

# Noc1 downregulation induces nucleolar stress and upregulates p53 isoforms, with a robust increase of the truncated p53E isoform in *Drosophila* wing discs

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Disruption of ribosome biogenesis triggers nucleolar stress, a conserved cellular response that activates p53. We previously demonstrated that depletion of Nucleolar Complex Protein 1 (Noc1) in *Drosophila* wing imaginal discs impairs rRNA maturation and ribosome assembly, resulting in elevated p53 levels and apoptosis, hallmarks of nucleolar stress. The *Drosophila* p53 gene produces four mRNA isoforms, yet their individual contributions to nucleolar stress responses remain poorly understood. Using newly designed isoform-specific qPCR primers, we found that although all p53 isoforms exhibit moderate transcriptional changes following Noc1 reduction, the truncated isoform p53E is robustly and preferentially upregulated. Notably, p53E lacks the N-terminal transactivation domain and has been reported to negatively regulate p53-induced apoptosis in specific tissues. Furthermore, our analyses indicate that  $\gamma$ -H2AV accumulation arises from caspase-dependent apoptosis rather than primary genomic lesions, suggesting the activation of a p53-dependent stress pathway distinct from canonical genotoxic pathways. Together, these findings suggest that p53E may be part of a novel mechanism activated during nucleolar stress, providing insight into how cells adapt to defects in ribosome biogenesis.

**Keywords:** p53 isoforms; nucleolar stress; Noc1; apoptosis;  $\gamma$ -H2AV; primer specificity; isoform primer design; mutant screen; *Drosophila* imaginal discs

## Introduction

Nucleolar stress is a cellular response to unbalanced ribosomal biogenesis (Boukoura and Larsen 2024). The nucleolus acts as a key stress sensor, responding to perturbations in ribosomal RNA (rRNA) processing and ribosome biogenesis by inducing the upregulation of p53 (Lindström et al. 2022).

*Drosophila* p53 exhibits significant structural and functional similarities with human p53 (Brodsky et al. 2000; Ollmann et al. 2000; Bourdon et al. 2005; Ingaramo et al. 2018). The p53 gene (FBgn0039044) maps to chromosome 3R and encodes four mRNA isoforms (p53A, p53B, p53C, and p53E). According to the current deep RNA sequencing-based genome annotation, these isoforms encode three distinct proteins (Boley et al. 2014). Notably, p53A, also known as  $\Delta$ Np53, and p53C differ in their 5' untranslated regions (5'UTRs), and while the protein product of p53C remains unconfirmed, its open reading frame (ORF) is predicted to be identical to that of p53A, and to encode a 44 kDa protein (Zhang et al. 2015; Chakravarti et al. 2022). This suggests that differential promoter regulation underlies their distinct functions in different cellular contexts. p53B is the longer isoform, carrying a unique N-terminal domain and encoding for a 56 kDa protein considered the closest homolog to human p53 (TP53) (Ingaramo et al. 2018). p53E is the shorter isoform that encodes a predicted 38 kDa protein lacking part of the N-terminal transactivation domain, and

is considered to act as a dominant negative in specific tissues and stress conditions (Wylie et al. 2022).

p53A and p53B are the most characterized protein isoforms. While their expression is comparable in the germline, they begin to exhibit distinct expression and regulation patterns across various developmental stages (Chakravarti et al. 2022). In somatic tissues, p53A mRNA and protein were found to be more abundant, likely reflecting differential roles during tissue development (Chakravarti et al. 2022). This may reflect their role in controlling apoptosis.

Initial studies demonstrated that overexpression of each p53 protein isoform leads to embryonic lethality. In addition, it was demonstrated that overexpression of p53A or p53B and of p53E induced apoptosis in the developing eye when driven by the GMR promoter, while in the ovary, overexpression of p53A and p53B proteins markedly enhanced apoptosis following irradiation, whereas p53E overexpression reduced the number of apoptotic cells, suggesting a dominant-negative role for this isoform in the ovary under irradiation-induced stress conditions (Zhang et al. 2015). Numerous studies have demonstrated that both p53 protein A and B isoforms play distinct roles in regulating stress responses and inducing apoptosis; indeed, p53A was shown to be both necessary and sufficient to activate the proapoptotic genes *rpr* and *hid* in response to DNA damage in somatic and germline tissues (Zhang et al. 2015; Chakravarti et al. 2022).

Although p53B can induce apoptosis when experimentally overexpressed, its behavior in normal tissues appears to be different. In several somatic tissues, the mRNA levels of p53B are only slightly lower than those of p53A, yet the amount of p53B protein detected is considerably lower. This discrepancy suggests that p53B may be translated less efficiently or undergo more rapid protein degradation (Zhang et al. 2014, 2015). These differences in protein abundance and activity among tissues may stem from competition between p53 isoforms that have opposing regulatory functions. Specifically, p53B primarily acts as a transactivator, promoting the expression of target genes, whereas p53A, which is more broadly expressed across tissues, can function as both a transactivator and a transrepressor (Wylie et al. 2022). Thus, the relative levels and functional properties of these isoforms likely contribute to the tissue-specific outcomes of p53 signaling. In support of this hypothesis, studies in photoreceptor neurons have shown that the balance between autophagy-mediated survival and apoptosis-driven cell death is governed by antagonistic regulatory inputs from p53A and p53B, whose mRNA levels shift differentially in response to oxidative stress (Robin et al. 2019).

Although p53 is best known for its conserved role in apoptosis triggered by DNA damage, it also exerts broad control over tissue physiology and metabolic homeostasis. Under nutrient-limiting conditions, p53-null mutant flies fail to sustain normal survival, revealing that p53 is essential for coordinating metabolic adaptation and efficient nutrient consumption during stress (Barrio et al. 2014). p53 also regulates growth and proliferation in a coordinated, non-autonomous manner and modulates apoptosis in response to stress, such as rRNA depletion (Mesquita et al. 2010). Furthermore, p53 is necessary to maintain imaginal disc plasticity and regenerative potential (Wells and Johnston 2012) and to act as a general sensor of competitive confrontation during MYC-induced cell competition (de la Cova et al. 2014). Notably, p53 isoform-specific protein functions have not been assessed in these experiments despite growing evidence that their function can differ significantly depending on the cellular context in which they are expressed. Thus, it may be important to consider that distinct signals and tissue-specific expression programs could drive isoform-specific roles in stress response and gene regulation.

