Transcriptional pausing factor M1BP regulates cellular homeostasis by suppressing autophagy and apoptosis in *Drosophila* eye

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Abstract

During organogenesis cellular homeostasis plays a crucial role in patterning and growth. The role of promoter proximal pausing of RNA polymerase II, which regulates transcription of several developmental genes by GAGA factor or Motif 1 Binding Protein (M1BP), has not been fully understood in cellular homeostasis. Earlier, we reported that M1BP, a functional homolog of ZKSCAN3, regulates wingless and caspase-dependent cell death (apoptosis) in the Drosophila eye. Further, blocking apoptosis does not fully rescue the M1BPRNAi phenotype of reduced eye. Therefore, we looked for other possible mechanism(s). In a forward genetic screen, members of the Jun-amino-terminal-(NH2)-Kinase (JNK) pathway were identified. Downregulation of M1BP ectopically induces JNK, a pro-death pathway known to activate both apoptosis and caspase-independent (autophagy) cell death. Activation of JNK pathway components can enhance M1BPRNAi phenotype and vice-versa. Downregulation of M1BP ectopically induced JNK signaling, which leads to apoptosis and autophagy. Apoptosis and autophagy are regulated independently by their genetic circuitry. Here, we found that blocking either apoptosis or autophagy alone rescues the reduced eye phenotype of M1BP downregulation; whereas, blocking both apoptosis and autophagy together significantly rescues the M1BP reduced eye phenotype to near wild-type in nearly 85% progeny. This data suggests that the cellular homeostasis response demonstrated by two independent cell death mechanisms, apoptosis and autophagy, can be regulated by a common transcriptional pausing mechanism orchestrated by M1BP. Since these fundamental processes are conserved in higher organisms, this novel functional link between M1BP and regulation of both apoptosis and autophagy can be extrapolated to humans.

Conflict of Interests
The authors declare no competing interests.

Keywords

Atg8a; caspase-independent; cell death; JNK; Promotor proximal pausing; Wg

Introduction

During organogenesis cellular processes like cell division, cell differentiation, and cell death are tightly regulated by the spatio-temporal expression of various genes that include transcription factors and signaling molecules [1]. Transcriptional regulation of gene expression is critical for patterning and cellular homeostasis [2–4]. Disruption of homeostatic processes by intrinsic or extrinsic signals ultimately disrupts the transcriptional process: either by directly affecting the transcription of target genes or by indirectly modifying the stability, abundance and activity of transcription factors [5]. A key response to homeostatic stress is activation of mechanisms that either lead to restoration of homeostasis by transcriptional mechanisms [6], or elimination of sub-optimal or less fit stressed cells by cell death mechanisms [7–9]. There are several types of cell death utilized by organisms to restore homeostatic balance [10]. Generally, cell death can occur by accident or by design. Accidental cell death like necrosis is often triggered by excessive and unexpected stress and is not directly controlled by specific genetic machinery. Alternatively, regulated cell death including apoptosis, autophagy, pyroptosis, entosis, parthanatos, etc. are controlled by specific genetic machinery and triggered by specific signals [10, 11]. Among these, apoptosis and autophagy may occur more frequently during development. Although these two processes occur independently, research shows that blocking apoptosis can induce increased autophagy in a compensatory fashion to trigger cell death [12, 13]. Although associations between aberrant cellular homeostasis to several diseases (for example, neurodegenerative diseases, cancers) have been made, mediators of the cellular response to homeostatic stress are not well understood.

Recently, transcriptional pausing mechanism(s) during transcription elongation have been implicated as another rate-limiting step alongside downstream events in transcription initiation [14, 15]. Therefore, these mechanism(s) serve as key regulators of gene expression during development as well as in response to homeostatic stress [16, 17]. During organogenesis, transcription factors facilitate RNA polymerase II (Pol II) recruitment to DNA resulting in dynamic spatio-temporal gene expression patterns [18, 19]. These transcription factors bind to enhancers, proximal promoters, or core promoters within genes to initiate Pol II-mediated transcription [20]. During transcription, Pol II usually gets recruited before the transcription start site (TSS) to promote transcription [14]. Promoter proximal pausing is conserved from flies to mammals [21, 22], and is known to regulate several developmentally controlled genes during *Drosophila* embryogenesis [23]. Sequence-specific transcription factors like GAGA factor (GAF) and Motif 1-Binding Protein (M1BP) regulate Pol II mediated transcriptional pausing [24]. M1BP is highly conserved across species and encodes a 55kDa protein containing a zinc-associated domain (ZAD) towards the N-terminus and five C_2H_2 zinc-finger domains towards the C-terminus. *Drosophila* M1BP is functionally homologous to vertebrate zinc-finger with SCAN and KRAB domain 3 (ZKSCAN3) transcription factors [25, 26]. M1BP functions by binding to Motif1 sequences on

target genes to regulate or pause their transcription and has been reported to block several genes during development [24]. Such transcriptional pausing mechanisms may be responsible for creating unique genetic signatures and regulating cellular processes [15, 24].

We showed that M1BP mediated transcriptional pausing can regulate evolutionarily conserved WNT/Wingless (Wg) signaling during *Drosophila* eye development [27]. In the *Drosophila* eye, M1BP regulates expression of Wg, a ligand for the WNT/Wg signaling pathway. Ectopic Wg signaling is known to suppress eye fate by activation of caspase-dependent cell death [28, 29]. Caspases mediate a majority of programmed cell death, or apoptosis, via formation of the apoptosome [30]. The initiator caspase, Dronc is activated by upstream pro-apoptotic genes like *reaper* (*rpr*), *head involution defective* (*hid*), and *grim aka RHG* [31–33]. RHG also negatively regulates Inhibitor of apoptosis proteins (IAPs) [34, 35]. Caspase-dependent cell death is triggered by activation of Dronc that further activates the executioner caspase Drice, and leads to apoptosis characterized by cell shrinkage, nuclear fragmentation, chromatin condensation and DNA fragmentation [36].

Loss of M1BP is linked to reduced chromatin accessibility, impaired cellular proliferation, and death [24, 37, 38]. Autophagy is a distinct form of cell death responsible for the elimination of damaged or excess cytoplasmic content by a catabolic lysosome mediated degradation system [39]. Autophagy is marked by phagophore formation, autophagosome formation, and autophagolysosomal degradation that are genetically controlled by evolutionarily conserved genes first identified in yeast [40, 41]. There are twenty highly conserved autophagy related genes (*Atg*) characterized in *Drosophila* [42]. Among them, the *Atg*8 gene family (*Atg8a* and *Atg*8b) is most widely studied, and its members serve as an excellent biomarker for autophagy [43]. Atg8a is the fly ortholog of human LC-3, an important component of autophagy machinery, is involved in phagophore formation, autophagosome formation and autophagolysosome formation, and serves as a reporter for autophagy [40]. During eye development various specialized cells-types are generated to form an organ. These specialized cell types are required for vision. During eye development, autophagy plays an important role in formation of organelle free zone (OFZ) cells to form the transparent lens. Autophagy also protects the retinal cells from external stressors such as aggregated proteins, damaged cell organelles etc. Autophagy is also required for recycling light sensitive pigments in the retina. Thus autophagy is critically involved in development and in function of retinal neurons [44]. Recent studies have linked autophagy to maintaining cellular homeostasis and development. Therefore, investigating how M1BP regulates gene expression and its targets in the developing eye may generate new insights on mediators of the cellular response to homeostatic stress during organogenesis.

In this context, we conducted a genetic modifier screen to identify interactors of M1BP. We tested transgenes expressing members of evolutionarily conserved signaling pathways in a M1BP loss-of-function background (Fig.1). Depletion of M1BP results in a reduced eye phenotype. In this screen, we identified the JNK pathway as a key mediator of M1BP function in maintaining cellular homeostasis. JNK, a member of the conserved MAP kinase super-family, is involved in cell proliferation and cell survival and is activated through a phosphorylation cascade via MAP kinases [45–48]. In *Drosophila*, JNK signaling is activated by binding of the Tumor necrosis factor (TNF) Eiger (Egr) to its receptors Wengen (Wgn)

and Grindelwald (Grnd), and through a conserved signaling cascade of kinases, which include Tak 1 (TGFβ activating kinase 1, a JNK kinase kinase JNKKK), *hemipterous (hep)* (JNK kinase), *basket (bsk)* (Jun kinase) and the effector *jun*. Activation of the JNK pathway results in triggering its transcriptional target *puckered (puc)* expression. Puc, a dual phosphatase, participates in a negative feedback loop to downregulate JNK activity [47, 48]. Current literature suggests that the JNK pathway acts as a hub for activating multiple modes of cell death including necroptosis, ferroptosis, pyroptosis, and autophagy [49].

Here we demonstrate that M1BP suppresses the JNK signaling pathway in *Drosophila* eye. Downregulation of M1BP results in a reduced eye to a "no-eye" phenotype. This phenotype is worsened upon activation of JNK signaling in this background that dramatically increases the frequency of the "no-eye" phenotype in progeny. Furthermore, modulation of JNK when M1BP levels are downregulated using *M1BP^RNAi* results in modulation of *wg*, induction of cell death, and autophagy. In addition, both JNK- and Wg- signaling are in a feedback loop to trigger cell death and autophagy when M1BP levels are reduced. Thus, M1BP suppresses both Wg and JNK signaling pathways. Our results demonstrate that during development, M1BP mediated transcriptional pausing mechanism regulates cellular response to homeostatic stress by regulating multiple cell death mechanisms like apoptosis and autophagy.

Results

Search for genetic modifiers of M1BP's reduced eye phenotype

The *Drosophila* adult eye is composed of 600–800 highly organized ommatidia or unit eyes arranged in a hexagonal lattice [50]. These develop from the larval eye-antennal imaginal disc [51]. In this study, the *ey*-Gal4 driver was used for targeted misexpression of our genes of interest in the developing eye as shown by UAS-GFP (*ey>GFP*, green) reporter expression (Fig. 1B). Targeted misexpression of the UAS-*M1BP^RNAi* transgene using *ey*-Gal4 driver *(ey>M1BP^RNAi)* shows a strong eye suppression phenotype of reduced-eye (Fig. 1A). This reduced eye phenotype is evident from pan-neuronal marker Elav staining that marks the nuclei of the retinal neuron. We employed a forward genetic screen to look for genetic modifiers of *ey>M1BP^RNAi* reduced eye phenotype to identify targets of M1BP mediated transcriptional regulation (Fig. 1A). Prior to initiating the screen, we addressed the issue, if *ey*-Gal4 driver has a diluting effect when we increased the number of UAS-constructs. We compared *ey-Gal4>UAS-M1BP^RNAi*, with *ey-Gal4>UAS-M1BP^RNAi*+ UAS-GFP and *ey-Gal4>UAS-M1BP^RNAi* +UAS-GFP (2 copies). We found that *ey> M1BP^RNAi* along with one or two more copies of UAS-GFP constructs exhibit same frequency of reduced eye phenotype (Supplementary Fig. 1).

Using a candidate gene approach, we individually misexpressed various inducible transgenes (UAS-X) of evolutionarily conserved signaling pathways in the *ey>M1BP^RNAi* background. We screened for candidates whose misexpression resulted in either worsening (enhancers) or rescue (suppressors) of the *ey>M1BP^RNAi* reduced eye phenotype (Fig. 1A). We identified *hemipterous (hep)* and *basket (bsk)*, members of the evolutionarily conserved JNK signaling pathways, as modifiers of *ey>M1BP^RNAi* eye phenotype. Quantification of the adult eye surface area reveals significant reduction to a "no-eye" phenotype along with reduced head

phenotype in $ey>M1BP^{RNAi}+hep^{Act}$ (n=627/916, ~69%) whereas in $ey>M1BP^{RNAi}+bsk^{DN}$ (n= 746/1092, ~68%) there is a significant rescue to near wild-type eye (Fig. 1A). In the developing larval eye disc, Wg is expressed in an antero-lateral fashion (Fig. 1C). Furthermore, downregulation of M1BP ($ey>M1BP^{RNAi}$) also results in a reduced eye field in the developing eye imaginal disc along with ectopic induction of Wg (Fig 1D). However, this eye field is significantly reduced or completely eliminated along with robust Wg induction in $ey>M1BP^{RNAi}+hep^{act}$ background (Fig 1E). Conversely, the eye field was significantly restored along with downregulation of Wg expression in $ey>M1BP^{RNAi}+bsk^{DN}$ background (Fig. 1F). These results suggest that *hep* and *bsk* are genetic modifiers of M1BP mediated reduced eye phenotype.

