

# The zebrafish *sf3b1*<sup>b460</sup> mutant reveals differential requirements for the *sf3b1* pre-mRNA processing gene during neural crest development

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**ABSTRACT** The functions of gene regulatory networks that control embryonic cell diversification occur on a background of constitutively active molecular machinery necessary for the elaboration of genetic interactions. The essential roles of subsets of such "housekeeping" genes in the regulation of specific aspects of development have become increasingly clear. Pre-mRNA processing is essential for production of functional transcripts by, for example, excision of introns. We have cloned the zebrafish *toast*<sup>b460</sup> locus and found that it encodes *splicing factor 3b, subunit 1 (sf3b1)*. The *sf3b1*<sup>b460</sup> mutation causes aberrant splicing of *sf3b1* resulting in functional and predicted non-functional transcripts and a 90% reduction in full-length Sf3b1 protein. The *sf3b1*<sup>b460</sup> mutation was isolated in a mutagenesis screen based on the absence of neural crest-derived melanophores. Further analysis revealed specific earlier defects in neural crest development, whereas the early development of other ectodermal populations appears unaffected. The expression of essential transcriptional regulators of neural crest development are severely disrupted in *sf3b1*<sup>b460</sup> mutants, due in part to defects in pre-mRNA processing of a subset of these factors, leading to defects in neural crest sublineage specification, survival and migration. Misexpression of a subset of these factors rescues aspects of neural crest development in mutant embryos. Our results indicate that although *sf3b1* is a ubiquitously essential gene, the degree to which it is required exhibits tissue-type specificity during early embryogenesis. Further, the developmental defects caused by the *sf3b1*<sup>b460</sup> mutation provide insights into genetic interactions among members of the gene regulatory network controlling neural crest development.

**KEY WORDS:** *neural crest, pre-mRNA, splicing, zebrafish*

## Introduction

The removal of introns from pre-mRNA is essential for the ultimate generation of functional proteins. The precise excision of introns is catalyzed by a dynamic protein complex called the spliceosome (Burge *et al.*, 1999). Major elements of the spliceosome include the U1, U2, U5 and U4/6 small nuclear ribonucleoprotein particles (snRNPs) that contain small nuclear RNAs, common Sm proteins and proteins unique to each snRNP (Kambach *et al.*, 1999). The U2 snRNP exists in an inactive 12S form and an active 17S form generated by interaction of the 12S snRNP with the splicing factors Sf3b and Sf3a that are themselves comprised of several subunits (Brosi *et al.*, 1993a, 1993b). Sf3b1 is the largest subunit

comprising the Sf3b complex (25 exons encoding a 1315 amino acid protein in zebrafish) and is required for prespliceosome assembly and splicing catalysis and is thus essential for pre-mRNA processing (Gozani *et al.*, 1996; Gozani *et al.*, 1998). Sf3b1 is also a component of the minor U12-dependent spliceosome (Will *et al.*, 1999). *sf3b1* is ubiquitously expressed in mouse during early embryogenesis (Isono *et al.*, 2001) and *sf3b1*<sup>-/-</sup> mouse embryos arrest at the 16-32 cell stage (Isono *et al.*, 2005) demonstrating its essential function. The ubiquitous expression and function of

*Abbreviations used in this paper:* hpf, hours postfertilization; BAC, bacterial artificial chromosome; *sf3b1*, splicing factor 3b, subunit 1; snRNPs, small nuclear ribonucleoprotein particles.

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*sf3b1* in eukaryotic cells defines *sf3b1* as one of a number of so-called “housekeeping” genes. However, more nuanced functional requirements for a number of housekeeping genes have been documented recently (for example, see Coutinho *et al.*, 2004; Nambiar and Henion, 2004; Nambiar *et al.*, 2007).

The possibility that *sf3b1* may also serve more graded functions in different tissues at different times has been suggested by several recent reports. For example, Isono and colleagues, employing *sf3b1*<sup>+/−</sup> mice, demonstrate the requirement for *sf3b1* in Polycomb group-mediated repression of *Hox* genes that regulate skeletal growth and patterning (Isono *et al.*, 2005). A prominent differential role in neurogenesis by stem cells in neurogenic compared to non-neurogenic regions of the adult mammalian brain for RNA processing genes, including *sf3b1*, has been suggested (Lim *et al.*, 2005). Also of interest is the observation that *sf3b1* levels can determine the alternative splicing pattern derived from the *BCL-x* gene, potentially regulating pro- or antiapoptotic responses of cells to external stimuli (Massiello *et al.*, 2006). Together, these studies suggest that in certain contexts *sf3b1* may differentially regulate developmentally relevant processes.

In a screen for mutations affecting neural crest development (Henion *et al.*, 1996), we isolated the *toast*<sup>b460</sup> (*tst*) mutant based on the absence of neural crest-derived pigment cells but presence of the non-neural crest-derived pigmented retinal epithelium, suggesting possible cell type-specific functions of the *tst* mutant locus during development. Later in development, *tst* mutant embryos die at approximately 40 hours postfertilization (hpf), presumably due to prominent dorsal CNS cell death and the lack of circulating blood.

Here we present the molecular identification and characterization of the *tst*<sup>b460</sup> locus and a detailed phenotypic analysis of the consequences of this mutation. Specifically, we identify *tst*<sup>b460</sup> as a hypomorphic mutant allele of *sf3b1* that results in a dramatic reduction in wild-type transcripts and protein. As a result of a T>G point mutation in the 5′ splice site of intron 4 of *sf3b1*, pre-mRNA splicing errors in *tst* mutant *sf3b1* result in the production of 3 aberrant, presumably non-functional transcripts as well as a minority proportion of wild-type transcripts. The primary consequences of the *tst*<sup>b460</sup> mutation in the ectoderm are selective defects in neural crest development during early embryogenesis. These defects initially present as defects in the levels of expression of key transcriptional regulators of early neural crest development. In specific cases, reduced expression levels result from aberrant pre-mRNA splicing producing severely truncated transcripts. Partial restoration of a subset of these transcriptional regulators by misexpression results in differential degrees of neural crest phenotype rescue. Our results demonstrate differential sensitivities to the levels of expression of the essential RNA processing gene *sf3b1* in different subdivisions of the ectoderm and among genes that control the early development of the neural crest. These results indicate that the status of specific elements of essential cellular machinery is an additional regulatory component in the control of neural crest cell diversification during embryogenesis.

## Results

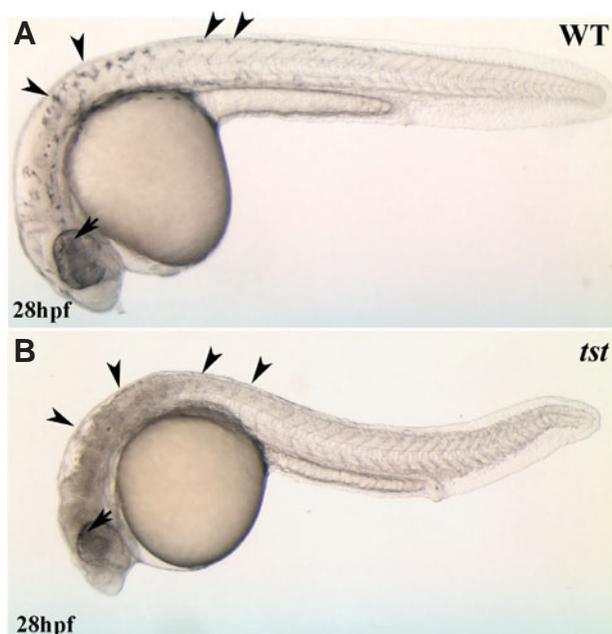
### Visible phenotype of live *tst*<sup>b460</sup> mutant embryos

The *tst*<sup>b460</sup> mutant was identified in an ENU-based mutagenesis screen for mutations that affect neural crest development (Henion

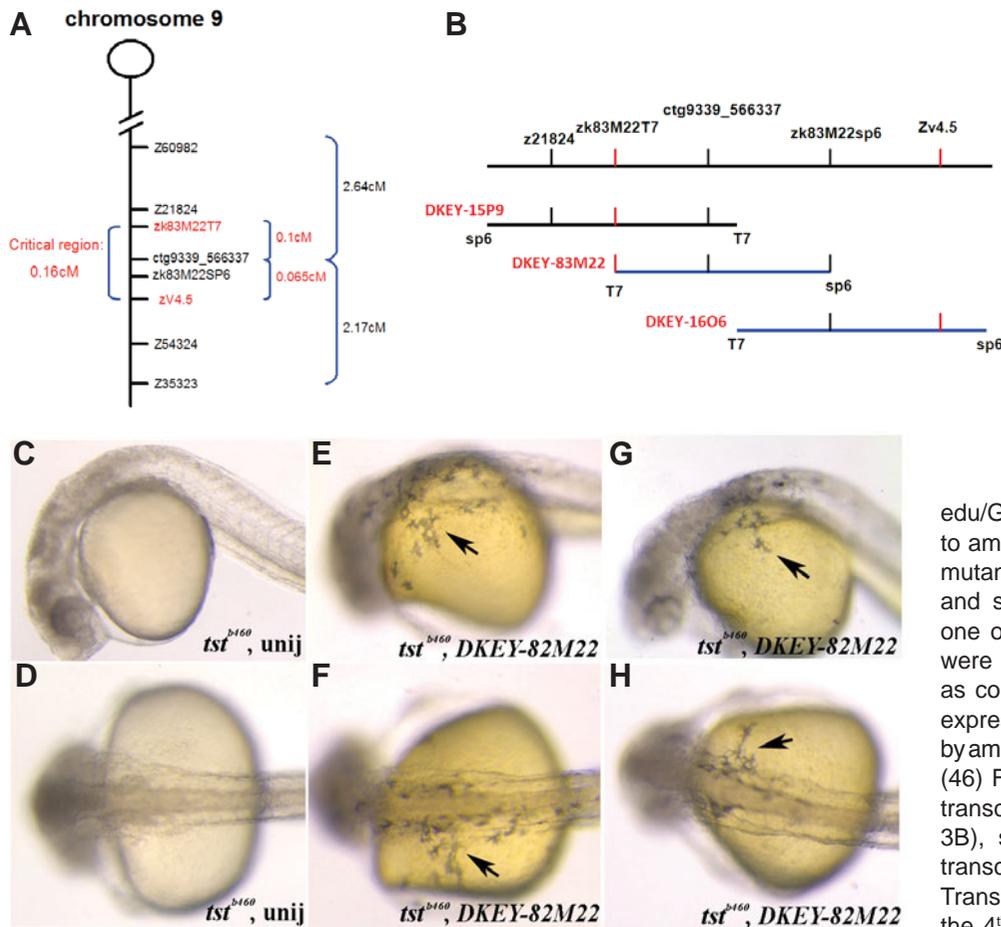
*et al.*, 1996). Differentiated neural crest-derived melanophores are completely absent in *tst*<sup>b460</sup> embryos (Fig. 1). In contrast, melanized pigmented retinal epithelial cells are present suggesting that, among pigment cells, the mutant phenotype is neural crest-specific. The *tst*<sup>b460</sup> mutation is recessive and *tst*<sup>b460</sup> mutants die by approximately 40 hpf most likely due to progressive dorsal CNS cell death and the absence of circulation.