Our previous study demonstrates that reducing the nucleolar gene *Noc1* (FBgn0036124) impairs rRNA processing and ribosome biogenesis, leading to apoptosis and the upregulation of p53 (Destefanis et al. 2022; Rambaldelli et al. 2025), both hallmarks of nucleolar stress. Using our newly designed p53 isoform-specific qPCR primers, we found that p53E is the most upregulated isoform following *Noc1* reduction. This suggests that nucleolar stress may induce a novel regulatory mechanism for p53E, whose strong upregulation upon nucleolar disruption points to a potential autoregulatory interaction in which one p53 isoform may influence the activity or stability of other isoforms. In addition,  $\gamma$ -H2AV phosphorylation accumulates because of caspase-dependent cell death rather than direct genomic lesions, consistent with the activation of a p53 stress pathway distinct from the canonical genotoxic pathway. Together, these findings suggest that p53E may be part of a novel mechanism activated during nucleolar stress, providing insight into how cells adapt to defects in ribosome biogenesis.

## Materials and methods

### Drosophila husbandry and lines

Animals were raised at low density in vials containing standard fly food, composed of 9 g/l agar, 75 g/l corn flour, 50 g/l fresh yeast,

30 g/l yeast extract, 50 g/l white sugar, and 30 ml/l molasses, along with nipagin (in ethanol) and propionic acid. The crosses and flies used for the experiments are kept at 25 °C, unless otherwise stated.

Line used: *w*<sup>1118</sup> from VDRC #6000, *Rotund-Gal4* BDSC#76179, *UAS-Noc1-RNAi* BDSC #25992, *UAS-p35* (Destefanis et al. 2022), *p53*<sup>5-1-4</sup> BDSC #6815, *p53*<sup>A2.3</sup> and *p53*<sup>B41.5</sup> from (Chakravarti et al. 2022).

### Alkaline comet assay

Using the *UAS-Gal4* system (Brand and Perrimon 1993), *Noc1* was downregulated in the entire imaginal disc using a temporally controlled expression system under the *Actin-Gal4* promoter in combination with the temperature-sensitive *Gal80-ts* allele that inhibits *Gal4* (Germani et al. 2018). Animals were left to develop for five days after egg laying (AEL) at the permissive temperature of 18 °C, then shifted to 30 °C for 60 h, during which *Gal4* was active and induced the ubiquitous expression of *Noc1-RNAi*. Wing imaginal discs were dissected and incubated in Ringer's solution, then enzymatically digested with Collagenase in TrypLE at 37–38 °C for 30 min following the protocol in (Pederzoli et al. 2025). After electrophoresis and Hoechst staining, photos of nuclei were imaged using a Zeiss Axio Imager fluorescent microscope with a 40× objective, and DNA damage was analyzed with CometScore 2.0.

### RNA extraction and RT-qPCR

RNA extraction was performed with the RNeasy Mini Kit (Qiagen). Five larvae or 30 wing discs were collected and placed in 250  $\mu$ l of lysis buffer, then processed following the manufacturer's instructions. The isolated RNA was quantified using a Nanodrop (Thermo Scientific) and stored at –80 °C for extended periods. RNA (1  $\mu$ g) was retro-transcribed into cDNA using the SuperScript IV VIL0 Master Mix (Invitrogen). DNase was added to the samples. To perform retro-transcription, the mix was incubated at 25 °C for 10 min, 50 °C for 10 min, and 85 °C for 5 min. The obtained cDNA was used to perform quantitative PCR (qPCR) with the SYBR Green PCR Kit (Qiagen). The final mix for each sample to be tested was composed by 8  $\mu$ l of cDNA, 2  $\mu$ l of primers (10  $\mu$ M), and 10  $\mu$ l of SYBR Green reaction mix. Reactions were carried out using a BioRad CFX96 machine with the following protocol: 50 °C for 2 min, 95 °C for 2 min, 95 °C for 15 s, and 60 °C for 30 s (repeated for 45 cycles), followed by 72 °C for 1 min. Analyses were done using Bio-Rad CFX Manager software. The relative abundance level for each transcript was calculated by subtracting the value of actin from the value of p53 and using the 2<sup>- $\Delta\Delta$ Ct</sup> method. For qPCR in wing discs analysis, the RNA was extracted from at least 20 wing imaginal discs, or from at least five larvae for each genotype. Data are expressed as arbitrary units (A.U.). The number of independent biological replicates is cited in the figure legend for each data set. The sequences of the primers used are in Table 1.

### Dissections and immunofluorescence

Larvae were collected at the third-instar stage, dissected in 1× phosphate-buffered saline (PBS), and fixed for 30 min in 4% paraformaldehyde (PFA) at room temperature (RT). After 15 min of tissue permeabilization with 0.3% Triton X-100, samples were washed in PBS with 0.04% Tween 20 (PBS-T) and blocked in 1% bovine serum albumin (BSA) for 1 h at RT. Samples were incubated overnight at 4 °C with primary antibodies in 1% BSA and, after washing, with Alexa-Fluor-488- or 555-conjugated secondary antibodies in BSA. During washing in PBS-T, nuclei were stained

**Table 1.** Sequences of the primers used in this work.

Target	Sequence	Reference
p53A/C	F: TATATCACAGCCAATGTCTGTG R: CCTCCGTGGAGTCATCC	This work
p53B	F: CCAACAAGGGCACTGATTC R: ATTCCGATCCCGATACTC	This work
p53C	F: GTGAGCAAATTCAAAAACGCG R: CCATTCGCCACGCAG	This work
p53E	F: GAGCGCAAACAATCCGT R: TGTGATTCTCTAGCTTGGGC	This work
p53 pan isoform	F: CCAAGCTAGAGAATCACAAACATCG R: TCGAGTACATCCAAAGAGACTTGG	(Zhang et al. 2014)

with Hoechst 33342 (Thermo Fisher Scientific). Imaginal discs were dissected from the carcasses and mounted on slides with Vectashield (LsBio-Vector Laboratories). Images were acquired using a Leica SP8 confocal microscope or Zeiss Axio Imager fluorescent microscope and analyzed with the Fiji program to measure fluorescence. Images were assembled using Adobe Photoshop 2025. Primary antibodies used were rabbit anti-cleaved Caspase3, 1:400 (Cell Signaling 9661); and anti p2HAV, 1:1000 (DSHB#UNC93-5.2.1)(Lake et al. 2013).

