**Conserved Regulation of p53 Network Dosage by MicroRNA–125b Occurs through Evolving miRNA–Target Gene Pairs**

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Conserved Regulation of p53 Network Dosage by MicroRNA–125b Occurs through Evolving miRNA–Target Gene Pairs

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Abstract

MicroRNAs regulate networks of genes to orchestrate cellular functions. MiR-125b, the vertebrate homologue of the Caenorhabditis elegans microRNA lin-4, has been implicated in the regulation of neural and hematopoietic stem cell homeostasis, analogous to how lin-4 regulates stem cells in C. elegans. Depending on the cell context, miR-125b has been proposed to regulate both apoptosis and proliferation. Because the p53 network is a central regulator of both apoptosis and proliferation, the dual roles of miR-125b raise the question of what genes in the p53 network might be regulated by miR-125b. By using a gain- and loss-of-function screen for miR-125b targets in humans, mice, and zebrafish and by validating these targets with the luciferase assay and a novel miRNA pull-down assay, we demonstrate that miR-125b directly represses 20 novel targets in the p53 network. These targets include both apoptosis regulators like Bak1, Igfbp3, Itch, Puma, Prkra, Tps3inp1, Tp53, Zac1, and also cell-cycle regulators like cyclin C, Cdc25c, Cdkn2c, Edn1, Ppp1ca, Sef11, in the p53 network. We found that, although each miRNA–target pair was seldom conserved, miR-125b regulation of the p53 pathway is conserved at the network level. Our results lead us to propose that miR-125b buffers and fine-tunes p53 network activity by regulating the dose of both proliferative and apoptotic regulators, with implications for tissue stem cell homeostasis and oncogenesis.

Introduction

MicroRNAs (miRNAs) are short non-coding RNA molecules that were first discovered as regulators of developmental timing, and later found to regulate complex networks of genes to orchestrate cellular functions. Lin-4 was the first miRNA gene to be discovered, and shown to regulate developmental timing by repressing its target genes at the post-transcriptional level [1]. Subsequently, miRNAs were found to regulate processes ranging from proliferation and apoptosis, to cell differentiation and signal transduction [2–4]. Several miRNAs are conserved in metazoan evolution, one prominent example being lin-4 whose vertebrate homologues comprise the miR-125a/b family [5]. Much like lin-4’s role of regulating the homeostasis of reiterative or self-renewing stem cells in C. elegans [6], recent studies have shown that miR-125a/b regulates mammalian neural stem cell commitment, as well as the mammalian hematopoietic stem cell (HSC) pool size [7–10]. Although Lin28 and Bak1 have been proposed as the critical targets of miR-125a/b for regulating these stem cell compartments [8,9], the hundreds of predicted targets for miR-125a/b suggest a more complex interplay between miR-125a/b and its targets in regulating proliferation and differentiation.

Depending on the cell context, miR-125b has been proposed to regulate both apoptosis and proliferation. miR-125b has been shown to downregulate apoptosis in many contexts, in some cases by repressing Tp53 and Bak1. Examples include mammalian hematopoietic stem cells, human leukemia cells, neuroblastoma cells, breast cancer and prostate cancer cells [9–18]. During zebrafish embryogenesis, loss of miR-125b leads to widespread apoptosis in a p53-dependent manner, causing severe defects in neurogenesis and somitogenesis [16]. On the other hand, miR-
Author Summary

MicroRNAs (miRNAs) are tiny endogenous RNAs that can regulate the expression of hundreds of genes simultaneously, thus orchestrating changes in gene networks and mediating cellular functions in both plants and animals. Although the identification of individual targets of miRNAs is of major importance, to date few studies have sought to uncover miRNA targets at the gene network level and general principles of miRNA regulation at the network level. Here we describe how miR-125b targets 20 apoptosis and proliferation genes in the p53 network. We found that, although each miRNA-target pair evolves rapidly across vertebrates, regulation of the p53 pathway by miR-125b is conserved at the network level. The structure of the miR-125b regulatory network suggests that miR-125b buffers and fine-tunes p53 network activity. This buffering feature of miR-125b has implications for our understanding of how miR-125b regulates oncogenesis and tissue stem cell homeostasis. We believe these findings on miR-125b support a new fundamental principle for how miRNAs regulate gene networks in general.

miR-125b can also downregulate proliferation in a variety of human cancer cell-lines [19–23] and one of its bona fide targets Lin28, also promotes cancer cell proliferation [24]. Therefore in different contexts, miR-125b appears to be able to regulate both apoptosis and proliferation.

