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## Inactivation of Hippo and cJun-N-terminal Kinase (JNK) signaling mitigate FUS mediated neurodegeneration *in vivo*

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### Abstract

Amyotrophic Lateral Sclerosis (ALS), a late-onset neurodegenerative disorder characterized by the loss of motor neurons in the central nervous system, has no known cure to-date. Disease causing mutations in human Fused in Sarcoma (FUS) leads to aggressive and juvenile onset of ALS. FUS is a well-conserved protein across different species, which plays a crucial role in regulating different aspects of RNA metabolism. Targeted misexpression of FUS in *Drosophila* model recapitulates several interesting phenotypes relevant to ALS including cytoplasmic mislocalization, defects at the neuromuscular junction and motor dysfunction. We screened for the genetic modifiers of human FUS-mediated neurodegenerative phenotype using molecularly defined deficiencies. We identified *hippo* (*hpo*), a component of the evolutionarily conserved Hippo growth regulatory pathway, as a genetic modifier of FUS mediated neurodegeneration. Gain-of-function of *hpo* triggers cell death whereas its loss-of-function promotes cell proliferation. Downregulation of the Hippo signaling pathway, using mutants of Hippo signaling, exhibit rescue of FUS-mediated neurodegeneration in the *Drosophila* eye, as evident from reduction in the

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Authors Contributions

N.G., A.S., A.S.M, N.R., P.D. performed the experiments, analyzed the data and provided comments to the manuscript.

N.G. screened the genetic modifiers and helped in manuscript writing.

A.S., U.P. and M.K.S. were involved in developing the concept, designing the experiments, analyzing the data and writing the manuscript.

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number of TUNEL positive nuclei as well as rescue of axonal targeting from the retina to the brain. The Hippo pathway activates c-Jun amino-terminal (NH<sub>2</sub>) Kinase (JNK) mediated cell death. We found that downregulation of JNK signaling is sufficient to rescue FUS-mediated neurodegeneration in the *Drosophila* eye. Our study elucidates that Hippo signaling and JNK signaling are activated in response to FUS accumulation to induce neurodegeneration. These studies will shed light on the genetic mechanism involved in neurodegeneration observed in ALS and other associated disorders.

## Keywords

Neurodegeneration; Amyotrophic Lateral Sclerosis (ALS) FUS; Fused in Sarcoma; TLS; *Drosophila* eye; Hippo pathway; JNK signaling; Cell death

## Introduction

Amyotrophic Lateral Sclerosis (ALS), a fatal late-onset neurodegenerative disorder, is characterized by degeneration of motor neurons in the brain and spinal cord (Brenner and Weishaupt, 2019; Riggs, 1985). One of the hallmarks of ALS is disruption in RNA metabolism, which results in degeneration of motor neurons (Daigle et al., 2013; Donnelly et al., 2013; Ling et al., 2013; Nussbacher et al., 2019; Ortega et al., 2020). Mutations in more than 30 genes including *superoxide dismutase 1 (SOD1)* (Rosen et al., 1993), *TAR DNA-binding protein-43 (TDP-43)* (Kabashi et al., 2008; Rutherford et al., 2008), *C9ORF72* (DeJesus-Hernandez et al., 2011; Renton et al., 2011), *ubiquilin-2 (UBQLN2)* (Deng et al., 2011), *optineurin (OPTN)* (Maruyama et al., 2010), and *fused in sarcoma (FUS)* (Kwiatkowski et al., 2009; Vance et al., 2009) results in ALS (White and Sreedharan, 2016). Of these, FUS is not only involved in causing ALS associated neurodegeneration (Chen et al., 2011), but also serves as a factor in the pathogenesis of other neurodegenerative disorders like polyglutamine diseases (Doi et al., 2008; Kino et al., 2016). FUS is a DNA/RNA binding protein that plays crucial role(s) in mRNA metabolism, nucleocytoplasmic RNA transport, and translation (Coyne et al., 2017; Kwiatkowski et al., 2009; Yang et al., 1998; Zinszner et al., 1997). Normally FUS shuttles between the nucleus and cytoplasm. Pathogenic mutations in FUS result in cytoplasmic mislocalization of a small fraction of the FUS protein. It suggests that cytoplasmic retention of FUS is likely causing cytotoxicity (Lanson et al., 2011; Zinszner et al., 1997). Further, deletion of Nuclear Export Signal (NES) disrupts the ability of this protein to enter the cytoplasm, which likely results in ALS neuropathy (Lanson et al., 2011). The RNA-binding ability of FUS has been suggested to play a major role in FUS mediated neurodegeneration and has been demonstrated by generating mutations in RNA recognition motifs (RRM) like R518K and R521C. However, the exact mechanism(s) that can block FUS-mediated neurodegeneration is unclear. Therefore, it is important to understand functional consequences of disease-causing mutations in FUS, which promote cytoplasmic accumulation of mutant FUS, in order to discern the mechanism of FUS-mediated neurodegeneration.

Since the genetic machinery is highly conserved, several animal models including mice, zebrafish, and fruit flies are in use to study the molecular mechanisms of ALS disease

(Casci and Pandey, 2015; Pandey and Nichols, 2011; Picher-Martel et al., 2016). Amongst these, *Drosophila melanogaster* a.k.a fruit fly, serves as a highly versatile and genetically tractable human disease model (McGurk et al., 2015; Olesnický and Wright, 2018; Pandey and Nichols, 2011; Sarkar et al., 2016; Singh and Irvine, 2012). The *Drosophila* eye serves as a useful model to study neurodegenerative disorders like ALS (Casci and Pandey, 2015) as the retinal neurons are sensitive to cellular insult(s) and are not crucial for the survival and reproductive ability of the fly (Sarkar et al., 2016; Tare et al., 2011). The adult compound eye of *Drosophila* develops from an epithelial bi-layer structure, which is housed inside the larva, called an eye-antennal imaginal disc. The larval eye imaginal disc develops into the adult compound eye comprised of 800 unit eyes called ommatidia (Kumar, 2018; Ready et al., 1976; Singh et al., 2005; Singh et al., 2012; Tare et al., 2013). Misexpression of wild-type FUS or mutant FUS, in the differentiating retinal neurons of *Drosophila* eye, using a GMR-Gal4 driver (Moses and Rubin, 1991) results in severe degeneration of retinal neurons. Targeted misexpression of FUS (FUS) or mutant FUS in motor neurons exhibits locomotor dysfunction (Casci and Pandey, 2015; Coyne et al., 2017; Daigle et al., 2013; Lanson et al., 2011; Pandey and Nichols, 2011; Xia et al., 2012).

We have used a well-characterized fruit fly model to screen for the genetic modifiers of FUS-mediated neurodegeneration using molecularly defined deletions. We identified a deficiency, which uncovers *hippo* (*hpo*) and other genes as a suppressor of FUS-mediated neurodegeneration. The Hippo pathway is an evolutionarily conserved growth regulatory pathway (Kango-Singh and Singh, 2009; Ma et al., 2019). The core components of the Hippo pathway are Ste20 family protein kinase *hpo* (Harvey et al., 2003; Pantalacci et al., 2003; Udan et al., 2003; Wu et al., 2003), the nuclear Dbf-2-related (NDR)-family kinase- *warts* (*wts*) (Justice et al., 1995; Xu et al., 1995), and *yorkie* (*yki*)- the *Drosophila* orthologs of the mammalian transcription co-activator yes-associated protein (YAP) (Huang et al., 2005). When Hippo pathway is activated, Hpo binds with and phosphorylates Salvador (*Sav*), a protein containing WW and coiled-coil domains, and phosphorylates the downstream Wts kinase (Kango-Singh et al., 2002; Kango-Singh and Singh, 2009; Tapon et al., 2002). The active, phosphorylated form of Wts binds with an adaptor protein Mob-as-tumor suppressor (Mats) (Lai et al., 2005) and in turn phosphorylates the transcriptional co-activator Yki. The phosphorylated form of Yki binds with 14-3-3 proteins, and undergoes 14-3-3 mediated proteasomal degradation in the cytoplasm (Dong et al., 2007; Ren et al., 2010; Wei et al., 2007). However, downregulation of the Hippo signaling leads to release of Wts-mediated repression of Yki, and the unphosphorylated Yki goes into the nucleus, binds to its partner(s) such as the TEAD family transcription factor Scalloped (*Sd*), and regulates expression of the downstream targets of the Hippo pathway like *expanded* (*ex*), *drosophila inhibitor of apoptosis* (*diap1*), *cyclins*, *bantam* (*ban*), etc. (Fulford et al., 2018; Kango-Singh and Singh, 2009; Yu and Guan, 2013). Loss-of-function of *hpo*, *sav*, *wts*, *mats* or overexpression of *yki* activity leads to cell proliferation and over growth, while gain-of-function of the Hpo signaling leads to apoptosis or cell death (Kango-Singh and Singh, 2009).

The Hippo pathway activates JNK, an evolutionarily conserved signaling pathway, to trigger cell death (Ma et al., 2015; Ma et al., 2017). JNK signaling is involved in a wide array of signaling events underlying tumorigenesis and is known to regulate cell proliferation,

invasive migration, and cell death (Dhanasekaran and Reddy, 2017). JNK, or stress activated kinase proteins of mitogen-activated protein kinase (MAPK) superfamily (Adachi-Yamada et al., 1999; Adachi-Yamada and O'Connor, 2002), induce cell death due to phosphorylation of transcription factors involved in cell death (Dhanasekaran and Reddy, 2008). JNK signaling pathway in *Drosophila* gets activated downstream of the Tumor Necrosis Factor (TNF) homolog Eiger (Egr) and its receptor Wengen (Wgn) by a conserved downstream signaling cascade. The core components of this pathway comprises of Tak1 (TGF- $\beta$ -activating kinase 1); JNK Kinase Kinase (JNKKK), Hemipterous (Hep; JNK Kinase- JNKK, which is closely related to mitogen-activated protein Kinase Kinases (MAPKKs) (Glise et al., 1995; Tournier et al., 1997). Other components are Basket (Bsk; a Jun kinase), D-Jun (Jun), and Puckered (Puc). *bsk* acts downstream of *hep* and is activated by constitutive *hep* phosphorylation (Sluss et al., 1996). Phosphorylation of Jun leads to activation of the JNK signaling pathway that functions as a dual specificity protein phosphatase (Martín-Blanco et al., 1998). *puc* functions as a transcriptional target of JNK signaling and is known to regulate JNK signaling through a negative feedback loop (Adachi-Yamada et al., 1999; Adachi-Yamada and O'Connor, 2002). Furthermore, ectopic activation of JNK signaling induces cell death during early eye imaginal disc development (Singh et al., 2006). Here we present evidences to show that targeted misexpression of FUS triggers neurodegeneration as seen in ALS models by activating the Hippo signaling pathway, which we identified in a forward genetic screen. We further investigated the genetic mechanism and found that activation of the Hippo signaling pathway due to gain-of-function of FUS in turn triggers the JNK signaling to induce neurodegeneration in the developing *Drosophila* eye.