Downregulation of M1BP results in JNK activation in the developing eye

We tested if the JNK signaling pathway is affected by M1BP levels in developing eye. Activation of JNK signaling triggers a signaling cascade by phosphorylating Tak1, *hep* (JNK kinase kinase), *bsk* (Jun kinase) and Jun (Fig. 2A). Activation of Jun upon phosphorylation triggers *puc* expression, which participates in a negative feedback loop with Bsk to regulate the JNK pathway [47, 48]. Since M1BP is a transcription pausing factor, we first employed a real-time qPCR approach to quantify transcript levels of *bsk* (JNK) gene and found a near 7-fold increase in *bsk* mRNA levels in $ey>M1BP^{RNAi}$ discs compared to *ey*-Gal4 controls (Fig. 2B). We validated our results by comparing p-JNK levels in $ey>M1BP^{RNAi}$ and *ey*-Gal4 control eye imaginal discs using a p-JNK antibody. Since JNK is expressed ubiquitously and $ey>M1BP^{RNAi}$ exhibits a highly reduced eye field (Fig. 2F, F') compared to the *ey*-Gal4 control (Fig. 2E, E'), we calculated p-JNK protein's e signal intensity. We found a significant increase in p-JNK signal intensity in $ey>M1BP^{RNAi}$ background (Fig. 2C) compared to controls. We also tested JNK pathway activation using the western blot analysis. We found a 2.5-fold increase in p-JNK signal in the $ey>M1BP^{RNAi}$ compared to *ey*-Gal4 controls (Fig. 1D) using the Empiria Studio® Software. Tubulin, a housekeeping gene, served as an internal control. To further validate our results, we tested the levels of *puc*-lacZ reporter, which serves as the functional read-out for JNK pathway activation [48]. In wild-type eye disc, *puc* is expressed in differentiated photoreceptor neurons of eye disc (Fig. 2G, G') whereas in $ey>M1BP^{RNAi}$ background, ectopic induction of *puc*-lacZ reporter was seen in the eye disc (Fig. 2H, H'), suggesting that JNK signaling is activated in the $ey>M1BP^{RNAi}$ background.

Modulation of JNK signaling levels affects M1BP mediated eye suppression

To test if the reduced eye (n=45/669, ~7%) or a "no-eye" phenotype (n=624/669, ~93%) in $ey>M1BP^{RNAi}$ discs are due to activation of JNK signaling, the levels of JNK signaling pathway members (Fig. 2A) were modulated using genetic approaches. JNK signaling status was determined by p-JNK staining in the third-instar eye imaginal discs. Downregulation of M1BP ($ey>M1BP^{RNAi}$) results in a near complete loss of photoreceptors in the eye disc (Fig. 3C, C') and almost "no-eye" phenotype(s) in adults (n=624/669, ~93%) (Fig. 3D). This phenotype is also accompanied with a significant increase in p-JNK levels (Fig. 3C, C', U) compared to the *ey*-Gal4 control (Fig. 3A, A', U). Blocking JNK signaling in the developing eye by targeted misexpression of a dominant negative *bsk*, bsk^{DN}, ($ey>bsk^{DN}$, Fig. 3E, E', F) and *puc* ($ey>puc$, Fig. 3I, I', J) alone served as controls and were similar to the

ey-Gal4 control (wild-type) as seen in the eye imaginal discs and adult eyes (Fig. 3A, A', B). Downregulation of M1BP along with JNK signaling by using bsk^{DN} (*ey>M1BP^{RNAi}+bsk^{DN}* (n=746/1092, ~68%, Fig. 3G, G', U) and *puc* (*ey>M1BP^{RNAi}+puc* (n=307/384, ~80%, Fig. 3 K, K', V), *rescued* M1BP mediated reduced eye or "no-eye" phenotype as evident from Elav staining in the eye imaginal discs (Fig. 3G, K, U) and adult eyes (Fig. 3H, L, V). However, activation of JNK signaling and downregulation of M1BP using an activated *hep*, hep^{Act}, in *ey>M1BP^{RNAi}+ hep^{Act}* (n=627/916, ~69%, Fig. 3 O, O', P) and activated *jun* in *ey>M1BP^{RNAi}+jun^{aspv7}* (n=207/353, ~59%, Fig. 3S, S', T) increased the severity of "no-eye" phenotype in the eye imaginal discs (Fig. 3O, S, U) and adult eyes (Fig. 3P, T, V). Furthermore, there were also pharate undeveloped pupae that were observed when JNK signaling was activated (Fig. 3V).This observation was supported by a significant increase in p-JNK levels in the eye imaginal discs (Fig. 3O, S, U) when compared to levels observed in *ey>M1BP^{RNAi}* (Fig. 3C). The controls used for activation of JNK signaling pathway were jun^{aspv7} (*ey>jun^{aspv7}*, Fig. 3M, M', N) and hep^{Act} (*ey>hep^{Act}*, Fig. 3Q, Q', R), which resulted in an eye suppression phenotype as compared to the wild-type *ey*-Gal4 control (Fig. 3A, A', B). These results suggest that M1BP is involved in regulating JNK signaling pathway levels to promote cell survival during eye development. Since M1BP also regulates Wg levels, it is important to understand the role of Wg and JNK signaling in of M1BP's reduced eye phenotype.

Modulation of JNK pathway affects *wingless* in M1BP loss-of-function background

During eye development, Wg serves as a negative regulator of eye fate and blocks progression of morphogenetic furrow (MF) [52–56]. To understand what happens to *wg*, a putative target of M1BP [27], when levels of the JNK pathway are modulated, we analyzed *wg* levels in the eye disc at the transcriptional level using a lacZ reporter and translational level using an antibody against Wg protein. Both *wg*-lacZ reporter and Wg protein are expressed in an antero-lateral fashion in the developing third instar eye disc of *ey*-Gal4 control (Fig. 4A, B). Downregulation of M1BP (*ey>M1BP^{RNAi}*) results in the ectopic induction of *wg*-lacZ as well as Wg protein (Fig. 4C, D) [27]. Downregulation of the JNK pathway using bsk^{DN} (*ey>bsk^{DN}*, Fig. 4E, F) and *puc* (*ey>puc*, Fig. 4I, J) alone in the developing eye disc exhibits the *wg*-lacZ reporter- and Wg protein- expression similar to the *ey*-Gal4 control (Fig. 4A, B). When JNK signaling and M1BP levels are downregulated together in the *ey>M1BP^{RNAi}+bsk^{DN}* (Fig. 4G, H) and *ey>M1BP^{RNAi}+puc* (Fig. 4 K, L) background, it exhibits a rescue of the "no-eye" phenotype and shows a dramatic reduction in *wg-lacZ and* Wg proteins levels compared to the *ey>M1BP^{RNAi}* discs (Fig. 4C, D). Activation of JNK signaling in the entire eye by using jun^{aspv7} (*ey>jun^{aspv7}*, Fig. 4Q, S), and hep^{Act} alone (*ey>hep^{Act}*, Fig. 4M, N) results in eye suppression along with induction of *wg* as compared to *ey*-Gal4 control (Fig. 4A, B). Note, that Wg level are not significantly affected in jun^{aspv7} alone compared to *ey*-Gal4 controls due to weakness of the transgene. Activation of JNK signaling along with downregulation of M1BP (*ey>M1BP^{RNAi}+jun^{aspv7}* (Fig. 4T, U) and *ey>M1BP^{RNAi}+hep^{Act}* (Fig. 4O, P) shows an increase in *wg* at the transcriptional as well as translational level when compared to *ey>M1BP^{RNAi}* alone (Fig. 4C, D). These results clearly suggest that modulation of the JNK pathway in the developing eye modulates *wg* levels. This observation supports the presence of a feedback loop during eye development [57]. Previous studies have shown that ectopic Wg or JNK are capable of inducing cell death

during eye development [27–29]. Therefore, it is important to test if reduced eye phenotype in M1BP downregulation is due to activation of Wg and JNK signaling mediated cell death.

Downregulation of M1BP induces *hid* expression and apoptotic cell death

We tested if M1BP downregulation (*ey>M1BP^RNAi*) can trigger apoptotic cell death by using *hid5'F-WT*-GFP, a transcriptional reporter for the *hid* gene [58]. *hid* is expressed in cells undergoing apoptosis and is generally seen in a few cells in control third instar eye-antennal imaginal disc [58]. To quantify the signal, we measured the GFP fluorescence intensity in uniform areas of 100X100 pixel boxes in the *ey*-Gal4 expression domain using a region of interest (ROI) approach [59]. A basal level of GFP (green) signal was observed in the eye region of *ey*-Gal4 control discs (Fig. 5A, A', B, B', E); whereas downregulation of M1BP in the developing eye resulted in dramatic induction of *hid*-GFP (green) in the eye disc (Fig. 5C, C', D, D', E). To test if the reduced eye phenotype in *ey>M1BP^RNAi* background is due to induction of caspase-dependent cell death, we downregulated *hid* levels using *hid^RNAi* and *hid^N14*, a N-terminal deletion construct, in the *ey>M1BP^RNAi* background. When compared to downregulation of M1BP alone (*ey>M1BP^RNAi*, Fig. 3C, D), downregulation of both *hid* and M1BP (*ey>M1BP^RNAi+hid^RNAi*, n=81/317, ~25%, Fig. 5G and *ey>M1BP^RNAi+hid^N14*, n=100/343, ~29%, Fig. 5I) showed rescue of the reduced eye or "no-eye" phenotype of *ey>M1BP^RNAi* alone as seen in the eye disc and the adult eye. Furthermore, we also tested levels of Wg and JNK signaling in this background to study their role. Downregulation of M1BP levels along with blocking of cell death by using *hid^RNAi* (*ey>M1BP^RNA+hid^RNAi*, Fig. 5L, L') and *hid^N14* (*ey>M1BP^RNAi+hid^N14*, Fig. 5N, N') exhibits mild restoration of Wg expression compared to the ectopic induction of Wg in *ey>M1BP^RNAi* alone (Fig. 4D). Also, the controls *ey> hid^RNAi* (Fig. 5K, K') and *ey> hid^N14* (Fig. 5N, N') show near wild-type expression of Wg. Additionally, we wanted to see pJNK levels, a read out of JNK signaling, which is known to activate caspase -dependent and -independent cell death. We observed that downregulation of M1BP levels along with *hid* by using *hid^RNAi* (*ey>M1BP^RNA+hid^RNAi*, Fig. 5P, P', S) and *hid^N14* (*ey>M1BP^RNAi+hid^N14*, Fig. 5R, R', S) resulted in a significant reduction in pJNK levels compared to M1BP (*ey>M1BP^RNAi*) alone (Fig. 2C, 5S). Although pJNK levels were slightly higher in *ey>M1BP^RNAi* than *ey*-Gal4 control (Fig. 2C, 5S). Other controls like *ey> hid^RNAi* (Fig. 5P, P') and *ey>hid^N14* (Fig. 5Q, Q') show near wild-type expression of pJNK. Thus, blocking cell death alone is not sufficient for rescuing M1BP mediated eye suppression phenotypes (Fig. 5J). Therefore, we wanted to assay other mechanisms of death.

