p53 directly represses human LINE1 transposons

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p53 is a potent tumor suppressor and commonly mutated in human cancers. Recently, we demonstrated that p53 genes act to restrict retrotransposons in germline tissues of flies and fish but whether this activity is conserved in somatic human cells is not known. Here we show that p53 constitutively restrains human LINE1s by cooperatively engaging sites in the 5'UTR and stimulating local deposition of repressive histone marks at these transposons. Consistent with this, the elimination of p53 or the removal of corresponding binding sites in LINE1s, prompted these retroelements to become hyperactive. Concurrently, p53 loss instigated chromosomal rearrangements linked to LINE sequences and also provoked inflammatory programs that were dependent on reverse transcriptase produced from LINE1s. Taken together, our observations establish that p53 continuously operates at the LINE1 promoter to restrict autonomous copies of these mobile elements in human cells. Our results further suggest that constitutive restriction of these retroelements may help to explain tumor suppression encoded by p53, since erupting LINE1s produced acute oncogenic threats when p53 was absent.

[**Keywords:** LINE1; p53; transrepression; tumor suppressors]

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p53, the most commonly mutated gene in human cancers, has been primarily studied in the context of cellular stress where it directs transcriptional programs specifying apoptosis, cell cycle arrest, and cellular senescence. Accordingly, stimulus-induced transactivation of effector genes is widely thought to account for p53-mediated tumor suppression (Lane and Levine 2010; Biegging et al. 2014; Chen 2016). However, mice collectively lacking essential p53 target genes (*p21*, *noxa*, and *puma*) remain tumor free and, likewise, p53 continues to function as a tumor suppressor even if decoupled from these cellular processes (Brady et al. 2011; Li et al. 2012; Valente et al. 2013). Recently, our laboratory demonstrated that p53 restrains mobile elements in the germline of flies and fish, raising the possibility that, if conserved, modalities related to this tonic activity might contribute to cancer prevention in longer-lived animals (Wylie et al. 2016a,b; Tiwari et al. 2017, 2018). Here, we tested whether retroelement repression by p53 is a conserved property that extends to human somatic cells.

LINE1 (L1) elements are considered the only active retrotransposon in the human genome (Lander et al. 2001). Although the majority of L1 copies are fragmented and/or mutated, ~100–150 full-length L1 elements are auto-

nous and retrotransposition competent (Moran et al. 1996; Brouha et al. 2003). These are all members of an evolutionarily young family of L1s designated L1 human specific (L1_{HS}). Intact L1_{HS} copies contain an internal promoter located within the 5'UTR (Swergold 1990; Alexandrova et al. 2012) and encode two open reading frames. L1 ORF1p is an RNA chaperone, while ORF2p possesses endonuclease and reverse transcriptase activities (Domroski et al. 1994; Feng et al. 1996; Martin et al. 2005; Martin 2010; Callahan et al. 2012). While L1s are normally silenced in somatic cells, derepression of these transposons is associated with certain cancers and some neuropathies (Muotri et al. 2010; Hancks and Kazazian 2012; Rodić et al. 2014; Wylie et al. 2016b; Burns 2017). Conceivably, L1 hyperactivity could drive pathologies by destabilizing genomes, promoting mutations, altering gene expression and triggering inflammatory responses (Hancks and Kazazian 2012, 2016; Lee et al. 2012; Tubio et al. 2014; Volkman and Stetson 2014; Chenais 2015; Kemp and Longworth 2015; Scott and Devine 2017; Briggs et al. 2018; Saleh et al. 2019; Rodriguez-Martin et al. 2020).

How p53 loss permits elevated retrotransposon activity is currently unknown, but several lines of evidence

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suggest that p53 could specify the deposition of epigenetic marks that silence mobile elements. For example, when a synthetic L1 was injected into Zebrafish, the L1-5'UTR was a target for p53-dependent H3K9 trimethylation (Wylie et al. 2016b) and consistent with this, in mouse fibroblasts treated with a DNA demethylating agent, intact p53 was needed to silence repetitive elements (Leonova et al. 2013). Furthermore, since informatic and empirical evidence for p53 binding to retroelements has been reported (Wang et al. 2007; Harris et al. 2009; Zemojtel et al. 2009; Cui et al. 2011), it is plausible that p53 could directly antagonize these elements.