### The *tst*<sup>b460</sup> locus encodes *sf3b1* and is a hypomorphic *sf3b1* mutant allele

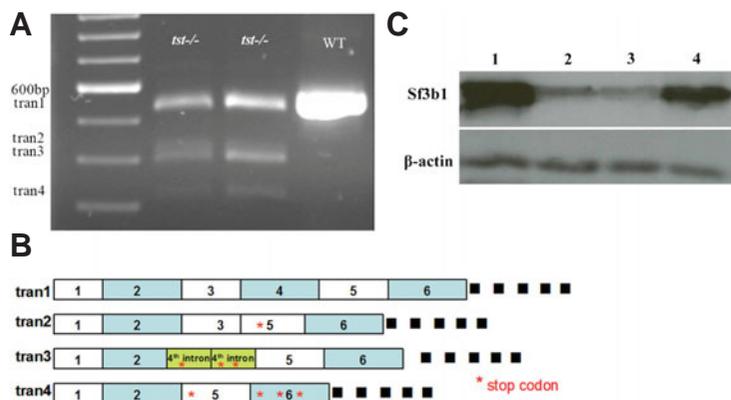
We mapped the *tst*<sup>b460</sup> mutation to chromosome 9 using standard linkage analysis. We then generated a mapping panel comprised of 2018 meioses to conduct recombination frequency analysis using several closely linked SSLP (z21824, z54324, and z35323) and SSCP (zk83M22T7, ctg9339\_566337, zK83M22SP6 and zv4.5) markers (see Materials and Methods). The result of this analysis defined a 0.164cM critical region containing the *tst*<sup>b460</sup> locus (Fig. 2A). Three BACs (DKEY-83M22, DKEY-16O6, DKEY-15P9) were found to map to this region (Fig. 2B; [http://www.sanger.ac.uk/cgi-bin/Projects/D\\_erio/mapsearch](http://www.sanger.ac.uk/cgi-bin/Projects/D_erio/mapsearch)). Because *tst*<sup>b460</sup> mutant embryos completely lack easily visualized neural crest-derived melanophores, we performed phenotype rescue experiments by misexpressing the DKEY-83M22 BAC in embryos from *tst*<sup>b460</sup> heterozygote crosses. All injected embryos were subsequently genotyped with tightly linked SSLP markers (z21824, z54324). We found that misexpression of BAC DKEY-83M22 resulted in robust melanophore phenotype rescue in *tst*<sup>b460</sup> mutants at efficiencies ranging from 57%–100% per experiment (Fig. 2 C–H and not shown). BAC DKEY-83M22 has been completely sequenced by the Sanger Center (<http://www.ebi.ac.uk/emb1>; <http://www.ncbi.nlm.nih.gov>). Based on this sequence and the regions of all three BACs that overlap, we identified several genes within this region using GENESCAN (<http://genes.mit>



**Fig. 1. Visible live phenotype of *tst*<sup>b460</sup> mutants.** Neural crest-derived melanophores; see arrowheads in (A) are absent in *tst*<sup>b460</sup> mutants; see arrowheads in (B). In contrast, melanized pigmented retinal epithelial cells are present in the mutants (arrow).



**Fig. 2. Fine mapping of the *tst* locus and BAC-mediated *tst*<sup>b460</sup> mutant phenotype rescue.** (A) Critical region containing the *tst*<sup>b460</sup> locus between closely linked SSLPs (z21824, z54324, z60982 and z35323) and SSCP (z83M22T7, ctg9339\_566337, z83M22SP6 and zV4.5) markers. (B) BACs (DKEY-83M22, DKEY-16O6, and DKEY-15P9) that map to the critical region containing *tst*<sup>b460</sup> locus. (C-H) Injection of DKEY-83M22 partially rescues the *tst*<sup>b460</sup> mutant melanophore phenotype. (C,D) Uninjected *tst*<sup>b460</sup> homozygous mutant embryos; (E-H) DKEY-83M22 injected *tst*<sup>b460</sup> homozygous mutant embryos, showing partial rescue of the melanophore phenotype (arrows).



**Fig. 3. Molecular consequences of *sf3b1* mutations (A,B).** Variant transcripts of *sf3b1* in *tst*<sup>b460</sup> mutants. (A) RT-PCR products were amplified between exon1 and exon6 of *sf3b1* with 20-24 hpf AB wildtype and *tst*<sup>b460</sup> homozygous mutant total RNA as template using *Sf3b1aa* (46)F and *Sf3b1aa* (579)R primers. The single AB wildtype RT-PCR product was 534 bp (tran1). *tst*<sup>b460</sup> homozygous mutants contained 4 different transcripts; tran1(534 bp), and variant transcripts tran2(412 bp), tran3(390 bp), and tran4(312 bp). (B) Diagram of variant transcripts in *tst*<sup>b460</sup> homozygous mutants. (C) Western blot analysis of *Sf3b1* protein levels in wildtype, *tst*<sup>b460</sup> mutant and hi3394a mutant embryos at 24 hpf. Lane 1: wildtype embryos; Lane 2: *tst*<sup>b460</sup> homozygous mutants; Lane 3: hi3394a homozygous mutants; Lane 4: hsp>*sf3b1* injected *tst*<sup>b460</sup> mutants with melanophore phenotype rescue.

edu/GENSCAN.html). We designed primers to amplify these candidate genes from *tst*<sup>b460</sup> mutant and wildtype embryos for RT-PCR and sequence analysis. We found that for one of these candidate genes, *sf3b1*, there were 4 different transcripts in *tst*<sup>b460</sup> mutants as compared to AB<sup>+</sup> wild-type embryos that express a single transcript (Fig. 3A) revealed by amplification using RT-PCR and the *Sf3b1aa* (46) F and *Sf3b1aa* (579) R primers. Of the transcripts expressed by *tst*<sup>b460</sup> embryos (Fig. 3B), sequence comparisons revealed that transcript 1 is the wild-type *sf3b1* mRNA. Transcript 2 is an aberrant transcript in which the 4<sup>th</sup> exon is missing. The 3<sup>rd</sup> transcript is missing the 3<sup>rd</sup> and 4<sup>th</sup> exons with the 4<sup>th</sup> intron aberrantly included between the 2<sup>nd</sup> and 5<sup>th</sup> exons. Transcript 4 is missing part of exon 2<sup>nd</sup> and all of exons 3 and 4. The splicing errors that result in the three aberrant transcripts all introduce premature stop codons that predict highly truncated proteins that lack the essential TP and RWD repeats that are believed to be essential functional domains for phosphorylation and protein-protein interactions, respectively (Seghezzi *et al.*, 1998; Gozani *et al.*, 1998), and thus non-functional or severely compromised products.

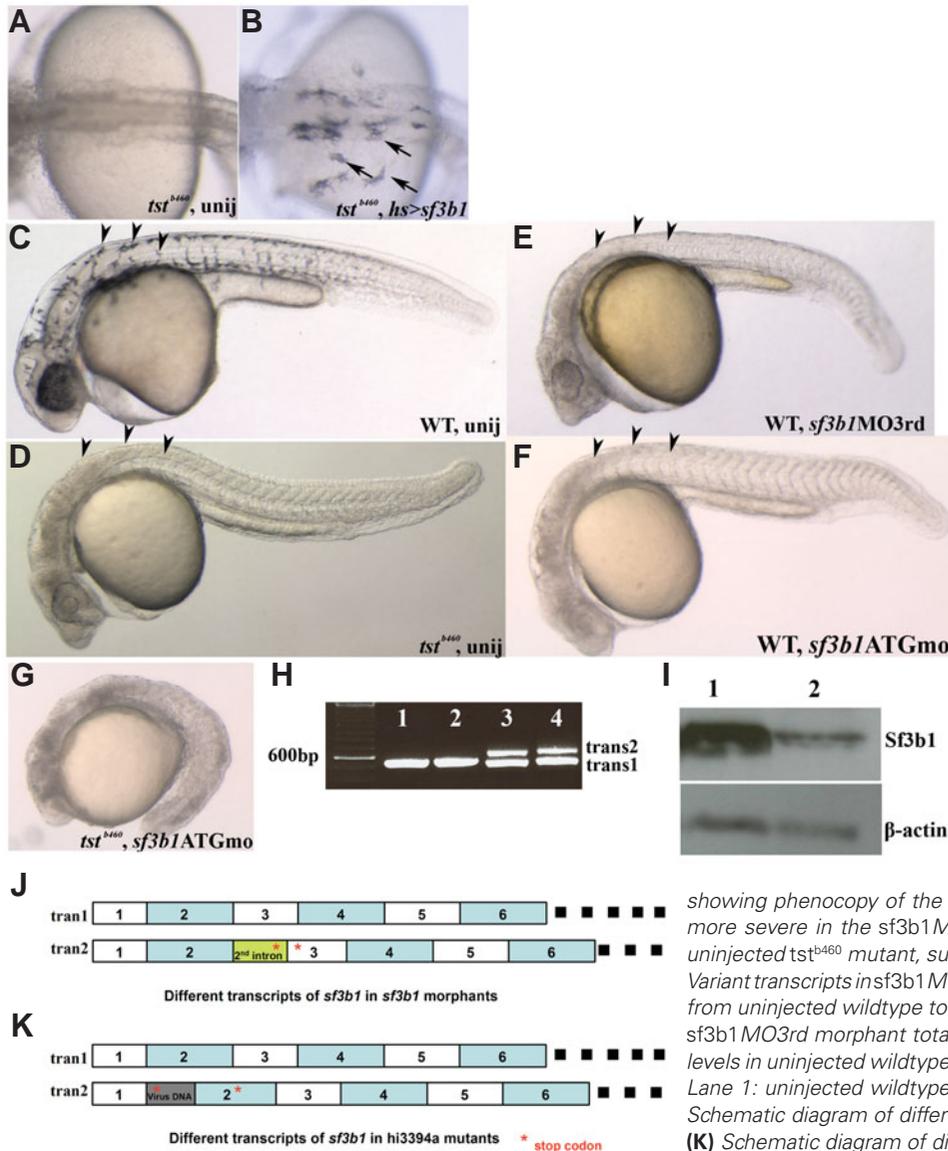
We employed a *Sf3b1* monoclonal antibody that recognizes a central portion of the *Sf3b1* protein (kindly provided by Dr. K. Isono, Chiba University; Horie *et al.*, 2003) and densitometer measurements of chemiluminescence detection in Western blots to obtain a quantitative determination of wildtype *Sf3b1* protein levels in *tst*<sup>b460</sup> mutant embryos compared to wildtype siblings. Actin protein expression was used as a control. We found that presumptive full length functional *Sf3b1* protein levels in pooled *tst*<sup>b460</sup> mutant embryos was dramatically reduced to 9.6 ± 4.2% of wildtype levels at 24hpf (Fig. 3C; n=10 different protein pools, each containing 20-30 embryos per pool). In addition, we found that the *Sf3b1* antibody employed did not recognize any products derived from the aberrant 5' truncated transcripts produced in *tst*<sup>b460</sup> mutants, as predicted by the antigen sequence used to produce the antibody (Horie *et al.*, 2003). These results suggest that the *tst*<sup>b460</sup> locus is likely to be *sf3b1* and that the allele is hypomorphic.

To further assess whether the *tst<sup>b460</sup>* mutant locus is *sf3b1*, we constructed the full-length wildtype cDNA for misexpression and phenotype rescue analysis (GenBank accession number pending). Because of the large size of the predicted full-length cDNA (4393 bp), we constructed the full-length cDNA from two fragments, one derived from EST clone fb99f09 and the other a RT-PCR fragment of the 5' region of the transcript generated from RT-PCR amplification from 18-24 hpf wildtype total RNA (see Methods). The presumptive full-length wildtype *sf3b1* cDNA was sequenced to confirm frame and sequence. The full-length cDNA was then ligated to the heatshock vector pCSHSP (Halloran *et al.*, 2000) to drive expression.

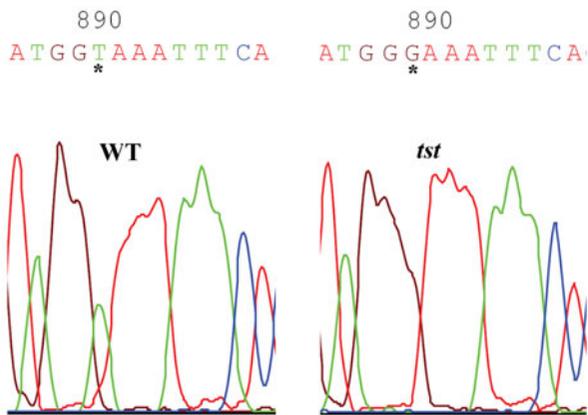
The *hsp>sf3b1* expression construct was injected into a single blastomere of 1-2 cell stage embryos derived from *tst<sup>b460</sup>* heterozygous parents and injected embryos were heat-shocked at 6-7hpf and 11-12hpf and raised to 27-36hpf. In a subset of experiments, embryos were heat-shocked an additional time at 23-24hpf. No phenotypic differences were observed between those embryos heat-shocked 2 or 3 times. Injected embryos

were examined for visible melanophores at 27- 36hpf and genotyped with the closely-linked SSLP markers z21824 and z54324. We found that misexpression of wildtype *sf3b1* by injection of *hsp>sf3b1* resulted in significant melanophore rescue in 80% of injected homozygous *tst<sup>b460</sup>* mutant embryos (Fig. 4 A,B; n=92 mutant embryos). Together with the efficient rescue exhibited by misexpressing the *sf3b1*-containing BACs and the severe reductions of wildtype Sf3b1 protein and full-length transcript levels in mutant embryos, these results provide further evidence that the *tst<sup>b460</sup>* locus is likely to be *sf3b1*.