Another molecular pathway that regulates both apoptosis and proliferation is the highly conserved p53 network [25–28]. Due to the central role of the p53 network in these two processes, and because we found that miR-125b regulates both human and zebrafish Tp53 but not mouse Tp53 [16], we sought to examine if miR-125b regulates the p53 network in a conserved manner in vertebrates. To address this question, we used a gain- and loss-of-function screen for miR-125b targets in different vertebrates, and validated these targets with the luciferase assay and a novel miRNA-target pull-down assay. We demonstrate that miR-125b directly represses 20 novel targets in the p53 network, including both apoptosis regulators like Bak1, Ighp3, Ich, Pana, Prka, Tp53inp1, Tp53, Zac1, and also cell-cycle regulators like cyclin C, Cdc25c, Cda2a, Edn1, Ppp1ca, Sct11. We found that although individual miRNA-target pairs were seldom conserved, regulation of the p53 network by miR-125b appears to be conserved at the network-level. This led us to propose that miR-125b buffers and fine-tunes p53 network dosage, with implications for the role of miR-125b in tissue stem cell homeostasis and oncogenesis.

Results

Identifying direct targets of miR-125b in the p53 network

To systematically identify direct targets of miR-125b in the p53 network of vertebrates, we first employed a bioinformatics approach by identifying all predicted miR-125b targets in the p53 network, followed by three complementary methods to screen and validate these targets for both direct binding and repression by miR-125b (Figure 1). Existing databases and prediction algorithms were used to shortlist a set of p53 network genes predicted to possess miR-125b-binding sites in their 3’ UTRs. We analyzed the Ingenuity Pathways Analysis™ (IPA) database and the p53 Knowledgebase [29,30] for a list of genes and proteins that participate in the p53 network, either by regulating p53 upstream, by direct interaction with p53 protein, or by serving as effectors of p53 function downstream. We then analyzed the TargetScan and MicroCosm Target databases [31,32] for genes that are predicted to possess miR-125b-binding sites in their 3’ UTRs, in three vertebrate genomes: human, mouse and zebrafish. The genes at the intersection of the predicted miR-125b target list and the list of p53 network genes constituted our list of predicted miR-125b targets in the p53 network (Table S1).

miR-125b gain- and loss-of-function screen in 3 vertebrates

Next we sought to screen our list of predicted targets for significant repression by miR-125b in cells, by performing a miR-125b gain- and loss-of-function screen. Gain-of-function (GOF) in miR-125b was achieved by transfection of miR-125b duplex into human SH-SY5Y or mouse N2A neuroblastoma cells, whereas loss-of-function (LOF) in miR-125b was achieved in human primary lung fibroblasts or mouse 3T3 fibroblasts by knocking down miR-125b with an antisense (AS) RNA (Figure 2A). We chose to perform a gain-of-function screen in human (SH-SY5Y) or mouse (N2A) neuroblastoma cells, because these cells possess low
levels of endogenous miR-125b (Figure S1A, S1B). For the loss-of-function screen, we chose human fetal lung (hLF) or mouse (3T3) fibroblasts because they possess high levels of miR-125b (Figure S1C, S1D). miR-125a/AS was co-transfected with miR-125b-AS to achieve a complete silencing of the miR-125a/ family, because miR-125a, which shares the same seed sequence and the same predicted targets as miR-125b, is also highly expressed in human and mouse fibroblasts (Figure S1C, S1D). Genes that were either significantly repressed by miR-125b or significantly derepressed by miR-125a/AS with fold-changes within the range of microRNA regulation (P<0.05, fold change > 1.3), were selected as candidate miR-125b targets (Figure 2B-2D). For zebrafish embryos, which possess high levels of miR-125b, the loss-of-function (LOF) screen was performed using an antisense morpholino cocktail that blocks the loop regions of all 3 pre-miR-125b hairpin precursors [16]. The gain-of-function (GOF) screen was performed by co-injecting miR-125b duplex with the morpholino (Figure 2A). All gene expression changes were measured with at least three biological replicates using qRT-PCR.