Here, we establish a role for p53 in the silencing of human retroelements and link retroelement derepression caused by p53 loss to oncogenic phenotypes. Using synthetic transposons together with CRISPR methods, we found that L1 activity and de novo transposition were significantly increased when p53 was eliminated. We also observed that p53 binds to the 5'UTR of L1s and established that constitutive occupancy at this internal promoter was required for deposition of repressive histone marks and for transcriptional suppression of these elements. Soon after p53 loss, chromosomal rearrangements associated with L1 sequences were observed. Likewise, without p53, immune response signatures were also detected and, using a retrotransposon antagonist, we demon-

strate that retroelement activity underlies this response. Together, these findings establish that tonic repression of L1s is a well-conserved property of p53 that safeguards human somatic cells against potential oncogenic threats presented by these transposons.

Results

p53 suppresses L1 expression in human cell lines

Using CRISPR editing methods a series of p53 knockouts were generated in diverse human cell lines and derepressed L1 activity was observed in both cancer derived (e.g., A375 and U2OS cell lines) as well as normal lung tissue derived HBEC3kt cells (Fig. 1). As shown in Figure 1A (top panel), a common exon found in all *p53* isoforms was targeted and genomic sequencing was performed to assign loss-of-function genotypes in derived clones (see Supplemental Table S1). Using Western blot analyses, absence of p53 was verified in these independent knockout clones derived from A375 (Fig. 1C), U2OS (Fig. 1D), HBEC3kt (Fig. 1E, and HCT116 cells (Supplemental Fig. S1E). Furthermore, where available, loss of *p53* transcripts was verified in RNA sequencing data sets, consistent with nonsense-mediated decay in independent p53 knockout lines (Fig. 1A, bottom panel). To further verify the specificity of our *p53*

knockout lines, we computationally identified 114 potential *p53* gRNA off target loci, which differed from the *p53* target sequence by up to four mismatches (Supplemental Fig. S1M). Less than half of these loci are located in gene bodies (Supplemental Fig. S1N) and, importantly, we confirmed that expression from these was not substantially different in cells exposed to the *p53* gRNA and the parental A375 cell line (Supplemental Fig. S1O).

Next, the effects of *p53* loss on L1 activity were tested by inspecting these elements in two ways. First, we examined ORF1p, a commonly used marker for the expression of intact L1 copies. As shown in Figures 1C–E and quantified in Supplemental Figure S1G–I, *p53* elimination triggered acute increases in L1 ORF1p expression in A375, U2OS, and HBEC3kt cells. Importantly, neither cas9 exposure alone (Fig. 1C–E; Supplemental Fig. S1G–I) nor introduction of a control gRNA targeting the huAAVS1 locus (Supplemental Fig. S1C,D) was sufficient to trigger ORF1p expression, confirming that *p53* loss caused ORF1p derepression in these experiments. As a second approach, we directly inspected L1-derived RNAs using RNA sequencing and/or droplet digital PCR (ddPCR) assays. Using methods adapted from Philippe et al. (2016) and as exemplified in Figure 1B and summarized in Supplemental Figure S1A,B, loss of *p53* was associated with elevated expression of individually identifiable L1_{Hs} RNAs in RNA sequencing data sets from A375 cells. Intriguingly, this effect was stochastic, and consistent with (Philippe et al. 2016), different L1_{Hs} elements were clearly derepressed in different clonal *p53* knockout cell lines (in Supplemental Fig. S1A,B note red and yellow dots indicating commonly and uniquely derepressed L1_{Hs} elements). Furthermore, the effect triggered by *p53* loss was restricted to L1_{Hs} transcripts since ancient L1-, L2-, Alu-, and ERV-derived sequences were not similarly impacted (see Supplemental Fig. S1J–L). Most L1_{Hs} genomic sequences lack a complete 5' end and are, for this reason, retrotransposition-incompetent. Therefore, to more directly focus on intact L1_{Hs} copies, we also developed a ddPCR assay that selectively targets L1_{Hs} RNAs containing the 5'UTR. Using this assay, 5'UTR L1_{Hs}-derived RNAs were increased approximately fivefold when *p53* is lost in A375 cells (Fig. 1F). Similarly, a fourfold to fivefold increase was observed in U2OS and, likewise, approximately fourfold elevation in HBEC3kt cells upon *p53* loss (Fig. 1G,H, respectively). Taken together, these results indicate that *p53*-mediated repression of intact and potentially active retroelements seen in model systems (Wylie et al. 2016b; Tiwari et al. 2017) is conserved in humans and detectable in numerous cultured cell lines. This pattern was not universal since, consistent with previous reports (Coufal et al. 2011), L1s were not activated in HCT116 cells, indicating that additional factors and/or cell-type specific contexts can influence the behavior of L1s when *p53* is lost (Supplemental Fig. S1E,F).