To further test whether the *tst<sup>b460</sup>* locus corresponds to *sf3b1*, we designed two antisense oligonucleotide morpholinos (MO) to knockdown Sf3b1 in wildtype embryos when injected at the 1-2 cell stage in wildtype embryos. We designed and tested *sf3b1*ATG MO, a start-site blocking MO and *sf3b1*3rdMO, a MO targeted to disrupt pre-mRNA splicing in a manner approximating the splicing errors found in *sf3b1* transcripts in *tst<sup>b460</sup>* mutant embryos. We confirmed by Western analysis that *sf3b1*ATG MO effectively knocked down Sf3b1 protein levels in injected wildtype embryos (Fig. 4I). Sequence analysis revealed that the *sf3b1*3rd MO causes a frequent failure in the splicing of the 3' splice site of the 2<sup>nd</sup> intron of *sf3b1* (Fig. 4 H,J) resulting in a premature stop codon in the 2<sup>nd</sup> intron. The injected *sf3b1*3rd MO phenocopied the melanophore and gross morphology phenotypes of *tst<sup>b460</sup>* mutants in injected wildtype embryos at 88-99% efficiency (Fig. 4E). The *sf3b1*ATG MO also resulted in phenocopy in 87-98% of groups of injected embryos



**Fig. 4. *sf3b1* gain- and loss-of-function analyses.** (A,B) The *tst<sup>b460</sup>* phenotype is partially rescued by *sf3b1* misexpression. (A) Melanophores are absent in uninjected *tst<sup>b460</sup>* mutant embryos. (B) *tst<sup>b460</sup>* homozygous mutant embryos were injected with *hs>sf3b1* and heat-shocked 2 or 3 times (6 hpf, 10 or 11 hpf, 22 or 23 hpf), 1h/each time. Several melanophores present in the dorsal trunk region and over the yolk region in injected *tst<sup>b460</sup>* mutant embryos (arrows). (C-F) Phenocopy of *tst<sup>b460</sup>* mutant embryos using morpholino knockdown analysis. (C) Lateral view of a live uninjected wildtype embryo at 30 hpf. (D) Lateral view of a *tst<sup>b460</sup>* homozygous mutant at 30 hpf. (E) Lateral view of a live *sf3b1*MO3rd morphant showing the absence of melanophores (arrowheads) and CNS (brain) cell death at 30 hpf. (F) Lateral view of a live *sf3b1*MOATG morphant showing phenocopy of the *tst<sup>b460</sup>* mutant at 30 hpf. (G) The phenotype was more severe in the *sf3b1*MOATG injected *tst<sup>b460</sup>* mutant as compared with uninjected *tst<sup>b460</sup>* mutant, suggesting that the *tst<sup>b460</sup>* allele is hypomorphic. (H) Variant transcripts in *sf3b1*MO3rd morphants. Lane 1 and lane 2: RT-PCR product from uninjected wildtype total RNA. Lane 3 and lane 4: RT-PCR product from *sf3b1*MO3rd morphant total RNA. (I) Western blot analysis of Sf3b1 protein levels in uninjected wildtype embryos and *sf3b1*MOATG morphants at 12 hpf. Lane 1: uninjected wildtype embryos; Lane 2: *sf3b1*MOATG morphants. (J) Schematic diagram of different *sf3b1* transcripts in *sf3b1*MO3rd morphants. (K) Schematic diagram of different *sf3b1* transcripts in *hi3394a* mutants.



**Fig. 5.** The nucleotide mutation at the 5' splice site of the 4<sup>th</sup> intron of *sf3b1* genomic DNA identified in *tst<sup>b460</sup>* mutants. In *tst<sup>b460</sup>*, T is changed to G.

(Fig. 4F). Injection of *tst<sup>b460</sup>* mutants with *sf3b1*ATG MO resulted in the arrest embryonic development during early somitogenesis (Fig. 4G), reminiscent of Sf3b1 null mice and consistent with the suggestion that *tst<sup>b460</sup>* is a hypomorphic allele. The phenocopy of *tst<sup>b460</sup>* mutants by two verified MOs targeting *sf3b1* provides further evidence that *tst<sup>b460</sup>* corresponds to *sf3b1*.

As mentioned previously, we isolated the full-length zebrafish *sf3b1* cDNA (Genbank accession number pending). Sequence analysis of the entire *sf3b1* coding sequence of wildtype and *tst<sup>b460</sup>* mutants failed to detect any nucleotide mutations in the *tst<sup>b460</sup>* coding sequence. Therefore, because *tst<sup>b460</sup>* mutants produce aberrant transcripts in the 5' end of the gene upstream of the 5<sup>th</sup> exon, we sequenced genomic sequences of wildtype and *tst<sup>b460</sup>* mutants upstream of the 6<sup>th</sup> exon of *sf3b1*. Our analysis revealed a T>G point mutation in the 5' splice site of the 4<sup>th</sup> intron in the *tst<sup>b460</sup>* *sf3b1* sequence (Fig. 5). The U1 snRNP recognizes the 5' splice site to form a complex that commits the premRNA to spliceosome assembly (reviewed by Nilsen, 1997). The recognition of the 5' splice site involves base-pairing between the conserved nucleotide at the 5' end of U1 and 5' intronic splice site. The specific nucleotide mutation at the 5' splice site of the 4<sup>th</sup> intron in *sf3b1* from *tst<sup>b460</sup>* mutant embryos is consistent with the production of the aberrant transcripts observed based on the current knowledge of the mechanics of pre-mRNA processing. The observed nucleotide mutation in *tst<sup>b460</sup>* mutant *sf3b1* combined with the evidence presented above strongly suggest that *tst<sup>b460</sup>* is a *sf3b1* mutant allele.

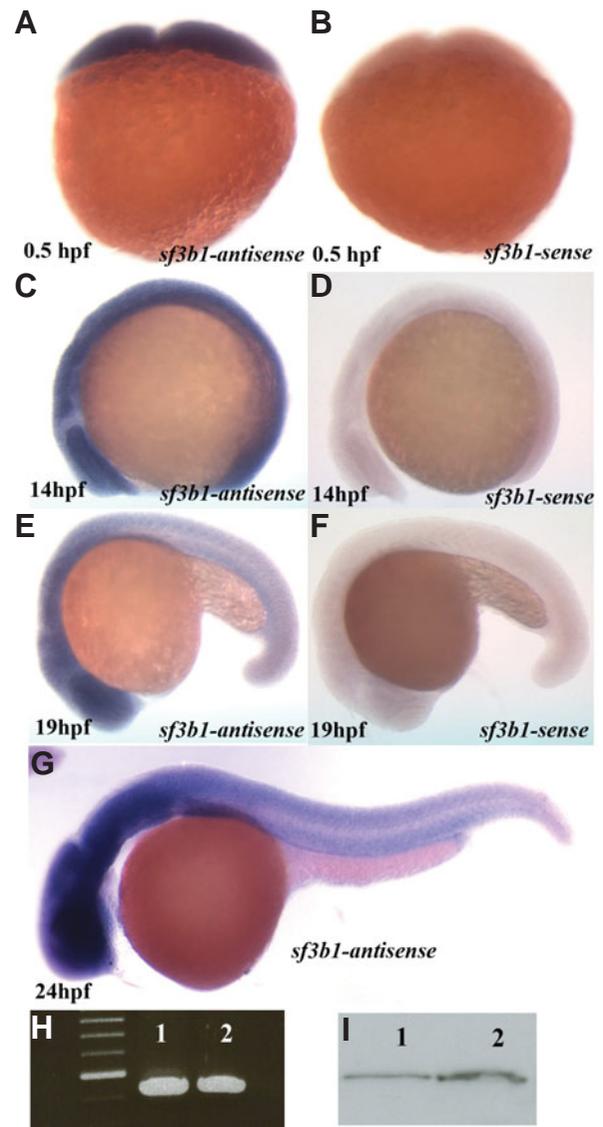
Lastly, as our analysis was proceeding, we learned that a viral insertion mutation in *sf3b1* was identified in a mutant screen (Amsterdam *et al.*, 2004). We obtained this mutant line, *hi3394a*,

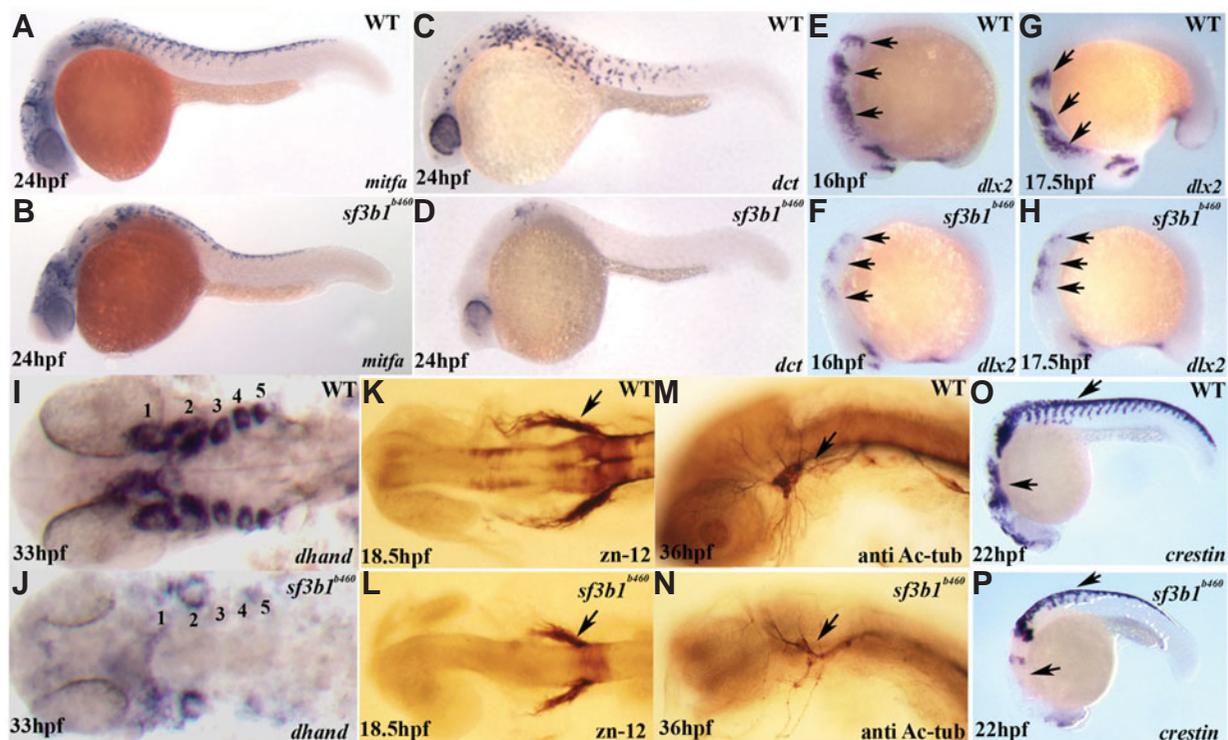
**Fig. 6.** Whole-mount *in situ* hybridization with *sf3b1* antisense probe and sense probes in wildtype embryos. (A,C) *sf3b1* is ubiquitously expressed at 0.5 hpf and 14 hpf. (E,G) The expression of *sf3b1* appears elevated in the brain, neural tube and ventral trunk region at 19 hpf and 24 hpf (lateral view, anterior to left). (B,D,F) *sf3b1* sense controls. (H) RT-PCR products using total RNA from 0.5hpf (lane1) and 24hpf (lane2) wildtype embryos as templates and *Sf3b1aa(46)F* and *Sf3b1aa(579)R* as primers. (I) Western blot analysis shows Sf3b1 protein present maternally. Lane 1: protein from 24 hpf wildtype embryos as control. Lane 2: protein from 1 hpf wildtype embryos.

and determined via sequence analysis that the viral insertion resulted in the production of two transcripts, a presumptive wild-type transcript and a second aberrant transcript in which the viral insertion between the 1<sup>st</sup> and 2<sup>nd</sup> exons results in the creation of a premature stop codon in the 2<sup>nd</sup> exon (Fig. 4K). We also found that full length Sf3b1 protein levels in *hi3394a* mutants was reduced to 13.5+/-6.2% of wildtype levels at 24hpf (see Fig. 3C). We determined that *hi3394a* live phenotype is very similar but slightly less severe than that of *tst<sup>b460</sup>* mutants particularly in the case of melanophores which are entirely absent in *tst<sup>b460</sup>* mutants whereas a small number of melanophores usually develop in *hi3394a* mutants (Amsterdam *et al.*, 2004). Critically, complementation analysis demonstrated that *tst<sup>b460</sup>* and *hi3394a* fail to complement. Taken together with all of the results described, we conclude that the *tst<sup>b460</sup>* locus corresponds to *sf3b1* and that it is a severe but hypomorphic allele. Thus, we designate *tst<sup>b460</sup>* as *sf3b1<sup>b460</sup>*.