Our GO/F/LOF screen revealed that in humans, out of 29 predicted targets in the p53 network, 13 genes were derepressed by miR-125b/AS in hLF cells and 20 genes were repressed by miR-125b in SH-SY5Y cells (Figure 2B). In mice, out of 22 predicted targets in the p53 network, 11 genes were derepressed by miR-125b/AS in 3T3 cells and 12 genes were repressed by miR-125b in N2A cells (Figure 2C). In zebrafish embryos, out of 20 predicted targets in the p53 network, 13 genes were derepressed by pre-miR-125b morpholino and 12 genes were repressed by the injection of miR-125b duplex (Figure 2D). In total, 22 human genes, 13 mouse genes and 14 zebrafish genes passed the gain- and loss-of-function qRT-PCR screen.

Direct binding interactions between miR-125b and mRNA targets from the p53 network

To assess which candidate miR-125b targets identified in the gain- and loss-of-function qRT-PCR screen are directly bound by miR-125b in cells, we employed a novel mRNA pull-down method developed by Lal et al. (manuscript in preparation). RNA transcripts bound to biotinylated-miR-125b were pulled down with streptavidin beads and quantified by qRT-PCR relative to mRNA bound to biotinylated-control miRNA (log2 fold change > 0.3, P<0.05). In this assay, biotinylated miRNAs were shown to be loaded into the RNA-induced silencing complex (RISC) and fully functional in repressing their target mRNAs (Lal et al., manuscript in preparation). This method provides a robust and complementary method for detecting miRNAs bound to endogenous target mRNAs, and serves as a useful approach for distinguishing direct and indirect targets in the same pathway (Lal et al., manuscript in preparation). Quantification of the pulled down mRNA targets in hLF cells revealed that 13 out of 22 gene transcripts, Bak1, Cdc25c, Edh1, Igbf3, Mue11a, Ppp1ca, Ppp2ca, Prka, Puma, Tdg, Tp53, Tp53inp1 and Zac1, were direct binding targets of miR-125b in human cells (Figure 3A). In mouse 3T3 cells, 11 out of 13 gene transcripts, Bak1, Hspa5, Ithc, Ppp1ca, Ppp2ca, Prka, Puma, Sell1, Spl, Tdg and Tp53inp1, were found to be direct binding targets of miR-125b (Figure 3B). In zebrafish embryos, 8 out of 14 gene transcripts, Cdc25c, Cdc26e, Gfzb1h, Hspa5, Ithc, Ppp1ca, Sell1, and Tp53, were pulled down by miR-125b (Figure 3C). Tp53 mRNA was pulled down by miR-125b only in human lung fibroblasts and zebrafish embryos but not in mouse fibroblasts, consistent with previously published results [16] and the Targetscan algorithmic prediction that miR-125b targets Tp53 in humans and zebrafish but not in mice.