## Materials and Method

### Fly Stocks

The fly stocks used in this study are listed at FlyBase (<http://flybase.bio.indiana.edu>). We used the transgenic fly strains that carry full length FUS (UAS-FUS)(Lanson et al., 2011) and mutant FUS, UAS-FUS R518K and UAS-FUS R521C(Casci et al., 2019). Other stocks used are UAS-*hpo* (Udan et al., 2003), *yw, hsf1p; UAS-hpo<sup>RNAi</sup>symp19/SM6-TM6B,Tb* (Pantalacci et al., 2003), UAS-*wts<sup>13f</sup>* (Kwon et al., 2015), UAS-*wts<sup>RNAi</sup>* (Fernández et al., 2011; Rauskolb et al., 2011), UAS-*yki<sup>RNAi (N+C)</sup>* (Zhang et al., 2008), a hyperactivated form that allows *yki* activation, UAS-*yki<sup>3SA</sup>* (Oh and Irvine, 2008), UAS-*Djun<sup>aspv7</sup>* (Treier et al., 1995), UAS-*bsk<sup>DN</sup>* (Adachi-Yamada et al., 1999), UAS-*hep<sup>Act</sup>* (Glise et al., 1995), UAS-*puc* (Martín-Blanco et al., 1998) and *diap1-4.3-GFP* (Ren et al., 2010). The *diap1-4.3-GFP* gene, contains a 4.3-kb *diap1* genomic fragment from +1.37 kb to +5.68 kb, which contains a Hippo response element that drives the Diap1 expression in response to Hippo pathway activity where GFP is a reporter for Yki activity. We used a molecularly defined deficiency *Df(2R)BSC782/+*, which is located on the right arm of the second chromosome, and uncovers  *$\beta$ Tub56D, par-1, CG16926, CG7744, CG15120, mei-W68, oseg6, TBCB, rep* and *hpo* genes (listed in FlyBase). We used the Canton-S (Wild-type) stock of *D. melanogaster* in this study. Fly stocks used in this study (Supplementary Table 1) were maintained at 25°C on cornmeal, molasses food.

## Genetic Crosses

We employed the Gal4/UAS system for targeted misexpression studies (Brand and Perrimon, 1993). The GMR-Gal4 enhancer directs expression of transgenes in the differentiating retinal precursor cells in the developing larval eye-antennal imaginal discs (Moses and Rubin, 1991; Tare et al., 2011). We used GMR-Gal4 to drive expression of transgenes. We used GMR-Gal4/ CyO; UAS-FUS / *TM3 Sb e Ser* (GMR>FUS), GMR-Gal4/ CyO; UAS-FUS R518K / *TM3 Sb e Ser* (GMR>FUS R518K), GMR-Gal4/ CyO; UAS-FUS R521K/ *TM3Sb e Ser* (GMR>FUSR521K) to sample FUS-mediated neurodegeneration in the eye. All fly stocks were maintained at 25°C. All Gal4/UAS crosses were maintained at 18°C, 25°C, or 29°C to sample different induction levels (Singh and Choi, 2003). The third instar larval eye-antennal imaginal discs were used for immunohistochemistry purposes.

## Immunohistochemistry

The eye-antennal imaginal discs were dissected from the wandering third-instar *Drosophila* larvae in 1X PBS (Phosphate Buffered Saline, pH 7.4) following the standard protocol (Sarkar et al., 2018a; Singh et al., 2002). Tissue samples were then fixed in 4% paraformaldehyde for 20 minutes and washed with 1X PBST (1XPBS + 0.2% TritonX-100), 3 times for 10 minutes each. Tissues were incubated with different combinations of primary and secondary antibodies. The primary antibodies used were: rabbit anti-FUS (1:500, Bethyl Laboratories, A300-302A), mouse anti-Dlg (1:100, Developmental Studies Hybridoma Bank-DSHB, 4F3), mouse anti-Wg (1:50, DSHB, 4D4), rat anti-Elav (1:100; DSHB, 7E8A10), mouse anti-Chaoptin, MAb24B10 (1:100, DSHB, 24B10) (Zipursky et al., 1984), and rabbit anti- $\beta$ -galactosidase (1:100, Promega, Z3781). Secondary antibodies (Jackson Immuno Research) used were donkey anti-rabbit IgG conjugated with FITC (dilution 1:200), donkey anti-mouse IgG conjugated with Cy3 (dilution 1:250), and goat anti-rat IgG conjugated with Cy5 (dilution 1:250). After primary and secondary antibody incubation, tissue samples were washed with 1XPBST (X3) for 10 minutes each to remove excess antibody bound to sample. Following washing, tissue samples were mounted in Vectashield mountant (Vector laboratories-H1000). The images were scanned using Olympus Fluoview-3000 Laser Scanning Confocal Microscope (LSCM) and final figures were made using Adobe Photoshop CS6 software (Sarkar et al., 2018a). The quantification of GFP intensity was done using built-in program in Olympus Fluoview 3000 confocal microscope. The statistical analysis of eye-antennal imaginal discs images was conducted using Microsoft Excel 2017 software, p-values were calculated using a Student's two-tailed t-test and the error bars represent the values of standard deviation calculated from the mean (Tare et al., 2016).

## Detection of Cell Death

The TUNEL assay marks the cells undergoing cell death. In TUNEL staining, the cleavage of double and single-stranded DNA is labeled by a Fluorescent tag (TMR Red) (McCall and Peterson, 2004; White et al., 1994). The fluorescently labeled nucleotides are added to 3' OH ends in a template-independent manner by Terminal Deoxynucleotidyl transferase (TdT). The dying cell's fragmented DNA is tagged with a fluorochrome and can be

detected using a fluorescence microscope or confocal microscope. Eye antennal discs after secondary antibody staining (Singh et al., 2006) were blocked in 10% normal donkey serum in Phosphate buffered saline with 0.2% Triton X-100 (PBT) and labeled for TUNEL assays using a cell death detection kit from Roche Diagnostics (12156792910). For negative controls, eye-antennal imaginal discs were treated with a cell death detection labeling mix without the TdT enzyme added.

The TUNEL positive nuclei were counted from five sets of eye imaginal discs for each experiment. The cell count number was used for statistical analysis using Microsoft excel 2013. The p-values were calculated using a Student's two-tailed t- test. The error bars represent the standard deviation from the mean (Steffensmeier et al., 2013).

### Adult Eye Imaging

The adult flies were prepared for imaging by freezing them at  $-20^{\circ}\text{C}$  for approximately 2 hours. Following this incubation, the flies were mounted on a dissection needle. The needle carrying the fly was placed horizontally over a glass slide using a putty. Images were captured on a MrC5 color camera mounted on an Axioimager.Z1 Zeiss Apotome using a Z-sectioning method (Sarkar et al., 2018a; Wittkorn et al., 2015). The final images were generated by compiling individual stacks from the Z-section using the extended depth of focus function of Axiovision software version 4.6.3.

### Scanning Electron Microscopy (SEM)

SEM was carried out using the standard protocol (Tare and Singh, 2009). The flies with phenotype were dehydrated through a series of increasing concentrations of acetone. The dehydrated fly samples were then incubated in 1:1 ratio of acetone and Hexamethyldisilazane (HMDS, Electron Microscopy Sciences) solution for 24 hours, followed by incubation in 100% HMDS alone. The flies were then air dried in the fume hood. The samples were mounted onto the conductive carbon tape attached to stubs (Electron Microscopy Sciences). The flies mounted on the stub were then subjected to sputter coating with gold using a Denton vacuum sputter coater (DV502). The samples were photographed using a Hitachi S-4800 High Resolution Scanning Electron Microscope (HRSEM) (Singh et al., 2019). The images were later analyzed using Adobe Photoshop CS6 software.

### Western Blot

The protein samples were extracted from heads of Wild-type, *GMR>FUS*, *GMR> FUS-R518K*, and *GMR>FUS-R521C* adult flies using a standardized protocol (Gogia et al., 2017). The primary antibodies used were Phospho-SAPK/JNK (Thr183/Tyr185) (81E11) (1:3000, Cell Signaling) and anti-FUS (1:1500, A300-302A, Bethyl laboratories). A signal was detected using a Horse Radish Peroxidase (HRP) conjugated goat anti-rabbit IgG (1:5000), and Super Signal West Dura Extended Duration Substrate (Thermo Scientific). Images were captured using the BioSpectrum® 500 imaging system and analyzed using Adobe Photoshop CS6 software. To measure the band intensity, statistical analysis was conducted on western blots using Microsoft Excel 2017 software. The p-values were

calculated using Student's two-tailed t-test and the error bars represent the values of standard deviation calculated from the mean (Sarkar et al., 2018b).

HEK293T cells (ATCC® CRL-3216™) were cultured in Advanced DMEM supplemented with 10% FBS and 1% Glutamax and grown at 37°C and 5% CO<sub>2</sub> (Casci et al., 2019). HEK293T cells were transiently transfected using Turbofect (Invitrogen) with pCI-neo-HA-FUS WT, pCI-neo-HA-FUS R518K and pCI-neo-HA-FUS R521C. Cells were lysed in RIPA buffer (150 mM NaCl, 1% NP40, 0.1% SDS, 1% sodium deoxycholate, 50 mM NaF, 2 mM EDTA, 1 mM DTT, 2.5 mM Na orthovanadate, 1x protease inhibitor cocktail [Roche 11836170001], pH 7.4] 24 hours post-transfection, and centrifuged down at 12000xg for 10 minutes. The supernatant was boiled in 1X NuPage LDS-Sample buffer (Invitrogen NP0007) at 95°C for 5 minutes. SDS-PAGE was performed using 4-12% NuPage Bis-Tris gels (Invitrogen), and the separated proteins were transferred onto nitrocellulose membranes using the iBlot2 system (Life Technologies 13120134). Nitrocellulose membranes were incubated overnight at 4°C with primary antibody (1:1000 LATS1, SCBT #398560; 1:500 Phospho-LATS1 (Thr1079), CST #8654; 1:8000  $\alpha$ -tubulin, Sigma T5168) followed by secondary antibody at room temperature for 1 hour. Membranes were imaged on Odyssey CLx (LI-COR Biosciences) and quantification of bands was performed using Image Studio (LI-COR Biosciences). Statistical analysis was performed using Graphpad Prism.