### Expression of zebrafish *sf3b1*

Analysis of the zebrafish genome indicates that the *sf3b1* gene





**Fig. 7. Neural crest progenitor and cranial ganglia phenotypes of *sf3b1*<sup>b460</sup> mutants.** There is reduced melanoblast expression of *mitfa* (A,B) and nearly absent melanoblast expression of *dct* (C,D) in *sf3b1*<sup>b460</sup> mutants. (E-J) Pharyngeal arches are dramatically reduced in *sf3b1*<sup>b460</sup> mutant embryos. (E-H) Lateral view of whole-mount *in situ* embryos with *dlx2* antisense RNA probe, arch progenitors denoted by arrows. (I,J) Dorsal view of flat-mounted *in situ* embryos with *dhand* antisense RNA probe, pharyngeal arches 1 and 2 are decreased and 3, 4 and 5 are absent in *sf3b1*<sup>b460</sup> mutants. (K-N) the cranial ganglia are also affected in *sf3b1*<sup>b460</sup> mutants. (K,L) Whole-mount antibody staining with *zn-12*. Arrows in (K,L) show the trigeminal ganglia. (M,N) Whole-mount antibody staining with anti-acetylated tubulin antibody. Arrows in (M,N) show the decreased and disorganized trigeminal ganglia in *sf3b1*<sup>b460</sup> mutants compared to wildtype siblings. (O,P) Crestin expressing cells are present in *sf3b1*<sup>b460</sup> mutant embryos. However, these cells are reduced in numbers, particularly in the cranial region and fail to migrate normally in the *sf3b1*<sup>b460</sup> mutant (P) compared to wildtype embryos (O).

is approximately 17 kb in size and that it is a single-copy gene. The full-length *sf3b1* is 4393 bp derived from 25 exons. The *sf3b1* cDNA predicts a protein comprised of 1315 amino acids. The amino acid sequence of zebrafish *Sf3b1* is highly conserved when compared to the human, mouse and *Xenopus* *Sf3b1* homologs (calculated 92% identity) and the carboxy-terminal three-fourths of *Sf3b1* shows even higher conservation (97.5%; see Supplemental Fig. 1). In addition, the amino acid sequences of *C. elegans* and *S. pombe* *Sf3b1* also show a high degree of conservation compared to vertebrates (Isono *et al.*, 2001).

To examine the expression pattern of zebrafish *sf3b1* we synthesized a 1.6kb antisense RNA probe complementary to the 2934-4393 bp 3' region of *sf3b1* and performed wholemount *in situ* hybridization (WISH) on embryos from 0-48 hpf (Fig. 6 A-G and not shown). We found using WISH that *sf3b1* is maternally expressed (Fig. 6 A,B) and confirmed this observation using RT-PCR and Western analyses (Fig. 6 H,I). Zytically, in general, we found that *sf3b1* mRNA is expressed ubiquitously at all stages examined (Fig. 6 C-G). Expression levels appear enhanced in the brain, dorsal neural tube and ventral trunk regions of later-stage embryos (Fig. 6 E-G), although this observation may be the result of differences in the cellular composition and architecture of these regions at these stages compared to other regions of the embryo. No qualitative differences in the pattern or level of *sf3b1* mRNA expression between wildtype and *sf3b1*<sup>b460</sup> mutant embryos were observed using the 3' riboprobe employed, as would be expected

based on the 5' locations of the splicing errors in *sf3b1*<sup>b460</sup> mutants.

#### The development of neural crest-derived cell subpopulations fails in *sf3b1*<sup>b460</sup> mutant embryos

As stated previously, a prominent visible phenotype of *sf3b1*<sup>b460</sup> mutant embryos is the complete absence of differentiated neural crest-derived melanophores (Fig. 1). Therefore, we examined the development of melanogenic precursors in *sf3b1*<sup>b460</sup> mutant embryos. To assess melanogenic sublineage development, we examined the expression of *mitfa*, a gene necessary for melanoblasts specification and an early marker of melanophore precursors (Opdecamp *et al.*, 1997; Lister *et al.*, 1999) and *dct*, whose expression is diagnostic of later stage melanoblasts (Kelsh *et al.*, 2000). The initial neural crest expression of *mitfa* in *sf3b1*<sup>b460</sup> mutant embryos is delayed by several hours and then significantly reduced in expression compared to stage matched wildtype embryos (Fig. 7 A,B). In contrast, the timing and pattern of PRE expression is qualitatively normal. The expression of neural crest *dct* is significantly reduced with only occasional *dct*<sup>+</sup> cells in the trunk of a minority of mutant embryos (Fig. 7 C,D). As is the case for *mitfa*, PRE *dct* expression in *sf3b1*<sup>b460</sup> mutants was not significantly different than wildtype embryos. Thus, we conclude that the specification and early development of the melanophore sublineage is severely disrupted in *sf3b1*<sup>b460</sup> mutants resulting in the absence of differentiated melanophores.

We also assessed the development of neural crest-derived

craniofacial cartilage progenitors by comparing the expression of *dlx2* (Akimenko *et al.*, 1994) and *dhand* (Yelon *et al.*, 2000) in the developing pharyngeal arches in wildtype and *sf3b1<sup>b460</sup>* mutant embryos. We found that neural crest expression of both genes in the pharyngeal arches was dramatically reduced in mutant embryos, being absent in posterior arches and strongly reduced in the first and second arches (Fig. 7 E-J). These results indicate that the development of the neural crest component of the pharyngeal arches is severely compromised in *sf3b1<sup>b460</sup>* mutants.

Another subpopulation of neural crest-derived cells that is specified and differentiates relatively early during embryogenesis is a subset of cranial ganglion neurons (Schilling and Kimmel, 1994). Other neurons that comprise the cranial ganglia are derived from the ectodermal placodes. In *sf3b1<sup>b460</sup>* mutants, the numbers of neurons and organization of trigeminal (Fig. 7 K-N) and posterior lateral line ganglia (not shown) are reduced and disorganized, respectively, compared to wildtype siblings. Although we cannot be sure that this phenotype results from a selective disruption of cranial neural crest development, as opposed to a disruption of placode development, the pattern of pan-neural crest *crestin* expression in the head at 22hpf (Fig. 7 O,P) is consistent with the possibility of a neural crest defect in the development of the cranial ganglia phenotype of *sf3b1<sup>b460</sup>* mutant embryos.

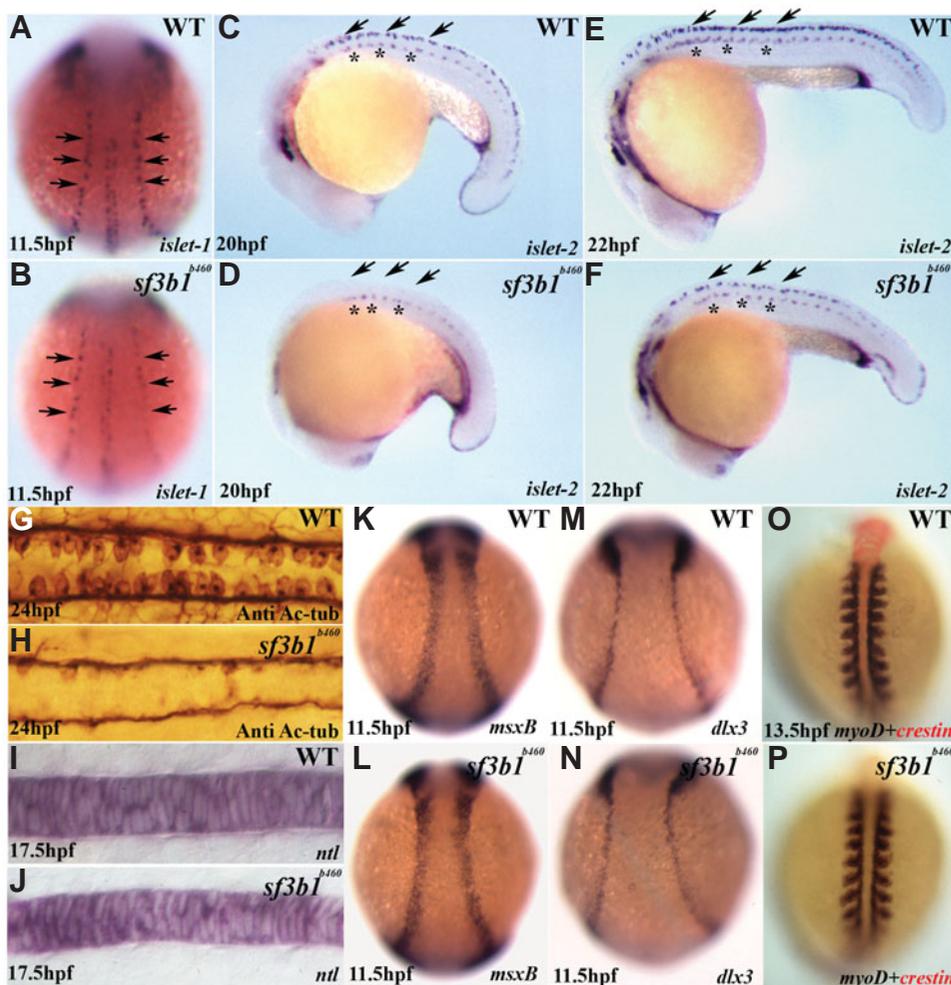
Within the developing neural plate border in zebrafish embryos, the neural crest and Rohon-Beard sensory neurons are thought

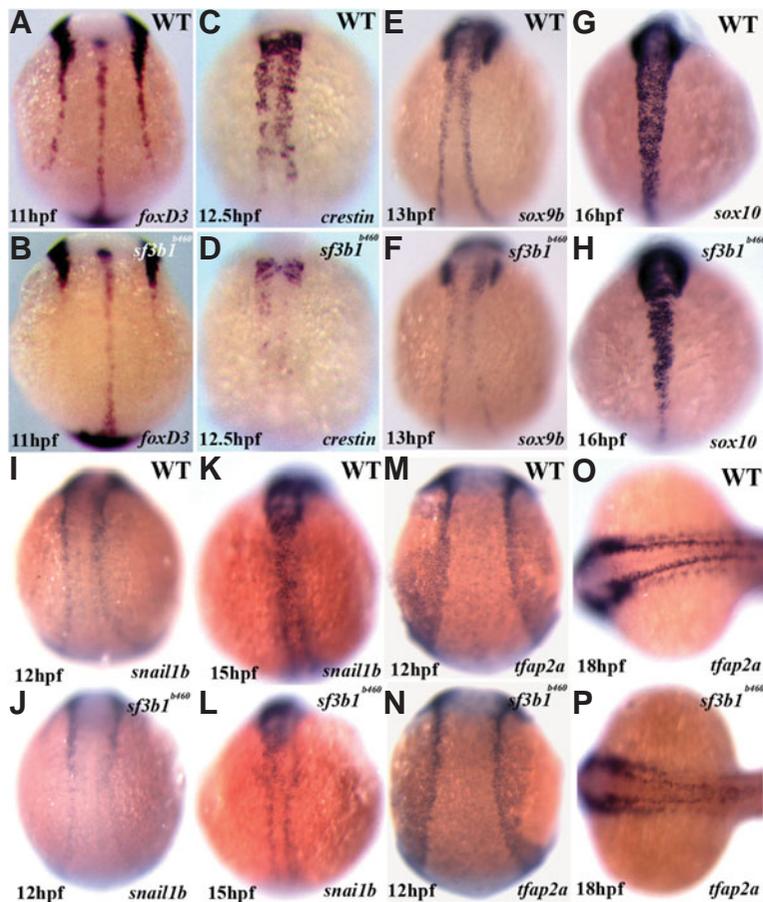
to comprise an equivalence group (Artinger *et al.*, 1999; Cornell and Eisen, 2000). At early somitogenesis stages, differentiated Rohon-Beard sensory neurons in the neural plate border can be identified based on the expression of *huC* as well as *islet1* (Artinger *et al.*, 1999; Cornell and Eisen, 2000). Examination of Rohon-Beard sensory neuron development at 11.5, 20 and 22hpf in *sf3b1<sup>b460</sup>* mutant and wildtype embryos revealed that the number of neurons was significantly reduced in mutant embryos compared to wildtype siblings (Fig. 8 A-F). Rohon-Beard sensory neuron numbers in *sf3b1<sup>b460</sup>* mutant embryos do not recover as revealed by immunostaining with acetylated tubulin and zn-12 antibodies at later developmental stages (Fig. 8 G,H). Thus, along with severe defects in the development of neural crest-derived cell subpopulations, *sf3b1<sup>b460</sup>* mutant embryos display abnormal development of another neural plate border cell subpopulation, Rohon-Beard sensory neurons. However, even given the early defects in neural crest and Rohon-Beard neuron development, the timing and pattern of the establishment of the neural plate border generally is unaffected in *sf3b1<sup>b460</sup>* mutant embryos as assessed by the expression of *msxB* and *dlx3* (Fig. 8 K-N). Further, the early development of somites (Fig. 8 O,P) and notochord (Fig. 8 I,J) were also normal in mutants. Lastly, development of primary motoneurons was also comparatively normal in mutant embryos (Fig. 8 C-F and not shown). Thus, early ectodermal developmental defects in *sf3b1<sup>b460</sup>* mutant embryos are by and large confined to

the neural crest/Rohon Beard neurons of the neural plate border.