Validation of miR-125b targets in the p53 network

As a final validation of the candidate miR-125b targets we have identified in the p53 network, we tested our candidate target genes with the luciferase reporter assay. Where cloning was successful, we cloned the entire 3′ UTR of selected candidate target genes into a Renilla luciferase reporter, and assayed luciferase expression following co-transfection of miR-125b duplex into HEK-293T cells. Transfection of miR-125b significantly suppressed 40-60% (P<0.01) of the luciferase activity of many 3′ UTR reporters of the miR-125b targets we analyzed, relative to transfection of the negative control miRNA (Figure 4A). For humans, the 3′ UTR reporters of Bak1, Cdc25c, Ppp1ca, Ppp2ca, Pkra, Puma, Tdg, Tp53, Tp53inp1, and Zac1 were significantly suppressed by miR-125b. In mice, the 3′ UTR reporters of Bak1, Ithc, Ppp1ca, Ppp2ca, Pkra, Puma, Sell1, Tdg, and Tp53inp1 were significantly suppressed by miR-125b (Figure 4B). In zebrafish, the 3′ UTR reporters of Cac, Cdc25e, Cdc26e, Gfzb1h, Hspa5, Ppp1ca, and Tp53 were significantly suppressed by miR-125b (Figure 4C). With the exception of zebrafish Cac, all genes tested were positive in the miR-125b-pull-down as well as the miR-125b gain- and loss-of-function screen. Amongst these targets, we found Ppp1ca, Pkra and Tp53 to be especially interesting from the evolutionary viewpoint, since all 3 vertebrate species possess these 3 genes, but each gene shows a different pattern of evolutionary conservation with respect to miR-125b repression. Ppp1ca is repressed by miR-125b in all 3 species, Pkra is repressed by miR-125b in humans and mice, while Tp53 is repressed in humans and zebrafish. To examine the sequence evolution of these mRNA-mRNA pairs in greater detail, we compared the Targetscan-predicted miR-125b binding sites of these genes in humans, mice and zebrafish. In Ppp1ca, the predicted binding site is 95% identical between humans and mice and 55% identical between humans and zebrafish, while the seed binding sequence is 100% conserved in all 3 species (Figure 4D). In Pkra, the predicted binding site is 94% identical between humans and mice, but only 26% identical between humans and zebrafish, while the seed binding sequence is completely absent in zebrafish (Figure 4D). In contrast, the predicted binding site in Tp53 is 64% identical between humans and zebrafish, and the seed binding sequence is 100% conserved between humans and zebrafish, but only 36% identical between humans and mice when the mouse seed binding sequence has acquired 2 point mutations (Figure 4D). The miR-125b-repression patterns we observed for each of these genes in the qPCR, pull-down and luciferase assays are consistent with these DNA sequence analyses, suggesting that evolution in the miRNA-mRNA binding is driving the evolution in miR-125b-repression patterns. Introduction of point mutations into the predicted seed binding sequences abrogated miR-125b-repression of each target 3′UTR luciferase reporter (P<0.05), validating the predicted miR-125b binding sites and confirming the miRNA-mRNA sequence evolution patterns we observed (Figure 4E).

Finally, we checked miR-125b regulation of protein expression in a subset of p53 network targets for which reliable Western blotting was possible. miR-125b significantly downregulated the protein levels of human BAK1, PP1CA, TP53INP1, PP2CA, CDC25C, and TP53 in SH-SY5Y neuroblastoma cells (Figure 4F). In mouse N2A neuroblastoma cells, miR-125b significantly downregulated mouse BAK1, PP1CA, PUMA, and ITCH protein (Figure 4G).

miR-125b regulation of the p53 network, but not individual miRNA-target pairs, is conserved

Our results reveal that miR-125b regulation of the p53 network is conserved at the network-level over the course of vertebrate evolution, but individual miRNA-target pairs are evolving rapidly.
Figure 2. GOF/LOF screen for p53 network genes regulated by miR-125b. (A) Loss-of-function (LOF) screens were performed in human primary lung fibroblasts (hLF) or mouse 3T3 fibroblasts by transfecting an antisense RNA against both miR-125a and miR-125b (miR-125a/b-AS), or by microinjecting morpholinos (MO) against pre-miR-125b hairpin precursors (all 3 isoforms) into zebrafish embryos. Gain-of-function (GOF) screens were performed in human SH-SY5Y and mouse N2A neuroblastoma by transfecting the miR-125b duplex into cells in culture, or by coinjecting the miR-125b duplex with the morpholinos against pre-miR-125b into zebrafish embryos. Fold changes in gene expression were measured by qRT-PCR twenty-four hours after transfection or injection, relative to the mock and negative control miRNA or morpholino, and shown as log2(fold change) using a heat-map. (B) Human: 13 genes were significantly derepressed by a loss of miR-125b, while 20 genes were significantly repressed by a gain of miR-125b, making a total of 22 genes that passed the screen (P < 0.05, fold change > 1.3, relative to mock control). (C) Mouse: 11 genes were significantly derepressed by a loss of miR-125b, while 12 genes were significantly repressed by a gain of miR-125b (P < 0.05, fold change > 1.3, relative to control MO), making a total of 14 genes that passed the screen. All experiments were performed with at least three biological replicates.

