1MB06487/Ask1MB06487 discs, 104.43±9.54, S.D. (n = 10); and of the Ask1MB06487/Def(3R)BSC636 88.37±13.03, S.D. (n = 10). Mean pixel intensity of the basal levels in interphase cells of the wild-type, 50.44±6.27, S.D. (n = 20); of the Ask1MB06487/Ask1MB06487, 41.65±4.80, S.D. (n = 10); and of the Ask1MB06487/Def(3R)BSC636, 27.22±5.73, S. D. (n = 18). (C) Inhibition of the endogenous Ask1 P-Ser83 levels after expressing the UAS-Ask1RNAi in the posterior compartment, using the hh-Gal4 driver. Scale bars: 5μm. (TIF)

S4 Fig. Supporting information for Fig 2. (A) Left: wild-type disc stained with P-Ser83. Right: disc in which apoptosis has been induced with the driver ptc-Gal4 UAS-rpr (ptc>rpr) and stained with P-Ser83 (n = 5). (B) SalE/Pv>hid discs stained with P-Thr (n = 12) and P-Ser83 (n = 5). YP-1: YO-PRO-1 nuclear staining. Purple: salE/Pv>rpr ablated zone. (C) Staining of Ask1 P-Ser83 after physical injury, (n = 6). Square:
magnified image. Scale bars: 50μm.

(TIF)

S5 Fig. Supporting information for Fig 3. (A) Design of the experiments in B, C and D with salE/Pv>rpr cell death (purple) and blocking Pi3K (ci>dp110DN) in the anterior compartment (black). (B-D) Staining of P-Akt (B), Ask1-P-Ser83 (C) and Ask1 P-Thr (D). White-dotted line indicates anterior-posterior boundary. TP3: TO-PRO-3 nuclear staining. Scale bars: 50μm.

(TIF)

S6 Fig. TUNEL assay and P-Thr of discs with simultaneous Sod1:Cat and rpr activation (salE/Pv>rpr, Sod1:Cat; low oxidative stress). Purple area shows where transgenes were activated.

(TIF)

S7 Fig. Supporting information for Fig 5. (A) P-JNK staining in wild type, salE/Pv>rpr, and two examples of salE/Pv>rpr ci>Ask1 RNAi. Thermal LUT was used to detect pixel differences. TP3: TO-PRO-3 nuclear staining. Purple indicates the dead zone. Colorized red indicates the ci>Ask1 RNAi zone. (B) Quantification of the P-JNK in raw images. A and P indicate the approximate areas measured for pixel intensity in n = 15 for each genotype. (C-E) Live imaging of a puc-Gal4, UAS-GFP used as a reporter of the JNK pathway. Discs were cut and cultured for 6 hours. White lines indicate the wound site. White arrows point to the endogenous puc expression zone at the tip of the notum. (C) Wild type. (D) Ask1MB06487 heterozygous. The nuclear marker NucRed Live647 was added in the culture medium. Scale bar: 50μm. (E) Quantification of puc-Gal4, UAS-GFP reporter at the wound site (n = 13 discs each genotype). Error bars indicate standard deviation. **P<0.01, ***P<0.001.

(TIF)

S1 Appendix. Computational verification that human Ser83 domain is missing on Drosophila melanogaster Ask1 gene loci. This file contains the plain text genewise alignment of the human M3K5_HUMAN protein sequence against the genomic segment of 5kbp upstream/downstream that is centered on the annotated fly Ask1 gene transcription start site. No alignment was reported for the first 126 residues of the human protein where the Ser83 RxRxx[ST] domain has been described.