### Image processing

Raw confocal images were processed with Fiji to obtain a z-stack projection (using the Average Intensity or Max Intensity options). Then, all images from the same experiment were processed, enhancing the brightness of all channels equally across all images.

## Results

### Reduction of Noc1 triggers DNA double-strand breaks, p53 upregulation, and apoptotic signaling that regulates $\gamma$ H2AV phosphorylation

We previously demonstrated that downregulation of the nucleolar factor Noc1 in *Drosophila* tissues leads to the accumulation of rRNA (Destefanis et al. 2022; Rambaldelli et al. 2025). Furthermore, Noc1-RNAi expression driven by the *rotund* promoter in cells of the wing imaginal discs induces DNA damage as evidenced by the comet-like morphology of the nuclei in the affected cells (Fig. 1a and b, and (Pederzoli et al. 2025)). These nuclei show a statistically significant increase in the comet tail length (or tail moment, calculated as tail length multiplied by the percentage of DNA in the comet head), indicating the presence of double-strand breaks absent in the nuclei of wild-type  $w^{1118}$  cells (Fig. 1c). Notably, reduction of Noc1 is accompanied by a significant increase in apoptosis in cells of the wing pouch (Fig. 1d–f), together with the upregulation of p53 at both transcriptional and protein levels (Fig. 1g and h).

To assess the impact of Noc1 downregulation on DNA damage signaling, we monitored  $\gamma$ H2Av phosphorylation, a well-established marker of double-strand breaks. Interestingly, blocking apoptosis prevented  $\gamma$ H2AV accumulation (Fig. 1i), indicating that in this context,  $\gamma$ H2AV phosphorylation is downstream of apoptotic signaling rather than a direct consequence of Noc1 loss. This suggests that apoptosis itself contributes to the observed DNA damage response in these tissues. Together, these findings are consistent with a cellular response to nucleolar stress resulting from impaired rRNA processing upon Noc1 reduction. Based on these data, we designed isoform-specific qPCR primers to determine which individual p53 isoforms respond to Noc1 reduction, as they may respond specifically to nucleolar stress.

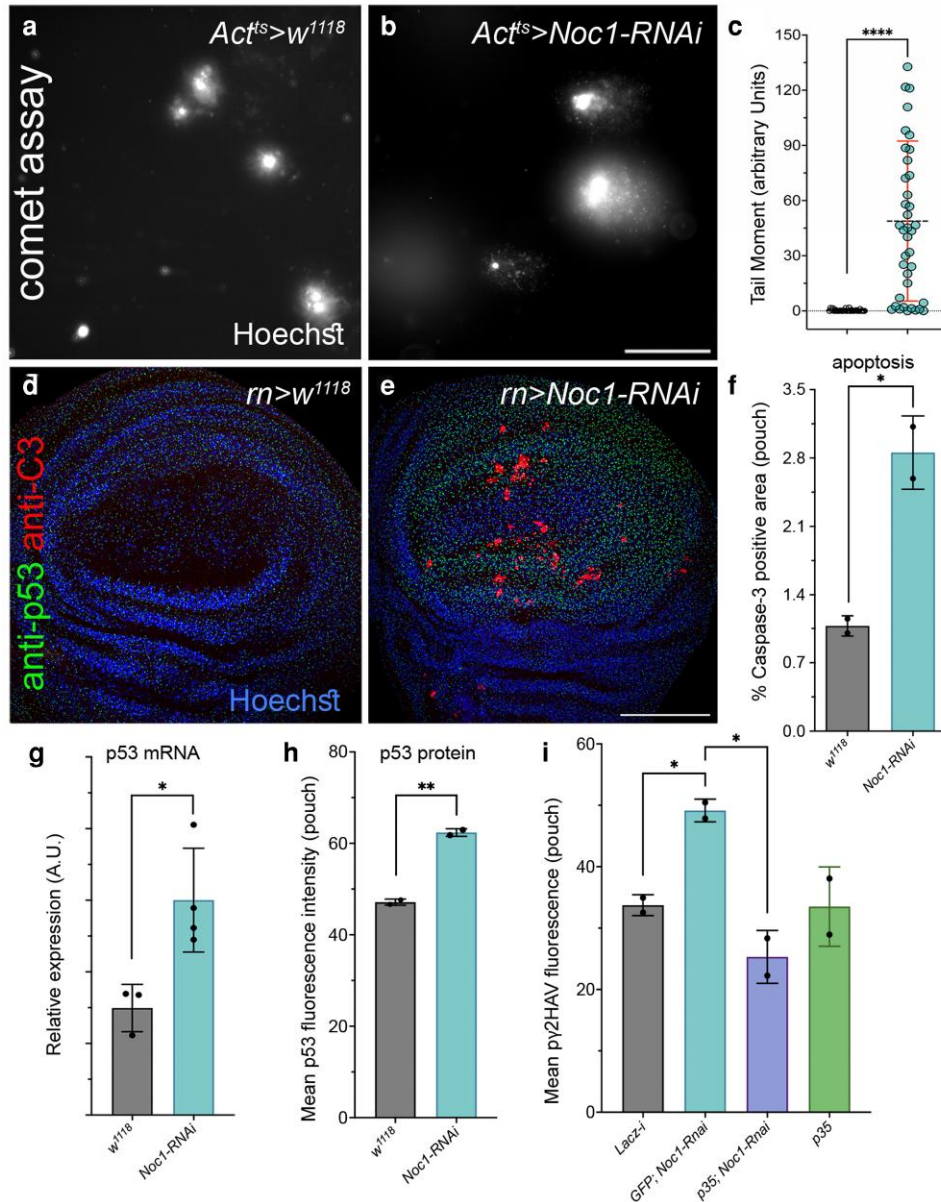
### P53 locus and novel primers for detecting distinct p53 isoforms