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To summarize our results, our list of predicted miR-125b targets in the p53 network (Table S1) was filtered and reclassified according to the results of the screen and validation assays (Figure 5). From the GOF/LOF screen we were able to identify mRNAs perturbed by miR-125b. However these results did not discriminate between direct or indirect targets. To supplement these experiments the pull-down assay was used to uncover mRNAs physically associated with miR-125b. Of note, the pull-down might not identify mRNA targets that are rapidly degraded, and as such the luciferase reporter assay can complement its shortcomings. Taken together...
Figure 4. Validation of miR-125b targets. Candidate p53 network genes that were positive in both the GOF/LOF screen and miR-125b pull-down were validated for targeting by miR-125b using the 3' UTR luciferase reporter assay and Western blots for protein expression. (A-C), Reporter genes containing the full-length 3' UTRs of each selected target gene were co-transfected with miR-125b duplex into 293T cells. Luciferase readings were obtained 48 hours after transfection and presented here as the average percentage of luciferase activity ± s.e.m. (n=3) relative to a scrambled duplex co-transfected control (100%). A reporter containing a 23-nucleotide-binding-site with perfect complementarity to miR-125b was used as the perfect match positive control, while the unmodified luciferase reporter was used as the empty negative control. (A) Human: 10 out of 13 candidate genes' 3' UTRs showed significant repression by miR-125b relative to the control (p<0.01). (B) Mouse: 9 out of 11 candidate genes' 3' UTRs showed significant repression by miR-125b relative to the control (p<0.01).
Conserved Regulation of p53 Network by miR-125b

significant repression by miR-125b relative to the control (p<0.01). (C) Zebrafish: 7 out of 8 candidate genes’ 3’ UTRs showed significant repression by miR-125b relative to the control (p<0.01). (D) Alignment of predicted miR-125b binding sites in the 3’UTRs of Ppp1ca, Prkra and Tp53 across three species. Seed-binding sequences are underlined. Bases conserved in two (blue) or three (black) species are highlighted. (E) The 3’UTR seed-binding sequences of 7 target mRNAs were mutated and assayed for direct binding to miR-125b using the luciferase reporter assay, relative to wild-type 3’UTR sequences. (F) The seed-binding sequences in the 3’UTR of 7 predicted target mRNAs were mutated and compared to wild-type sequences for binding to miR-125b using luciferase reporter assay. (G) Western blot analysis of protein expression of selected target genes two days after a transfection of miR-125b duplex, miR-125b antisense (AS) or negative control duplex or negative control antisense. (F) Western blots showed that miR-125b repressed Bak1, PPP1CA, PUMA, and ITCH levels in mouse N2A neuroblastoma cells. Abbreviations: h, human; m, mouse; z, zebrafish.

The three assays provide a powerful means to identify direct miR-125b targets. In order to minimize false positives, we counted the number of assays for which each gene target was positive, and gene targets that failed to pass at least 2 assays in at least one vertebrate species were filtered out. Predicted targets that passed 3 assays (red), 2 assays (orange), 1 assay (yellow), or predicted targets that failed all assays but whose orthologues in other species passed 3 assays of direct regulation by miR-125b (pink), were colored as indicated (Figure 5). Using our conservative estimate of miR-125b targets in the p53 network, we found that in all three vertebrates we examined – humans, mice and zebrafish – miR-125b regulates multiple p53 network genes. This shows that miR-125b regulation of the p53 network is conserved at least at the network level. However very few individual gene targets of miR-125b in the p53 network were conserved across all three vertebrates (Figure 5; Figure 6A-6C). Instead, conserved miR-125b regulation of the p53 network appears to occur through evolving miRNA-target pairs in the three vertebrates – zebrafish (Figure 6A), mouse (Figure 6B), and humans (Figure 6C). In general, we observe miR-125b regulating 2 general classes of genes in the p53 network: (i) apoptosis regulators like Bak1, Kyph3, Ick, Puma, Prkra, Tp53inp1, Tp53, and Zac1, and (ii) cell-cycle regulators like cyclin C, Cdc25c, Cdb2c, Edn1, Ppp1ca, and Sel1l.