(PDF)

S2 Appendix. Complete alignment of the selected 50 MAP3K5/Ask1 protein orthologs that illustrates high conservation across the sequence. Many positions are preserved from sponges to humans, including serine/threonine residues, whilst other show a dual pattern of conservation with specific substitutions between chordate and invertebrate species. From the whole set of 338 sequences on 164 species that define the EggNOG (http://eggno gdb.embl.de) homologs cluster KOG4279, comprising human MAP3K5 and its counterpart in fly Ask1-PC, we filtered out 49 proteins that were confidently annotated as MAP3K5 orthologs on different taxa plus fly Ask1-PB isoform downloaded from FlyBase, which are listed here (names shown in parentheses are those used on the alignment sequence labels and refer to UniProt identifiers when possible): Homo sapiens ENSP00000351908 (M3K5_HUMAN), Pan troglodytes ENSPTRP00000043317 (K7CFQ9_PANTR), Gorilla gorilla ENSGGOP0000022651 (G3S3K1_GORGO), Macaca mulatta ENSMMUP00000021508 (F7G2W1_MACMU), Pongo abelii ENSPPYP00000019077 (H2PKF0_PONAB), Otoblemur garnettii ENSOGAP0000000498 (H0WGN9_OTOGA), Callithrix jacchus ENSCJAP00000015659 (F7G2H9_CALJA), Rattus norvegicus ENSRNOP00000051338 (D3W27_RAT), Mus musculus ENSMUSB00000093485 (M3K5_MOUSE), Cavia porcellus ENSCPO0000003829 (H0V2Q4_CAVPO), Bos taurus
ENSBTAP00000011949 (F1MXH6_BOVIN), Pteropus vampyrus ENSPVAP00000009127 (ens9127_PTEVA), Mustela putorius furo ENSMUP00000014726 (G9K9I6_MUSPP), Loxodonta africana ENSLAFP00000000494 (G3UDL3_LOXAF), Echinops telfairi ENSTEP00000012543 (ens12543_ECHTE), Canis lupus familiaris ENSCAFP00000000362 (F1PA02_CANFA), Ailuropoda melanoleuca ENSAMEP00000010579 (D2H0V2_AILME), Tursiops truncatus ENSTTPR000000002388 (ens2388_TURTR), Dasypus novemcinctus ENSDNOP00000014966 (ens14966_DASNO), Sarcophilus harrisii ENSSHAP00000000495 (G3WHQ0_SARHA), Gallus gallus ENSGALP00000022482 (F1NYS9_CHICK), Meleagris gallopavo ENSMGAP00000015731 (G1NK21_MELGA), Taeniopygia guttata ENSTGUP00000011202 (H0ZKT1_TAEGU), Xenopus tropicalis ENSXETP000000005271 (F7A415_XENT), Gasterosteus aculeatus ENSGACP000000009628 (G3NWA9_GASAC), Oreochromis niloticus ENSONIP00000007016 (I3JDS6_ORENI), Danio rerio ENSDARP000000006740 (E7F2L3_DANRE), Takifugu rubripes ENSTRUP00000002505 (H2RQT9_TAKRU), Tetraodon nigroviridis ENSNIP00000022322 (Q4RFA2_TETNG), Branchiostoma floridae JGI1219597 (C3XTE7_BRAFL), Drosophila melanogaster FBpp0089256 (Ask1-PB_DROME) and FBpp0289484 (Ask1-PC_DROME), D. ananassae FBpp0120816 (B3LXS9_DROAN), Culex quinquefasciatus CPIJ008487-PA (BOWNO2_CULQO), Anopheles gambiae AGAP000747-PB (QEA6_ANOGA), Aedes aegypti AAELO08311-PA (Q16Z55_AEDAE), Bomyx mori GB14238-PA (H9KAX6_APMIE), Nasonia vitripennis NV13729-PA (K7IXA9_NASVI), Acyrthosiphon pisum ACYPI009039-PA (J9K801_ACYPI), Tribolium castaneum TC000430-PA (TCOG-S2_TRICA), Daphnia pulex DappuP328832 (P32883_DAPP), Caenorhabditis elegans F59A6.1 (NY-1_CAEEL), C. japonica CIA14063 (K711N3_CAEJA), C. briggsae CBG0299 (A8WNU1_CAEBR), Strongylocentrotus purpuratus SPU_010872r (H319Y9_STRPU), Amphimedon queenslandica PAC_15725247 (15725247_AMFQE), Trichoplax adhaerens TriadP10277 (B3RYN2_TRIAD).

(PDF)

S3 Appendix. Data for the graphics of Fig 4.

(DOCX)

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