The *Drosophila* genome contains a single p53 locus (Fig. 2a, FBgn0039044 and NCBI gene ID: 2768677) on chromosome 3R, which produces four distinct mRNA transcripts through alternative promoter usage and splicing, resulting in four protein isoforms (Fig. 2b). The canonical p53A (also referred to as  $\Delta$ Np53) lacks part of the N-terminal Transactivation Domain (TAD), producing a protein of approximately 44 kDa. The p53B transcript retains the full TAD, encoding for a protein of about 56 kDa, while the rest of the sequence is identical to that of p53A mRNA. p53C protein shares the same amino acid sequence as p53A, but their mRNAs originate from an alternative transcription start site and are predicted to produce a longer transcript. Finally, p53E is the shortest isoform, predicted to encode a 38 kDa protein that lacks the N-terminal TADs of p53A and p53B, but retains the DNA-binding domain. This isoform has been shown to inhibit the apoptotic response to ionizing radiation, suggesting that it may act as a dominant-negative regulator of the other p53 isoforms (Zhang et al. 2015; Wylie et al. 2022).

Although primer sets have previously been reported for distinguishing among p53 mRNA isoforms (Supplementary Fig. 1), none have been fully optimized to reliably differentiate between p53A and p53B. Additionally, primers capable of specifically discriminating between p53B and p53C, or uniquely amplifying p53C, remain unavailable. To overcome these limitations, we designed and optimized a novel set of primers to enable precise and isoform-specific analysis of p53 transcripts (Table 1).

To ensure isoform specificity and accurate validation, primers were designed using the sequences of the available p53<sup>5A-1-4</sup> null lines and the isoform-specific p53A and p53B mRNA mutant lines. Briefly, the p53A null line carries a 23 bp deletion and 7 bp insertion at the end of the first exon which expands also in the first intron (Robin et al. 2019); the mutant p53B null line has a 14 bp deletion and 1 bp insertion in the second exon, resulting in multiple early STOP codons (Chakravarti et al. 2022); and the p53<sup>5A-1-4</sup> null line carries a large deletion of approximately 3.3 Kb that removes four exons, preventing the transcription of a functioning mRNA, resulting in a complete loss of p53 protein expression (Fig. 2c) (Rong et al. 2002).

The new primers for the p53A isoform span the exon-exon junction between exons 1 and 2 (Fig. 2d-i, shown as light green arrows in the panel, and Table 1), providing enhanced specificity compared to earlier designs (shown as arrows in pink, and Table 1). Like the previously available primers, this set can be used to assess p53A expression in the p53A null line as the forward primer anneals upstream of the deleted region, while the reverse primer anneals right after the deletion, allowing the amplification of a specific region of the mutant (Fig. 2d-ii). A limitation of the



**Fig. 1.** Noc1 downregulation induces DNA double-strand breaks (DSBs), leading to p53 upregulation and apoptosis, which contributes to  $\gamma$ H2AV phosphorylation. (a–c) Comet assay showing fluorescent images of nuclei stained with Hoechst (white) from cells of the wing imaginal discs of *w<sup>1118</sup>* third instar larvae (a), and from animals in which *Noc1* was transiently reduced for 60 h under the temporal control of the *Act-Gal80<sup>5</sup>* system. b), scale bar is 50  $\mu$ m. c) Quantification of single-cell tail moment from the comet assay performed on nuclei from the genotypes shown in (a and b). These data are obtained from pooling at least 20 wing imaginal discs of each genotype, and one representative experiment is shown. Statistical analysis was performed using the non-parametric Kruskal-Wallis as described in (Pederzoli et al. 2025). Experiments in (d–i) are from wing imaginal discs from third instar control larvae (*m > w<sup>1118</sup>*) or from larvae in which expression of the transgene was reduced using the indicated Rnai lines in the wing pouch under the control of the *rotund* (*m*) promoter. d and e) Confocal images of wing imaginal discs from third instar larvae immunostained for the expression of p53 protein (green) and for Caspase3, as a marker of apoptosis (red). Nuclei were stained with Hoechst (blue). Scale bar is 100  $\mu$ m. f) Quantification of Caspase3 by immunofluorescence in cells from the pouch of wing discs from third instar larvae of the indicated genotype. g) qPCR analysis from RNA extracted from at least 20 wing imaginal discs from larvae of the indicated genotype, p53 mRNA expression was analyzed using p53 pan isoform primers (Table 1). Data are expressed as arbitrary units (A.U.) (H). Quantification of p53 by immunofluorescence was analyzed in the pouch of wing imaginal discs from animals of the indicated genotype. i) Quantification of  $\gamma$ H2AV phosphorylation by immunofluorescence was analyzed in the wing pouch from larvae of the indicated genotype. In (c), the asterisks represent the *P*-values from Student's *t*-test \*\*\*\**P* < 0.0001, in one out of three independent experiments performed, where the dots represent the number of nuclei analyzed. In (f), (g), (h), and (i), the dots represent the number of independent biological replicates, with at least five imaginal discs analyzed for each genotype. The asterisks represent the *P*-values from Student's *t*-test \**P* < 0.05 and \*\**P* < 0.01; the error bars indicate the standard deviations.

new primers, as with previously published ones, is their inability to distinguish between *p53A* and *p53C* transcripts, due to the high sequence similarity between them. To design primers capable of distinguishing *p53B* from *p53C* isoform, we took advantage of the difference in the length of their 5' untranslated region

(5'UTR) (Fig. 2e). While the reverse primer anneals in a sequence common between *p53B* and *p53C*, the forward primer for *p53B* anneals at the end of the second exon and extends into the beginning of exon 3 (green arrows in Fig. 2e-i and 2e-iii). This design maximizes specificity, ensuring amplification of only *p53B*. Moreover,

this set of primers is suitable for screening p53B in p53B null lines, as they do not anneal within the deleted region (Fig. 2e-ii).