p53 inhibits L1-retrotransposition

Results in Figure 1 demonstrate a strong inverse relationship between *p53* status and the expression of L1_{Hs} RNA

and protein. To determine whether retrotransposition rates are similarly affected, we stably transfected parental and *p53* knockout A375 and U2OS cells with a synthetic L1 designated 99-gfp-LRE3 (Ostertag et al. 2000; Kopera et al. 2016). This indicator (see Fig. 2A) encodes a retrotransposition competent copy of L1_{Hs} engineered to permit direct measurements of de novo retrotransposition rates. As illustrated in Figure 2A EGFP expression

Figure 2. *p53* loss is permissive for human L1 de novo retrotransposition. (A) Schematic depiction of the 99-gfp-LRE3 retrotransposition indicator. eGFP fluorescence indicates a complete retrotransposition cycle. Representative confocal images (obtained at 63 $\times$; scale bars, 20 μ m) of GFP fluorescence in stable wild-type, cas9-treated control, and *p53*^{-/-} A375 (B) and U2OS (D) 99-gfp-LRE3 integrant cell populations counterstained with DAPI. Quantification ($n=3$) of the percentage of GFP-positive cells by flow cytometry for A375 (C) and U2OS (E) cells, error bars represent the standard error of the mean. (*) P -value ≤ 0.05 , (**) P -value < 0.005 . Droplet digital PCR (ddPCR) quantification ($n=2$) of de novo LINE1 integration events in A375 (F) and U2OS (G) cells, normalized to single-copy gene *puma* abundance. Primer pair 2 (barbed arrows shown in A) specifically detects spliced GFP (indicating a complete retrotransposition life cycle) in genomic DNA. Bar graphs are averages of two biological replicates and error bars are the standard deviation.

produced by this synthetic element requires a completed retrotransposition life cycle, including transcription, splicing, reverse transcription, and integration. Using this tool, we observed significantly increased retrotransposition when p53 was removed (Fig. 2B–E) and, importantly, CRISPR/cas9 exposure alone did not trigger increased retrotransposition (Fig. 2B–G). When quantified by flow cytometry, *de novo* insertions were increased fivefold to sixfold in U2OS p53 knockout cells (Fig. 2E) and increased twofold to threefold in p53 knockout A375 cells (Fig. 2C) relative to parental counterparts or a control transposition indicator lacking the endonuclease, LRE3-H230A-GFP (EN⁻) (Supplemental Fig. S2C,D). To verify this conclusion, we directly inspected the genomes of wild-type and p53 knockout cells using a ddPCR assay that measures *de novo* retro-insertions. Importantly, starting copy numbers of the integrated 99-gfp-LRE3 plasmid were similar across genotypes, as verified by genomic PCR using primers that detect unspliced *eGFP* (Supplemental Fig. S2A,B). As seen in Figure 2, F and G, normalized values of spliced *eGFP* copies, indicating a complete cycle of retrotransposition, were considerably higher in the genomes of p53 mutant A375 and U2OS cells relative to their wild-type counterparts. Therefore, p53-dependent differences seen in Figure 2, B–E, reflect genuine differences in derived copies of 99-gfp-LRE3 retroinserts, establishing that human p53 restricts retrotransposition in these cells.

The L1-5'UTR is sufficient for p53 repression

Human L1 elements carry an internal RNA polymerase II promoter within their ~900-bp 5'UTR (Swergold 1990). Previously, we showed p53-dependent deposition of re-

pressive histone marks occurred at this 5'UTR when 99-gfp-LRE3 plasmids were injected into zebrafish embryos (Wylie et al. 2016b), suggesting that p53-dependent suppression of L1s may require the L1-5'UTR. To determine whether the 5'UTR is sufficient for p53 repression, we placed the L1-5'UTR upstream of an eGFP cassette to create an L1 expression reporter (L1-5'UTR-eGFP). Importantly, for these studies, we integrated the L1-5'UTR-eGFP expression reporter at the AAVS1 safe harbor site (Fig. 3A) and, in this way, avoided confounders associated with transient transfections, copy number differences, and position effects. We used genomic PCR to verify single-copy insertions of L1-5'UTR-eGFP at the AAVS1 safe harbor site in our edited A375 and U2OS cells (Supplemental Fig. S3A–C). Using flow cytometry, we monitored L1-5'UTR-eGFP activity and detected significantly higher eGFP expression levels in the p53 knockout A375 and U2OS cells compared with wild-type cells (Fig. 3B,C, Supplemental Fig. S3F,G; Supplemental Tables S2, S3 for the quantification). Likewise, *eGFP* transcript levels were similarly elevated in p53 knockout A375 and U2OS cells (Supplemental Fig. S3D,E), suggesting that p53 repression occurs at the level of transcription. Together, Figure 3, B and C, and Supplemental Figure S3, D and E indicate that p53 antagonizes the production of L1 RNAs by operating through DNA sequences within the 5'UTR of L1 elements.