A number of transcription factors, including *foxd3*, *tfap2a*, *snai1b*, *sox10* and *sox9b*, have been shown to play important roles in the fate-specification, survival, migration and/or differentiation of distinct and overlapping neural crest sublineages in multiple species (Gammill

**Fig. 8. Early development of ectodermal and mesodermal derivatives in *sf3b1<sup>b460</sup>* mutants.** (A,B) Whole-mount in situ hybridization with *islet-1* antisense RNA probes shows that the number of Rohon-Beard precursor cells (arrows) is decreased in *sf3b1<sup>b460</sup>* mutants (dorsal view, anterior to top). (C-F) Whole-mount in situ hybridization with *islet-2* antisense RNA probe shows deficiency in the development of Rohon-Beard sensory neurons (arrows), whereas the development of primary motor neurons is comparatively less affected in mutant embryos (asterisks). (G,H) Dorsal view of spinal cords labeled with anti-acetylated tubulin (Anti Ac-tub), anterior to left, showing that the Rohon-Beard sensory neuron phenotype persists at later stages in *sf3b1<sup>b460</sup>* mutants. (I,J) Induction of the neural plate border occurs normally in *sf3b1<sup>b460</sup>* mutants. Whole-mount in situ hybridization with *msxB* (K,L) and *dlx3* (M,N). (I,J,O,P) The derivatives from paraxial mesoderm and chordamesoderm are not affected in *sf3b1<sup>b460</sup>* mutants. (I,J) Lateral view of whole-mount in situ with *ntl*, anterior to left. (O,P) Dorsal view of double in situ with *crestin* (red) and *myoD* (blue), anterior to top.





**Fig. 9. Significant abnormalities in trunk neural crest expression of critical transcriptional regulators as well as *crestin* at early developmental stages in *sf3b1<sup>b460</sup>* mutants, but grossly normal expression in the cranial neural crest.** Whole-mount in situ hybridization with *foxd3* (A,B), *crestin* (C,D), *sox9b* (E,F), *sox10* (G,H), *snail1b* (I-L), and *tfap2a* (M-P) antisense RNA probes. All embryos are dorsal view, anterior to top with the exception of (O,P) which is anterior to left.

and Bronner-Fraser, 2003). Therefore, because of the miscues exhibited in NC development in *sf3b1<sup>b460</sup>* mutant embryos, we examined the neural crest expression of these genes in *sf3b1<sup>b460</sup>* mutants. We also examined the expression of the neural crest marker gene *crestin* (Luo *et al.*, 2001).

At 12.5hpf, the expression of *crestin* is nearly absent in the trunk of *sf3b1<sup>b460</sup>* mutant embryos, whereas expression is present but reduced in cranial neural crest (Fig. 9 C,D). Later in development, at 22hpf, *crestin* is expressed by trunk neural crest in mutant embryos although by noticeably fewer cells and, other than in the most rostral somitic regions, is confined to the dorsal neural tube suggesting defective trunk neural crest migration. At the same stage, and qualitatively in contrast to earlier embryonic stages, cranial neural crest expression of *crestin* is nearly absent in *sf3b1<sup>b460</sup>* mutant embryos (Fig. 7 O,P).

At 11hpf, the expression of *foxd3* is essentially absent in trunk neural crest in *sf3b1<sup>b460</sup>* mutant embryos, whereas expression by cranial neural crest is very similar to that observed in wildtype embryos (Fig. 9 A,B). The expression of *sox9b* at 13hpf is also very strongly reduced in trunk neural crest and is reduced to a lesser extent in cranial neural crest in *sf3b1<sup>b460</sup>* mutant embryos

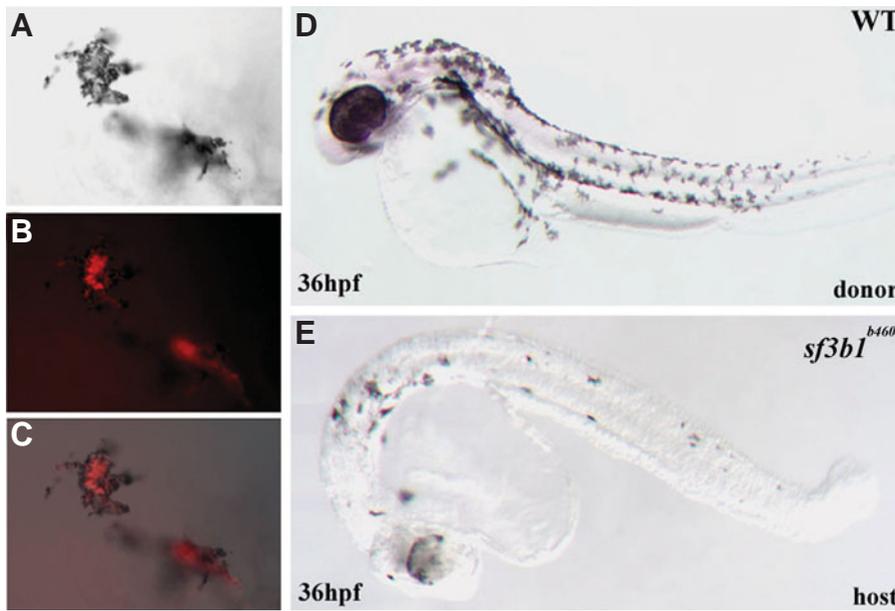
(Fig. 9 E,F), an abnormal neural crest expression pattern similar to that of *foxd3*. Strong trunk neural crest as well as cranial neural crest expression of *snail1b* at 12 hpf and 15 hpf and *sox10* at 16hpf is evident in wildtype embryos (Fig. 9 G,I,K). The expression of both genes in *sf3b1<sup>b460</sup>* mutant embryos was markedly reduced in trunk neural crest and was similar but slightly reduced in cranial neural crest (Fig. 9 H,J,L). At 12hpf, neural crest expression of *tfap2a* was indistinguishable between *sf3b1<sup>b460</sup>* mutant and wildtype embryos (Fig. 9 M,N). In contrast, a noticeable reduction in neural crest expression of *tfap2a*, particularly in the trunk, is evident in *sf3b1<sup>b460</sup>* mutant embryos at 18hpf (Fig. 9 O,P). The abnormal neural crest expression patterns of *foxd3*, *sox9b*, *snail1b* and *sox10* observed at earlier developmental stages in *sf3b1<sup>b460</sup>* mutant embryos are maintained at 18hpf (not shown). The fact that neural crest *tfap2a* expression is initially normal indicates that neural crest induction occurs normally in *sf3b1<sup>b460</sup>* mutants. Thus, the neural crest expression of multiple other transcriptional regulators of neural crest development are disrupted in *sf3b1<sup>b460</sup>* mutant embryos in an asynchronous manner and share a comparatively more severe trunk neural crest as opposed to cranial neural crest phenotype. As these transcription factors have been implicated as regulators of neural crest sublineage specification, survival and/or migration, these results suggest that the severe deficiencies in later neural crest development in *sf3b1<sup>b460</sup>* mutants is, at least in part, a consequence of the dysregulated expression of early transcriptional regulators of neural crest development.

#### ***sf3b1* is required cell autonomously for neural crest development**

To determine whether or not *sf3b1* function is required cell autonomously for neural crest development, we analyzed genetically mosaic embryos generated by cell transplantation between mutant and wildtype embryos. Because *sf3b1<sup>b460</sup>* mutant embryos completely lack neural crest-derived melanophores, we scored mosaic embryos for the presence or absence of donor-derived melanophores. Embryos generated from crosses of *sf3b1<sup>b460</sup>* heterozygous adults were separated into two groups, half of which were injected at the 1-4 cell stage with lyseinated rhodamine dextran (LRD) to label all cells. Cells from LRD-labeled embryos were transplanted to unlabeled siblings at the late blastula stage (Ho and Kane, 1990) and according to the zebrafish embryo fate map (Kimmel *et al.*, 1990). Host and donor embryos were grown to 24-36hpf when the visible phenotype of absent melanophores in *sf3b1<sup>b460</sup>* mutant embryos is readily apparent. All embryos were also genotyped with linked markers as described previously. We found that wildtype donor neural crest cells readily generate melanophores in *sf3b1<sup>b460</sup>* mutant hosts (Fig. 10; n=23) while we never observed mosaic embryos in which *sf3b1<sup>b460</sup>* mutant neural crest cells generated melanophores in wildtype hosts (not shown; n=29). We conclude that *sf3b1* is required cell autonomously for neural crest development.

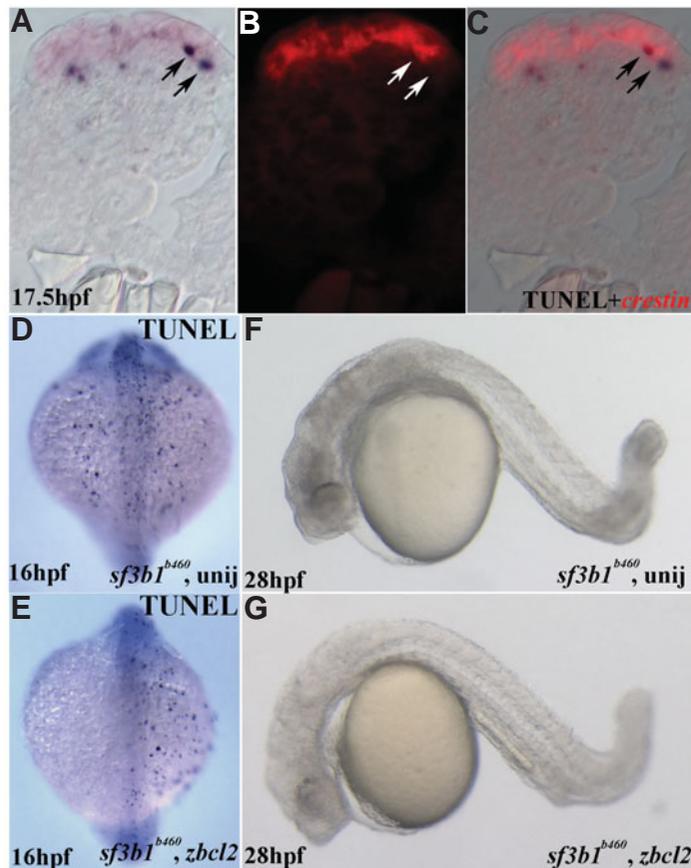
#### **Increased neural crest apoptosis and migratory defects in *sf3b1<sup>b460</sup>* embryos**

Because the transcription factors *foxd3*, *tfap2a*, *snail1b*, *sox9b* and *sox10* have been shown to be required to differing degrees for



**Fig. 10. Genetic mosaics of *sf3b1<sup>b460</sup>* mutants and wildtype embryos.** (A) Melanophores derived from wildtype donor cells on the yolk of a *sf3b1<sup>b460</sup>* mutant host (DIC image). (B) Fluorescence image of the same (donor-derived) cells. (C) (A,B) merged. (D) A wildtype donor. (E) An *sf3b1<sup>b460</sup>* mutant host with wildtype donor-derived melanophores.

crucial aspects of neural crest development including the specification of neural crest sublineage fates, neural crest cell survival and neural crest migration and the expression of all of these genes is disrupted in *sf3b1<sup>b460</sup>* mutants, we further investigated general aspects of neural crest development in mutant embryos. As described earlier, the specification of neural crest sublineage fates is disrupted in *sf3b1<sup>b460</sup>* mutants, at least in the case of the melanophore and craniofacial cartilage sublineages.



to wildtype embryos.

We then examined cell death during development in *sf3b1<sup>b460</sup>* mutant embryos using TUNEL. At early stages of trunk neural crest development (12-14hpf) the numbers of dying cells in mutants is moderately increased compared to wildtype embryos, particularly in the anterior CNS (not shown). Subsequently, numerous TUNEL-positive cells along the dorsal neural tube of mutant embryos, but not wildtype embryos, are observed. We employed WISH with the neural crest marker *crestin* of 17.5hpf embryos previously processed by TUNEL and examined cryosectioned samples of the trunk region. We found that many of the TUNEL-positive cells co-labeled with *crestin* indicating increased neural crest cell death in *sf3b1<sup>b460</sup>* mutants compared to wildtype siblings (Fig. 11 A-C). At subsequent stages of development, in whole mount, TUNEL processed embryos, we observed a progressive wave, from rostral to caudal, of substantial dorsal CNS cell death (not shown) that presumably leads to the demise of *sf3b1<sup>b460</sup>* mutant embryos by 40 hpf.