Through detailed analysis of the 5'UTR sequence of p53C, we identified a region unique to this isoform (highlighted in orange in Fig. 2f-i). Thus, we designed primers that specifically target this region, enabling the selective detection of p53C mRNA and providing a distinct primer set for this isoform. Furthermore, the availability of p53C-specific primers allows for a more accurate interpretation of results obtained with the p53A primer set, which cannot distinguish between the p53A and p53C transcripts.

In addition to p53A, p53B, and p53C, we developed isoform-specific primers for the truncated p53E isoform. These primers target the unique 5' sequence generated by the p53E transcription start site (Fig. 2g), providing selective amplification and preventing cross-detection with other isoforms.

Primer specificity was tested by performing qPCRs on RNA extracted from imaginal discs of  $w^{1118}$  third-instar larvae. All primers amplified a single product, as evidenced by a single band observed in the gel (Fig. 2h) and confirmed by melting curve analysis, which showed a distinct peak for each primer (Supplementary Fig. 2).

### Validation of isoform-specific primers in wild-type and p53 mutant background

To further assess the specificity and to complete the characterization of the newly designed primer sets, we conducted qPCR analysis on RNA from wild-type control  $w^{1118}$  and p53 mutant lines using primers targeting the single p53 mRNA isoforms (Table 1).

As shown in Fig. 3a, the p53 pan primers, which target a region common to all the p53 isoforms (Fig. 2b-i), detect p53 transcript levels in the p53A null line that are comparable to those of control  $w^{1118}$ . In contrast, the p53B null line exhibits a small but not significant increase in p53 mRNA expression, suggesting a potential compensatory upregulation in response to the loss of p53B.

Using the p53A/C primers, we observed that the p53A/C mRNA levels were significantly reduced by approximately 50% in the p53A null line (Fig. 3b). In contrast, the p53B null line showed a significant twofold increase in p53A/C mRNA expression. Consistent with the data in Fig. 3a, these findings may suggest the presence of a compensatory response to the other isoforms when p53B is eliminated in null animals, whereas loss of p53A has minimal effect on overall transcript abundance.

In the p53B null line (Fig. 3c), we observed an unexpected significant upregulation of p53B mRNA. However, since the mutation does not prevent transcription and the primers amplify regions downstream of the deletion, it is likely that the mutant transcript can still be detected by qPCR. Overall, this finding suggests that compensatory or autoregulatory feedback mechanisms may act to increase p53B transcription or stabilize the mutant mRNA in response to reduced levels of p53B protein.

Finally, analysis of the p53C isoform, using primers that specifically detect the transcript, reveals a significant reduction in its expression in both p53A and p53B null lines (Fig. 3d), suggesting that both p53A and p53B isoforms, directly or indirectly, participate in the regulation of p53C expression.

Importantly, the complete absence of p53 transcripts in the  $p53^{5A-1-4}$  null line confirms the specificity and reliability of the isoform-targeted primer set used in this study.

### p53E is the predominantly upregulated isoform upon Noc1 downregulation

Analysis of p53 expression in cells of the wing imaginal disc in which Noc1 was reduced in the pouch revealed a significant upregulation of p53 mRNA levels when assessed using pan-isoform

primers (Figs. 4a and 1g). Because these primers recognize a conserved region shared by all four p53 isoforms, as illustrated in the magnified view (Fig. 2b-i), this initial analysis did not allow discrimination between individual isoform contributions. Given that Noc1 reduction leads to rRNA accumulation and increased p53 levels (Fig. 1b and c), and considering that p53 isoforms can be expressed at different levels under various stress conditions, we therefore designed isoform-specific primers to determine which p53 isoform(s) are preferentially upregulated in response to Noc1-RNAi-induced nucleolar stress.

Among the isoforms, p53E exhibited the strongest induction ( $P < 0.001$ ), whereas all the other isoforms showed a moderately significant increase ( $P < 0.05$ ). Notably, p53E has been reported to act as a dominant negative in a tissue-specific manner. Thus, our analysis suggests that p53E may contribute to the repression of specific target genes during the nucleolar stress response.