Downregulation of M1BP triggers autophagy

To test if autophagy is responsible for the reduced eye phenotype of *ey>M1BP^RNAi*, we first assessed the levels of *Atg8a-mCherry*, a reporter for autophagic cell death, in the third instar eye discs. Atg8a is an important component of autophagy machinery and is involved during all stages of the process like in phagophore formation, autophagosome formation and autophagolysosome formation. Therefore, Atg8a is a widely used as a marker to assay autophagy [43, 60, 61]. We assayed *Atg8a* as an indicator of the entire process of autophagy and also downregulated *Atg8a* to block autophagy. Downregulation of M1BP (Fig. 6C, C') in the entire eye results in a nearly a two-fold increase in *Atg8a-mCherry* levels as compared to *ey*-Gal4 control discs (Fig. 6B, B', D). These observations suggest that M1BP may

also be involved in suppressing autophagy during eye development. Therefore, we further validated these results using real-time quantitative PCR and semi-quantitative western blot. Downregulation of M1BP (*ey>M1BP^{RNAi}*) resulted in a significant upregulation of *Atg8a* transcripts when compared to the *ey*-Gal4 control discs (Fig. 6E). Furthermore, western blot revealed a significantly higher Atg8a protein levels in *ey>M1BP^{RNAi}* when compared to *ey*-Gal4 (Fig. 6F). We tested if blocking autophagy is sufficient to rescue M1BP mediated eye suppression. In *Atg8a^1* mutant's heterozygous background, eye development is near wild-type as observed in the adult eye (Fig. 6A, G) and eye imaginal disc (Fig. 6H). However, in the *ey>M1BP^{RNAi}* background (*ey>M1BP^{RNAi}+Atg8a^{1−/+}*, n=126/470, ~27%, Fig. 6J) a significant rescue of the "no-eye" or reduced eye phenotype of *ey>M1BP^{RNAi}* alone to near wild-type eye is seen in nearly 27% of the flies (Fig. 6J, K). Heterozygous *Atg8a^{1−/+}* mutants show normal Wg expression in the antero-lateral margins (Fig. 6F, F'). Reducing M1BP levels in *Atg8a^1* mutant's heterozygous background (*ey>M1BP^{RNAi}+ Atg8a^{1−/+}*, Fig. 6K, K') reveals restoration of Wg expression to normal levels in the eye discs when compared to *ey>M1BP^{RNAi}* alone. pJNK levels were studied in these eye discs and signal intensity was quantified. Interestingly, downregulation of M1BP in *Atg8a^1* mutant's heterozygous background (*ey>M1BP^{RNAi}+ Atg8a^{1−/+}*, Fig. 6L, L', N) shows pJNK levels similar to that of *ey>M1BP^{RNAi}*. These results suggest that JNK signaling is upstream of *Atg8a* and regulates autophagy.

Downregulation of M1BP exhibits JNK signaling mediated activation of autophagy

JNK signaling is emerging as a node for multiple modes of cell death [49]. We tested if activation of autophagy is mediated through JNK signaling. To do this, we assayed the levels of an autophagy marker while modulating the JNK pathway in a background where M1BP was downregulated. Downregulation of M1BP in entire eye disc (*ey>M1BP^{RNAi}*, Fig. 7B, B', K) ectopically enhances *Atg8a-mCherry* expression in the eye disc when compared to *ey*-Gal4 control (Fig. 7A, A', K). In the developing eye disc, downregulation of both M1BP and JNK signaling together by using *bsk^{DN}* (*ey>M1BP^{RNAi}+bsk^{DN}*, Fig. 7D, D', K) and *puc* (*ey>M1BP^{RNAi}+puc,* Fig. 7 F, F', K), result in a significant rescue in the number of Elav positive cells; however, the *Atg8a* levels are reduced compared to *ey>M1BP^{RNAi}* (Fig. 7A, A', K). The modulation of controls like *bsk^{DN}* (*ey>bsk^{DN}*, Fig. 7C, C') and *puc* (*ey>puc*, Fig. 7E, E') alone in the entire developing eye show minimal or highly reduced *Atg8a-mCherry* expression compared to *ey*-Gal4 control (Fig. 7A, A'). Activation of JNK signaling in the entire eye disc by using *jun^{aspv7}* (*ey>jun^{aspv7}*, Fig. 7I, I') or *hep^{Act}* (ey>*hep^{Act}*, Fig. 7G, G', K) showed increased *Atg8a-mCherry* expression compared to *ey*-Gal4 control (Fig. 7A, A', K). When we activated JNK signaling along with downregulating M1BP in *ey>M1BP^{RNAi}+hep^{Act}* (Fig. 7H, H', K) and (*ey>M1BP^{RNAi}+jun^{aspv7}* (Fig. 7J, J', K), we see a significant increase in *Atg8a-mCherry* expression in the eye discs compared to *ey>M1BP^{RNAi}* alone (Fig. 7B, B', K). Thus, during eye development M1BP is required to suppress JNK signaling and in turn suppresses autophagy. Interestingly, blocking caspase-dependent (apoptosis) alone or caspase-independent (autophagy) alone rescued the "no-eye" phenotype to near wild-type eyes in nearly 25% of the flies but not all flies. Therefore, we tested if blocking both cell death and autophagy along with M1BP downregulation can exhibit a stronger rescue.

M1BP blocks both caspase-dependent and caspase-independent cell death

To test this hypothesis, we genetically misexpressed P35 and reduced *Atg8a* levels to half using a heterozygous combination along with downregulation of M1BP function using M1BP^{RNAi} (*Atg8$^{1-/+}$*; *ey> M1BP^{RNAi} + P35*) in the same fly. As expected, we observed a very strong rescue to near wild-type adult eye (Fig. 8C) compared to the "no-eye" phenotype of *ey>M1BP^{RNAi}* (Fig. 8B) alone flies. Furthermore, this rescue was observed in ~ 85% (n=308/363) of the flies (Fig. 8D). We wanted to see how JNK signaling may play a role in this rescue. We stained for pJNK and quantified its intensity. Our results clearly showed an increase in pJNK expression when M1BP is downregulated (Fig. 8I, I', K). Additionally, our results showed that blocking apoptosis and autophagy alone in *ey>M1BP^{RNAi}* background did not rescue JNK pathway activation (Fig. 8K). Interestingly, upon blocking both apoptosis and autophagy, we did not see a decrease in JNK signaling levels to that of wild-type (Fig. 8J, J', K). We also checked the expression of Wg, a negative regulator of eye development that is normally expressed in the antero-lateral margins shown in *ey*-Gal4 control (Fig. 8E, E'). We see a significant restoration of Wg expression levels when both caspase -dependent (apoptosis) and -independent (autophagy) are blocked in *ey>M1BP^{RNAi}* background (Fig. 8G, G'). This data suggests that transcriptional pausing mechanism mediated through M1BP regulates both apoptosis and autophagy during development (Fig. 8L).

Discussion

During organogenesis, cell death occurs at basal levels to maintain cellular homeostasis. Multicellular organisms typically produce excess cells that are eliminated to maintain terminally differentiated cells and organ size during morphogenesis. This strictly regulated cell death is referred to as "Programmed Cell Death" (PCD) [10], and requires appropriate signals due to activation of molecular programs. Generally, dying cell(s) are replaced by new cell-type(s) whereas in other cases they are replaced by brand new cell(s) of the same type.

In the developing *Drosophila* eye, cellular homeostasis plays an important role in organogenesis. Most cell death utilizes caspase-dependent cell death, also called as apoptosis. Autophagy, a lysosome mediated self-degradation process observed in eukaryotic cells, is contextual and eliminates unfit cells to ensure macromolecule turnover and maintains cellular homeostasis. It can function as a prodeath or pro-survival process depending on context dependent manner. Autophagy delivers cytoplasmic materials to the lysosome for degradation [11]. The genetic machinery involved in autophagy is highly conserved. In *Drosophila*, a recent study has demonstrated that several autophagy genes are active in the developing eye[62]. Loss of just one does not induce a severe phenotype while loss of multiple genes shows a spectrum of reduced eye phenotypes [62]. Interestingly, autophagy (cellular "self-eating") often accompanies apoptosis [13]. Therefore, both apoptosis and autophagy are required to remove excess cells to regulate the cell number of terminally differentiated cells and form the highly organized adult compound eye [62–65].These cellular processes are also crucial for organ sculpting and morphogenesis during development [7, 8]. There are also genetic evidences for autophagy-dependent cell death in mammalian cells that are unable to undergo apoptosis[12]. This suggests that autophagy, which often accompanies

apoptosis, can compensate for apoptosis when apoptosis is blocked or inhibited. During embryonic development, removal of interdigital web cells is delayed in apoptosis-deficient Bax/Bak KO mice, which gets further delayed in Atg5/ Bax/ Bak triple KO mouse [12]. Thus, a compensatory backup mechanism involving apoptosis and autophagy ensures that unwanted cells get deleted by non-apoptotic (autophagy) means.

Aberrant activation of several signaling pathways and extrinsic stress signals are known to regulate these processes by triggering developmental cell death [28, 29, 41, 66, 67]. Since these two processes often accompany each other but are regulated by their distinct genetic machinery, it is important to understand how these two processes are regulated during development.

We found that Pol II pausing, which dynamically regulates gene expression of 10–40% of all genes in *Drosophila* [23, 68], plays an important role in regulating apoptosis and autophagy. Thus, the promoter-proximal stalling of developmentally important genes plays a crucial function in development [69]. M1BP is expressed ubiquitously in all cells. It binds to Motif1 sequences on target genes and subsequently recruits other factors that regulate transcription negatively. This negative regulation is facilitated by association with negative elongation factor (NELF) and potentially by the presence of 1+ nucleosomes downstream of target genes [24]. In gene ontology, M1BP regulated genes for basic biological processes such as cell cycle, metabolism, cell viability and the cytoskeleton are overrepresented [15, 24]. Previously, we have demonstrated that M1BP downregulates *wg* by a transcriptional pausing mechanism that blocks apoptosis in the developing *Drosophila* eye [27]. However, the underlying molecular genetic mechanism by which M1BP regulates cellular homeostasis during development is not clear. We therefore studied how this transcriptional pausing factor can contribute at a molecular level to regulate cellular homeostasis. The premise is based on our observation that blocking caspase-dependent cell death (apoptosis) did not fully rescue M1BP's downregulation phenotype of "no-eye" or reduced eye. To further dissect the M1BP mediated eye suppression function, we performed a forward genetic screen to look for genetic modifiers of M1BP downregulation phenotype of "no-eye" or reduced eye. We identified *hemipterous (hep)* and *basket (bsk),* the two members of JNK signaling pathway in the screen (Fig. 1). To validate the activation of JNK pathway in M1BP loss-of-function background, we investigated three functional readouts of the JNK pathway - (i) the transcriptional status of *jnk* using qRT PCR, (ii) *puc*-lacZ reporter expression, and (iii) pJNK protein levels using an immunohistochemical approach (Fig. 2). Our qPCR data showed a 7-fold increase in *jnk* mRNA levels. Additionally, *puc*-lacZ reporter expression and pJNK protein levels were significantly upregulated when M1BP is downregulated in the developing eye (Fig. 2). Puc, a dual phosphatase is the effector of JNK signaling and reflects activation status of the pathway [48, 67, 70]. Downregulation of M1BP clearly activates JNK signaling as seen via evidences mentioned above. These clearly suggest that a transcriptional pausing mechanism can regulate JNK itself and the activation status of the signaling pathway. Furthermore, we performed MEME analysis[20, 71, 72], a bioinformatics approach to screen for potential M1BP binding sites (YGGTCACACTR) in JNK pathway members. There are five binding sites on *puc* indicating that M1BP can bind at these sites to suppress *puc* expression by transcriptional pausing (Supplementary Fig. 2).

We found that M1BP can repress JNK signaling in the developing eye (Fig. 2). Additionally, modulation of JNK signaling levels can affect M1BP loss-of-function phenotype. Thus, the transcriptional pausing function of M1BP suppresses JNK signaling to promote eye development (Fig. 3). Recently, *wg* has been reported as a putative target of M1BP with Motif1 sequences before the transcription start site (TSS) of *wg* [27]. We used *wg*-lacZ reporter expression and Wg protein expression patterns that respectively corresponds to the transcriptional status and translational levels. We found that *wg* responds to the modulation of JNK signaling in the developing *Drosophila* eye (Fig. 4, 8). Furthermore, alteration of JNK signaling in M1BP loss-of-function background affects *wg* expression, which is suggestive of the existence of feedback loop during eye development [57].