The elevated levels of neural crest cell death in *sf3b1<sup>b460</sup>* mutant embryos raised the possibility that the abnormal development of the neural crest in *sf3b1<sup>b460</sup>* mutants results from neural crest cell apoptosis as a consequence, directly or indirectly, of abrogated Sf3b1 function. To test this possibility, we first investigated whether misexpression of *bcl2* mRNA in *sf3b1<sup>b460</sup>* mutant embryos significantly attenuates cell death as detected by TUNEL. We found that injection of 1-4 cell stage embryos resulted in a dramatic reduction in TUNEL-positive cells in *sf3b1<sup>b460</sup>* mutant embryos (Fig. 11 D,E)

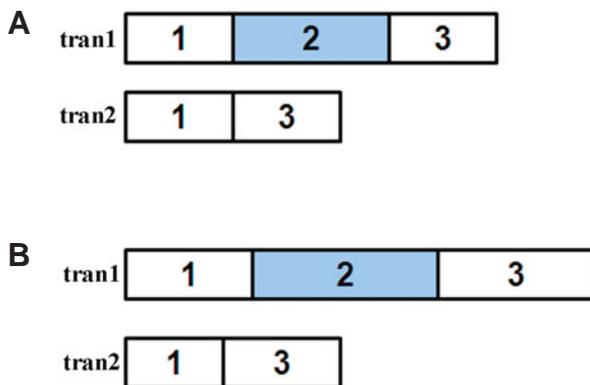
**Fig. 11. Analysis of cell death in *sf3b1<sup>b460</sup>* mutants.** (A-C) Double labeling with TUNEL (NBT/BCIP) and *crestin* whole-mount in situ hybridization (fast red). Transverse section is from the trunk region of a *sf3b1<sup>b460</sup>* mutant, dorsal is up. (A) DIC image/TUNEL; (B) fluorescence image/*crestin* of the same section; (C) (A,B) merged. (D) Dorsal view of uninjected *sf3b1<sup>b460</sup>* mutant at 16 hpf with TUNEL staining. (E) Dorsal view of *zbc12* mRNA injected *sf3b1<sup>b460</sup>* mutant at 16 hpf with TUNEL staining. Reduced TUNEL staining is evident on the left side. (F) Lateral view of live uninjected *sf3b1<sup>b460</sup>* mutant at 28 hpf. (G) Lateral view of live *zbc12* injected *sf3b1<sup>b460</sup>* mutant at 28 hpf, showing less CNS cell death as compared with uninjected *sf3b1<sup>b460</sup>* mutants but no melanophore rescue.

including cells along and in the immediate vicinity of the neural tube previously shown to be neural crest cells (see above). While the extent of CNS cell death observed in *sf3b1<sup>b460</sup>* mutants, confirmed by genotyping, was dramatically reduced (Fig. 11 D,E), we did not observe any rescue of neural crest-derived tissues such as craniofacial cartilage anlagen or in neural crest-derived cells such as melanophores when embryos were observed between 28 (Fig. 11 F,G) and 40hpf (not shown). Thus, although we cannot conclude that apoptosis plays no role in the development of the neural crest phenotype of *sf3b1<sup>b460</sup>* mutant embryos, these results strongly suggest that much of the abnormal neural crest development observed in mutant embryos results from miscues in the function of other key regulatory components of neural crest development that are dependent upon normal Sf3b1 function.

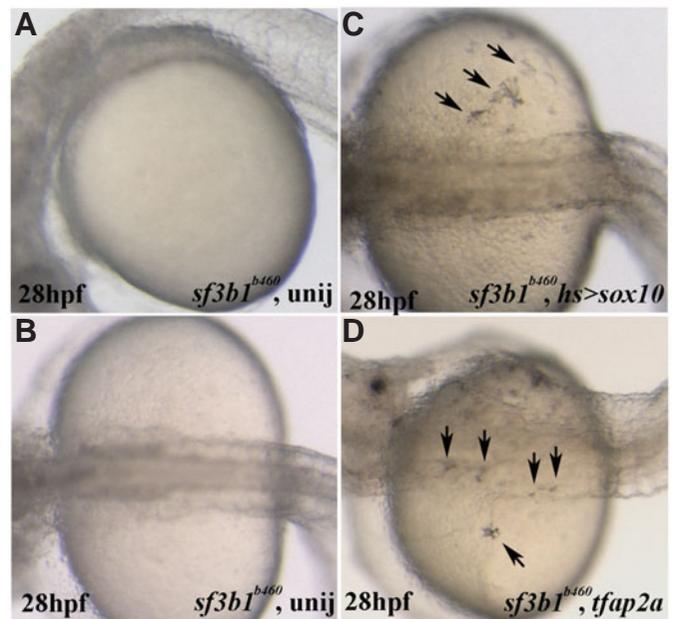
#### Aberrant pre-mRNA processing of a subset of transcriptional regulators of neural crest development

Because Sf3b1 is essential for pre-mRNA processing, and wild-type Sf3b1 protein levels in *sf3b1<sup>b460</sup>* mutant embryos are reduced by 90% due to aberrant pre-mRNA processing of *sf3b1* itself (Fig. 3C; see above), we examined the pre-mRNA processing of the essential transcriptional regulators of neural crest development whose neural crest expression is abnormal in *sf3b1<sup>b460</sup>* mutant embryos (Fig. 9; see above). To do so, we isolated RNA from 18-24hpf *sf3b1<sup>b460</sup>* mutant embryos and performed RT-PCR using gene-specific primers. Samples were then sequenced and compared to published sequence data. We found that only normal (identical to wildtype in size and sequence) transcripts for *foxd3*, *tfap2a* and *sox10* were present in *sf3b1<sup>b460</sup>* mutants suggesting that the relatively low levels of predicted functional Sf3b1 protein present in mutant embryos is sufficient for the correct pre-mRNA processing of these gene products. In contrast, we found that the pre-mRNA processing of *snai1b* and *sox9b* transcripts was severely disrupted in *sf3b1<sup>b460</sup>* mutants (Fig. 12).

In *sf3b1<sup>b460</sup>* mutant embryos, we were unable to detect nor-



**Fig. 12. Abnormal pre-mRNA processing in *sf3b1<sup>b460</sup>* mutant embryos.** (A) diagram of the exons of *snai1b*. tran1: shows the normal *snai1b* transcript. tran2: shows the abnormally spliced transcript in *sf3b1<sup>b460</sup>* mutant embryos, in which exon2 is deleted. Both transcripts were detected in mutant embryos. (B) diagram of the exons of *sox9b*. There are two types of transcripts, wildtype transcript (tran1) and the abnormally spliced transcript (tran2) whose partial exon1, whole exon2 and partial exon3 were deleted in *sf3b1<sup>b460</sup>* mutant embryos. Only the aberrant transcript was detected in mutant embryos.



**Fig. 13. Misexpression of genes that regulate neural crest development results in melanophore rescue in *sf3b1<sup>b460</sup>* mutant embryos.** (A) Lateral view of live uninjected *sf3b1<sup>b460</sup>* mutant at 28 hpf (anterior to left). (B) Dorsal view of live uninjected *sf3b1<sup>b460</sup>* mutant at 28hpf (anterior to left). (C) *hs>sox10* injected *sf3b1<sup>b460</sup>* mutant embryo at 28 hpf (dorsal view), showing partial melanophore rescue (arrows). (D) *tfap2a* mRNA injected *sf3b1<sup>b460</sup>* mutant embryo at 28hpf (dorsal view), showing partial melanophore rescue (arrows).

mal *sox9b* transcripts (GenBank Accession No. NM131644 and AF277097). Instead the only *sox9b* transcript detected was devoid of the 138-1379 bp coding region and thus not expected to encode any protein product (Fig. 12). While we cannot rule out the presence of small amounts of normal *sox9b* transcripts in *sf3b1<sup>b460</sup>* mutants present at levels below the limit of detection of methods employed, these results suggest that *sf3b1<sup>b460</sup>* mutants lack *sox9b* functional activity. The ability to detect *sox9b* mRNA by WISH in *sf3b1<sup>b460</sup>* mutants is likely due to the fact that the antisense probe used consists of 541 bp of the 3'-UTR present in the abnormal transcripts in addition to 404 bp corresponding to missing coding sequence. Thus, some specific hybridization may be expected in *sf3b1<sup>b460</sup>* mutants albeit at reduced efficiency.

We detected two species of *snai1b* transcripts in *sf3b1<sup>b460</sup>* mutants (Fig. 12). One species corresponded to the wildtype transcript (GenBank Accession. No. NM130989) whereas the other species contained a 513 bp deletion (106-618 bp) of coding sequence corresponding to a region encoding three of the five zinc finger motifs found in *snai1b* (Thisse *et al.*, 1995) and as such predicting that if a protein product is produced from the aberrant transcript that its function would be severely compromised. The antisense riboprobe used for WISH detection of *snai1b* is approximately 400 bp (Thisse *et al.*, 1995) corresponding to 5'-UTR sequence and coding sequence up to the first zinc finger. Thus, the riboprobe used would be predicted to recognize both normal and abnormal transcripts in *sf3b1<sup>b460</sup>* mutants. Therefore, the reduced levels of neural crest *snai1b* expression observed in *sf3b1<sup>b460</sup>* mutants using WISH is likely to underestimate actual functional Snai1b protein levels. In any case, the aberrant pre-mRNA processing of

*snai1b* in *sf3b1<sup>b460</sup>* mutant embryos strongly suggests that *Snai1b* function during neural crest development in *sf3b1<sup>b460</sup>* mutants is severely disrupted.

#### **Differential *sf3b1<sup>b460</sup>* mutant neural crest phenotype rescue activity of transcriptional regulators of neural crest development**

Because the neural crest expression of the transcription factors *foxd3*, *tfap2a*, *sox10*, *sox9b* and *snai1b* are severely disrupted in *sf3b1<sup>b460</sup>* mutants and because each of these transcription factors have been shown to be differentially required for aspects of early neural crest development in multiple species including zebrafish, we asked whether partial restoration of the function of each of these genes could rescue the neural crest phenotype of *sf3b1<sup>b460</sup>* mutants when misexpressed. To do so, we individually misexpressed each transcription factor via mRNA or inducible cDNA injected into 1-4 cell stage *sf3b1<sup>b460</sup>* mutant embryos. We allowed injected embryos to develop until 36hpf and examined each for the presence or absence of differentiated melanophores, which are entirely absent in uninjected mutant embryos, followed by genotyping using linked markers.

We found that misexpression of *hs>sox10* resulted in efficient (41% of injected *sf3b1<sup>b460</sup>* mutant embryos) melanophore rescue in *sf3b1<sup>b460</sup>* mutants (Fig. 13). In the vast majority of individual injected mutant embryos, multiple melanophores were observed in the head and trunk as well as over the yolk. Similarly, we observed a similar melanophore rescue when *tfap2a* mRNA was misexpressed in mutant embryos (Fig. 13), although qualitatively fewer melanophores were present in each rescued mutant embryos compared to *hs>sox10* injected mutants and rescue was slightly less efficient (34%). As was the case with *hs>sox10* injected mutants, melanophores in *tfap2a* injected mutants were observed in the head, trunk and over the yolk. In contrast to the melanophore phenotype rescue activities of *sox10* and *tfap2a* in *sf3b1<sup>b460</sup>* mutant embryos, we did not observe any differentiated melanophores in mutant embryos in which *foxd3*, *snai1b* or *sox9b* mRNAs were misexpressed.

## **Discussion**

In a screen for ENU-induced mutations that result in defects in neural crest development in zebrafish (Henion *et al.*, 1996), we isolated the *tst<sup>b460</sup>* mutant based on the absence of neural crest-derived melanophores. *tst<sup>b460</sup>* was found to be a recessive, embryonic lethal mutation with embryos dying at approximately 40hpf presumably due to prominent CNS cell death and/or the lack of circulation.