### Real Time Quantitative Polymerase Chain Reaction

Real time Quantitative Polymerase Chain Reaction (RT-qPCR) was performed according to the standardized protocol (Mehta and Singh, 2017; Mehta et al., 2019). Total RNA was extracted in 500  $\mu$ l of TRIzol Reagent (Thermo Fisher, Cat. No. # 15596926) from twenty pairs of third instar eye-antennal imaginal discs (n = 40), which were dissected from wild-type (WT), GMR>FUS, GMR>FUS-R518K and GMR>FUS-R521C background larvae. The quality of isolated RNA was determined by using the Nanodrop 2000 spectrophotometer (Thermo Scientific). Good quality samples had A260/A280 ratio greater than 2 and a peak at 260 nm. cDNA was produced from total RNA through RT-PCR using the first-strand cDNA synthesis kit (GE healthcare, Cat# 27926101). RT-qPCR was performed using iQ™ SYBR Green Supermix (Bio-Rad) and Bio-Rad iCycler (Bio-Rad) following the kits protocol for 25  $\mu$ l. Primers used for Jun-N-terminal kinase (JNK) are: (fwd: CCAACCGTCCGAAACTATGT); rev: CCGGCGGCTATTCTGATTATTA). The expression level of Glyceraldehyde 3-phosphate dehydrogenase (GAPDH) was used as an internal control to normalize the results (fwd: CAATGGATTTGGTCGCATCG; rev: CCGTTGACCACCAGGAAACC). The fold change was calculated relative to the expression level of the respective control (WT, Canton-S).

## Results

### Search for genetic modifier of human-FUS mediated neurodegeneration.

The larval eye imaginal discs develop into an adult compound eye comprised of nearly 800 ommatidia arranged in a stereotypical fashion (Fig. 1 A, F). We stained the eye-antennal imaginal discs with Disc large (Dlg; Green, a membrane specific marker) and pan neural marker Embryonic Lethal Abnormal Vision (Elav; Red), which marks the

nuclei of the photoreceptor neurons (Fig. 1A). Targeted misexpression of full length human-FUS (FUS) in differentiating retinal neurons using a Glass Multiple Repeat (GMR) eye specific promoter (GMR-Gal4/+), exhibits a strong retinal degeneration phenotype in the eye imaginal disc (n=10, Fig. 1C) and highly reduced adult eye with loss of eye pigmentation (n=200, Fig S1A, Fig. 1H). The control GMR-Gal4 driver alone does not show any neurodegenerative phenotype as seen in the eye imaginal disc and the adult eye (Fig. 1B, G, S1A). We tested other transgenic alleles of mutant FUS as well. Targeted misexpression of FUS R518K (Fig. 1D, I), and FUS R521C (Fig. 1E, J) also exhibits strong neurodegenerative phenotypes both in eye imaginal discs and the adult eye (Fig. S1A).

We verified if these transgenes were expressing human FUS protein (GMR>FUS, in heterozygous combination) in the *Drosophila* eye by using an antibody against human FUS protein. We found that FUS expression was restricted to the GMR expression domain of the developing third instar eye imaginal disc of GMR>FUS, GMR>FUS R518K, and GMR>FUS R521C (Fig. S2A-C, G). These results confirmed that the neurodegeneration seen in the eye-antennal imaginal discs and the adult fly eyes is due to misexpression of the human FUS protein. We also calculated the intensity of FUS signals as seen by immunohistochemistry detected by FUS antibody in eye-antennal imaginal disc and found that there was no significant difference of FUS levels in all three alleles (Fig. 1G). The protein extracts from these samples showed a significant accumulation of FUS protein in a semi-quantitative western blot (Fig. S2 H, H').

In order to understand the genetic basis of the FUS gain-of-function mediated neurodegeneration as seen in ALS, we employed a forward genetic screen using the molecularly defined deficiencies. We identified a deficiency line Df(2R)BSC782, which in heterozygous combination (Df(2R)BSC782/+) can rescue the GMR>FUS mediated neurodegeneration as seen in the eye imaginal disc and the adult eye (Fig. 1N, S). This molecularly defined *Df(2R)BSC782/+* deficiency is located on the right arm of the second chromosome, and uncovers *βTub56D*, *par-1*, *CG16926*, *CG7744*, *CG15120*, *mei-W68*, *oseg6*, *TBCB*, *rep*, and *hpo* genes (Fig. 1K). Targeted misexpression of other FUS transgenes in Df(2R)BSC782/+ background, which includes GMR>FUS R518K+*Df(2R)BSC782/+* (Fig. 1O, T) and GMR>FUS R521C + *Df(2R)BSC782/+* (Fig. 1P, U), exhibit a stronger rescue with a near complete wild-type phenotype in both the eye-antennal imaginal discs as well as the adult eyes. The heterozygous combination of deficiency Df(2R)BSC782/+ alone (Fig. 1L, Q) and GMR>Df(2R)BSC782/+ (Figure 1M, R) respectively, which served as controls, exhibit a wild-type phenotype in the eye-antennal imaginal disc and the adult eye. We also tested the levels of FUS expression levels in eye-imaginal discs from GMR>FUS, *Df(2R)BSC782/+*, GMR>FUS R518K+*Df(2R)BSC782/+* and GMR>FUS R521C + *Df(2R)BSC782/+*, where FUS-mediated neurodegenerative phenotypes are rescued (Fig.1). Interestingly, in this background, FUS levels were not significantly affected (Fig.S2D-G).

### ***hpo* acts as a genetic modifier of FUS mediated neurodegeneration.**

To narrow down the gene, which is responsible for modifying the FUS mediated neurodegeneration phenotype, we individually tested the genes uncovered by



Df(2R)BSC782 deficiency (Fig. 1K), using gain-of-function and loss-of-function approaches. Of these, modulation of *hippo* (*hpo*), one of the genetic loci uncovered by Df(2R)BSC782, affects the FUS neurodegenerative phenotype (Fig. 2). In comparison to the wild-type adult eye (Fig. 2A), and control GMR-Gal4/+ eye (Fig. 2B), gain-of-function of FUS as seen in GMR>FUS/+ (Fig. 1H, 2C), GMR>FUS R518K/+ (Fig. 1I, 2D), and GMR>FUS R521C/+ (Fig. 1J, 2E) results in a strong neurodegenerative phenotype (Lanson et al., 2011). These eyes are of reduced size and exhibit a rough eye phenotype, with disorganized lenses that clearly marks the onset of neurodegeneration. Gain-of-function of *hpo* in various backgrounds like FUS (GMR>FUS + *hpo*) (Fig. 2F), FUS R518K (GMR>FUSR518K+*hpo*; Fig. 2G), and FUS R521C (GMR>FUS R521C+*hpo*; Figure 2H) exhibit further enhancement in the severity of neurodegenerative phenotypes of FUS gain-of-function in the eye. On the contrary, loss-of-function of *hpo* by misexpressing *hpo*<sup>RNAi</sup>, with FUS (GMR>FUS+*hpo*<sup>RNAi</sup>), FUS R518K (GMR>FUS R518K+*hpo*<sup>RNAi</sup>), or FUS 521C (GMR>FUS R521C+*hpo*<sup>RNAi</sup>), rescues FUS mediated neurodegenerative phenotype (Fig. 2 I, J, K). These phenotypes were statistically significant (Fig. S1B). Our data strongly suggests that the *hpo* gene plays a role in triggering FUS mediated neurodegeneration in the *Drosophila* eye.

### Hippo signaling downregulation can rescue FUS mediated neurodegeneration

Hpo, a kinase, is one of the core components of an evolutionarily conserved Hippo signaling pathway. We therefore tested if this neuroprotective function is exclusive to *hpo* gene or if it is dependent on the Hippo signaling pathway. We therefore tested other members of the Hippo signaling pathway. Ectopic expression of FUS, FUS R518K, and FUS R521C in differentiating retinal neurons using a GMR driver, which results in a neurodegenerative phenotype in the eye-antennal imaginal discs (Fig.1) and the adult fly eye (Fig. 1, 2C, D, E). In comparison to the controls (Fig. 3A, B), activation of Hippo signaling via misexpression of *wts* and *yki*<sup>RNAi</sup> along with FUS (GMR>FUS+*wts*, GMR>FUS+*yki*<sup>RNAi</sup>), results in a strong enhancement of neurodegenerative phenotypes (Fig. 3F, I) as compared to the controls (Fig.3A, B, C). Similarly, activation of Hippo signaling in the gain-of-function of FUS R518K (GMR>FUS R518K+*wts*, Fig. 3G; GMR>FUS R518K+*yki*<sup>RNAi</sup>, Fig. 3J), or FUS R521C (GMR>FUS R521C+*wts*, Fig. 3H; and GMR>FUS R521C+*yki*<sup>RNAi</sup>, Fig. 3K) backgrounds in the GMR domain of eye worsens the FUS mediated neurodegeneration phenotype (Fig. 3C, D, E).

We further verified our hypothesis by downregulation of Hippo signaling using *wts*<sup>RNAi</sup> and *yki* misexpression in the FUS gain-of-function backgrounds. Downregulation of Hippo signaling in FUS (GMR>FUS+*wts*<sup>RNAi</sup>, Fig. 3L GMR>FUS+*yki*<sup>3SA</sup>, Fig. 3O), FUS R518K (GMR>FUS R518K+*wts*<sup>RNAi</sup>, Fig. 3M; GMR>FUS R518K+*yki*<sup>3SA</sup>, Fig. 3P), or FUS R521C (GMR>FUS R521C+*wts*<sup>RNAi</sup>, Fig. 3N; and GMR>FUS R521C+*yki*<sup>3SA</sup>, Fig. 3Q) results in overgrowth and in the case of *wts* showed rescue of the neurodegenerative phenotype. These phenotypes were statistically significant (Fig.S2C).

### Downregulation of Hippo signaling can block FUS mediated neuronal cell death.

We decided to test if Hippo signaling downregulation rescues FUS mediated neurodegeneration by blocking cell death. We used TUNEL staining, which marks the

fragmented DNA, to mark the nuclei of the dying neurons (Cutler et al., 2015; Gogia et al., 2017; McCall and Peterson, 2004; Sarkar et al., 2018b; Tare et al., 2011; White et al., 1994). We performed TUNEL staining in third instar larval eye-antennal imaginal discs of the wild-type larvae (Figure. 4A), GMR>FUS (Fig. 4B). The number of TUNEL positive cells, shown as white dots, were quantified from five eye-antennal imaginal discs (n=5) of each genotype. We took the mean, standard deviation, standard error, and p-values of this data and recorded it on a graph (Fig. 4I). Note that there are fewer TUNEL positive dying cells nuclei in the wild-type eye-antennal imaginal discs, which serves as control, as compared to the experiments (Fig. 4 A, I). Targeted misexpression of FUS (GMR>FUS) exhibits an almost two-fold increase in the number of TUNEL positive cells as compared to the control (Fig. 4 B, I). We also tested other components of the Hippo signaling pathway. Activation of Hippo signaling as seen in GMR>FUS+*hpo* (Fig. 4C), GMR>FUS+*wts* (Fig. 4E), GMR>FUS+*yki*<sup>RNAi</sup> (Fig. 4G) results in the enhancement of the neurodegenerative phenotype of GMR>FUS alone (Fig. 4I). Furthermore, activation of Hippo can result in a nearly two-fold increase in cell death as compared to the GMR>FUS alone (Fig. 4I). Inactivation of Hippo signaling in GMR>FUS background as seen in GMR>FUS+*hpo*<sup>RNAi</sup> (Fig. 4D), GMR>FUS+*wts*<sup>RNAi</sup> (Fig. 4F), GMR>FUS+*yki*<sup>3SA</sup> (Fig. 4H) resulted in the suppression of GMR>FUS mediated neurodegeneration. This suppression is accompanied by a significant decrease in the number of TUNEL positive cells or dying nuclei/ cell death in differentiating eye neurons compared to the wild-type (Fig. 4I). These observations confirm that downregulation of Hippo signaling blocks cell death in the differentiating photoreceptor neurons and thus plays a primary role of rescuing FUS mediated neurodegeneration.