The evolutionarily conserved JNK signaling cascade, which can be triggered by various signals [70], is a key regulator of many cellular processes including proliferation, differentiation, and cell death [49]. Several studies have shown that JNK is a hub for activating multiple modes of cell death including apoptosis as well as autophagy during development. Modulation of JNK signaling has been linked to organ size control of the wing, leg and antenna in *Drosophila* [73]. Furthermore, our data reveals that JNK signaling is ectopically induced when M1BP is downregulated (Fig. 2) and modulation of JNK result in modulation of the eye phenotype (Fig. 3). Thus, we assayed two important cell death pathways such as apoptosis and autophagy that are activated by the JNK pathway. While, the role of JNK is widely studied and known in activation of apoptosis and autophagy, here, we wanted to focus on its potential role in transcriptional pausing mechanism in regulating these two processes during development. The rationale was that M1BP can regulate JNK signaling. We found that M1BP's downregulation phenotype of reduced eye or "no-eye" is due to induction of cell death as evident from the ectopic induction of apoptotic marker *hid*-GFP (Fig. 5). Surprisingly, not only caspase-dependent apoptosis alone but also caspase-independent autophagy is involved as evident from ectopic induction of autophagic marker *Atg8a-mCherry* (Fig. 6). We also validated the activation of autophagy in *ey>M1BP^{RNAi}* background by detecting autophagosome-lysosome flux using tandem mCherry-GFP-Atg8a reporter. mCherry- and GFP marks the presence of autophagosomes, and GFP is used to monitor flux as it will get quenched in the acidic environment of the lysosome. Whereas mCherry, unaffected by the acid remains active until the protein is degraded. Downregulation of M1BP resulted in reduced GFP in acidic environment indicating autophagolysosome activation (Supplementary Fig. 3). We found that blocking either apoptosis (Fig. 5) or autophagy (Fig. 6) alone in *ey>M1BP^{RNAi}* background rescued the "no-eye" phenotype in nearly 30% and 20% flies respectively. However, no complete rescue was observed in either scenario and frequency remained low. This led us to hypothesize that during development M1BP mediated transcriptional pausing of JNK regulates both caspase-dependent cell death and autophagy. To validate this hypothesis, we checked the levels of JNK pathway readout using pJNK levels. We observed pJNK levels similar to that of *ey>M1BP^{RNAi}* when we downregulated either the levels of autophagy (*ey>M1BP^{RNAi}+Atg8a^{+/−}*) or apoptosis (*ey>M1BP^{RNAi}+hid^{RNAi}* or *ey>M1BP^{RNAi}+hid^{N14}*). Since JNK signaling levels were not affected by modulation of either apoptosis or autophagy in *ey>M1BP^{RNAi}* background suggests that JNK is upstream of both apoptosis as well as autophagy. Based on these results, we propose a model

showing that M1BP regulates cell survival by suppressing JNK pathway levels which in turn triggers apoptosis and autophagy (Fig. 8L). Therefore, the two distinct and independent cell homeostasis mechanisms are being commonly regulated by JNK signaling and can explain the compensatory relation between apoptosis and autophagy during development.

We also wanted to understand if these two cell death processes could be controlled simultaneously. Therefore, we blocked both caspase-dependent apoptosis and caspase-independent autophagy in a background where M1BP levels have been downregulated and found a significant rescue to near wild-type adult eye phenotypes in ~ 85% of flies (Fig. 8B, C, D). Therefore, it is possible that a transcriptional pausing mechanism mediated through M1BP is required to suppress both apoptosis and autophagy by regulating the expression of both *wg* and JNK signaling in the developing *Drosophila* eye. There are several studies that demonstrate how apoptosis and autophagy are independently regulated[10, 11, 49]. In this study, we show for the first time how a common transcription pausing mechanism is regulating signaling components that can in turn regulate cellular homeostasis to suppress both autophagy and apoptosis. Also, we have provided multiple evidences for the activation of autophagy when M1BP is downregulated (Figure 6). We have also shown the link of M1BP downregulation with JNK modulation (Figure 7).

In order to understand if M1BP mediated regulation of cellular homeostasis through regulation of apoptosis and autophagy is seen in other tissues, we assayed the markers for cell death, autophagy, and JNK signaling in M1BP loss-of-function background in fly eye- and wing imaginal disc (Supplementary Fig. 4). Downregulation of M1BP in *bi*-Gal4 expression domain, which marks the pouch region of wing (*bi>M1BP^{RNAi}*), results in ectopic pJNK, Dcp1, and *Atg8a-mCherry* expression in the pouch region of the wing imaginal disc. This finding suggests that the transcription pausing function of M1BP may have similar target(s) in the eye and wing imaginal discs.

In conclusion, here we describe the regulation of cellular homeostasis by a transcription pausing factor M1BP during development. Furthermore, the evidence for *wg* and JNK as targets of M1BP mediated regulation of cellular homeostasis is clear from our results, however, there are still flies that show the M1BP loss-of-function phenotype of reduced eye when both these processes are suppressed. It is based on the fact that both the extent of rescue as well as frequency of rescue phenotypes among the flies where both autophagy and apoptosis are blocked in *ey>M1BP^{RNAi}* background, was significantly higher than when autophagy or apoptosis was blocked independently. However, the rescue was not complete. Therefore, it is possible that M1BP might be playing a role in modulating other mechanism(s) of cellular homeostasis. It is known that there are many other types of cellular homeostasis mechanisms. Since the current study has restricted only to the two common mechanisms, this warrants further research to explore the regulation achieved by transcriptional pausing mechanisms during development.

Several studies suggest that the regulatory mechanism in Pol II pausing during fly development are also conserved in higher organisms. Regulation of transcriptional elongation by positive or negative factors have been shown to have a role in maintaining cellular homeostasis [74]. Since *Drosophila* genes are highly conserved in humans, results

from this study can be extrapolated to the maintenance of cellular homeostasis in higher organisms. Additionally, JNK signaling and *wg* are known to play an important role in apoptosis and autophagy during development. While there are several signaling nodes that can trigger these cell death pathways, here we show that M1BP regulates both these processes in a JNK-dependent manner during development. Therefore, it is crucial to add to the existing research about how M1BP, a ubiquitously expressed transcriptional pausing factor, can regulate other signaling pathways to achieve optimum cellular homeostasis by suppressing autophagy. During normal eye development, autophagy is required to drive tissue heterogeneity and determine their structure. For example, in mammals, autophagy is required to produce the organelle free zone (OFZ) cells to form the transparent lens. Dysregulated autophagy has been associated with several ocular diseases like cataracts, corneal dystrophy, Glaucoma, diabetic retinopathy, age-related macular degeneration etc. [44, 75]. Therefore, regulation of different types of cell death seems to be a conserved mechanism used by organisms.

Materials and methods

Fly stocks

Fly stocks used in this study are described in Flybase (http://flybase.bio.indiana.edu). We used Canton-S as wild-type, *ey*-Gal4 [76], and UAS-*M1BP*RNAi (BL41937) [24, 27]. Other stocks used include UAS-*jun*aspv7 [77], an activated form of *Drosophila jun*, UAS-*hep*Act [78], an activated form of *hep*, UAS-*bsk*DN, a dominant negative form of *bsk* [79], UAS-*puc* [48] a negative regulator of *bsk*, *wg*-lacZ [76], *puc*E69 [48], UAS-*wg*RNAi (BL31249, BL33902) [80], UAS-*P35* [81], *hid 5'F-WT* GFP [58], UAS-*hid*RNAi (hidGD1673) (VDRC v8269) [82], UAS-*hid* N14 [83], a N-terminal deletion construct lacks the sequences encoding the N-terminal DIAP1 binding domain that is required for proper *hid* function [83–85]. Other stocks used were UAS-*mCherry-Atg8a*, and *Atg8a*1 mutant [86]. Fly stocks were maintained at 25°C on regular cornmeal, yeast, molasses food medium.

Genetic crosses

The Gal4/UAS system was used to misexpress the gene of interest in a spatiotemporal manner [87]. All crosses were maintained at 18°C and 25°C, unless specified, to check different induction levels [88]. We have used the *ey*-Gal4 driver in this study to target misexpression of transgene(s) in the entire developing eye domain of larval eye-antennal imaginal disc.

Immunohistochemistry

Eye-antennal discs of wandering third instar larvae were dissected in 1X phosphate buffered saline (PBS), fixed in 4% paraformaldehyde in PBS (fixative) for 20 minutes and washed in PBST (three times). The tissues were stained using a combination of antibodies following the standard protocol [89, 90]. Primary antibodies used were mouse anti-β-GAL (1:100; Developmental Studies Hybridoma Bank, DSHB); rabbit anti-pJNK (1:250; Cell Signaling Technologies, CST), rat anti-Elav (1:100), mouse anti-Wg (1:100; DSHB), mouse anti-Dlg (1:100, DSHB). Discs were washed in PBST thrice for 10 minutes each. Secondary antibodies used were donkey anti-rat IgG conjugated to Cy5 (1:250), donkey anti-rabbit

IgG conjugated to Cy3 (1:300) or goat anti-mouse IgG conjugated to FITC (1:200) (Jackson Laboratories). Discs were mounted in Vectashield and photo-documented on a Fluoview 3000 Laser Scanning Confocal Microscope [57, 91]. We took the images at 20X and 40X magnification unless stated otherwise. We analyzed and prepared the final figures using Adobe Photoshop CS6 software.

Quantification of intensity using Fiji/ ImageJ software

Fiji/ ImageJ software was used for quantification of intensity from images [59, 92]. The split channel function was used to select the appropriate channel for analysis. ROI tool was used to select regions of a uniform area 100X100pixels boxes in the eye disc and measurements were obtained. Graphs were plotted using the mean intensity per unit area (μm^2). The values were analyzed and plotted as mean +/− Standard Error of the Mean (SEM) using GraphPad Prism. For each genotype, 5 images were used to perform quantification.

Adult eye imaging

For capturing adult eye images, flies were first frozen at −20°C for ~4 hours. The flies were then mounted onto needles after removing the wings and legs and imaged at 10X magnification. The needle with the fly was embedded in a clay putty on a glass slide to stably position it to get the lateral view of the fly eye/head. Images were taken on a MrC5 color camera mounted on an Axioimager.Z1 Zeiss Apotome using a Z-sectioning function of Axiovision software 4.6.3 [27, 66, 93–95]. The final images were prepared using Adobe Photoshop CS6 software.

Frequency of eye phenotype

Three independent sets of two hundred flies were screened (200 X 3= 600) and the frequency of eye phenotype(s) were calculated for each genetic cross. The eye phenotypes were categorized as no-eye, reduced-eye, or normal eye. Analyses were performed and graphs were plotted in GraphPad Prism.

Quantitative analyses of area of the eye

Image J software was used to quantify the area (μm^2). The region of interest (ROI) was drawn along the perimeter of the adult eye and the surface area of the eye was measured and the graph was plotted in GraphPad Prism.

Real time quantitative PCR

60–80 eye-antennal imaginal discs were used for RNA isolation [96, 97]. Tissue was collected and homogenized in TRIzol reagent (Invitrogen, Cat# 15596026). Total RNA in the aqueous phase was transferred to RNA Clean & Concentrator-5 (Zymo research, Cat# R1013) columns and eluted in about 15–20 μL DNase/RNase-free water. Quality and quantity of RNA was determined by Nanodrop 2000 spectrophotometer (Thermo Scientific). The cDNA was synthesized from 250 ng of total mRNA through Reverse Transcription reaction using first-strand cDNA synthesis kit (GE healthcare, Cat# 27926101). Real Time-qPCR was performed using BioRad iQ SYBR Green Supermix (Bio-Rad, Cat# 1708860) according

to the standard protocol [96, 97]. Fold change was calculated using comparative CT method ($2^{-\Delta\Delta CT}$ method). The primers used were:

GAPDHFw5'-GGCGGATAAAGTAAATGTGTGC-3'

GAPDHRev5'-AGCTCCTCGTAGACGAACAT-3'

JnkFw5'-CCAACCGTCCGAAACTATGT-3'

JnkRev5'-CCGGCGGCTATTCTGATTATTA-3'

Atg8aFw5'- GGTCAGTTCTACTTCCTCATTCGC-3' [62]

Atg8aRev5'- ATAGTCCTCGTGATGTTCCTG-3' [62]

Western blotting

Entire adult fly heads were used for sample preparation from *ey*-Gal4 and *ey>M1BP^RNAi* flies. Protein was extracted in 2X lamellae buffer in the presence of phosphatase band protease inhibitor using a standardized protocol [57, 98]. The protein samples were resolved in a 10% gel and transferred onto a nitrocellulose membrane. The membrane was washed with 1X TBST, blocked in 5% w/v BSA in 1X TBST and incubated overnight at 4°C with primary antibody rabbit Phospho-SAPK/JNK (1:1000) (Cell Signaling Thr183/Tyr185) (81E11) antibody, rabbit anti-Atg8a (1:10000) (Millipore Sigma, Cat. # ABC974) and mouse anti-α-Tubulin antibody (1:12000) (SIGMA, Cat. # T5168) diluted in 5% w/v BSA in 1X TBST. After three washes with 1X PBST, the blot was incubated for 1hour in secondary antibody: horseradish peroxidase conjugated goat anti-rabbit IgG-HRP (1:5000) and goat anti-mouse IgG-HRP (1:5000) (Santa Cruz Biotechnology, Cat. # Sc-2005). Signal was detected using Horse Radish Peroxidase (HRP) conjugated goat anti–rabbit IgG using chemiluminescence substrate (LICOR). Images were captured using the LICOR Odyssey XF Imaging System. Empiria Studio software was used to quantify and normalize the signal. Relative pJNK levels were then statistically quantified and normalized by using LI-COR Image Studio lite 5.2 software and graph was plotted in GraphPad Prism.