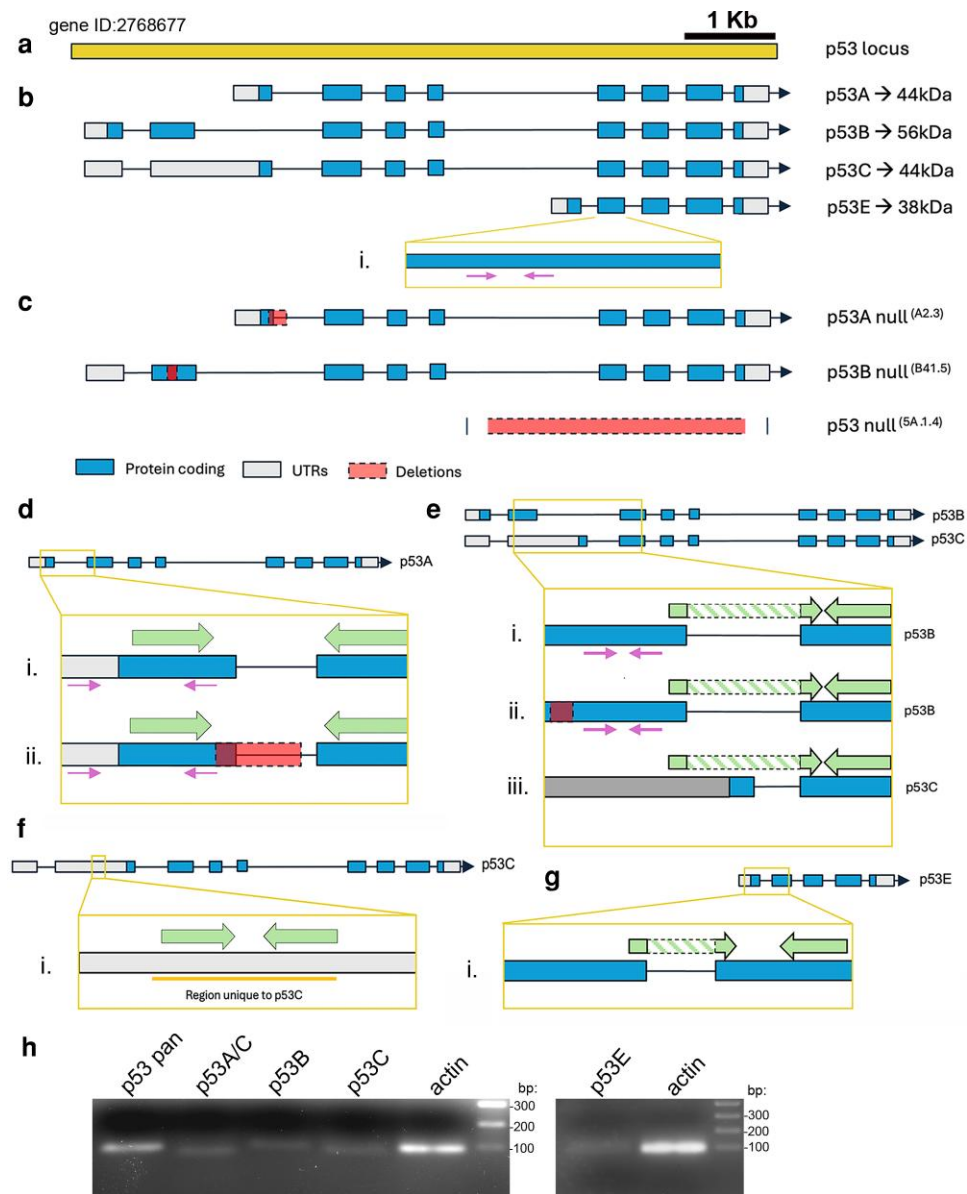
Collectively, these findings highlight an isoform-specific regulation of p53 in response to nucleolar stress, with p53E emerging as a potential novel key mediator of transcriptional repression. The differential upregulation of p53 isoforms suggests a fine-tuned cellular strategy, where transrepressor isoforms may selectively modulate gene expression to orchestrate an appropriate stress response, while other isoforms potentially maintain canonical p53 functions. This underscores the importance of considering isoform-specific roles in understanding p53-mediated pathways under nucleolar stress conditions.

## Discussion

In this study, we present an optimized and validated set of isoform-specific primers designed for the accurate detection and quantification of *Drosophila* p53 transcript variants. Here, we also designed primers to specifically detect the unique p53C isoform, enabling precise isoform discrimination. We first utilized p53 mutant lines to validate the specificity of these primers. These analyses show that the p53A null line retains some of p53A and p53C transcript levels (Fig. 3b and d). Moreover, despite the introduction of premature stop codons within the coding sequence of the p53B mutation, our qPCR analysis revealed an upregulation of p53B-mRNA in p53A/C and B mutant lines (Fig. 3d and c), which may suggest the presence of a compensatory mechanism, to buffer the loss of p53B function, or to compensate for the reduction of p53A/C protein, highlighting isoform-specific plasticity of the p53 network, where alternative isoforms may adjust expression to preserve critical stress response functions.

Our analysis of isoform-specific p53 expression under nucleolar stress induced by Noc1 reduction (Milkereit et al. 2001; Destefanis et al. 2022) (Fig. 1) reveals that all p53 isoforms were upregulated ( $P < 0.05$ ), with p53E showing the most robust increase ( $P < 0.01$ ), marking its first correlation with nucleolar stress and its role in stress-induced apoptosis. This truncated variant, characterized by a unique N-terminal domain, has been shown to counteract p53-induced apoptosis in eye imaginal discs (Zhang et al. 2015).

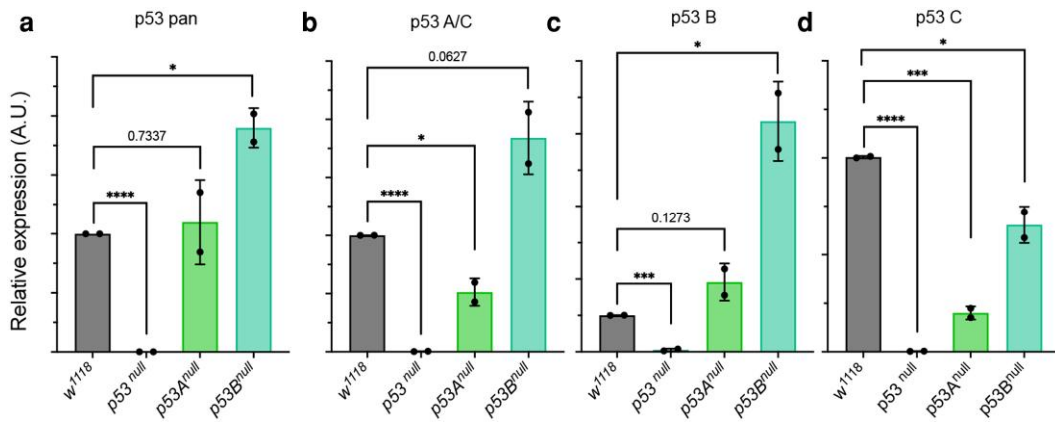
Indeed, the transcriptional repression activity of p53 isoforms is highly context-dependent. While p53A was shown to act as both an activator and a repressor in salivary glands, p53E functions only as a transrepressor and is dispensable for transactivation in irradiated embryos, where p53A was essential (Wylie et al. 2022). However, our study specifically addresses nucleolar stress, which is mechanistically distinct from stress induced by DNA damage. In wing imaginal discs subjected to Noc1 depletion, p53E may be selectively induced to engage stress-specific or compensatory mechanisms that modulate the nucleolar stress response in a manner that