H3K9me3 and H3K27me3 are repressive histone marks characteristic of heterochromatin (Bannister and Kouzarides 2011). These marks are also enriched at repetitive sequences including LINE1 elements (Molaro and Malik 2016; Yang and Wang 2016), but efforts to inspect individual L1 copies are confounded by the absence of uniquely

Figure 3. p53-mediated suppression of LINE1s operates at the level of transcription through the 5'UTR. (A) The integrated L1 reporter construct, L1-5'UTR-eGFP, is detectable with primers indicated (barbed arrows). Flow cytometry of GFP in WT, cas9-treated control, and p53^{-/-} A375 (B) or U2OS (C) cells containing the L1-5'UTR-eGFP exposes bright GFP-positive subpopulations in p53^{-/-} cells (see Supplemental Tables S2, S3 for gating parameters). Additional biological replicates are shown in Supplemental Tables S2 and S3; for unstained controls, see Supplemental Figure S3, F and G. ChIP results (D,E) showing H3 normalized H3K27me3 (D) and H3K9me3 (E) ratios at the L1-5'UTR, *p21*, and *puma* in wild-type (black) and p53^{-/-} (red) cells. Note that changes in H3K27me3 and H3K9me3 enrichment are specific to the 5' UTR of the L1-5'UTR-eGFP and p53 dependent. Error bars represent 95% confidence intervals.

assignable sequences. Therefore, to ask whether p53 might regulate deposition of these marks, we took advantage of the integrated L1-5'UTR-eGFP reporters with primers that uniquely detect this template (Fig. 3A) and, using these, we conducted ChIP experiments to interrogate internal L1 promoter sequences. As seen in Figures 3D and E, when p53 was absent H3K27 trimethylation was reduced by ~70% and, similarly, H3K9 trimethylation of L1-5'UTR-eGFP was reduced by 50% across the same region. Importantly, these p53-dependent patterns were specific to L1s, since we did not observe differential histone methylation or expression at the canonical p53 target genes *p21* and *puma* (Fig. 3D,E; Supplemental Fig. S3H,I).

p53 directly binds and represses L1-5'UTR

To determine how p53 regulates the deposition of repressive histone marks at the L1-5'UTR, we first assessed the impacts of p53 on known retroelement repressive pathways. Importantly, expression levels of epigenetic modifiers (Supplemental Fig. S4A,C), small RNA effectors (Supplemental Fig. S4B,D), and piRNA precursors (Supplemental Fig. S4E,F) known to repress transposons were not changed in p53 knockout cell lines. Indeed, the piRNA pathway component *piwil4* was modestly induced in p53 knockout cells, consistent with a potential adaptive response to hyperactive L1s in these cells. These results indicate that transcriptional regulation of target genes

coding for known effectors does not underlie p53 mediated L1 silencing, suggesting that p53 may directly bind and repress these retrotransposons.

The L1 5'UTR contains three predicted p53 binding sites (Smeenk et al. 2008; Harris et al. 2009; Heinz et al. 2010), which we designated as sites I–III (see Fig. 4A). Using in vitro assays, Harris et al. (2009) demonstrated that purified p53 protein bound a synthesized stretch of the L1 5'UTR (bases 427–523) containing both sites I and II. Site III was informatically predicted (see the Materials and Methods). To inspect potential p53 binding to the L1 5'UTR, we conducted ChIP assays in A375 cells containing the uniquely identifiable L1-5'UTR-eGFP reporter. For assay validation, we confirmed robust p53 binding signals at benchmark target genes (e.g., *puma* and *p21*) that were present in wild-type samples but absent from p53 knockout cells (Supplemental Fig. S4G,H). As seen in Figure 4B, in these same lysates we detected significant enrichment of p53 signal at the L1 5'UTR (Harris et al. 2009). For these assays, we used primer pairs specific for the L1 expression reporter in the L1 5'UTR and a negative control region at the eGFP 3' end (see Fig. 4B). Importantly, signals detected in the 5'UTR in wild-type cells were absent from p53^{-/-} cells, while signal detected at the negative control was similar in both cell types. To identify specific sequences needed for p53 binding, we produced a series of L1-5'UTR-eGFP variants that separately eliminated sites I, II, or III (as illustrated in Fig. 4A). Furthermore, like the parental reporters,