Because miR-125b represses both pro-apoptosis and anti-apoptosis genes, as well as both proliferation and cell-cycle arrest genes in all three vertebrates (Figure 5), miR-125b appears to modulate the p53 network on the whole through an incoherent feedforward loop (FFL) [33,34] acting on the cellular processes of apoptosis and cell proliferation (Figure 6D). An incoherent type-2 FFL is a regulatory pattern in which X represses a target Z and also represses Y, another repressor of Z (Figure 6D). Incoherent FFLs have been found in the transcription factor networks of human embryonic stem cells and hematopoietic stem cells, and have been shown to modulate E2F1 dosage in the Myc-E2F1 pathway [35-37]. Besides accelerating responses and acting as amplitude filters [38-40], the incoherent FFL motif is also a noise buffering motif that reduces the variance of network dosage [41-43]. Thus our finding that incoherent FFLs fit the overall structure of network relationships between miR-125b and the p53-mediated processes, suggests that miR-125b is fine-tuning and buffering p53 network dosage.

Discussion

In this study, we sought to identify direct targets of miR-125b in the p53 network of humans, mice and zebrafish, to better understand how miR-125b regulates the p53 network throughout evolution and how that might relate to its conserved role in regulating tissue stem cells.

We identified 20 direct targets of miR-125b in the p53 network, including 15 novel targets like Zac1, Puma, Ick and Cdc25c, and also targets like Bak1 and Tp53 that were identified in previous studies [9,16-18]. In general, we found that miR-125b directly represses 2 classes of genes: apoptosis regulators and cell-cycle regulators. With the exception of Ppp1ca, Ick and Edn1, very few individual targets were strictly conserved throughout vertebrate evolution. Instead, we found that only the network-level of regulation was conserved, and miR-125b-regulation of individual apoptosis and proliferation regulators appears to be evolving rapidly from species to species. This observation suggests that, at least within the vertebrates, the 3’ UTR sequences of each gene target is evolving rapidly via neutral genetic drift. In other words, the loss or gain of a single miR-125b-binding site in the 3’ UTR of most genes appears to have a relatively insignificant effect on the fitness of an organism. On the other hand, the strict conservation of miR-125b-regulation at the network-level in humans, mice and zebrafish, suggests that natural selection acts on the network-level rather than the gene-level with regard to miRNA-target evolution. It will be interesting to see if this novel paradigm applies to other microRNAs or gene networks as well.

Previous studies on miRNA evolution have suggested that a relatively poor conservation of individual miRNA-target pairs but strong conservation of a miRNA-gene network relationship is consistent with miRNAs’ role as buffers of gene expression [42,44,45]. Our observation that an incoherent FFL-like network motif fits the overall structure of the miR-125b – p53 network models with respect to apoptosis and cell proliferation, lends further support to this idea since incoherent FFL network motifs are well-adapted for noise filtering [41,43,46]. It is thought that miRNAs are at least partially responsible for the phenomenon of developmental or phenotypic stability within each species [41,42], termed “canalization” by C. H. Waddington [47]. These studies suggest that miRNAs have a conserved role in regulating the overall stability of pathways/networks, a role which is relatively unaffected by the loss or gain of individual miRNA-targets over the course of evolution. A network buffering function has also been suggested for the regulation of muscle development by miR-1 – throughout evolution, regulation of the Wnt pathway by miR-8, and fine-regulation of Pten dosage by a variety of miRNAs [48-50]. Our findings suggest that the fine-tuning of p53 network dosage by miR-125b is another example of this paradigm.

Fine-regulation of p53 network dosage by miR-125b may also explain miR-125b’s conserved role in regulating tissue stem cell homeostasis. In C. elegans, loss-of-function mutations in lin-4 lead to a delay in differentiation and thus expansion of vulval precursor cells, seam stem cells in the lateral hypodermis and mesoblasts, causing multiple defects in larval development [6]. In zebrafish, loss of miR-125b leads to widespread p53-dependent apoptosis with consequent defects in early embryogenesis, especially in neurogenesis and somitogenesis [16]. Overexpression of miR-125a/b causes an expansion of mammalian hematopoietic stem cells (HSCs) and aberrant differentiation, leading to myeloid leukemia [9,10] and also lymphoid leukemia if miR-125b is overexpressed in fetal liver HSC-enriched cells [12]. However, the molecular underpinnings of miR-125a/b’s regula-
tion of tissue stem cell homeostasis had remained unclear largely due to the complex nature of microRNA regulation of gene networks. The 2 classes of miR-125b targets in the p53 network, and the incoherent FFL network motifs that we found, may at least partially explain how miR-125b regulates tissue stem cells in vertebrates. By fine-tuning both apoptosis regulators and cell-cycle regulators, miR-125b may fine-tune the p53 network dosage to drive the self-renewal of tissue stem cells. It could explain how overexpression of miR-125b leads to an expansion of self-renewing hematopoietic stem cells while loss of miR-125b leads to aberrant apoptosis and proliferation, with consequent defects in tissue differentiation.