Statistics

Statistical analysis was performed using GraphPad Prism. For all eye frequency data, multiple genotypes with subgroups were compared with each other using 2-way ANOVA. Statistical comparisons with *ey*Gal4 and *ey>M1BP^RNAi* were depicted using # and * respectively. For mean integrated intensity quantification data, we have used one-way ANOVA to compare >2 groups. Comparison between two groups was performed using Mann-Whitney test. Graphs were plotted as mean +/− SEM with the individual values. The error bars represent Standard error from mean (SEM). Statistical significance is set at 95% CI and is shown by p-value: **** $P<0.0001$; *** $P<0.001$; ** $P<0.01$; * $P<0.05$.

Supplementary Material

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Acknowledgments

Abbreviations

hid	head involution defective
rpr	reaper
DIAP	Drosophila inhibitor of apoptosis proteins
RD genes	Retinal Determination genes
MF	Morphogenetic Furrow
PCD	Programmed cell death
ER	Endoplasmic reticulum
Wg	Wingless
JNK	c-Jun amino-terminal (NH2) Kinase
MAPKs	Mitogen-activated protein kinases
TNF	Tumor necrosis factor
Egr	Eiger
Wgn	Wengen
Tak 1	TGFβ activating kinase 1
JNKKK	JNK kinase kinase
Hep	Hemipterous
JNKK	JNK kinase
Bsk	Basket
Puc	Puckered
Pol II	RNA polymerase II
DNA	Deoxyribonucleic acid
TSS	Transcription start site

ESCs	Embryonic stem cells
Hop	Hopscotch
Dpp	Decapentaplegic
GAF	GAGA factor
M1BP	Motif 1 Binding Protein
ZAD	Zinc-associated domain
ZKSCAN3	Zinc finger with a SCAN and a KRAB domain 3 (ZKSCAN3)
UAS	Upstream activation sequence
ELAV	Embryonic lethal abnormal vision
RT-qPCR	Real time quantitative polymerase chain reaction
LOF	Loss of function
GOF	Gain of function
GFP	Green fluorescent protein
WT	Wild-type
PBS	Phosphate buffered saline
PFA	Paraformaldehyde
PBST	Phosphate Buffered Saline with Triton X-100
β-GAL	Beta-galactosidase
DSHB	Developmental Studies Hybridoma Bank
CST	Cell Signaling Technologies
Dlg	Discs large
IgG	Immunoglobulin G
FITC	Fluorescein isothiocyanate
RNA	Ribonucleic acid
cDNA	Complementary DNA
Ct	cycle threshold
GAPDH	glyceraldehyde-3-phosphate dehydrogenase
SAPK	Stress-activated protein kinases
HRP	Horse Radish Peroxidase

ROI	Region Of Interest
SEM	Standard Error of the Mean
CI	Confidence Intervals

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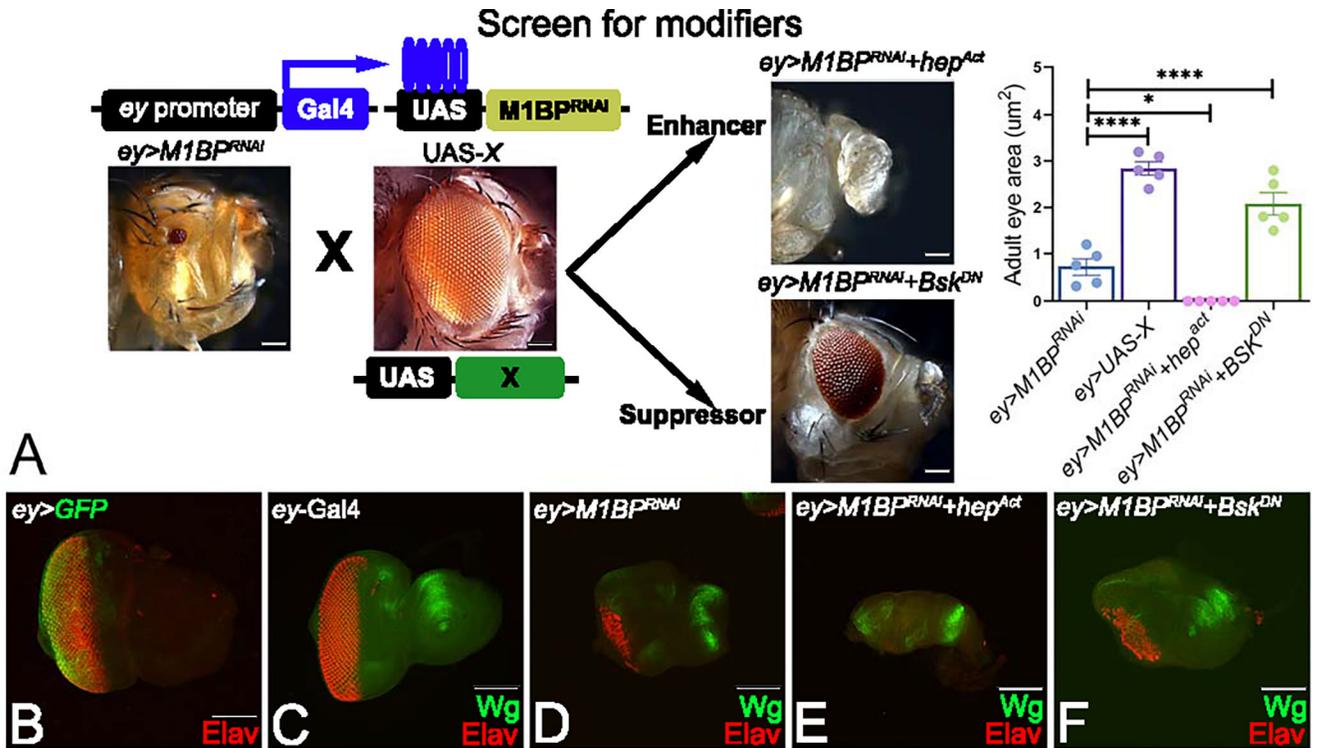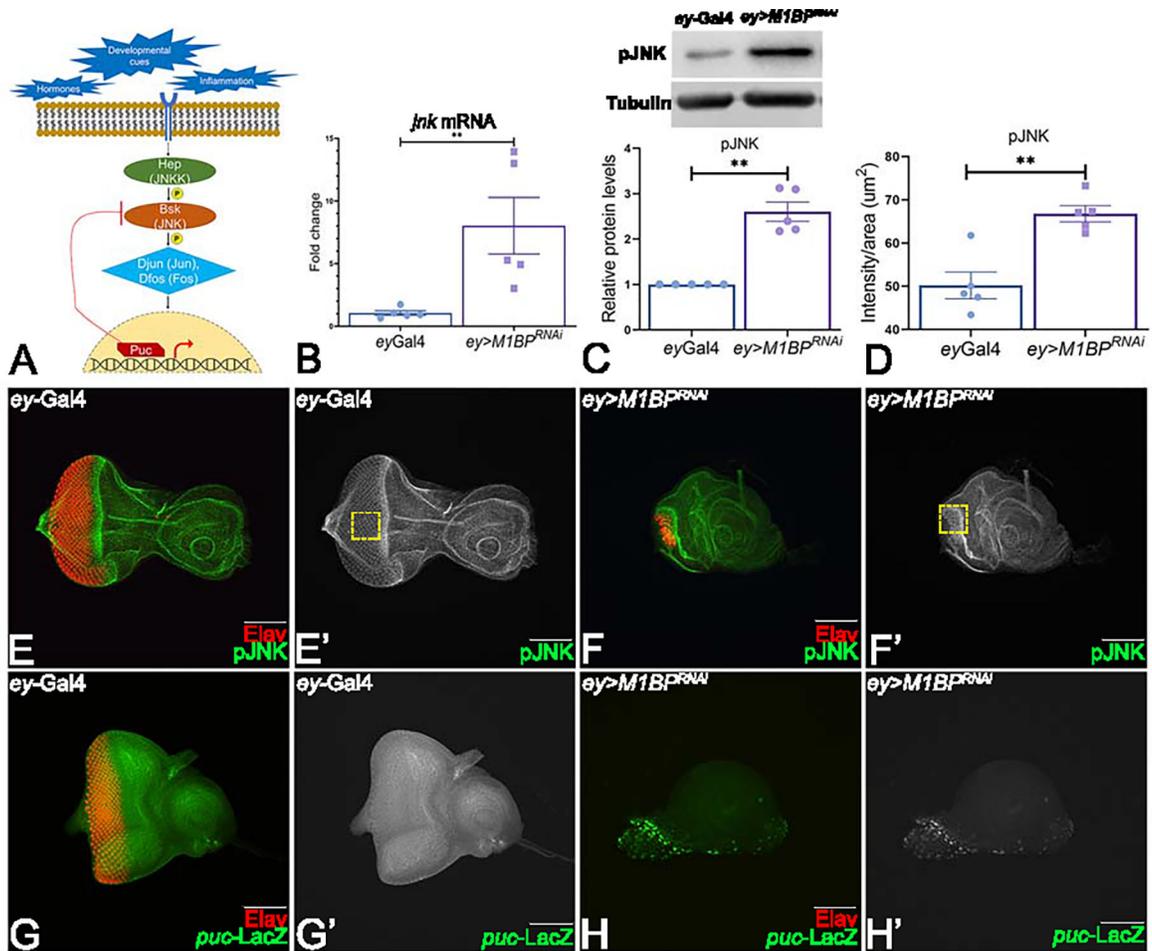
Figure 1: Forward genetic screen to identify modifiers of M1BP mediated eye suppression.
(A) Schematics of forward genetic screen for identifying genetic modifiers of M1BP mediated eye suppression phenotype where various UAS-X/transgene lines were individually misexpressed using the *ey*-Gal4 driver along with downregulation of M1BP. Modifiers were classified into enhancers and suppressors based on whether they enhance or suppress the *ey>M1BP^{RNAi}* reduced eye phenotype. Quantification of adult eye surface area (μm^2) using Fiji/ImageJ software (NIH) to assay differences in eye phenotype. (B) *ey>GFP* where the GFP transgene reporter marks the *ey*-Gal4 driver expression domain in the developing third instar eye antennal disc. Note that the eye antennal imaginal disc is stained with pan-neuronal marker Elav (red) that marks the nuclei of retinal neurons. (C, D, E, F) Eye antennal imaginal discs stained for Wg (green) and pan-neuronal marker Elav (red). (D) Downregulation of M1BP in the entire eye (*ey>M1BP^{RNAi}*) results in suppression of eye fate and increased Wg expression compared to (C) *ey*-Gal4 control. Eye discs showing (E) enhancement (*ey>M1BP^{RNAi}+hep^{Act}*) and (F) suppression (*ey>M1BP^{RNAi}+bsk^{DN}*) of the *ey>M1BP^{RNAi}* phenotype. The orientation of all imaginal discs is identical with posterior to the left and dorsal up. The magnification of all eye-antennal imaginal discs is 20X and adult eyes are 10X unless specified.
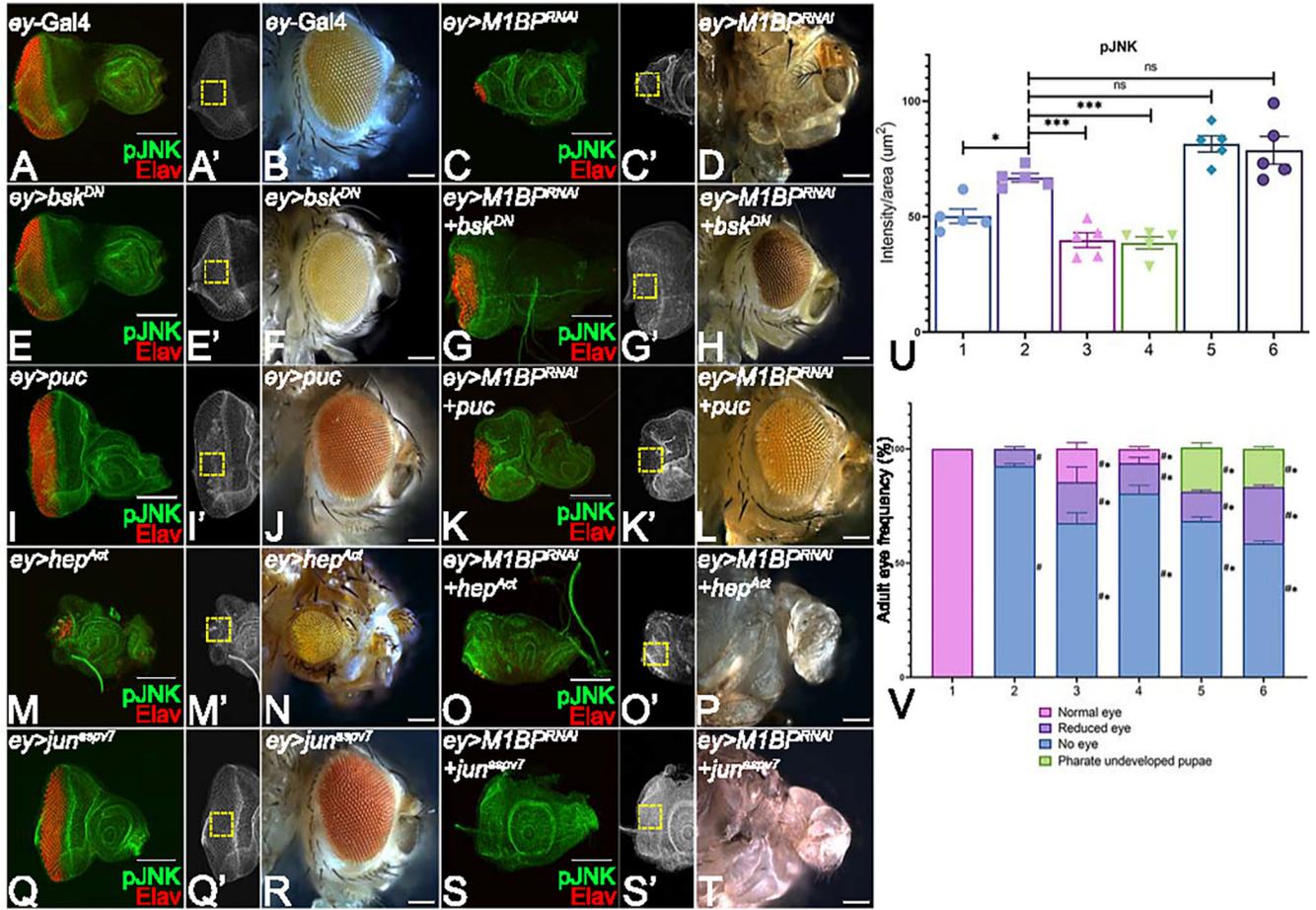
Figure 2: Downregulation of M1BP activates JNK signaling in the developing eye.
(A) Schematic representation of c-Jun NH (2)-terminal kinase (JNK) signaling pathway.
(B) Relative expression of *jnk* at the transcriptional level using quantitative real time PCR
(qRT-PCR) in *ey>M1BP^RNAi* and *ey*-Gal4 control eye-antennal imaginal discs. (C) Levels
of phospho-JNK (pJNK) in a semi-quantitative Western Blot shows higher levels of JNK
signaling in *ey>M1BP^RNAi* compared to *ey*-Gal4 control. The tubulin bands serve as control
for normalization. The quantification of p-JNK band intensity reveals a significantly higher
levels in *ey>M1BP^RNAi* background as compared to the control *ey*-Gal4. (D) Quantification
of pJNK intensity using Fiji/ImageJ software (NIH). (E-H) Eye antennal imaginal disc of
third instar larvae stained for pan-neuronal marker Elav (red) and (E, F) pJNK (green)
and (G, H) puc-lacZ (green). Eye antennal imaginal disc showing split channel for (E′,
F′) pJNK staining, (G′, H′) lacZ staining for (E, E′, G, G′) *ey*-Gal4 and (F, F′, H, H′)
ey>M1BP^RNAi. Graphs were plotted with mean +/− SEM. Statistical significance in each
graph is shown by p-value: ****p<0.0001, ***p<0.001; **p<0.01; *p<0.05. The orientation
of all imaginal discs is identical with posterior to the left and dorsal up. The magnification of
all eye-antennal imaginal disc is 20X unless specified.