### Loss of Hippo signaling restores impaired axonal targeting due to FUS expression.

During *Drosophila* eye development, the axonal projections from photoreceptor neurons innervates the medulla and lamina layers of the brain's optic lobe. We used a sensory neuron marker 24B10 (Chaoptin, DSHB), which marks the photoreceptor neurons and their axons (Fig. 5A) (Zipursky et al., 1984). Axonal targeting studies using Chaoptin staining is a qualitative approach to determine the functionality of the retinal neurons. We asked if the neurodegenerative phenotype seen in GMR>FUS also involves disruption of axonal targeting from the retina to the brain. We reasoned that if the downregulation of Hippo signaling that rescues the GMR>FUS phenotype in larval eye-antennal disc, it also may rescue the axonal targeting from the retina to the brain. In the wild-type eye-antennal imaginal discs, which serve as control, R1-R6 axons of the respective photoreceptors of an ommatidium innervates the lamina of the brain while R7 and R8 axons innervate the medulla (Meinertzhagen and Hanso, 1993; Newsome et al., 2000). It is known that when axonal targeting is disrupted it impairs axonal transport and leads to neurodegeneration observed in neurodegenerative diseases (Sarkar et al., 2018b). We investigated the effect of modulating *hpo* levels in the full-length FUS background on the axonal projections of photoreceptor retinal axons. In comparison to the wild-type eye-antennal imaginal disc (Fig. 5A), ectopic expression of FUS (GMR>FUS; Fig. 5B), FUSR518K (GMR> FUSR518K; Fig. 5C) and FUSR521C (GMR>FUSR521C; Fig. 5D) in the GMR domain resulted in a disorganized, defective and aberrant axonal targeting. Furthermore, gain-of-function of *hpo* along with FUS (GMR>FUS+*hpo*; Fig. 5E), FUS R518K (GMR> FUS R518K+ *hpo*; Fig.

5F) and FUS R521C (GMR>FUS R521C+ *hpo*; Fig. 5G) in the GMR domain severely impaired axonal targeting in larval eye-antennal imaginal discs as seen in (Fig. 5B, C, D). The axonal projections significantly decrease, and fail to innervate the lamina and medulla regions of the optic lobe (Fig. 5E, F, G). In contrast, loss-of-function of *hpo* signaling by misexpressing *hpo*<sup>RNAi</sup> in FUS (GMR>FUS+*hpo*<sup>RNAi</sup>; Fig. 5H), FUS R518K (GMR>FUS R518K+ *hpo*<sup>RNAi</sup>; Fig. 5I) and FUS R521C (GMR>FUS R521C+ *hpo*<sup>RNAi</sup>; Fig. 5J) background significantly restores retinal axonal targeting in the developing eye antennal imaginal discs.

We further extended our analysis to other members of Hippo signaling pathway to determine their role. Downregulation of Hippo signaling by misexpressing *wts*<sup>RNAi</sup>, *yki*<sup>3SA</sup> with FUS or mutant human FUS (GMR>FUS+*wts*<sup>RNAi</sup>, GMR>FUS R518K+*wts*<sup>RNAi</sup>, GMR>FUS R521C+*wts*<sup>RNAi</sup>, GMR>FUS+*yki*<sup>3SA</sup>, GMR>FUS R518K+*yki*<sup>3SA</sup> and GMR>FUS R521C+*yki*<sup>3SA</sup>), in the GMR domain of the developing eye also showed rescue of axonal targeting (Fig. S3). In contrast, activation of Hippo signaling by misexpressing *wts* or *yki*<sup>RNAi</sup> with FUS or mutant FUS (GMR>FUS+*wts*, GMR>FUS R518K+ *wts*, GMR>FUS R521C+*wts*, GMR>FUS+*yki*<sup>RNAi</sup>, GMR>FUS R518K+*yki*<sup>RNAi</sup> and GMR>FUS R521C+*yki*<sup>RNAi</sup>), in the GMR domain of the eye showed enhanced loss of the axonal targeting phenotype as compared to the controls (Fig. S3). These results confirm that downregulation of Hippo signaling in the FUS or mutant FUS backgrounds restores axonal targeting from the retinal neuron to the brain.

### Misexpression of FUS activates Hippo signaling

A transcriptional target of Yki, *death*- associated inhibitor of apoptosis (*diap1*), serves as a functional read out of the Hippo signaling pathway (Ren et al., 2010). *Drosophila inhibitor of apoptosis protein 1* (Diap1), is the direct transcriptional target of the Hippo pathway. The *diap1-4.3-GFP* gene, contains a Hippo response element, which drives the Diap1 expression in response to Hippo pathway activity where GFP is a reporter to study Hpo activity. When Hippo signaling is downregulated, which corresponds to the higher levels of Yki in the nuclei, activation of *diap1-4.3-GFP* levels (Ren et al., 2010) is triggered. In *diap1-4.3-GFP*, the GFP reporter expression is used to study Hpo activity.

Thus, when Hippo signaling is activated, *diap1-4.3-GFP* levels are decreased in the eye disc. We checked for *diap1-4.3-GFP* expression to further validate if Hippo signaling is activated in GMR>FUS background. The wild-type expression of *diap1-4.3-GFP* (tagged with a GFP reporter) is mostly restricted to the photoreceptor neurons present towards the posterior end of eye discs, and in the antennal region of eye-antennal imaginal disc (Fig. 6A, A'), as well as serves as the control. The GMR domain extends from the posterior margin of the eye disc to the Morphogenetic furrow (MF) which is marked by a proneural marker, ELAV (Fig. 6A, A', white arrowheads). Increasing the levels of FUS (GMR>FUS+*diap1-4.3-GFP*, Fig. 6B, B'), FUS R518K (GMR>FUS R518K+*diap1-4.3-GFP*, Fig. 6C, C') and FUS R521C (GMR>FUS R521C+*diap1-4.3-GFP*; Fig. 6D, D') in the GMR domain resulted in significant reduction of Diap1 levels (Fig. 6E). These results confirm that an increase in levels of FUS protein does not affect the expression of downstream targets of Hippo signaling.

To determine if Hippo signaling is activated in mammalian cells, we checked the levels of LATS1 and phosphorylated LATS1, one of the components of Hippo signaling cascade, in HEK293T cells ectopically expressing either FUS-WT or ALS-associated mutants, FUS-R518K and FUS-R521C. We observed that while the total levels of LATS1 protein were unchanged between the groups (Fig. 6F,F’), the levels of phosphorylated LATS1 were increased in the mutant FUS-expressing cells when compared to the untransfected control (Fig. 6F,F’), suggesting a hyper-activation of upstream kinase cascade. Ectopic expression of FUS-WT alone did not change the phosphorylated LATS1 levels (Fig 6F, F’), indicating that activation of Hippo signaling was specific to mutant FUS.

### Modulating levels of JNK signaling pathway modulates FUS mediated neurodegeneration

It is known that Hippo regulates JNK signaling (Ma et al., 2015; Ma et al., 2017), which is involved in neurodegeneration and cell death (Flerdegen et al., 1997; Sarkar et al., 2016; Tare et al., 2011). We tested if modulating JNK signaling levels can block FUS-mediated neurodegeneration. JNK signaling comprises of a series of kinases, which in turn regulates expression of *puckered* (*puc*) (Fig. 7A). *puc*, a dual phosphatase, regulates JNK signaling through a negative feedback loop (Martin-Blanco et al., 1998). The phospho-Jun kinase enzyme can phosphorylate its downstream substrate Jun in its N-terminal domain. The phosphorylated Jun is used to quantify levels of JNK activation using an antibody against p-Jun (Tare et al., 2011). Initially we tested JNK activation levels by quantifying the JNK levels with quantitative-PCR (qPCR) (Fig. 7B) and then quantifying levels of the p-JNK protein in semi-quantitative western blot (Fig. 7C, C’). In GMR>FUS background, JNK gene expression levels increased two fold, whereas in other constructs like GMR>FUS R518K or GMR>FUS R521C, the JNK expression levels were 10 and 4 folds respectively (Fig. 7B). We also employed a semi-quantitative approach to compare the amount of phospho-Jun kinase (p-JNK) levels in the wild type to that of FUS (GMR>FUS) and mutant FUS (GMR>FUS R518K and GMR>FUS R521C). In comparison to the wild type, p-JNK levels were upregulated in GMR> FUS, GMR> FUS R518K and GMR>FUS R521C backgrounds. Tubulin bands served as a control in the western blot. We calculated the intensity of the bands and presented them as a graph (Figure 7C, C’).

To further test if FUS mediated neurodegeneration can be suppressed by modulating the levels of JNK signaling, we initially activated JNK signaling by misexpressing *Djun<sup>aspv7</sup>* and constitutively active *hemipterous* (*hep<sup>Act</sup>*) in FUS and mutant FUS (FUS R518K and FUS R521C) backgrounds. In comparison to the wild-type eye imaginal disc and the adult eye (Fig. 7D, D’), we analyzed their resultant phenotypes in both eye-antennal imaginal discs (Fig. 7 F, G, H, J, K, L) and the adult fly eyes (Fig. 7 F’, G’, H’ J’,K’,L’). Targeted misexpression of *D-jun* and *hep* alone in the GMR domain (GMR>*jun* and GMR>*hep*), which served as control, led to a strong neurodegenerative phenotype in eye-antennal imaginal discs (Fig. 7E, I), adult fly eyes (Fig. 7 E’, I’). Moreover, activation of the JNK signaling by misexpressing *hep<sup>Act</sup>* along with FUS or mutant FUS in the GMR domain worsens the neurodegeneration phenotype (Fig. 7J-L’) as seen in eye imaginal discs (Fig. 7D) and adult fly eyes (Fig. 7D’). Similarly, misexpression of *Djun<sup>aspv7</sup>* with FUS or mutant FUS in the GMR domain (GMR>FUS+*Djun<sup>aspv7</sup>*, GMR>FUS R518K+*Djun<sup>aspv7</sup>* and GMR>FUS R521C+*Djun<sup>aspv7</sup>*), resulted in a small, rough eye phenotype with loss of

pigment cells (Fig. 7F-H'). We further validated our results by downregulating the JNK signaling in FUS gain-of-function backgrounds. Misexpression of *bsk<sup>DN</sup>* (*GMR>bsk<sup>DN</sup>*) or *puc* (*GMR>puc*) alone, which serve as controls, exhibits near wild-type eye imaginal discs (Fig. 7M,Q) and adult fly eyes (Fig. 7 M' Q'). Downregulating JNK by misexpressing *bsk<sup>DN</sup>* with FUS in GMR domain (*GMR>FUS+bsk<sup>DN</sup>*) or FUS R518K (*GMR>FUS R518K+bsk<sup>DN</sup>*), *GMR>FUS R521C* (*GMR>FUS R521C+bsk<sup>DN</sup>*), exhibits a strong rescue of the FUS mediated neurodegenerative phenotype (Fig. 7N-P'). Similarly, when *puc* was misexpressed with FUS or mutant FUS (*GMR>FUS+puc*) or FUS R518K (*GMR>FUS R518K+puc*), *GMR>FUS R521C* (*GMR>FUS R521C+puc*) as seen in the eye imaginal discs (Fig. 7 R, S, T) and adult fly eyes (Fig. 7 R', S', T). Thus, activation of JNK signaling is responsible in FUS mediated neurodegeneration.