Figure 4. p53 physically binds to repress the L1-5'UTR. The schematic in A shows empirically and computationally defined p53 binding sites in the 5'UTR of L1-5'UTR-eGFP. Note that site I, site II, and site III were independently targeted for deletion. (B) ChIP for p53 and control IgG performed with illustrated primer pairs (1 and 2) that uniquely detect binding at the L1 reporter. Note the absence of signal in p53^{-/-} cells and at the L1-GFP negative control region spanning vector-derived sequences (Ctrl primer pair). Error bars represent 95% confidence intervals. (C) ChIP testing of p53-binding site deletions indicated in A along with the intact wild-type 5'UTR interval. Note, site I and site II contribute to p53 binding, but site III had no effect. Error bars represent 95% confidence intervals. In D, flow cytometry was used to detect for GFP intensities in the intact and mutant L1-5'UTR-eGFP reporters in A375 cells. Note that sites I and site II are required for p53 repression of the L1 expression reporter (see Supplemental Fig. S4L for another biological replicate). (E) Shows that corresponding eGFP mRNA transcript levels mirror GFP intensities seen in these same cells. Bar graphs are averages of three biological replicates. Error bars represent standard error of the mean.

these were similarly targeted to the identical AAVS1 site and confirmed to a single copy of the L1-5'UTR-eGFP reporter (Supplemental Figs. S3A, S4J). As seen in Figure 4C, deletion of sites I and II led to the complete loss of p53 enrichment at the L1-5'UTR, while deletion of site III had no effect on p53 levels. p53 enrichment at the negative control eGFP 3' end was similar across all cell lines (Supplemental Fig. S4I).

To determine the functional relevance of p53 binding to the 5'UTR we inspected L1-5'UTR-eGFP expression in these same cells, using flow cytometry. As seen in Figure 4D and Supplemental Figure S4L, site I or site II deletions in the L1 5'UTR produced eGFP peaks in wild-type cells that were similar to those observed in p53^{-/-} cells. Deletion of site III, in contrast, showed no such effect, producing eGFP peaks comparable with those observed in wild-type cells expressing the intact reporter (Fig. 4D; Supplemental Fig. S4L). Similarly, deletion of sites I and II led to increased eGFP RNA (Fig. 4E) and protein (Supplemental Fig. S4K), while deletion of site III had no effect on p53 repression. Together, these data expose *cis* sequences within the 5'UTR of L1 elements that mediate binding to p53 as well as transcriptional repression by p53.

De novo rearrangements in p53^{-/-} cells are associated with retroelements

Genomic instability is a hallmark of cancer, and is commonly associated with loss or mutation of p53 (for review, see Eischen 2016). Using RNA sequencing data sets, we

assessed the frequency of gene fusion transcripts, which may be indicative of genomic rearrangements, in A375 wild-type and p53 knockout cell lines. Importantly, early passage p53 knockout cells (<20 doublings) were used for these experiments, enabling us to ask whether genomic instability indicative of p53 mutant human cancers is rapidly induced by the loss of p53. We identified nine fusion transcripts in p53 knockout A375 cell lines that were absent in the parental strain, together with 13 fusion transcripts present in both parental and derived cell lines (Fig. 5A). Both intrachromosomal ($n = 5$) and interchromosomal ($n = 4$) fusions were observed (Supplemental Table S4) indicating that genomic stability is rapidly compromised following loss of p53.

To ask whether genomic rearrangements observed in p53 knockout cells were associated with retroelement activity, we inspected rearrangement junctions for the presence of repeat elements. Half ($n = 9$ of 18) of the *de novo* rearrangement breakpoints identified in A375 p53 knockout cells spanned an annotated retroelement, yet none of the pre-existing rearrangements were associated with retroelement sequences (Fig. 5B). Intriguingly, virtually all ($n = 8$ of 9) of these were SINEs or LINEs (Fig. 5C; Supplemental Fig. S5A), suggesting that these elements may have promoted the *de novo* rearrangements seen after p53 loss. Importantly, these analyses also showed that transposable element-initiated transcripts were not elevated in p53 knockout cell lines (Supplemental Fig. S5B), indicating that p53 does not globally impact transcriptional properties or chromatin features.

Figure 5. *De novo* rearrangements detected after p53 loss are associated with retroelements. (A) Circos plot of genomic rearrangements in A375 cells. Genomic rearrangements emerging after p53 loss are indicated (red lines) along with pre-existing rearrangements in the parental A375 cell line (gray lines). Repeat element proximity is indicated by red tiles (outer gray ring) along with annotations for rearrangement junctions (inner gray ring). (B) Genomic rearrangements occur at repeat elements in p53 knockout ($n = 9$ of 18 breakpoints) but not wild-type ($n = 0$ of 26 breakpoints) cells. (C) p53 knockout associated genomic rearrangements are commonly associated with LINE and SINE ($n = 8$ of 18 breakpoints), but not LTR or DNA repeat elements ($n = 1$ of 18 breakpoints).