**Figure 3: Modulation of JNK signaling affects eye suppression phenotype of M1BP downregulation.**

(A, C, E, G, I, K, M, O, Q, S) Eye antennal imaginal disc of third instar larvae stained for pJNK (green) and pan-neuronal marker Elav (red). (A, A') *ey*-Gal4 control discs showing pJNK expression. (A', C', E', G', I', K', M', O', Q', S') Eye antennal imaginal disc showing split channel for pJNK staining. Quantification was performed using standard 100X100 pixel ROI. ROI is shown as yellow dashed boxes. Adult eye images of control (B) *ey*-Gal4, downregulation of *M1BP* (D) *ey>M1BP^{RNAi}*, downregulating JNK pathway: (F) *ey>bsk^{DN}*, (H) *ey>M1BP^{RNAi}+bsk^{DN}*, (J) *ey>puc,* (L) *ey>M1BP^{RNAi}+puc*, upregulating JNK pathway: (N) *ey>hep^{Act}*, (P) *ey>M1BP^{RNAi}+hep^{Act}*, (R) *ey>jun^{aspv7}*, (T) *ey>M1BP^{RNAi}+jun^{aspv7}*. Eye-antennal imaginal discs of control (A, A') *ey*-Gal4, downregulation of *M1BP* (C, C') *ey>M1BP^{RNAi}*, downregulating JNK pathway: (E, E') *ey>bsk^{DN}*, (G. G') *ey>M1BP^{RNAi}+bsk^{DN}*, (I, I') *ey>puc,* (K, K') *ey>M1BP^{RNAi}+puc*, upregulating JNK pathway: (M, M') *ey>hep^{Act}*, (O, O') *ey>M1BP^{RNAi}+hep^{Act}*, (Q, Q') *ey>jun^{aspv7}*, (S, S') *ey>M1BP^{RNAi}+jun^{aspv7}*. Note that (H, L) downregulation of JNK signaling showed significant rescues in adult eye phenotypes; whereas (P, T) activation of JNK signaling enhanced the eye suppression phenotype of (D) *ey>M1BP^{RNAi}*. (U) Quantification of pJNK intensity using Fiji/ImageJ software (NIH). The genotypes depicted in the graph are 1: *ey*-Gal4, 2: *ey>M1BP^{RNAi}*, 3: *ey>M1BP^{RNAi}+bsk^{DN}*, 4: *ey>M1BP^{RNAi}+puc*, 5: *ey>M1BP^{RNAi}+hep^{Act}* and 6: *ey>M1BP^{RNAi}+jun^{aspv7}*. (V) Graphical representation of

adult eye phenotype frequency. The genotypes depicted in the graph are 1: *ey*-Gal4, 2: *ey>M1BP^{RNAi}*, 3: *ey>M1BP^{RNAi}+bsk^{DN}*, 4: *ey>M1BP^{RNAi}+puc*, 5: *ey>M1BP^{RNAi}+hep^{Act}* and 6: *ey>M1BP^{RNAi}+jun^{aspv7}*. For adult eye frequency, statistical significance with *ey*Gal4 and *ey>M1BP^{RNAi}* are depicted using # and * respectively. Graphs were plotted with mean +/− SEM. For quantification, statistical significance in each graph is shown by p-value: ****p<0.0001, ***p<0.001; **p<0.01; *p<0.05. The orientation of all imaginal discs is identical with posterior to the left and dorsal up. The magnification of all eye-antennal imaginal discs is 20X and adult eyes are 10X unless specified.
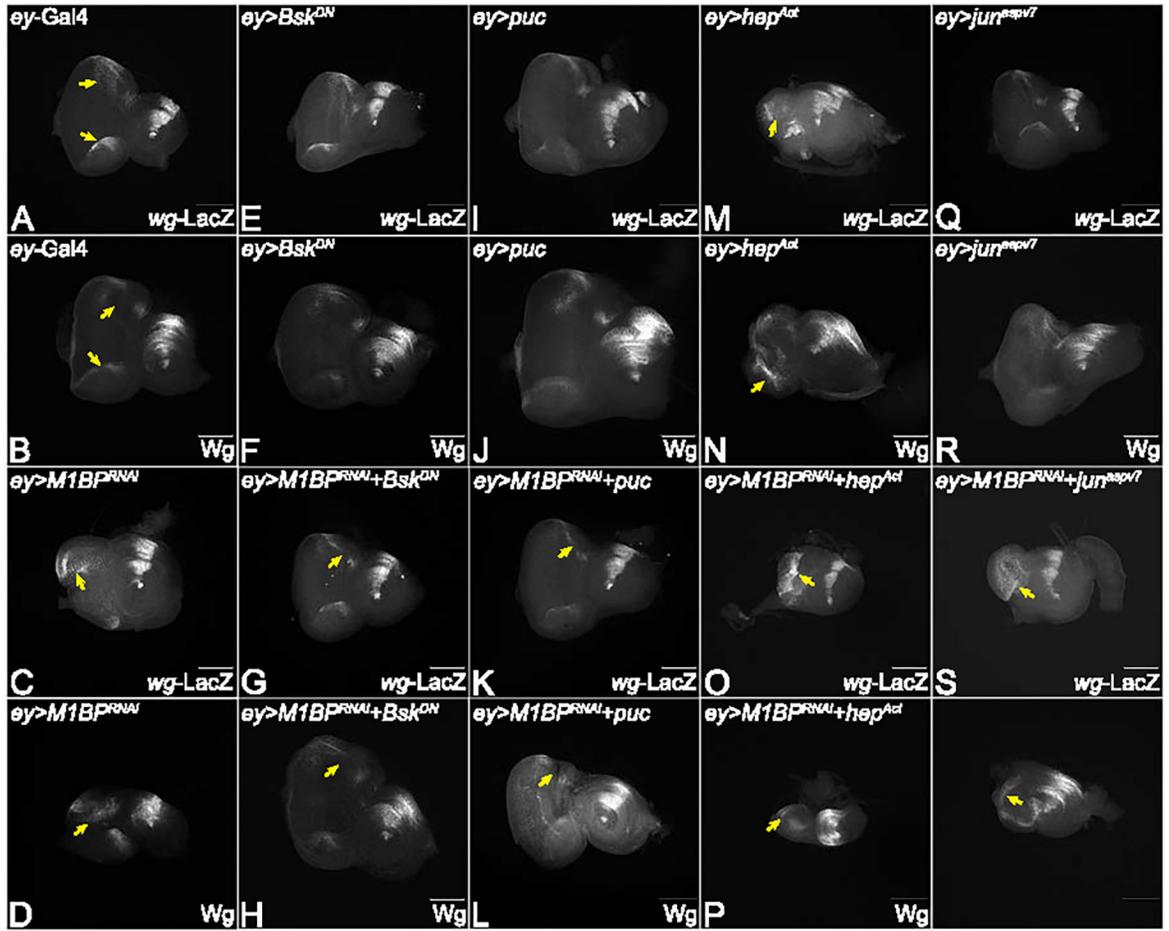
Figure 4: Modulation of JNK pathway in M1BP LOF background affects *wingless* levels.
Eye-antennal imaginal discs from third instar larvae of (A) *ey*-Gal4, downregulation
of *M1BP* (C) *ey>M1BP^RNAi*, downregulating JNK pathway: (E) *ey>bsk^DN*, (G)
ey>M1BP^RNAi+bsk^DN, (I) *ey>puc*, (K) *ey>M1BP^RNAi+puc*, upregulating JNK pathway:
(M) *ey>hep^Act*, (O) *ey>M1BP^RNAi+hep^Act*, (Q) *ey>jun^aspv7*, (T) *ey>M1BP^RNAi+jun^aspv7*
were assessed for changes in *wg*-lacZ expression shown in grayscale. (B, D, F, H, J,
L, N, P, R, S, U) Eye-antennal imaginal discs from third instar larvae of (B) *ey*-Gal4,
downregulation of *M1BP* (D) *ey>M1BP^RNAi*, downregulating JNK pathway: (F) *ey>bsk^DN*,
(H) *ey>M1BP^RNAi+bsk^DN*, (J) *ey>puc*, (L) *ey>M1BP^RNAi+puc*, upregulating JNK pathway:
(N) *ey>hep^Act*, (P) *ey>M1BP^RNAi+hep^Act*, (R) *ey>jun^aspv7*, (S) *ey>M1BP^RNAi+jun^aspv7*
were assessed for changes in Wg expression shown in grayscale. (A, B) *ey*-Gal4 control
discs showing expression in the anterolateral margins of the eye disc. (C, D) Loss of
M1BP in the entire developing eye (*ey>M1BP^RNAi*) results in ectopic induction of *wg*
expression. Downregulation of JNK pathway in the background of loss of M1BP in the
entire eye (G, H) *ey>M1BP^RNAi+bsk^DN* and (K, L) *ey>M1BP^RNAi+puc*. Activation of JNK
pathway in the background of loss of M1BP (O, P) *ey>M1BP^RNAi+hep^Act* and (T, U)
ey>M1BP^RNAi+jun^aspv7. Yellow solid arrows indicate the induction or suppression of Wg.
The orientation of all imaginal discs is identical with posterior to the left and dorsal up. The
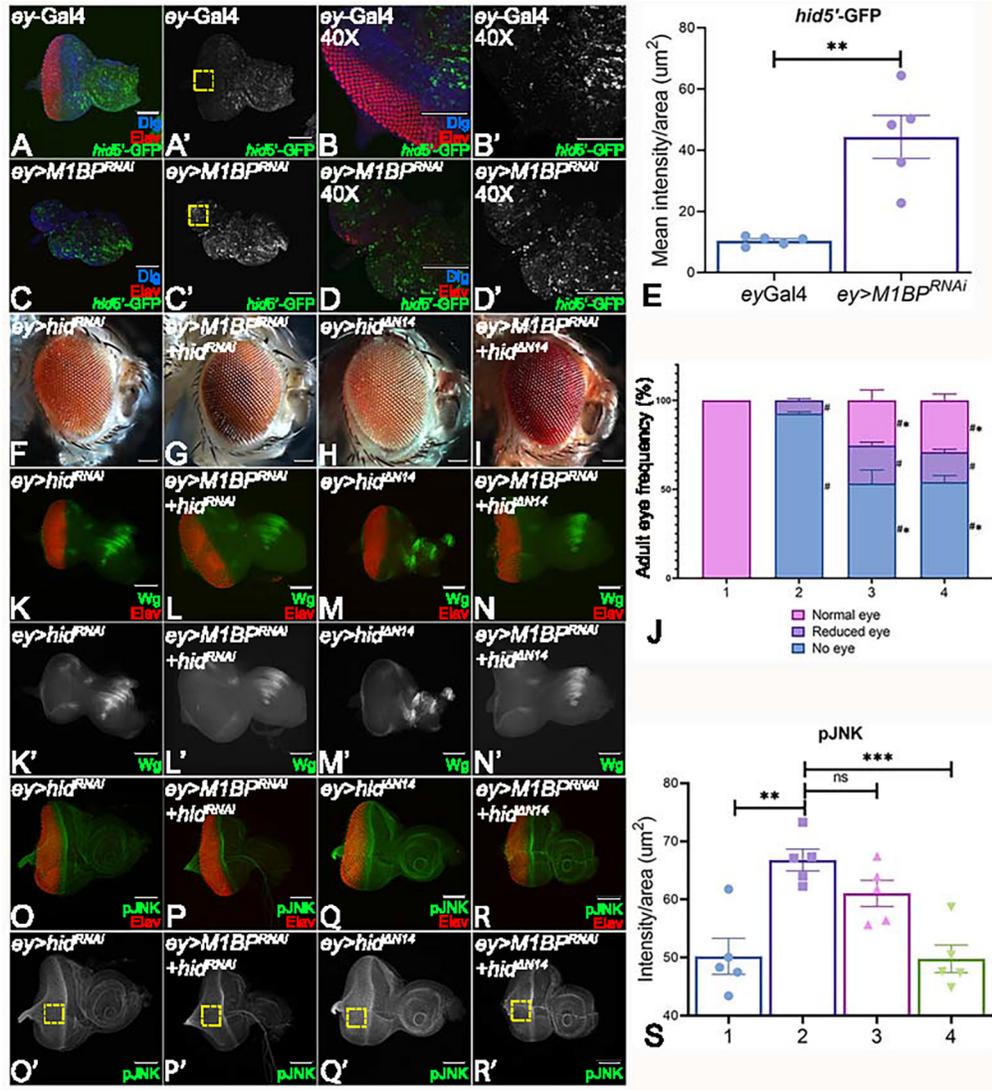magnification of all eye-antennal imaginal discs is 20X unless specified.