## Discussion

ALS, a progressive neurodegenerative disorder, is caused by mislocalization of FUS in cytoplasm which impairs the RNA metabolism (Lanson et al., 2011). However, the molecular genetic basis of this disease is elusive. It is possible that mislocalization of FUS proteins trigger(s) aberrant signaling, which results in the onset of the progressive neurodegenerative phenotype. Therefore, there is a need to identify the downstream targets of signaling pathways activated by the FUS mislocalization. Generation of animal model systems like *Drosophila*, which are genetically tractable, have facilitated the use of genetic screens to identify the modifiers of neurodegenerative phenotypes (Casci and Pandey, 2015; Pandey and Nichols, 2011; Sarkar et al., 2016).

Targeted misexpression of FUS and its mutants in the differentiating retinal neurons using transgenic approaches exhibited strong neurodegenerative phenotypes in the *Drosophila* adult eye (Daigle et al., 2013; Lanson et al., 2011). In a forward genetic screen using the molecularly defined deficiencies, we identified a deficiency line *Df(2R)BSC782*, which can rescue FUS mediated neurodegeneration. This deficiency uncovers *hpo* and other genes (Fig.1). In a candidate gene approach, where we tested individual genes uncovered by the deficiency *Df(2R)BSC782*, we identified *hpo* as one of the genetic modifiers of ALS phenotypes (Fig. 2). Interestingly, it has been shown that FUS/ LATS1/2 inhibits hepatocellular carcinoma (HCC) via activating Hippo pathway (Bao et al., 2018). Gain-of-function of *hpo* results in cell death whereas *hpo* loss-of-function results in over proliferation and overgrowth (Kango-Singh and Singh, 2009; Udan et al., 2003). We found that modulating the levels of *hpo* can modify FUS mediated neurodegeneration phenotype. Loss-of-function of *hpo* can significantly rescue the neurodegenerative phenotype of FUS (*GMR>FUS*) and other mutant (*GMR>FUS R518K* and *GMR>FUS R521C*) accumulation (Fig. 2). We further confirmed that the Hippo pathway is activated in FUS overexpressing retinal neurons (*GMR>FUS*) to trigger neurodegeneration in the *Drosophila* eye (Fig. 3).

### Role of Hippo in neurodegeneration

Studies from ours and other labs have shown that other components of Hippo pathway like Fat (Ft), Teashirt (Tsh), and Crumbs (Crb) induce neurodegeneration in models of Alzheimer's Disease (Moran et al., 2013; Napoletano et al., 2011; Sarkar et al., 2016;

Steffensmeier et al., 2013). In addition, the Hippo pathway may also be involved in neural development (Jiang et al., 2009; Wittkorn et al., 2015). The major function of the Hippo signaling pathway has been in growth regulation and cancer (Kango-Singh and Singh, 2009; Ma et al., 2019; Snigdha et al., 2019). Recently, Hippo signaling has been implicated in many disease models where it plays role in apoptosis, autophagy, regeneration and cell survival (Calamita and Fanto, 2011; Ma et al., 2019). Thus, it is interesting to find a role for Hippo pathway in FUS mediated neurodegeneration.

Our genetic analysis suggested that Hippo signaling acts downstream of FUS accumulation (Fig. S2). In genetic backgrounds where FUS and its mutant proteins were misexpressed in the eye, we found a robust accumulation of FUS protein in the eye imaginal disc. In *GMR>FUS, Df(2R)BSC782/+* background, FUS accumulation was unaltered suggesting that the rescue of the neurodegeneration phenotype occurred by a mechanism downstream of FUS accumulation (Fig.S1). Our data shows that FUS accumulation triggers downstream signaling events involving Hippo and JNK signaling. We tested the levels of *diap1-4.3-GFP*, which serves as a reporter for Hippo signaling, in *GMR>FUS* and *GMR>FUS R518K* and *GMR>FUS R521C* background. When Hippo signaling is activated it triggers cell death and *diap1-4.3-GFP* levels are downregulated (Zhang et al., 2008). We found that *diap1-4.3-GFP* levels were downregulated in these backgrounds, which suggests that Hippo signaling is activated (Fig. 6). Earlier studies have shown that Hippo can regulate JNK signaling (Ma et al., 2015; Ma et al., 2017). We also tested the reporters of JNK signaling and found that JNK signaling is activated in gain-of-function of FUS (Fig. 7).

Furthermore, if we block these signaling pathways, even though the neurodegenerative phenotype caused by accumulation of FUS is rescued, it does not affect the accumulation of FUS protein suggesting that JNK signaling doesn't block expression or turnover of toxic FUS protein. Interestingly, JNK signaling is dependent on timing (temporal scale). We have assayed the effect of JNK signaling in the mature third instar eye-imaginal disc. Based on our results, we propose a model that accumulation of FUS triggers Hippo and JNK signaling, which acts downstream of FUS accumulation. Activation of Hippo signaling and JNK signaling triggers neurodegeneration (Fig. 8).

Thus, we demonstrate that inactivation of Hippo and JNK signaling can rescue FUS-mediated neurodegeneration in *Drosophila* eye. Our data suggests that this neuropathy can be alleviated by modulating levels of Hippo signaling and JNK signaling pathways (Fig. 2, 3, 7). Both pathways are crucial for normal development and in disease backgrounds for onset and progression. The members of these two pathways seem to be interesting therapeutic targets, which can mitigate the onset or rapid progression of neurodegenerative diseases like ALS.

MST, the mammalian homologue of *hpo*, a kinase, is involved in many biological processes from cell proliferation to cell death, patterning and growth (Kango-Singh and Singh, 2009; Yu and Guan, 2013). There are two *hpo* homologues MST1 and MST2 in the mammalian genome (Creasy and Chernoff, 1995). In addition, *hpo* is involved in other neurodegenerative models like the superoxide dismutase SOD1 G93A mouse model of ALS (Lee et al., 2013), and in VAPB models in *Drosophila* (Sanhueza et al., 2015). The SOD1

G93A mouse model of ALS is linked to decrease in motor neuron survival. It has been shown that downregulation of MST1, fly homolog of *hpo*, exhibits neuroprotective function in this mouse ALS model (Lee et al., 2013). Hippo signaling dysregulation has been identified in Huntington's disease brain and neuronal stem cells (Mueller et al., 2018). Thus, MST1/ Hippo can be an interesting link between cancer and neurodegenerative disorder like ALS (Yamamoto et al., 2019). The members of Hippo pathway can serve as potential therapeutic target for ALS as well as other neurodegenerative disorder.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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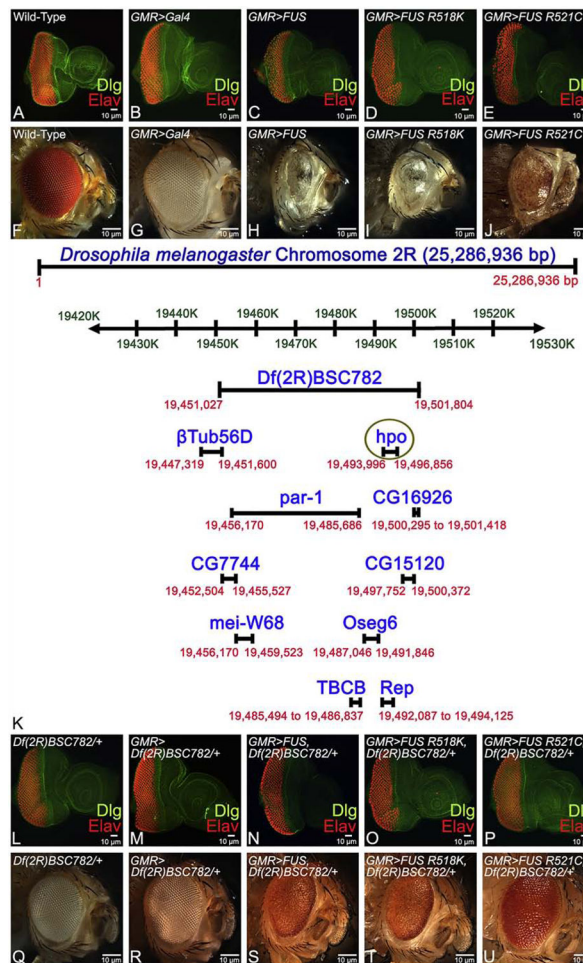
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**Figure 1. Identification of a molecularly defined deletion as a genetic modifier of human FUS mediated neurodegeneration.**

Eye imaginal disc and the adult eye of (A, F) Wild-type and (B, G) *GMR>Gal4/+* (an eye specific promoter) that serve as control. (A) Third instar larval eye antennal imaginal disc, which develop into (F) the adult compound eye. Note that the eye-antennal imaginal discs are stained with a membrane specific marker Disc large (Dlg: Green) and pan neural marker Embryonic Lethal Abnormal Vision (Elav: Red), which marks the nuclei of the photoreceptor neurons. In comparison to controls, misexpression of (C, H) FUS (*GMR>Gal4/CyO; UAS-FUS/ TM3Sb e Ser, GMR>FUS*) or mutant FUS (D, I) FUS R518K (*Gal4/CyO; UAS-FUS R518K/ TM3Sb e Ser, GMR>FUS R518K*), and (E, J) FUS R521C (*Gal4/CyO; UAS-FUS R521C/ TM3Sb e Ser, GMR>FUS R521C*) in differentiating retinal neurons using *GMR>Gal4* driver results in strong neurodegeneration in retinal neurons as seen in (C, D, E) eye imaginal discs and the (H, I, J) adult eye. Note that phenotype worsens as the (C, D, E) larval eye-antennal imaginal discs transforms into (H, I, J) adult eyes. In a deficiency screen, *Df(2R)BSC782* was identified as a genetic modifier of FUS mediated neurodegenerative phenotype. (K) The map shows the region and genes (including *hpo*) covered by deficiency line (*Df(2R)BSC782/+*). (L, Q) *Df(2R)BSC782/+* (M, R) *GMR>Df(2R)BSC782/+* served as controls. In a heterozygous background of (*Df(2R)BSC782/+*), misexpression of (N, S) FUS (*GMR>FUS/+; Df(2R)BSC782/+*), or