Multiple lines of evidence indicate that the pre-mRNA splicing factor *Sf3b1* is the locus mutant in *tst<sup>b460</sup>*. We found that; *tst<sup>b460</sup>* maps to a critical region of chromosome 9 containing *Sf3b1*; a BAC containing *Sf3b1* displayed phenotype rescue activity when misexpressed in mutant embryos; *tst<sup>b460</sup> Sf3b1* contains a T>G point mutation in the 5' splice site of the 4<sup>th</sup> intron and that *tst<sup>b460</sup>* embryos produce 3 aberrant, likely non-functional transcripts resulting from defective pre-mRNA processing that are consistent with this mutation; *Sf3b1* protein levels are reduced by approximately 90% in mutant embryos; knockdown of *Sf3b1* in wildtype embryos using two different morpholinos phenocopies *tst<sup>b460</sup>* mutants; morpholinos can increase phenotype severity in mutant embryos suggesting that *tst<sup>b460</sup>* is hypomorphic; misexpression of *Sf3b1* in *tst<sup>b460</sup>* mutant

embryos results in melanophore phenotype rescue; *tst<sup>b460</sup>* failed to complement a previously identified (Amsterdam *et al.*, 2004) viral insertion mutant in *Sf3b1*. Taken together, we conclude that *tst<sup>b460</sup>* is a strong hypomorphic *Sf3b1* mutant thus designated *Sf3b1<sup>b460</sup>*.

A prominent phenotype of *Sf3b1<sup>b460</sup>* mutants is the absence of neural crest-derived melanophores that is preceded by the abnormal neural crest expression of genes required for the development of and/or diagnostic of melanoblasts. This phenotype is in turn preceded by abnormal neural crest expression of transcription factors known to be required for early aspects of neural crest development (Gammill and Bronner-Fraser, 2003). These include *foxd3*, *tfap2a*, *snai1b*, *sox9b* and *sox10*. A subset of these, *sox9b* and *snai1b*, undergo defective pre-mRNA processing while a different subset, *sox10* and *tfap2a*, display neural crest rescue activity when misexpressed in *Sf3b1<sup>b460</sup>* mutants.

In zebrafish, mutants for *foxd3*, *tfap2a*, *sox9b* and *sox10* have been isolated and characterized (Dutton *et al.*, 2001; Knight *et al.*, 2003; Barrallo-Gimeno *et al.*, 2004; Yan *et al.*, 2005; Carney *et al.*, 2006; Montero-Balaguer *et al.*, 2006; Stewart *et al.*, 2006; Arduini *et al.*, 2009). Mutants for each of these transcription factors display both discreet and overlapping neural crest development phenotypes. The neural crest phenotypes of *Sf3b1<sup>b460</sup>* mutant embryos with respect to the expression patterns of these transcription factors, errors in pre-mRNA processing of a subset of these and the melanogenesis rescue activity of a different subset of these transcription factors provides multiple insights into the functional interactions among these genes in regulating early neural crest development and the role of *Sf3b1* in this context.

Specifically, neural crest expression of *sox10* is reduced, particularly in the trunk, of *Sf3b1<sup>b460</sup>* mutants and misexpression of *sox10* rescues melanogenesis in mutant embryos. Further, while *sox10* pre-mRNA appears to be processed normally in mutant embryos, melanoblast specification is severely disrupted ultimately resulting in the complete absence of differentiated melanophores in *Sf3b1<sup>b460</sup>* mutants. Because *sox10* is known to be globally necessary for melanogenesis, including in zebrafish (Kelsh and Eisen, 2000; Dutton *et al.*, 2001), it appears that the reduction in neural crest *sox10* expression levels in *Sf3b1<sup>b460</sup>* mutants is sufficient to prevent melanogenesis. But, since *sox10* pre-mRNA processing is normal in mutants, the role of *Sf3b1* in the regulation of *sox10* expression is indirect. Importantly, the neural crest expression of *sox9b*, *foxd3* and *tfap2a* are also reduced in *Sf3b1<sup>b460</sup>* mutants. Neural crest expression of *sox10* has been shown to be reduced in both *sox9b* (Yan *et al.*, 2005) and *foxd3* (Montero-Balaguer *et al.*, 2006; Stewart *et al.*, 2006) mutants and entirely abolished in *foxd3;tfap2a* double mutants (Arduini *et al.*, 2009). These results indicate that *sox9b* and *foxd3* regulate neural crest *sox10* expression while combined *foxd3* and *tfap2a* function also synergistically regulates *sox10*. However, *foxd3* and *tfap2a* transcripts are normally generated in *Sf3b1<sup>b460</sup>* mutants, albeit at reduced levels or fail to be maintained, respectively. Importantly though, *sox9b* pre-mRNA processing is severely disrupted in *Sf3b1<sup>b460</sup>* mutants leading to the absence of functional transcripts and *sox9b* has been shown to regulate the neural crest expression of both *foxd3* and *sox10* (and *snai1b*, see below; Yan *et al.*, 2005). In turn, maintenance of neural crest *tfap2a* expression is dependent upon *foxd3* (Stewart *et al.*, 2006). Taken together, these results suggest the possibility that failure in the *Sf3b1*-dependent pre-mRNA processing of *sox9b* in *Sf3b1<sup>b460</sup>* mutants results in downregulation of neural crest *foxd3*

and *sox10* expression, and that the resulting reduction in *foxd3* expression levels results in decreased neural crest expression of *tfap2a*. Thus, *sf3b1*-dependent pre-mRNA processing of *sox9b* is necessary for the establishment and/or maintenance of the transcription factor network that ultimately promotes *sox10*-dependent melanogenesis.

We also found that misexpression of *tfap2a* rescued melanogenesis in *sf3b1<sup>b460</sup>* mutants. While neural crest *tfap2a* expression is initially normal, expression is not maintained at wildtype levels. Analysis of transcripts indicates that pre-mRNA processing of *tfap2a* is normal in *sf3b1<sup>b460</sup>* mutants. In zebrafish *tfap2a* mutants (Knight *et al.*, 2003; Barrallo-Gimeno *et al.*, 2004), melanogenesis is significantly delayed, although appears to recover by 48-72 hpf. Lethality in *sf3b1<sup>b460</sup>* mutants (40 hpf) corresponds to the same timeframe in which melanogenesis is delayed in *tfap2a* mutants. Further, in addition to the delay in overt melanophore differentiation (pigmentation), the expression of both *mitfa*, and to a greater extent the levels and pattern of the expression of *c-kit* are also delayed in *tfap2a* mutants. In the latter case, *c-kit* is known to be required for melanoblast/melanophore migration and survival in zebrafish (Parichy *et al.*, 1999), suggesting that the reduction in *tfap2a* expression in *sf3b1<sup>b460</sup>* mutants could contribute to a delay in *mitfa*-dependent melanoblast specification as well as *c-kit*-dependent melanophore migration/survival thus contributing to the *sf3b1<sup>b460</sup>* mutant melanophore phenotype that can be partially rescued upon *tfap2a* misexpression.

Although misexpression of *snai1b* in *sf3b1<sup>b460</sup>* mutants and of itself failed to rescue the melanogenesis phenotype of mutant embryos, the fact that *snai1b* pre-mRNA processing is severely disrupted in mutants resulting in a significant reduction in the levels of neural crest *snai1b* expression may contribute to the overall neural crest phenotype of *sf3b1<sup>b460</sup>* mutants. Studies in multiple species indicate a role for *snai1b* in epithelial-mesenchymal transitions (EMT), including that which generates the premigratory neural crest population (Gammill and Bronner-Fraser, 2003; Nelms and Labosky, 2010). It is plausible that reduced neural crest *snai1b* expression in *sf3b1<sup>b460</sup>* mutants, due at least in part to aberrant pre-mRNA processing, could disrupt neural crest EMT and contribute to the increased levels of neural crest apoptosis observed in mutant embryos. A consequence could be a reduction in the overall neural crest population that could contribute to the derivative phenotypes observed in *sf3b1<sup>b460</sup>* mutants.

It should also be noted that while our discussion concerning neural crest derivatives has focused on melanophores with respect to the *sf3b1<sup>b460</sup>* mutant phenotype, the disruptions in the neural crest expression of the transcription factors analyzed as a consequence, directly or indirectly of disruption of Sf3b1 function, may also be involved in the development of the other neural crest derivative phenotypes including craniofacial progenitors and cranial ganglion neurons. In addition, there is a strong possibility that additional genes that regulate neural crest development, whose expression was not analyzed, may also be disrupted directly or indirectly in *sf3b1<sup>b460</sup>* mutants and contribute to the expression defects in the transcription factors analyzed and/or neural crest development.

Our results provide new insights into *sf3b1* function during development. For example, we have shown that the 90% reduction in Sf3b1 levels in *sf3b1<sup>b460</sup>* mutants results in selective defects in development rather than global early developmental arrest observed in mouse null mutants and when *sf3b1* is further knocked down

in *sf3b1<sup>b460</sup>* mutants. These results are similar to previous reports (Isono *et al.*, 2005; Lim *et al.*, 2006; Massiello *et al.*, 2006) and to recent reports indicating that inhibition of Sf3b1 function can lead to selective effects, in this case anti-proliferative effects of a Sf3b1 inhibitor due to selective aberrant pre-mRNA processing of cell cycle genes as opposed to a global disruption of pre-mRNA processing (Corrionero *et al.*, 2011; Folco *et al.*, 2011). Our results are also similar to recent findings indicating selective defects in early neural crest development when the functions of genes thought to be generalized regulators of transcriptional elongation are disrupted (Nguyen *et al.*, 2010; White *et al.*, 2011).

Lastly, we found that the development of neural crest is differentially affected in *sf3b1<sup>b460</sup>* mutants, potentially revealing new functional insights. We previously elucidated a framework gene regulatory network regulating neural crest cell diversification in which *foxd3* and *tfap2a* are globally and synergistically required for neural crest cell diversification and this function is due in large part to their regulation of the SoxE family genes *sox9a*, *sox9b* and *sox10* (Arduini *et al.*, 2009). Our results here have revealed new levels of interactions within this network and with additional genes (e.g. *snai1b*). Taken together, our results and those of others suggest that alterations in the function of the ubiquitous and essential pre-mRNA processing gene *sf3b1* may have implications for specific aspects of development as well as clinically relevant conditions in humans.

## Materials and Methods

### Zebrafish strains and maintenance

Zebrafish embryos and adults were raised and maintained in the Ohio State University zebrafish facility as described (Westerfield, 1993), and staged according to Kimmel *et al.*, 1995. The *toas<sup>b460</sup>* (*tsf*) mutation was induced in the wild-type \*AB genetic background by ENU (ethyl-N-nitrosourea) mutagenesis (Henion *et al.*, 1996). *tsf*WIK lines were obtained by outcrossing with wild-type WIK background fish for genetic mapping and genotyping. Heterozygous mating pairs (\*AB background or WIK background) were used to produce embryos (wild-type and mutant) for all experiments. The *hi3394a* (*sf3b1*) mutant line was kindly provided by the Hopkins's lab from MIT (Amsterdam *et al.*, 2004).

### Whole-mount *in situ* hybridization and immunohistochemistry

Whole-mount single and double *in situ* hybridization were performed as described (Jowett and Lettice, 1994; Jowett, 2001) with minor modifications. Embryos were fixed overnight in 4% PFA (paraformaldehyde) in 1X PBS (phosphate-buffered saline solution) at 4°C. Embryos older than 24 hpf were treated with PTU (1-phenyl-2-thiourea) to prevent melanin synthesis. Prehybridization and hybridization were carried out at 65°C. The following digoxigenin (DIG) labeled antisense RNA probes were used in single *in situ* hybridization: *crestin* (Luo *et al.*, 2001), *foxd3* (Odenthal and Nusslein-Volhard, 1998), *snai1b* (*snail-2*, Thisse *et al.*, 1995), *sox10* (Dutton *et al.*, 2001), *tfap2a* (Knight *et al.*, 2003), *sox9b* (Chiang *et al.*, 2001), *dlx2*, *dlx3* (Akimenko *et al.*, 1994), *islet-1* (Inoue *et al.*, 1994), *islet-2* (Appel *et al.*, 1995), *huC* (Park *et al.*, 2000), *dct* and *mitfa* (Lister *et al.*, 1999), *nkx2.5* (Lee *et al.*, 1996), and *myoD* (Weinberg *et al.*, 1996). Whole-mount double *in situ* hybridization was performed by using DIG labeled *myoD* antisense RNA probe and fluorescein labeled *crestin* antisense RNA probe as described (Jowett, 2001) with minor modifications.

*sf3b1* antisense and sense RNA probes were synthesized corresponding to 3'-*sf3b1* around 1.6 kb (2934-4393 bp) from a *PCR/Isf3b1c* construct linearized with *NotI* or *BamHI* separately and transcribed with SP6 or T7 separately. Other *sf3b1* antisense and sense RNA probes were made corresponding to 5'-*sf3b1* around 1.6 kb (1-1659 bp) from a *PCR/Isf3b1a* construct linearized with *NotI* and *HindIII* and transcribed with SP6 and

T7 separately.