Figure 5: Downregulation of M1BP triggers cell death.

Eye-antennal imaginal discs stained for pan-neuronal marker Elav (red), Dlg (blue), a membrane specific marker, to mark outline of the tissue and (A-D) *hid5'F-WT*-GFP (green). Eye antennal imaginal disc showing split channel for (A'-D') *hid5'F-WT*-GFP for (A, A', B, B') *ey*-Gal4 and (C, C', D, D') *ey>M1BP^RNAi* in (A, A', C, C') 20X and (B, B', D, D') 40X. (E) Quantification of *hid5'F-WT*-GFP intensity using Fiji/ImageJ software (NIH) using standard 100X100 pixel ROI. ROI is shown as yellow dashed boxes. Caspase-dependent cell death was blocked in the entire eye (F, K, K', O, O') *ey>hid^RNAi*, (H, M, M', Q, Q') *ey>hid^N14*. Blocking hid mediated apoptotic cell death in *ey>M1BP^RNAi* background (G, L, L', P, P') *ey>M1BP^RNAi+hid^RNAi*, (I, N, N', R, R') *ey>M1BP^RNAi+hid^N14* shows rescue when compared to *ey>M1BP^RNAi*. (J) Graphical representation of adult eye phenotype frequency. The genotypes depicted in the graph are 1: *ey*-Gal4, 2: *ey>M1BP^RNAi*, 3: *ey>M1BP^RNAi+hid^RNAi* and 4: *ey>M1BP^RNAi+hid^N14*. Statistical significance with *ey*Gal4 and *ey>M1BP^RNAi* are depicted using # and * respectively. Eye antennal imaginal discs of indicated genotypes stained for (K-N) Wg

(green), (O-R) pJNK (green) and pan-neuronal marker Elav (red). Eye-antennal imaginal discs showing split channel for (K'-N') Wg and (O'-R') pJNK staining. (S) pJNK intensity quantification using Fiji/ImageJ software (NIH). The genotypes depicted in the graph are 1: *ey*-Gal4, 2: *ey>M1BP^{RNAi}*, 3: *ey>M1BP^{RNAi}+hid^{RNAi}* and 4: *ey>M1BP^{RNAi}+hid ^{N14}*. Quantification was performed using standard 100X100pixels ROI. ROI is shown as yellow dashed boxes. Graphs were plotted with mean +/− SEM. Statistical significance in each graph is shown by p-value: ****$p<0.0001$, ***$p<0.001$; **$p<0.01$; *$p<0.05$. The orientation of all imaginal discs is identical with posterior to the left and dorsal up. The magnification of all eye-antennal imaginal discs is 20X and adult eyes are 10X unless specified.
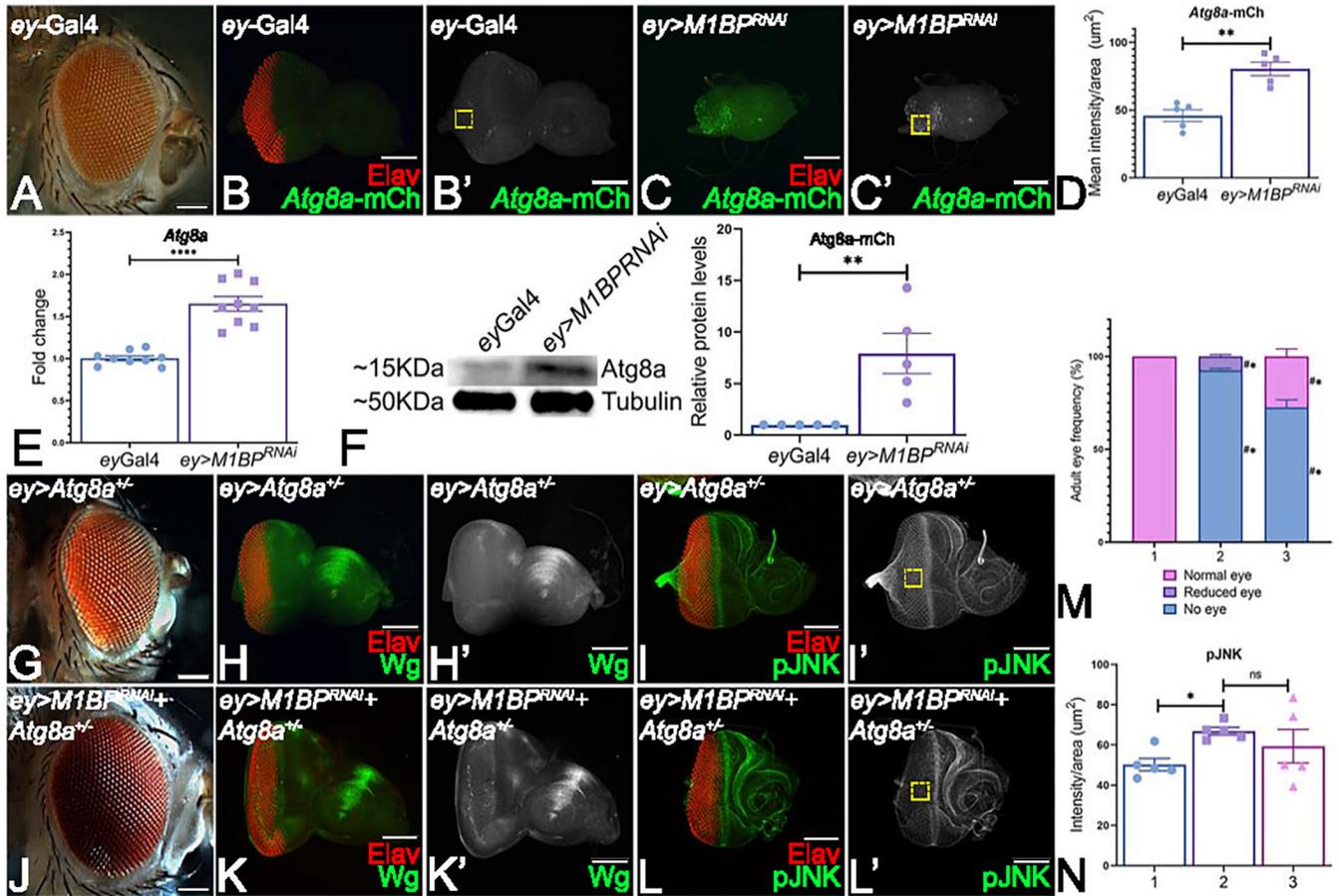
Figure 6: Downregulation of M1BP triggers autophagy.
(A) *ey*-Gal4 adult eye. (B, B', C, C') Eye-antennal imaginal discs stained with pan neuronal marker Elav (red) and assessed for changes in expression of *Atg8a-mCherry* (green) reporter. (B', C') Eye antennal imaginal disc showing split channel for *Atg8a-mCherry* expression. (B, B') *ey*-Gal4 control discs. (C, C') Downregulation of M1BP in the entire developing eye (*ey>M1BP^RNAi*). (D) Quantification of *Atg8a-mCherry* intensity conducted using Fiji/ImageJ software (NIH) using standard 100X100 pixel ROI. ROI is shown as yellow dashed boxes. (E) Relative expression of *Atg8a* at the transcriptional level using quantitative real time PCR (qRT-PCR) in *ey>M1BP^RNAi* and *ey*-Gal4 control eye-antennal imaginal discs. (F) Levels of Atg8a in a semi-quantitative Western Blot shows higher levels of autophagy activation in *ey>M1BP^RNAi* compared to *ey*-Gal4 control. The tubulin bands serve as internal loading control for normalization. The quantification of Atg8a band intensity reveals a significantly higher level in *ey>M1BP^RNAi* when compared to *ey*-Gal4. (G, H, H', I, I') Blocking autophagy in the entire eye using *Atg8a* mutant (*ey>Atg8a^+/−*) results in a near wild-type eye. Blocking autophagy using *Atg8a* mutant in *ey>M1BP^RNAi* background (H, I, I', J, J') *ey>M1BP^RNAi+Atg8a^+/−* can rescue the eye suppression phenotype. Eye antennal imaginal discs of indicated genotypes stained for (H, K) Wg (green), (I, L) pJNK (green) and pan-neuronal marker Elav (red). Eye-antennal imaginal discs showing split channel for (H', K') Wg and (I', L') pJNK staining. (M) Graphical representation of adult eye phenotype frequency. The genotypes depicted in the