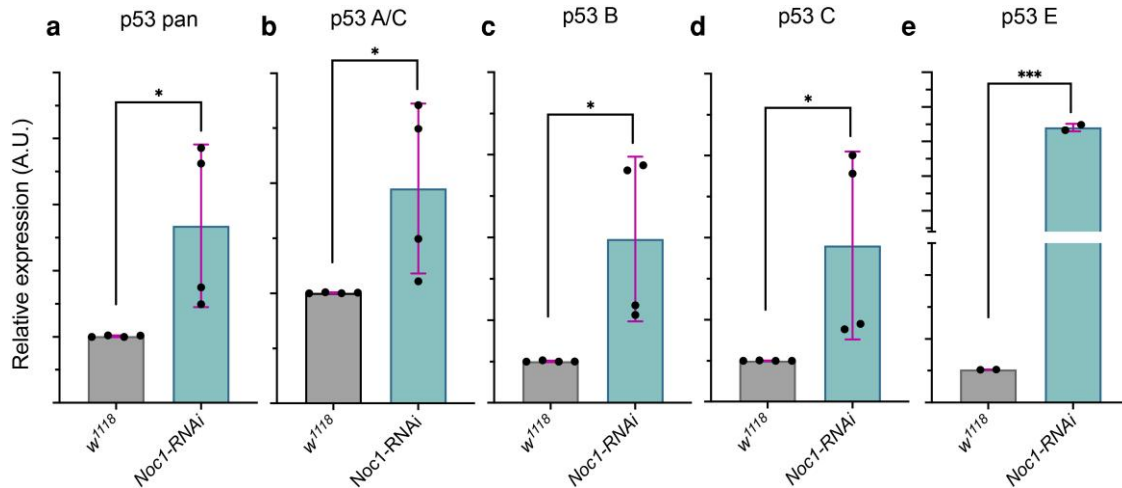
**Fig. 2.** Schematic representation of the p53 locus and primer design used to detect specific isoforms. a and b) Schematic representation of the p53 locus (gene ID 2768677) and of the p53 isoforms A, B, C, and E. (b-i), magnification of a common exon with the representation of the region where the primers that capture all p53 isoforms anneal (Zhang et al. 2015). These primers detect mRNA transcripts for all isoforms and were used in Fig. 1f; they will be referred to as p53 pan. c) Schematic representation of the p53<sup>A2.3</sup>, p53<sup>B41.5</sup>, and p53<sup>5A.1-4</sup> null mutant alleles. The p53<sup>A2.3</sup> null line was generated with a 23 bp deletion and a 7 bp insertion spanning from the end of the first exon until the first base of the first intron (in red), generating multiple early STOP codons (Supplementary Fig. 3a). This disrupts protein production from the p53A and p53C isoforms, but mRNA may still be detectable if primers anneal outside the deleted region. p53<sup>B41.5</sup> null line carries a 14 bp deletion and a 1 bp insertion in exon 2, resulting in a frameshift and premature stop codons leading to a truncated, non-functional p53B protein (Supplementary Fig. 3b)(Chakravarti et al. 2022). p53<sup>5A.1-4</sup> null line, contains a 3.3 kb deletion that removes the common 3' coding region (in red), eliminating protein expression from all isoforms. d) p53A isoform. (d-i and ii). Magnification of exons 1 and 2 of p53A and p53<sup>A2.3</sup> null isoform, respectively, showing where the primers designed in this study anneal (large arrows in green) compared to previous primers (in pink). As shown in (d-ii), the new primers can also be used to detect p53A mRNA in the p53<sup>A2.3</sup> null mutant, because the forward primer anneals outside the deleted region. In contrast, the previous reverse primer (small arrows in pink) may fail to detect the transcript in the p53A null line. The new primers also detect p53C mRNA and will therefore be referred to as p53A/C. e) p53B and p53C isoforms. (e-i, ii, and iii). Magnification of exons 2 and 3, shared by both isoforms, showing where the primers designed in this study anneal (in green). The new primers specifically detect p53B mRNA, whereas the old primers (pink) also detect p53C mRNA. Moreover, the new pair is also suitable for analyzing the p53B null line because the forward primer does not anneal in the deleted region marked in red (e-ii). This pair of primers will be referred to as p53B. f) p53C isoform. (f-i). Magnification of the region in exon 2 that includes the sequence of 78 bp, from 440 to 518 nucleotides from the beginning of the p53C transcript (in orange), unique to this isoform. Optimized primers (green) detect p53C mRNA and are referred to as p53C. g) p53E isoform. (g-i). Magnification of the region between exons, highlighting the amplified region specific to this isoform. h) Agarose gel (2%) showing qPCR-amplified products from RNA extracted from wing imaginal discs of *w<sup>1118</sup>* third-instar larvae, using the indicated primers and actin as a control. A single band is observed for each primer pair, demonstrating the specificity for mRNA detection.

differs across tissues and developmental stages. p53E could influence the relative abundance and composition of p53 oligomeric complexes, thereby fine-tuning p53 signaling output.

In this context, p53E may alter the balance of p53A-p53B-containing complexes. Depending on the cellular milieu, p53A can affect the formation of homo- or hetero-oligomers, ultimately



**Fig. 3.** Expression analysis of p53 mRNA isoforms by RT-qPCR in p53 mutant lines using isoform-specific primers. (a–d) Relative expression of p53 mRNA isoforms compared to *actin5C*, used as a control. The relative p53 isoforms analyzed with the specific primers are indicated at the top of each graph. The genotype of the lines is shown below. Relative expression of p53 mRNA is shown as arbitrary units (A.U.). p53 pan detects all p53 isoforms; p53A/C specifically detects p53A and the predicted p53C isoforms; p53B, p53D, and p53E detect only the respective isoform; *w<sup>1118</sup>* animals are used as control. The asterisks indicate the *P*-values obtained from Student's *t*-test analysis from two independent biological experiments. RNA was extracted from five whole larvae of the respective genotype for each experiment. \**P* < 0.05, \*\*\**P* < 0.001, and \*\*\*\**P* < 0.0001, and the error bars indicate the standard deviations.