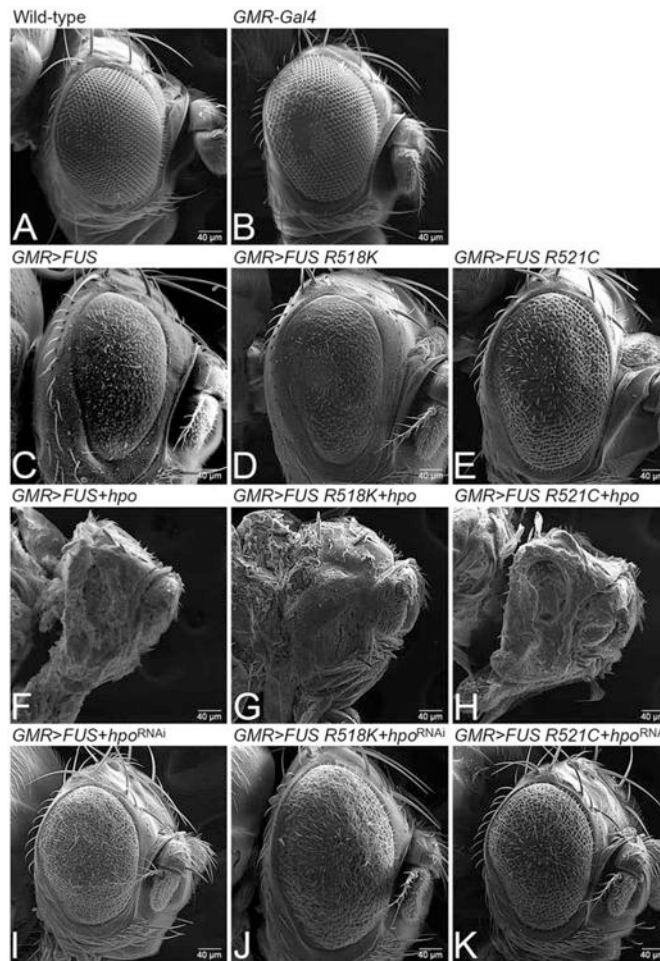
mutant FUS, (O, T) FUS R518K (GMR>FUS R518K/+ ; *Df(2R)BSC782/+*) and (P, U) FUS R521C (GMR>FUS R521C/+; *Df(2R)BSC782/+*), results in the significant rescue of FUS mediated neurodegeneration as seen in (N, O, P) the eye-antennal imaginal discs and (S, T, U) the adult compound eyes of *Drosophila* respectively. The neurodegeneration phenotype in the eye was significantly rescued and pigment cells were restored. The orientation of all imaginal discs in the figure is posterior to left and dorsal up. Magnification of all eye discs is 20X.

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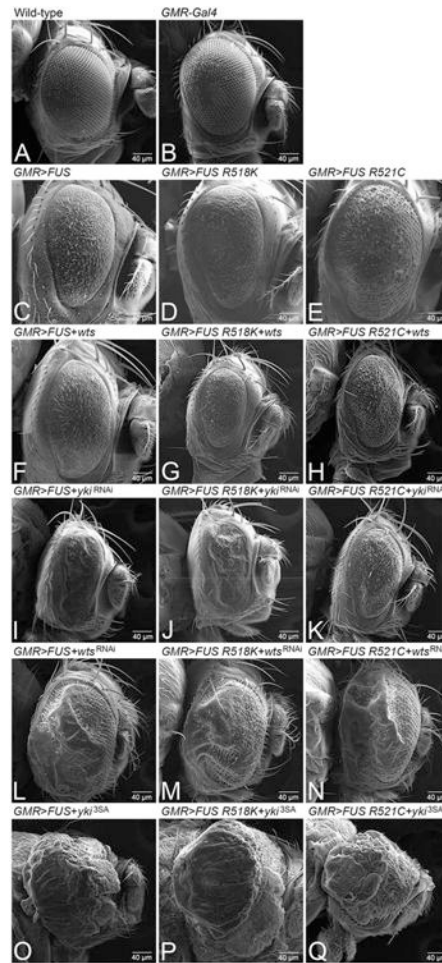
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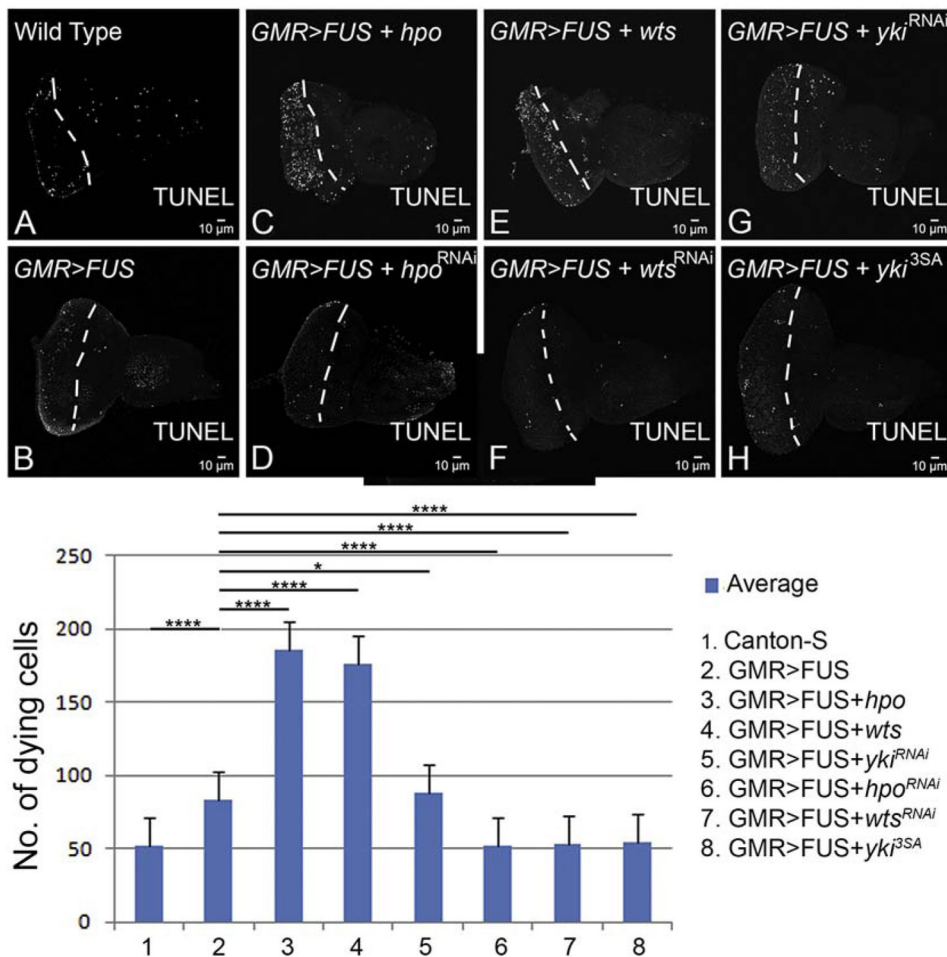
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**Figure 2. *hpo* is the genetic modifier of FUS mediated neurodegeneration in *Drosophila* eye.** (A) Wild-type and (B) *GMR-Gal4*/*+* adult fly eyes serve as controls. Targeted misexpression of FUS or mutant FUS R518K and FUS R521C in the GMR domain of eye (C) *GMR>FUS* (D) *GMR>FUS R518K* (E) *GMR>FUS R521C* results in reduced and rough eye phenotype due to neurodegeneration. Activation of Hippo signaling by misexpressing *hpo* along with FUS or mutant FUS (FUS R518K and FUS R521C) in the GMR domain of eye using GMR driver, (F) *GMR>FUS + hpo*, (G) *GMR>FUS R518K+ hpo*, (H) *GMR>FUS R521C +hpo*, further enhances the FUS mediated neurodegeneration phenotype as compared to misexpressing FUS or mutant FUS in GMR domain alone. Downregulation of Hippo signaling by misexpressing *hpo<sup>RNAi</sup>* along with FUS (I) *GMR>FUS+ hpo<sup>RNAi</sup>*, or mutant FUS (J) *GMR>FUS R518K+ hpo<sup>RNAi</sup>*, (K) *GMR>FUS R521C+hpo<sup>RNAi</sup>* in the GMR domain rescues the FUS mediated neurodegeneration, as seen in SEM images of the adult eyes of FUS or mutant FUS in GMR domain alone. Note that all transgenes are in heterozygous combination. Magnification of all the scanning electron microscopic images of adult fly eyes is 180X.



**Figure 3. Downregulation of Hippo signaling rescues FUS mediated neurodegeneration.** (A) Wild-type and (B) GMR-Gal4/+ adult fly eyes as controls. Targeted misexpression of WT FUS or mutant FUS R518K and FUS R521C in the GMR domain of eye (C) GMR>FUS (D) GMR>FUS R518K (E) GMR>FUS R521C results in rough eye or neurodegeneration phenotype. Activation of Hippo signaling by misexpressing *wts* or *yki<sup>RNAi</sup>* along with full length wild-type FUS or mutant FUS (FUS R518K and FUS R521C) in the GMR domain of eye using GMR driver, (F) GMR>FUS+ *wts*, (G) GMR>FUS R518K+ *wts*, (H) GMR>FUS R521C+ *wts*, (I) GMR>FUS+ *yki<sup>RNAi</sup>*, (J) GMR>FUS R518K+ *yki<sup>RNAi</sup>*, (K) GMR>FUS R521C+ *yki<sup>RNAi</sup>*, further enhances or worsens the FUS or mutant FUS alone mediated neurodegeneration phenotype. In-activation of Hippo signaling by misexpressing *wts<sup>RNAi</sup>*, *yki<sup>3SA</sup>* along with full length wild-type FUS (L) GMR>FUS+ *wts<sup>RNAi</sup>*, (O) GMR>FUS+ *yki<sup>3SA</sup>*, or mutant FUS (M) GMR>FUS R518K+ *wts<sup>RNAi</sup>*, (N) GMR>FUS R521C+ *wts<sup>RNAi</sup>*, (P) GMR>FUS R518K+ *yki<sup>3SA</sup>*, (Q) GMR>FUS R521C+ *yki<sup>3SA</sup>* in GMR domain rescues the FUS mediated neurodegeneration, as seen in scanning electron microscopic (SEM) images of adult fly eyes of FUS alone or mutant FUS. Note that all transgenes are in heterozygous combination. Magnification of all the SEM images of the adult fly eyes is 180X.