graph are 1: *ey*-Gal4, 2: *ey>M1BP^RNAi*, 3: *ey>M1BP^RNAi+Atg8a^+/−*. Statistical significance with *ey*Gal4 and *ey>M1BP^RNAi* are depicted using # and * respectively. (N) pJNK intensity quantification using Fiji/ImageJ software (NIH). The genotypes depicted in the graph are 1: *ey*-Gal4, 2: *ey>M1BP^RNAi*, 3: *ey>M1BP^RNAi+Atg8a^+/−*. Quantification was performed using standard 100X100pixels ROI. ROI is shown as yellow dashed boxes. Graphs were plotted with mean +/− SEM. Statistical significance in each graph is shown by p-value: ****p<0.0001, ***p<0.001; **p<0.01; *p<0.05. The orientation of all imaginal discs is identical with posterior to the left and dorsal up. The magnification of all eye-antennal imaginal discs is 20X and adult eyes are 10X unless specified.
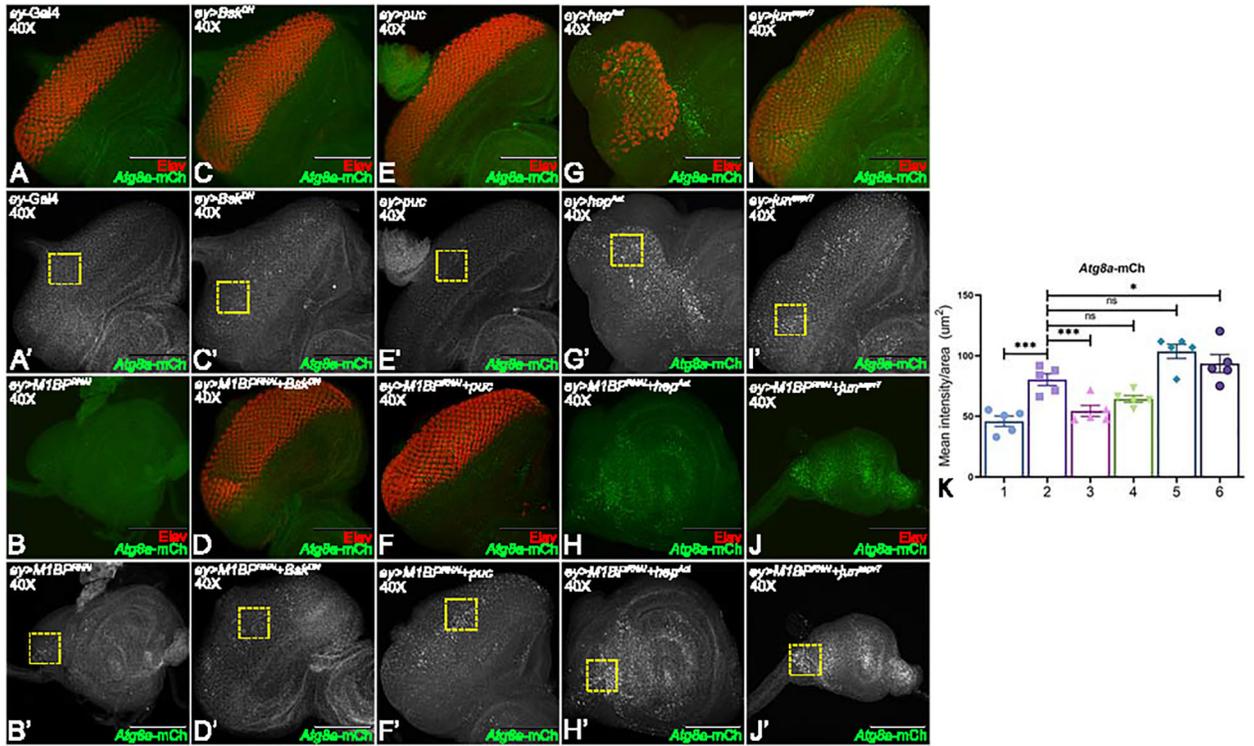
Figure 7: Activation of JNK signaling in the background of M1BP downregulation promotes autophagy.

(A-J) Eye-antennal imaginal discs of (A, A') *ey*-Gal4 control, (B, B') *ey>M1BP^RNAi*, (C, C') *ey>bsk^DN*, (D, D') *ey>M1BP^RNAi+bsk^DN*, (E, E') *ey>puc,* (F, F') *ey>M1BP^RNAi+puc*, (G, G') *ey>hep^Act*, (H, H') *ey>M1BP^RNAi+hep^Act*, (I, I') *ey>jun^aspv7*, (J, J') *ey>M1BP^RNAi+jun^aspv7* were assessed for changes in *Atg8a-mCherry* (green) and pan neuronal marker Elav (red) expression. (A'-J') Eye antennal imaginal disc showing split channel for *Atg8a-mCherry* expression. (C-F) JNK pathway was downregulated using *bsk^DN* (C, C') *ey>bsk^DN* and *puc* (E, E') *ey>puc*. (G-J) JNK pathway was activated using *hep^Act* (G, G') *ey>hep^Act* and *jun^aspv7* (I, I') *ey>jun^aspv7*. (K) Quantification of *Atg8a-mCherry* intensity using Fiji/ImageJ software (NIH). Quantification was performed using standard 100X100 pixels ROI. ROI is shown as yellow dashed boxes. Graph was plotted with mean +/− SEM. The genotypes depicted in the graph are 1: *ey*-Gal4, 2: *ey>M1BP^RNAi*, 3: *ey>M1BP^RNAi+bsk^DN*, 4: *ey>M1BP^RNAi+puc*, 5: *ey>M1BP^RNAi+hep^Act* and 6: *ey>M1BP^RNAi+jun^aspv7*. Statistical significance in each graph is shown by p-value: ****p<0.0001, ***p<0.001; **p<0.01; *p<0.05. The orientation of all imaginal discs is identical with posterior to the left and dorsal up. The magnification of all eye-antennal imaginal discs is 40X.
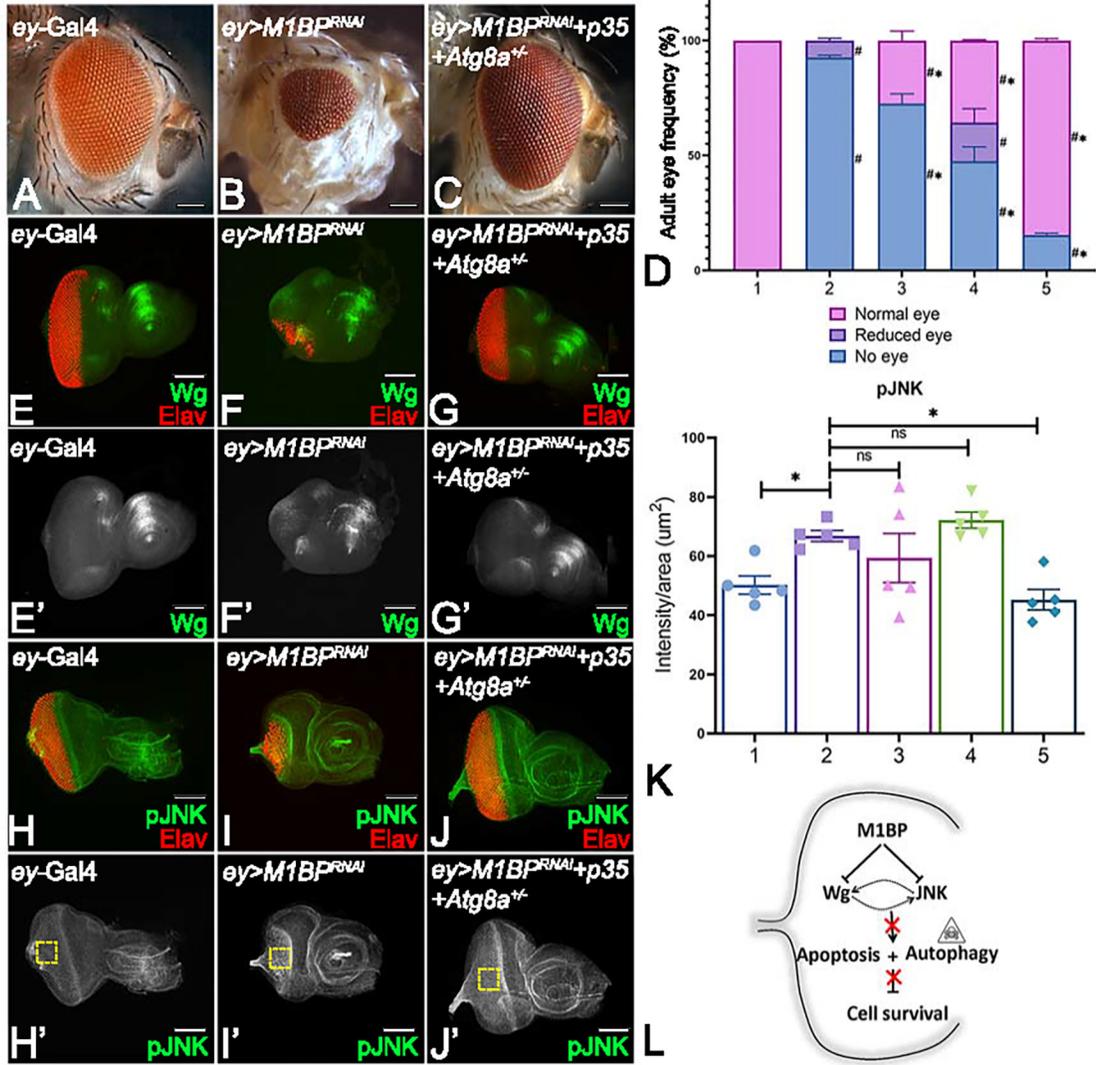
Figure 8: M1BP is required to block both caspase-dependent and caspase-independent cell death during development.

Adult eye of (A) *ey*-Gal4 showing normal arrangement of photoreceptors and (B) *ey>M1BP^{RNAi}* showing reduced eye phenotype. (C) Blocking caspase -dependent and -independent (autophagy) cell death in the background of M1BP downregulation *ey>M1BP^{RNAi}+p35+Atg8a^{+/−}* eye shows significant rescue. (D) Graphical representation of adult eye phenotype frequency. Graphs were plotted with mean +/− SEM. The genotypes depicted in the graph are 1: *ey*-Gal4, 2: *ey>M1BP^{RNAi}*, 3: *ey>M1BP^{RNAi}+Atg8a^{+/−}*, 4: *ey>M1BP^{RNAi}+p35*, 5: *ey>M1BP^{RNAi}+p35+Atg8a^{+/−}*. Statistical significance with *ey*Gal4 and *ey>M1BP^{RNAi}* are depicted using # and * respectively. Eye-antennal imaginal discs of the indicated genotypes were stained for (E-G) Wg (green), (H-J) pJNK (green) and pan-neuronal marker Elav (red). Eye-antennal imaginal discs showing split channel for (E'-G') Wg and (H'-J') pJNK staining. (K) pJNK intensity quantification using Fiji/ImageJ software (NIH). Quantification was performed using standard 100X100pixels ROI. ROI is shown as yellow dashed boxes. The genotypes depicted in the graph are 1: *ey*-Gal4, 2: *ey>M1BP^{RNAi}*,

3: *ey>M1BP^{RNAi}+Atg8a^{+/−}*, 4: *ey>M1BP^{RNAi}+p35*, 5: *ey>M1BP^{RNAi}+p35+Atg8a^{+/−}*. (L) Model for M1BP mediated regulation of cellular homeostasis. M1BP suppresses JNK-mediated apoptosis and autophagy to promote cell survival in the developing eye. Statistical significance in each graph is shown by p-value: ****$p<0.0001$, ***$p<0.001$; **$p<0.01$; *$p<0.05$. The orientation of all imaginal discs is identical with posterior to the left and dorsal up. The magnification of all eye-antennal imaginal discs is 20X and adult eyes are 10X unless specified.