**Fig. 4.** Upregulation of p53 isoforms in Noc1-downregulated cells. (a–e) qPCRs from RNA extracted from wing imaginal discs of *w<sup>1118</sup>* animals and expressing *Noc1-RNAi* in the wing pouch using the *rotund* promoter. Data are expressed as relative expression of p53 mRNA isoforms as arbitrary units (A.U.) compared to *actin5C*, used as a control. The asterisks represent the *P*-values obtained from Student's *t*-test analysis: \**P* < 0.05, and \*\*\**P* < 0.001. The error bars indicate the standard deviations. The dots represent the number of independent biological experiments. In each biological experiment, 30–40 wing imaginal discs were dissected per genotype and pooled for RNA extraction.

shaping transcriptional responses and cell fate decisions. Consistent with this model, shorter p53 isoforms have been shown to repress the activity of longer isoforms through heterotetramer formation, a conserved mechanism of p53 isoform regulation (Anbarasan and Bourdon 2019). Supporting isoform-specific regulation in vivo, Chakravarti et al. (Chakravarti et al. 2022) used isoform-specific mutants with the *hid-GFP* reporter in the germline, demonstrating differential transcription outputs among p53 isoforms. Together, these studies highlight the importance of heterotetramer formation among p53 isoforms (Zhang et al. 2014) and suggest a potential role for post-translational modifications (PTMs) in modulating isoform-specific functions. Further investigation will be required to determine how these regulatory mechanisms integrate under distinct cellular stress conditions, including nucleolar stress.

PTMs also contribute to isoform-specific regulation of p53. Indeed, ubiquitination, phosphorylation, sumoylation, and

acetylation may differ among p53 isoforms due to variations in their amino acid sequences. Lok/Chk2-dependent phosphorylation at serine 4 activates DNA repair and apoptotic responses (Peters et al. 2002; Brodsky et al. 2004). The E3 ligase dSyx5 acts as a negative regulator of p53 protein levels, while the role of the deacetylase dSir2 remains unclear (Bauer and Helfand 2009). Sumoylation at lysines 26 and 302 is essential for its full transcriptional and apoptotic activity (Mauri et al. 2008).

Moreover, our findings also highlight key mechanistic differences in p53 regulation between flies and humans. In vertebrates, nucleolar stress primarily activates p53 post-translationally through protein stabilization, mediated by ribosomal proteins that inhibit the E3 ubiquitin ligase MDM2 (Lindström et al. 2022). In contrast, *Drosophila* lacks a clear MDM2 homolog with E3 ligase activity, suggesting an alternative regulatory mechanism. The gene *companion of reaper* (*Corp*) has been proposed as a structural and partial functional analog of mammalian MDM2. Corp binds

to p53 and negatively regulates its levels through a feedback loop, similar to MDM2; however, it lacks an E3 ligase domain and thus cannot mediate canonical p53 degradation (Chakraborty et al. 2015). These distinctions point to a divergence in the evolution of p53 regulatory circuits, where transcriptional control may play a more prominent role in flies. This difference emphasizes the evolutionary divergence in p53 regulation and raises important questions about whether isoform-specific transcriptional control might also serve as a stress adaptation mechanism in vertebrates, including humans.

Further supporting the evolutionary conservation of nucleolar stress responses, our previous data show that downregulation of the human homolog of Noc1, called CEBPZ (Rambaldelli et al. 2025), also activates p53 both transcriptionally and at the protein level, consistent with our findings in *Drosophila* (Destefanis et al. 2022). Noc1 functions as part of a conserved nucleolar heterodimeric complex with Noc2 and Noc3, which is essential for pre-rRNA processing and 60S ribosomal subunit assembly, as demonstrated in both yeast and *Drosophila* (Milkereit et al. 2001; Destefanis et al. 2022). Our finding that Noc1 depletion triggers p53 upregulation in both flies and humans highlights the evolutionary conservation of nucleolar stress responses. This conserved activation highlights the broader relevance of nucleolar stress-induced p53 signaling and underscores the value of isoform-specific tools in elucidating the underlying regulatory mechanisms.

In conclusion, our analysis, enabled by isoform-specific primers and p53 mutant lines, reveals previously unrecognized residual activity and compensatory regulation, underscoring the intricate and multifaceted control of p53 in *Drosophila*. Our results indicate that p53E may participate in the apoptotic response triggered by nucleolar stress, suggesting it may have an isoform-specific role in this process. Furthermore, the observation that  $\gamma$ -H2AV phosphorylation reflects caspase-dependent apoptosis rather than direct genomic lesions indicates activation of a p53 stress response distinct from canonical genotoxic pathways. Collectively, these findings suggest that p53E is a component of a novel nucleolar stress-responsive mechanism that influences cellular adaptation to defects in ribosome biogenesis.

While our findings are consistent with potential transcriptional regulation of stress responses, further studies will be required to determine the relative contributions of transcriptional and post-translational mechanisms in maintaining cellular homeostasis.

## Data availability

All data are available in the published manuscript and its online Supplementary Files: [Supplementary Fig. 1](#), which reports the sequences of all the primers published for p53 isoforms, along with their corresponding recognition region in the p53 locus; [Supplementary Fig. 2](#), which includes the melting curve of the primer sets used; and [Supplementary Fig. 3](#). Sequences showing the mutation present in the p53<sup>A2,3</sup>, p53<sup>B41,5</sup> mutant lines.

Supplemental material available at [G3](#) online.

## Acknowledgments

We thank the Bloomington Stock Center (NIH P40OD018537).

## Funding

Department CIBIO Core Facilities is supported by the European Regional Development Fund (FESR) 2021–2027. The input from members of the COST Action CA21154 TRANSLACORE, supported

by COST (European Cooperation in Science and Technology and by the Dipartimento di Eccellenza 2023-2027, Legge 232/2016 project n 40613, funded by the MUR. This document includes language and clarity improvements made using ChatGPT (OpenAI).

## Conflicts of interest

None declared.

## Author contributions

AVC performed the experiments, conceived the study, and contributed to the writing of the manuscript. SB contributed to the final version of the manuscript. VM performed the experiments and analyzed the data. PB participated in the design, coordinated and supervised the team, and contributed to the writing and editing of the final manuscript. All authors have read and agreed to the published version of the manuscript.

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Editor: D. Bergstrahl