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Figure 7. A LINE1 antagonist, 3TC, prevents the inflammatory program triggered by p53 loss (A) Schematic of the experimental workflow. Expression of selected immunity (B) and inflammation response (C) genes was quantified using RT-ddPCR. Likewise, targets with overlapping immune pathway designations (D) and 5' intact L1_{Hs} (E) were similarly analyzed by RT-ddPCR. Wild-type and two p53 knock-out lines treated in parallel with vehicle, DMSO, only (filled bars) or 3TC (open bars). Transcript levels were normalized to β -actin and their fold change was calculated relative to their wild-type values (see Supplemental Fig. S6). For B–D, bar graphs plot the average of three biological replicates and error bars represent SEM. For Figure E, bar graphs represent the average of three biological replicates and error bars indicate the standard deviation. Two tailed *t*-tests were performed for all samples. (*) *P*-value < 0.05, (**) *P*-value < 0.005, (***) *P*-value < 0.0005, (ns) not significant. Note that treatment with the reverse transcriptase inhibitor, 3TC, prevented induction of immune and inflammatory gene sets but did not impact L1_{Hs} RNAs.

Accordingly, we further showed that loss of this repressive activity was permissive for L1 eruptions that triggered destabilized genomes and inflammatory responses commonly seen in p53 mutant cancers (Gudkov et al. 2011; Eischen 2016). Using CRISPR methods we generated a series of p53 knockouts in diverse human cell lines and observed generally derepressed L1 activity in both cancer derived (e.g. A375 and U2OS cell lines) as well as normal lung tissue derived HBEC3kt cells (Fig. 1). In p53 deficient cells, elevated L1_{Hs} derived RNAs were also tightly correlated with elevated ORF1 protein and de novo retroinserts (Fig. 2). This effect impacted native and synthetic L1 variants that were integrated and engineered for unique detection. Notably, derepression of individually identifiable L1 retroelements was stochastic and varied between p53 mutant cell lines (Supplemental Fig. S1A,B). Therefore, while p53 loss is a condition that is permissive for L1 eruptions, unknown inciting factors also govern individual elements in this context (Wylie et al. 2016b). Conceivably, p53 restriction could target any step in the L1 retrotransposon lifecycle, but several lines of evidence indicate that suppression oc-

curs at the level of transcription. First, acute induction of these transcripts occurred when p53 was removed (Fig. 1) yet known retroelement silencers were not affected (Supplemental Fig. S4A–F). Second, the internal L1_{Hs} promoter region conferred p53 regulation when fused to reporter templates (Fig. 3B,C). Third, the deposition of repressive histone marks at the internal L1_{Hs} promoter region was clearly p53 dependent (Fig. 3D,E). Together, these results strongly indicate that p53 prevents L1 expression by preventing L1 transcription.

To test the possibility of direct transrepression by p53 at the internal L1 promoter region, we functionally tested three distinct p53 binding sites by independently eliminating each. Two sites, (sites I and II) were clearly essential for binding and repression while a third site (site III) was dispensable for both (Fig. 4C). Surprisingly, loss of either site I or site II completely disrupted binding and completely abolished repression, suggesting that sites I and II are concurrently needed (Fig. 4D,E; Supplemental Fig. S4K, L). This “all-or-none” property suggests that cooperative p53 binding could mediate direct transrepression at the

internal L1 promoter. Interestingly, Harris et al. (2009) inspected these same p53 sites and found that site I, but not site II, was needed for stimulus-dependent transcriptional induction. Therefore, a single site can produce opposing outputs and it is likely that additional unknown factors specify the direction of activity *in vivo*. Several technical considerations could also account for our departures from Harris et al. (2009). For example, Harris et al. (2009) used transient cotransfections to follow episomal reporters as p53 was overexpressed, yet here we tracked integrated reporters after p53 was removed. Second, the reporters in Harris et al. (2009) lack ~100 bp of the L1 promoter region we used here. Third, Harris et al. (2009) conducted their assays in HCT116 cells, which, for unknown reasons, was the only line of four tested here that did not show L1 derepression after p53 loss. Despite these differences, p53 clearly engages the internal promoter of these transposons and, accordingly, future studies may identify instructive mechanisms that specify inductive versus repressive outcomes.