Immunohistochemistry was performed as described (Henion *et al.*, 1996). Embryos were fixed in 4% PFA for 2 h at room temperature or overnight at 4°C. Antibodies were used at the following dilutions: zn-12 (Trevarrow *et al.*, 1990), 1:4000; acetylated tubulin (Sigma), 1:200; goat anti-mouse, 1:200; PAP, 1:200.

#### TUNEL assays

Whole-mount TUNEL staining was performed on zebrafish embryos fixed overnight with 4% PFA in 1X PBS at 4°C, permeabilized with methanol, and rehydrated. TdT/digoxigenin or fluorescein-dUTP reactions (Roche) were done for 1 h on ice, followed by 1 h at 37°C. After labeling with digoxigenin or fluorescein-dUTP, anti-digoxigenin or anti-fluorescein-AP (Roche) was used to bind digoxigenin or fluorescein. NBT/BCIP was used for color development. For whole-mount *in situ* hybridization and TUNEL double labeling, the TUNEL assay was performed first by using fluorescein-dUTP and NBT/BCIP for color development. After TUNEL staining, the embryos were incubated in 2mg/ml glycine (pH 2.9) for 10 min at room temperature to inactivate anti-fluorescein-AP, washed with 1X PBS to recover pH and then, WISH was performed as described. Fast red was used for detection.

#### Genetic mosaic assays

Genetic mosaic assays were performed by transplanting cells between wild-type and mutant embryos at the late blastula stage as described (Ho and Kane, 1990). The embryos obtained from *sf3b1<sup>b460</sup>* heterozygous carriers were divided into two groups, a donor group and a host group. Donor embryos were injected with rhodamine dextran (10<sup>3</sup> MW; Molecular Probes) at the 1- to 4-cell stage. Rhodamine dextran-labeled cells from donor embryos were transplanted into different regions of host embryos at the same developmental stage (dome stage 3.7-4 hpf) according to the zebrafish embryo fate map (Kimmel *et al.*, 1990). The host and donor embryos were grown under the same conditions (28.5°C). After 24 hpf, wild-type and *sf3b1<sup>b460</sup>* mutant embryos were identified based on phenotype. Since *sf3b1<sup>b460</sup>* mutants completely lack melanophores, it allowed us to easily assess the terminal cell fate of transplanted donor cells. Transplanted cells were visualized using fluorescence microscopy on a Zeiss Axioplan/DIC microscope.

#### Genetic mapping and positional cloning

*ts<sup>b460</sup>* heterozygous carriers (\*AB background) were crossed to WIK wild-type to produce a mapping line. Female heterozygous F1 generation fish were used for the *in vitro* production of parthenogenetic diploid F2 embryos. Homozygosity at most loci was achieved by suppressing the second meiotic division by application of hydrostatic pressure (Westerfield, 1993). Based on live phenotype, F2 diploid embryos were divided into two groups, homozygous mutants and wild-type embryos including homozygous and heterozygous wild-type embryos. The *ts<sup>b460</sup>* locus was mapped to its corresponding chromosome using linkage analysis with simple sequence length polymorphisms (SSLPs). Through recombination frequency analysis using haploid F2 mutants and WT embryos, several closely linked SSLPs and simple-strand conformation polymorphisms (SSCPs) were identified and a critical region was defined. Bacterial artificial chromosomes (BAC, commercially available from RZPD) and expressed sequence tags (ESTs, commercially available from RZPD) were searched for within this critical region through the zebrafish genome database at Wellcome Trust Sanger Institute ([http://www.sanger.ac.uk/Projects/D\\_rerio/](http://www.sanger.ac.uk/Projects/D_rerio/)). SSCP and SSCP markers that were used for fine genetic mapping:

ctg9339 566337  
 5'-CTGGAACCCCATGTCAAA-3'  
 5'-AGGTGCAAGGCAGAGTTAGC-3'  
 zk83M22SP6  
 5'-GGTTCATAGTAGCTGAAGACACCTG-3'  
 5'-GTCAGTACATTCAGCAGAACATTTCG-3'  
 zk83M22T7  
 5'-GGCATGCAAGCTTCTCGTTTGTAT-3'

5'-TCCACAACAGCTTCTAAATTCCGC-3'  
 zV4.5

5'-CCACGTCGATCTCCGAGTA-3'  
 5'-GTGCCTCTTCACGCCTACAT-3'

Zebrafish Microsatellite markers used were z60982, z21824, z54324, z35323.

#### BAC microinjection and phenotype rescue analysis

BAC DNA clones overlapping with the defined critical region used for injection were isolated using a Qiagen protocol for large plasmid isolation with Qiagen midiprep kit. For phenotype rescue assays, the BAC DNA was microinjected into a single blastomere of 1- to 2-cell stage embryos obtained from heterozygote intercrosses. The BAC DNA was diluted to 100-120 ng/μl as a working stock in sterile 1% phenol red. Injected embryos were scored according to live phenotype. Since *ts<sup>b460</sup>* mutant embryos completely lack melanophores and differentiated melanophores are easily observed in live embryos after 24 hpf, this allowed us to easily and accurately assess phenotypic rescue. Potential rescued mutants were genotyped with highly linked SSLP markers (z21824, z54324) for definitive confirmation. The embryos for injection of BACs were obtained by intercrossing heterozygous parents that are generated by crossing *ts<sup>b460</sup>* heterozygotes on \*AB background to WIK. Therefore, *ts<sup>b460</sup>* homozygous mutants could be identified by genotyping with polymorphic SSLP markers.

#### cDNA sequencing analysis in *ts<sup>b460</sup>* mutant and wild-type embryos

Total RNA from 18-24 hpf AB wild-type embryos and *ts<sup>b460</sup>* mutants was isolated using TRIzol reagent (Sigma). Qiagen one-step RT-PCR kit was used for RT-PCR with \*AB wild-type and *ts<sup>b460</sup>* mutant total RNA as templates. Primers used for *sf3b1* cDNA fragments are:

sf3b1(1-2703 bp)  
 5'-AAATGGCGCAGATCGCCAAA-3'  
 5'-CGGCACCAAGGTTACCCATGATTT-3'  
 sf3b1a(1-1659 bp)  
 5'-AAATGGCGCAGATCGCCAAA-3'  
 5'-ACCAAACCTCCCTGGCCTTATCTGT-3'  
 sf3b1b(1486-3159 bp)  
 5'-TGCTGGTCTGAAGTTGATGAGTCCA-3'  
 5'-TGCAGTTCTCCTGCACCTTCTCAT-3'  
 sf3b1c(2934-4393 bp)  
 5'-AAGCTTATGGCCATTTGGGTGTG-3'  
 5'-AAGAGCGAAGCTTGACAGACCAGGA-3'  
 sf3b1aa (46)F  
 5'-ATGACATTGAGGCTCAGATCCTGG-3'  
 sf3b1aa (579)R  
 5'-TCACTGCTTTCAGCTCTCCAGCTT-3'

The RT-PCR programs were as described in the Qiagen protocol (OneStep RT-PCR kit). The fresh RT-PCR products were cloned into PCR II vector by TA cloning (Invitrogen). The cDNAs obtained from RT-PCR were sequenced by the Ohio State University Plant-Microbe Genomics Facility (PMGF). The sequencing analysis was performed using Clustal X 1.81 and sequence analysis program from <http://www.ncbi.nlm.nih.gov>.

#### *sf3b1* cDNA cloning and misexpression analysis

The wild-type full length cDNA of *sf3b1* was assembled with wild-type cDNA fragment 1-2703 bp and EST clone *fb99f09*. The EST clone *fb99f09* (RZPD) was sequenced by PMGF at Ohio State University and it contains a 3'-*sf3b1* cDNA fragment which overlaps with the *sf3b1* cDNA fragment 1-2703 bp. The 2.5 kb fragment from *sf3b1* cDNA fragment 1-2703 bp in PCR II vector (digested with *XhoI/Clal*) and the 1.9 kb fragment from *fb99f09* EST clone (digested with *Clal/XbaI*) were cloned into the *XhoI/XbaI* site of *pCS2+* vector as *pCS2+sf3b1*. The *sf3b1* full length cDNA from construct *pCS2+sf3b1* with *XhoI* (blunt)/*XbaI* site was cloned into *Clal* (blunt)/*XbaI* site of the heatshock vector *pCSHSP* (Halloran *et al.*, 2000; Dutton *et al.*, 2001) as *hsp>sf3b1*. DNA of *hsp>sf3b1* was diluted to a final concentration of 25-30 ng/μl in 0.5% phenol red. *ts<sup>b460</sup>* mutant embryos and their wild-type

siblings from intercrosses of the *tst* WIK line were injected with *hsp>sf3b1* at the 1- to 2- cell stage and raised at 28.5°C. Embryos were heatshocked 2 or 3 times at 6-7 hpf, 11-12 hpf, and 23-24 hpf time points for 1 hour at 37°C. Embryos were genotyped as described previously.

*hs>sox10* (Dutton *et al.*, 2001) was injected into a single blastomere of *tst* mutant embryos at 1- to 2- cell stages and their wild-type siblings from intercrosses of *tst* WIK line. The injected embryos were heatshocked as described (Dutton *et al.*, 2001). The injected embryos were scored as in the BAC DNA injection experiments, and genotyped by the highly linked SSLP markers (z21824 and z54324).

Synthetic sense-polyA-capped mRNAs were transcribed *in vitro* from linearized templates using the mMESSEGE mMACHINE kit (Ambion). After transcription, the mRNAs were recovered as described (mMESSEGE mMACHINE kit manual). *pCS2+foxd3* (Stewart *et al.*, 2006), *pCS2+tfap2a* (Knight *et al.*, 2003), *pCS2+sox9a*, *pCS2+sox9b* (Yan *et al.*, 2005) and *pCS2+bcl2* (Langenau *et al.*, 2005), were digested with *NotI*. The linearized DNA was used as template for *in vitro* transcription by SP6 polymerase. The RNA for injection was diluted to a final concentration range from 50-100 ng/μl and injected into a single blastomere of *ts<sup>b460</sup>* mutant and wild-type embryos from intercrosses of *tst* WIK line at the 1- to 4- cell stage. The injected embryos were scored and genotyped as described above. The embryos injected with *bcl2* were fixed at 17 hpf and used for the TUNEL assay to detect programmed cell death as well as subsequently (see above) for phenotype rescue assessment.

#### Morpholino phenocopy analysis

Two Morpholino antisense oligonucleotides (MO) were designed and acquired from Gene Tools: *sf3b1*MO3rd that targets pre-mRNA splicing at the 3rd exon (ATGAATCCTCGTCATCATCCTAAA), and *sf3b1*ATGmo that targets the start site of *sf3b1* (GGCGATCTGCGCCATTTTCGTGCTG). Each morpholino was injected separately into one blastomere of wild-type embryos at the 1- to 4- cell stage to interfere with either pre-mRNA splicing of *sf3b1* or to block translation, respectively. Wild-type embryos for morpholino injections were divided into two groups: morphants (embryos injected with morpholinos) and uninjected wild-type embryos. Efficiency of *sf3b1*MO3rd against the splice site was evaluated by RT-PCR with primers *sf3b1*aa (46)F and *sf3b1*aa (57)R. Western blot with mouse *Sf3b1* monoclonal antibody (Horie *et al.*, 2003) was used to determine the efficiency of *sf3b1*ATGmo.

#### Western blotting

Western blot analysis was performed as described (Monani *et al.*, 2003) with minor modifications. Dechorionated zebrafish embryos (12 hpf-24 hpf) were dissolved in blending buffer (10% SDS, 50mM Tris HCl, pH 6.8, 10mM EDTA, μl/each embryo) and incubated in boiling water for 5-10 min. BAC protein assay kit (PIERCE) was used for measuring protein concentration. 80 μg of protein from different embryos (Wild-type embryos, *ts<sup>b460</sup>* homozygous mutants, *hi3394a* homozygous mutants, *sf3b1*ATGmo morphants, *hsp>sf3b1* injected *ts<sup>b460</sup>* mutant embryos with phenotypic rescue) mixed with an equal volume of sample buffer (62.5mM Tris HCl, pH 6.8, 10% glycerol, 0.1% Bromophenol blue, 10% β-mercaptoethanol) was loaded and electrophoresed on 8% polyacrylamide gels. Then, samples were transferred to nitrocellulose membranes (Whatman GmbH). TBS Blotto A (Santa cruz biotechnology, Inc.) was used as a blocking reagent. Primary antibodies for western blotting are mouse monoclonal antibodies, anti-*Sf3b1* (kindly provided by Dr. K. Isono; Horie *et al.*, 2003) and anti-actin as protein loading control (Abcam Inc.). HRP-conjugated secondary antibody was used to bind primary antibody and detected by chemiluminescence system (ECL western blotting analysis system, Amersham Bioscience).

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