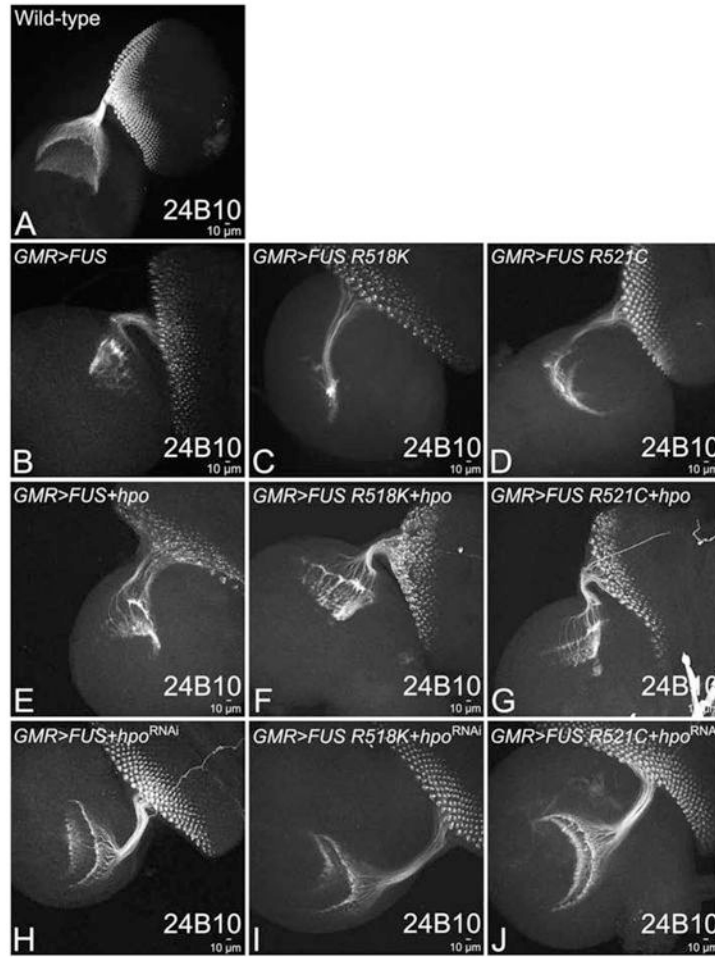


**Figure 4. Downregulation of Hippo signaling blocks cell death in the differentiating neurons to rescue FUS mediated neurodegeneration.**

TUNEL, marks the fragmented DNA within the nuclei of dying cells (also called as TUNEL positive cells). TUNEL staining was carried out in (A) wild-type third instar larval eye-antennal imaginal disc, displaying randomly distributed TUNEL positive dying cells shown as white dots. The area between the white dotted line and posterior margin of the eye imaginal disc is the domain of *GMR*-Gal4 driver. The number of TUNEL positive cells were counted from five eye-antennal imaginal discs ( $n=5$ ) of all the genotypes in the *GMR*-Gal4 domain. (B) Misexpression of *FUS* in the differentiating photoreceptor neurons of the eye using *GMR*-Gal4, eye specific promoter (*GMR>FUS*), show elevated levels of TUNEL positive cells, indicating increased frequency of cell death in photoreceptor neurons, as compared to the wild-type control (A,I). Note that the number of dying cells (TUNEL positive cells) increase nearly ~0.5 fold in (B) *GMR>FUS* as compared to the (A) wild-type eye-antennal imaginal discs. Activation of Hippo signaling by misexpressing *hpo*, *wts*, *yki<sup>RNAi</sup>* in WT *FUS* background, in *GMR* domain of eye in (C) *GMR>FUS+hpo*, (E) *GMR>FUS+wts*, and (G) *GMR>FUS+yki<sup>RNAi</sup>*, results in further increase in the number of dying retinal neurons (or TUNEL positive cells). On the other hand, downregulation of Hippo signaling by misexpressing *hpo<sup>RNAi</sup>*, *wts<sup>RNAi</sup>*, *yki<sup>3SA</sup>* in full length wild-type *FUS* background in (D) *GMR>FUS+hpo<sup>RNAi</sup>*, (F) *GMR>FUS+wts<sup>RNAi</sup>*,

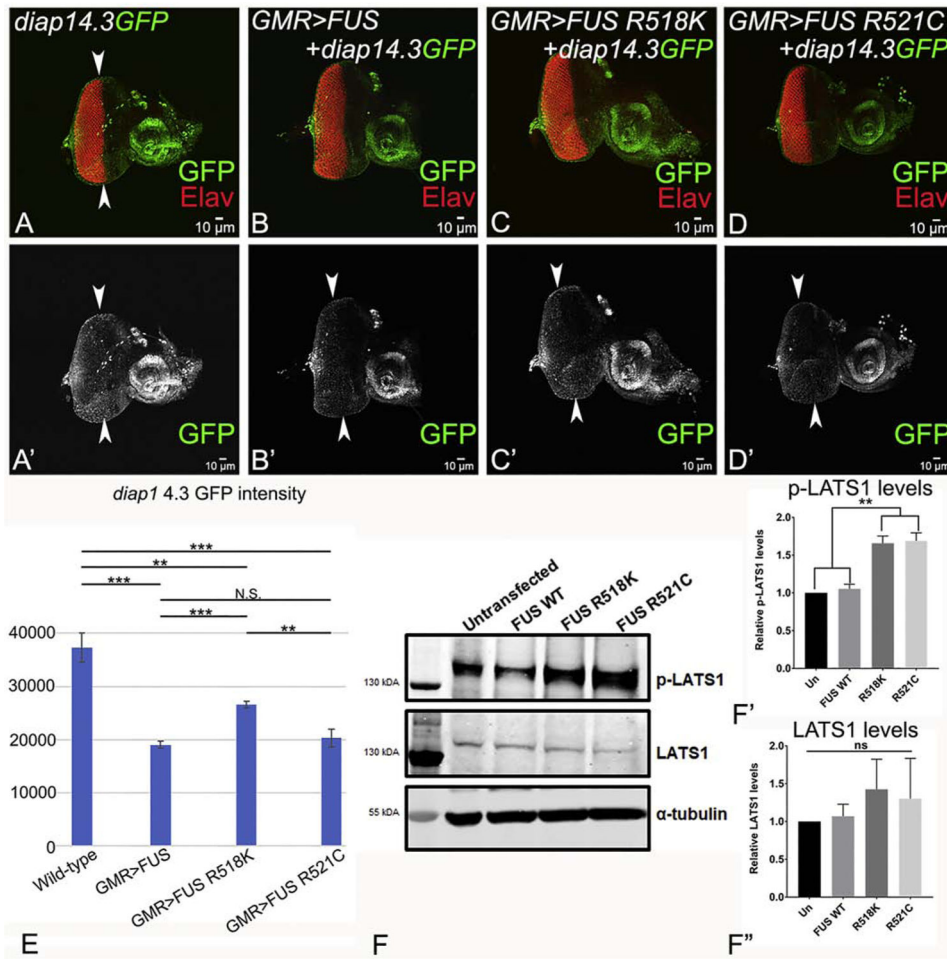


and (H)  $GMR>FUS+yki^{3SA}$ , results in significant reduction in the dying retinal neurons. (I) A graph representing the Quantitative analysis (statistics). Comparing the number of dying nuclei of neurons, when FUS is ectopically expressed alone, and when FUS is expressed together with components of Hippo pathway in the eye, clearly shows that downregulation of Hippo signaling significantly rescues FUS mediated neurodegeneration by blocking neuronal cell death. The p-values for the estimation of cell death were calculated between  $GMR>FUS$  and all the components of Hippo pathway using Student's two-tailed t-test (n=5) in Microsoft excel software.  $GMR>FUS$  phenotype was found to be statistically significant with respect to Wild-type (\*\*\*\*,  $p<0.0001$ ). The genotypes including  $GMR>hpo+FUS$  (\*\*\*\*,  $p<0.0001$ ),  $GMR>FUS+hpo^{RNAi}$  (\*\*\*\*,  $p<0.0001$ ),  $GMR>FUS + wts$  (\*\*\*\*,  $p<0.0001$ ),  $GMR>FUS + wts^{RNAi}$  (\*\*\*\*,  $p<0.0001$ ) and (G)  $GMR>FUS + yki$ , (\*\*\*\*,  $p<0.0001$ ),  $GMR>FUS + yki^{RNAi}$  (\*,  $p<0.05$ ). Note that all transgenes are in heterozygous combination. Magnification of all eye-antennal imaginal discs is 20X.



**Figure 5. Loss of function of Hpo restores axonal projections (impaired by gain of function of FUS) from retina to the brain.**

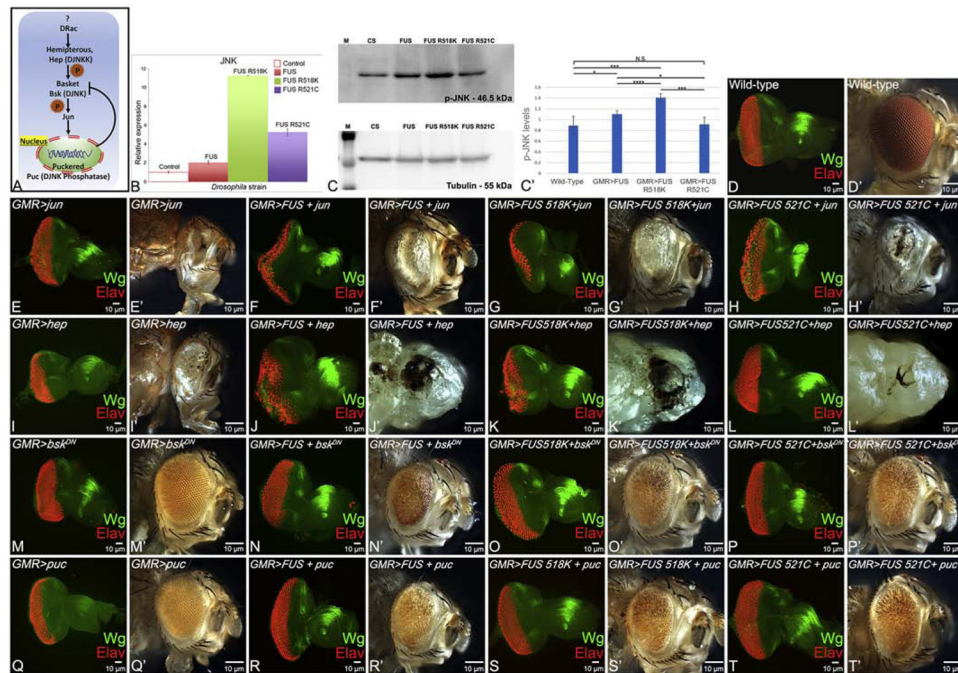
MAb24B10 (Chaoptin), marks the retinal axons that innervate lamina and medulla regions of the brain. Top panel shows, (A) wild-type larval eye-antennal imaginal disc, and (B) GMR>FUS (C) GMR>FUS R518K, and (D) GMR>FUS R521C showing expression of the retinal axons marked by 24B10 in control and when FUS or mutant FUS (FUS R518K and FUS R521C) are misexpressed in the GMR domain respectively. Activation of Hippo signaling by misexpressing *hpo*, along with FUS or mutant FUS results in impaired targeting of retinal axons to the brain (E) GMR>FUS+*hpo*, (F) GMR>FUS R518K+*hpo*, (G) GMR>FUS R521C+*hpo*, as compared to the wild-type (A). In-activation of Hippo signaling by misexpressing *hpo*<sup>RNAi</sup>, along with FUS and mutant FUS results in restoration of targeting of retinal axons to the brain (H) GMR>FUS+*hpo*<sup>RNAi</sup>, (I) GMR>FUS R518K+*hpo*<sup>RNAi</sup>, (J) GMR>FUS R521C+*hpo*<sup>RNAi</sup>. Note that all transgenes are in heterozygous combination. Magnification of all the eye-antennal imaginal discs is 20X.