p53 is commonly studied as a stress-responsive transactivator (Lane and Levine 2010; Chen 2016) and, within this framework, down-regulation of target genes is widely considered to be a secondary effect (for review, see Sullivan et al. 2018). In certain stimulus-dependent contexts, the occlusion of positive transcription factors has been suggested to account for p53 suppression (Bargonetti et al. 1997; Ori et al. 1998; St Clair et al. 2004) but, to our knowledge, the findings presented here constitute the first example of direct and constitutive transrepression mediated by p53 in unstressed cells. Indeed, our data showing that p53 binds and recruits repressive histone marks to LINE1s but not to canonical transactivation targets such as *p21* and *puma* (Fig. 3D,E; Supplemental Figs. S3H,I, S4G,H) are inconsistent with a simple occlusion model of p53 gene repression. Further work is needed to establish precise mechanisms and explore perhaps whether regulatory RNAs and/or physical interactions with epigenetic modifiers are involved. Since direct tonic repression accounts for p53-mediated suppression of LINE1s, and since these transposons appear to constitute drivers that remodel cancer genomes and incite immune responses (Lee et al. 2012; Tubio et al. 2014; Rodriguez-Martin et al. 2020), these observations could have important implications for understanding p53 tumor suppression.

Consistent with this hypothesis, studies in Figures 5–7 illustrate how derepressed retrotransposons present acute oncogenic threats that are typically associated with p53 loss. The prevalence of homologous recombination seen in Figure 5 is one compelling example since, unlike pre-existing alterations, the chromosomal rearrangements occurring after p53 removal were highly associated with retroelements at their junctions (Fig. 5). Furthermore, these *de novo* rearrangements occurred within 20 cell doublings after CRISPR editing, raising the possibility that p53 continually acts to suppress homologous recombination between distal retrotransposon sequences. Consistent with this possibility, chromatin features are indeed important determinants of recombination frequency (Stapley et al. 2017) and, as seen in Figure 3, D and E, we showed that

p53 acts to maintain repressive chromatin marks at LINE1s. Hence, facilitated recombination mediated by derepressed retroelements could represent important early events that destabilize genomes after p53 loss.

Other oncogenic threats posed by L1s relate to inflammatory responses (Leonova et al. 2013; Brégnard et al. 2016; Saleh et al. 2019) and two lines of evidence here show that hyperactive L1s elicit acute inflammatory transcriptional programs in the absence of p53. First, consistent with previous reports (Gudkov et al. 2011) retroelement derepression and immune gene induction both occurred rapidly and simultaneously after p53 loss (Fig. 6). Second, treatment with 3TC definitively prevented the induction of nearly all inflammatory genes (Fig. 7). Since 3TC is a potent inhibitor of the L1 reverse transcriptase (Jones et al. 2008; Dai et al. 2011; Banuelos-Sanchez et al. 2019), these results indicate that L1s can drive inflammatory genes when p53 is abolished. Furthermore, these results also suggest that the trigger eliciting immune signatures lies downstream from the LINE1 enzyme and could possibly involve RNA–DNA hybrids and/or the integration process itself. Taken together, these studies establish direct transrepression of human L1 transposons as a constitutive p53 function that, if compromised, promotes oncogenic features seen in p53 driven cancers.

Materials and methods

Generation of p53 knockout cells using CRISPR Cas9

CRISPR cas9 editing (pX458) was used to generate p53 knockout cell lines. p53 knockout clones were identified by Western blot using p53 monoclonal DO-1 (sc-126) and Sanger sequencing was performed to determine the nature of each p53 knockout allele. L1-ORF1p expression levels were examined using anti-L1-ORF1p antibodies. Fiji software (Schindelin et al. 2012) was used to measure the band intensities of L1ORF1p and β -actin from Western blot experiments.

Generation of huAAVS1 targeted L1-5'UTR-EGFP reporter cells

The WT L1 5'UTR and 5'UTR sites I, II, and III deletion mutants were cloned at the FseI and AfeI sites of the huAAVS1-egfp-Hyg donor plasmid (kind gift from Joshua Mendel laboratory, University of Texas Southwestern). Cells were transfected with huAAVS1-TALEN1 (Addgene 59025), huAAVS1-TALEN2 (Addgene 59026), and donor plasmids. Three days after transfection, GFP positive single cells were sorted into 96-well plates by FACS. Once confluent, targeted knock-in was verified using PCR primers indicated in Supplemental Table S5. Detailed methods are in the Supplemental Material.