**Figure 6. Targeted misexpression of FUS in the developing eye of *Drosophila*, transcriptionally activates Hippo signaling.**

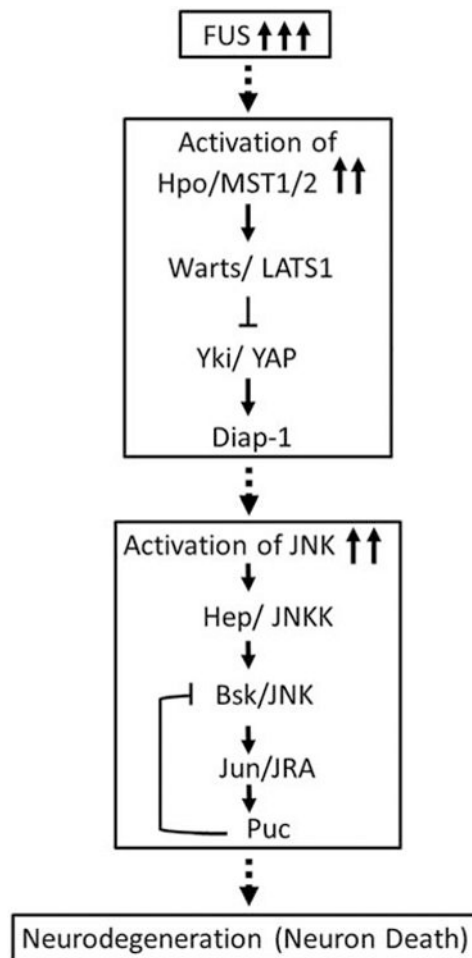
*diap1*-4.3-GFP (*diap1*, tagged with GFP reporter) shows the transcriptional activity of Hippo signaling pathway. Gain-of-function of *hpo* triggers cell death and result in downregulation of *diap1*-4.3-GFP levels. Panels shows *diap1*-4.3-GFP (green) and Elav (red) expression in the eye-antennal imaginal discs from larvae of all the genotypes. (A, A') Wild-type expression of *diap1*-4.3-GFP in eye-antennal imaginal discs, which serve as control. The white arrowheads marks the boundary of domain of GMR expression domain on dorsal and ventral margins of the eye disc. Misexpression of FUS or mutant FUS (R518K and FUS R521C) along with *diap1*-4.3-GFP in the GMR domain of eye (B, B') GMR>FUS+ *diap1*-4.3-GFP, (C, C') GMR>FUS R518K+ *diap1*-4.3-GFP, (D, D') GMR>FUS R521C+ *diap1*-4.3-GFP, exhibits downregulation of *diap1*-4.3-GFP reporter. (E) Quantitative analysis, comparing the GFP intensity levels in the eye imaginal disc of (A) *diap1*-4.3-GFP, which serve as control, (B) GMR>FUS, (C) GMR>FUS R518K, and (D) GMR>FUSR521C. The quantification of GFP intensity was done using in built program in Olympus Fluoview laser scanning confocal microscope. The p- values for the estimation of *diap1* 4.3-GFP intensity levels were calculated from five individual eye-imaginal disc for each genotype using a Student's t- test in MS excel and a graph was plotted. (A, A', B', C', D') The *diap1* 4.3-GFP signal intensity was calculated only in the GMR domain, which

extends from the posterior margin of the eye imaginal disc to the MF which is marked by white arrowheads. The data exhibits that GFP levels are reduced in the backgrounds where FUS targeted expression occurs. Genotypes including, GMR>FUS+ *diap1-4.3-GFP* (\*\*\*, p<0.001), GMR>FUS R518K+ *diap1-4.3-GFP* (\*\*, p<0.01) and GMR>FUS R521C+ *diap1-4.3-GFP* (\*\*\*, p<0.001) were found to be statistically significant from *diap1-4.3-GFP* (control). GMR>FUS+ *diap1-4.3-GFP* was found to be statistically significant from GMR>FUS R518K+ *diap1-4.3-GFP* (\*\*\*, p<0.001) while non-significant from GMR>FUS R521C+ *diap1-4.3-GFP* (N.S., p>0.05). GMR>FUS R518K+ *diap1-4.3-GFP* was found to be statistically significant as compared to GMR>FUS R521C+ *diap1-4.3-GFP* (\*\*, P<0.01). Note that all transgenes are in heterozygous combination. The magnification of all the eye-antennal imaginal discs is 20X. (F, F', F'') Semi-quantitative western blot (the tubulin bands in western blot, serve as controls) and (F', F'') quantitative analysis of western blot represented in the form of a graph (provides the levels of (F') p- LATS1 and (F'') LATS1 levels in untransfected HEK293T cells (ATCC® CRL-3216™) (negative control), HEK293T cells (ATCC® CRL-3216™) cells transfected with WT FUS, mutant FUS (FUS R518K and FUS R521C). The intensity of the (F') p-LATS1, and (F'') LATS1 bands was quantified, normalized by image studio ( Li-COR Biosciences), and the graph was made using Graphpad Prism. The p-values for the estimation of p-JNK levels in all combinations in Western Blot were calculated in a set of three (n=3), using one-way ANOVA with Tukey's multiple comparisons test. The p-values for untransfected versus FUS R518K = 0.0015 and p-value for untransfected versus R521C = 0.0011. Errors bar indicate s.e.m.



**Figure 7. Modulating the levels of JNK signaling affects FUS mediated neurodegeneration.** Panel shows, (A) Schematic representation of JNK signaling pathway, (B) levels of phospho-JNK (p-JNK) in quantitative-PCR (q-PCR), (C) semi-quantitative western blot (the tubulin bands in western blot, serve as controls) and (C') quantitative analysis of western blot represented in the form of a graph (provides the levels of JNK in WT FUS and mutant FUS (FUS R518K and FUS R521C) background). (C') The p-JNK bands intensity was quantified, normalized and the graph was made using Image J software (NIH). The p-values for the estimation of p-JNK levels in all combinations in Western Blot were calculated in a set of three (n=3), using Student's t-test in MS Excel software. GMR>FUS was found to be statistically significant from 1. Wild-type ( $p < 0.05$ , \*), 2. GMR>FUS R518K ( $p < 0.0001$ , \*\*\*\*), and 3. GMR>FUS-R521C ( $P < 0.001$ , \*\*\*). Genotypes including GMR>FUS, GMR>FUS-R518K were found to be statistically different ( $p < 0.05$ , \*) and ( $p < 0.001$ , \*\*\*) respectively, while GMR>FUS-R512C was found to be non-significant ( $p > 0.05$ , N.S.) as compared to Wild-type (Canton-S, Control). (B, C, C') When compared to wild-type controls, higher levels of p-JNK were observed when FUS or mutant FUS are misexpressed in GMR domain of the eye (GMR>FUS, GMR>FUS R518K and GMR>FUS R521C) in q-PCR and western blot approaches. (D) Wild-type eye-antennal imaginal disc and (D') adult fly eye, serve as the control. Note the expression of Wg (green) and Elav (red) in the eye-antennal imaginal discs from third instar larvae of all the genotypes. Activation of JNK signaling by misexpressing activated *hemipterous* (*hep*, GMR>*hep*<sup>Act</sup>), and activated *Djun* (GMR>*jun*<sup>aspv7</sup>) alone in GMR domain of the eye, results in few number of dying cells in (I,E) eye imaginal discs and (I',E') adult fly eye, respectively. Furthermore, activation of JNK signaling by misexpressing *hep* and *jun* in FUS or mutant FUS background using GMR-Gal4 driver, (J,J') GMR>FUS+*hep*<sup>Act</sup>, (K,K') GMR>FUS R518K+*hep*<sup>Act</sup> and (L,L') GMR>FUS R521C+*hep*<sup>Act</sup>, (F,F') GMR>FUS+*jun*<sup>aspv7</sup>, (G,G') GMR>FUS R518K+*jun*<sup>aspv7</sup> and (H,H')

GMR>FUS R521C+*jun<sup>aspv7</sup>*, results in (J,K,L,F,G,H) dramatic increase in dying cell population in the eye-antennal imaginal discs, leading to (J',K',L',F',G',H') no-eye or small eye with loss of pigmentation phenotype in the adult fly eyes. However, downregulation of JNK signaling by misexpression of dominant negative *bsk*, and *puc* (a dual phosphatase) alone in GMR domain (GMR>*bsk<sup>DN</sup>* and GMR>*puc*), results in near wild-type (M,Q) eye-antennal imaginal discs and (M',Q') adult fly eyes respectively. However, downregulating JNK signaling by misexpressing *bsk*, and *puc* in FUS or mutant FUS background (N,N') GMR>FUS+*bsk<sup>DN</sup>*, (O,O') GMR>FUS R518K+*bsk<sup>DN</sup>*, (P,P') GMR>FUS R521C+*bsk<sup>DN</sup>* and (R,R') GMR>FUS+*puc*, (S,S') GMR>FUS R518K+*puc* and (T,T) GMR>FUS R521C+*puc*, results in significant reduction in the dying cell population in (N,O,P,R,S,T) eye imaginal discs, thereby leading to a strong rescue of the (N',O',P',R',S',T') adult fly eye phenotype. The magnification of all eye-antennal imaginal disc is 20X.



**Figure 8. Model representing the mechanism by which FUS leads to neurodegeneration and cell death.**

Increase in the levels of human FUS or mutant FUS, FUS R518K and FUS R521C activates Hippo signaling, which in turn leads to the activation of JNK signaling. This aberrant Hippo and JNK signaling due to accumulation of FUS triggers cell death in the retinal neurons of eye. Thus, downregulation of Hippo and JNK signaling in FUS background, rescues FUS-mediated neurodegeneration.