FACS analysis

All FACS analyses were performed on an LSR Fortessa SCC using equal numbers of cells. Live–dead gating was performed based on forward scatter versus side scatter profile. Data analysis and percent GFP-positive cell counts were performed in FloJo. When analyzing GFP signals, untransfected (see Supplemental Fig. S3F,G) cells were used for the baseline gating to identify EGF-positive cells. EGFP fluorescence intensity of WT-L15'UTR-EGFP cells was used to create a second gating to identify bright GFP-positive

cells in p53^{-/-} L15'UTR-EGFP samples (see Supplemental Tables S2, S3).

3TC experiment

WT and p53 knockout A375 cells were treated with DMSO or 10 μ M 3TC for 72 h.

RNA expression

Total RNA was extracted with TRIzol (Invitrogen), followed by DNase treatment using Turbo DNase (Ambion). Droplet digital PCR (ddPCR) was performed using Bio-Rad's EvaGreen system. Gene expression was normalized to β -actin expression, and fold induction was calculated relative to the wild type. See Supplemental Table S5 for the primer sequences used in ddPCR experiments.

Retrotransposition assay

Cells (2×10^5) were transfected with equal amounts of 99-GFP-LRE3 plasmid, as described previously (Ostertag et al. 2000). After 72 h, cells were selected for puromycin resistance. Flow cytometry was used to determine the percentage of EGFP-positive cells 9–10 d after transfection. For GFP integration assays, 2 μ g of genomic DNA from each sample was digested for 1 h with EcoRI (NEB) followed by ddPCR using primers detecting unspliced GFP as indicated in Supplemental Table S5. To calculate relative retrotransposition rate, ddPCR was performed on genomic DNA using primers detecting spliced EGFP (see Supplemental Table S5) and normalized against the single-copy gene *puma*. Detailed methods are in the Supplemental Material.

Statistical analysis

All quantitative data were collected from experiments performed in two to three biological replicates. Two-tailed Student's *t*-tests were performed to identify statistically significant difference (cutoff $P < 0.05$).

RNA sequencing and data analysis

RNA was extracted from A375 wild-type or A375 p53^{-/-} cells using Trizol (Invitrogen) followed by DNase treatment using TurboDNase (Ambion). Samples with RNA integrity scores >9.0 , was used for sequencing. Sequencing libraries were prepared following the manufacturer's instructions (NEB E7750S with poly (A enrichment module NEB E7490S) and sequenced on an Illumina NextSeq to an average sequencing depth of 40 million 150-bp paired-end reads per sample. Reads were aligned to the hg38 assembly of the human genome with STAR (Dobin et al. 2013). HTseq counts tables were used as input to perform normalization and differential gene expression analysis with Deseq2 (Love et al. 2014; Anders et al. 2015). STAR-Fusion was used to identify potential gene fusion products (Haas et al. 2017). Gene ontology annotations were performed using the DAVID online resource (Huang et al. 2009a,b). GSEA analyses were performed using the h.all.v7.0.symbols.gmt gene set (Subramanian et al. 2005). Retroelement expression levels were determined using a customized pipeline adapted from (Wylie et al. 2016b). LIONS analytical pipeline was used to measure transposon initiated transcription (Babaian et al. 2019). RNA sequencing data sets were deposited with the Gene Expression Omnibus (GEO) database under accession number GSE159134. Detailed methods are in the Supplemental Material.

p53-binding site prediction

Putative binding sites within the L1 5'UTR were bioinformatically identified using the p53scan program (Smeenk et al. 2008) and Homer's scanMotifGenomeWide module (Heinz et al. 2010). Individual p53 motif files for right and left p53 consensus decamers were used for motif detection.

ChIP ddPCR

Chromatin immunoprecipitation protocol was adapted from Carey et al. (2009). Briefly, A375 cells were fixed with 1% formaldehyde, quenched with glycine, and nuclear lysates were prepared. DNA was sheared to an average of 300–700 bp with a Covaris Sonicator (5 min, 200 cpb, intensity 5). Sonicated lysate was diluted 1:10, aliquoted, and incubated overnight with antibody. Antibody-bound chromatin was precipitated with 25 μ L of protein A/G beads (EMD Millipore IP0515ML). Eluted chromatin was treated with RNase and proteinase K, cross-links were reversed, and samples were purified by standard phenol:chloroform extraction and ethanol-precipitated. ChIP enrichment was quantified by ddPCR with EvaGreen (Bio-Rad 1864034) on a QX200 droplet reader (Robin et al. 2014; Wylie et al. 2016b). Antibodies used in these experiments included p53 ChIP (Santa Cruz Biotechnology sc-126), IgG (Diagenode C15400001), H3 (Abcam ab1791), H3K27m3 (Active Motif 39155), and H3K9m3 (Diagenode C15410058). Detailed materials and methods are in the Supplemental Material.

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p53 directly represses human LINE1 transposons

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