Several studies have implicated miR-125b as an oncogene in a variety of mammalian tissue compartments, e.g. leukemia, neuroblastoma, prostate cancer and breast cancer [9–18]. These studies have ascribed miR-125b’s anti-apoptotic effect as an oncogene to its direct suppression of Bak1 or Tp53 [9,16-18]. On the other hand, several research groups have also reported miR-125b’s role as a

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**Figure 5. Summary of genes in p53 network that are directly targeted by miR-125b.** Only targets that passed ≥ 2 validation assays, in at least one species, are shown. Red: predicted targets validated by 3 assays; Orange: predicted targets validated by 2 assays; Yellow: predicted targets validated by 1 assay; Pink: predicted targets not validated by any assay, but validated by 2 assays in another species.

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<td>ITCH</td>
<td>anti-apoptosis</td>
<td>ubiquitinates p63 and p73 for degradation to prevent apoptosis</td>
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<td>CCNC</td>
<td>cell cycle</td>
<td>binds to CDK3 to phosphorylate Rb and promote G0/G1 transition</td>
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<tr>
<td>CDC25C</td>
<td>cell cycle</td>
<td>repressed by p53; activates cyclin B1/CDK1 complex to promote G2/M transition</td>
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<td>EDN1</td>
<td>cell cycle, anti-apoptosis</td>
<td>repressed by p53; co-mitogenic factor with IGF1 and PDGF to promote G1/S transition and downregulate apoptosis</td>
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<tr>
<td>SEL1L</td>
<td>cell cycle arrest</td>
<td>p53 target; upregulates Pten to downregulate PI3K signaling and mediate G1/S arrest</td>
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<tr>
<td>CDKN2C</td>
<td>cell cycle arrest</td>
<td>inhibits CDK4 to mediate G1/S arrest</td>
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<tr>
<td>PPP1CA</td>
<td>cell cycle, cell cycle arrest; pro-apoptosis</td>
<td>inactivates p33 to promote proliferation; activates Rb for cell cycle arrest and apoptosis; activates Bad to induce apoptosis</td>
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<td>PPP2CA</td>
<td>pro- and anti-apoptosis</td>
<td>activates Bad and inactivates Bcl-2 to induce apoptosis; inactivates p53 to prevent apoptosis</td>
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<td>BAK1</td>
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<td>p53 target; mediates apoptosis</td>
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<td>PUMA</td>
<td>pro-apoptosis</td>
<td>p53 target; mediates apoptosis</td>
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<tr>
<td>IGF1B3</td>
<td>pro-apoptosis; cell cycle arrest</td>
<td>p53 target; downregulates IGF1R-PI3K signaling to promote apoptosis and cell cycle arrest</td>
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<tr>
<td>ZAC1</td>
<td>pro-apoptosis; cell cycle arrest</td>
<td>p53 target; binds and coactivates p53 transcription</td>
</tr>
<tr>
<td>TP53</td>
<td>pro-apoptosis; cell cycle arrest</td>
<td>Tumor suppressor protein p53</td>
</tr>
<tr>
<td>SP1</td>
<td>pro-apoptosis; cell cycle arrest</td>
<td>binds and coactivates p53 transcription</td>
</tr>
<tr>
<td>PRKRA</td>
<td>pro-apoptosis; stress response</td>
<td>p53 target; activates PKR to promote apoptosis</td>
</tr>
<tr>
<td>TP53NI</td>
<td>pro-apoptosis</td>
<td>p53 target; mediates apoptosis</td>
</tr>
<tr>
<td>GTF2H1</td>
<td>stress response; DNA repair</td>
<td>binds to p53 to modulate nucleotide excision repair of DNA</td>
</tr>
<tr>
<td>HSPA5</td>
<td>stress response</td>
<td>binds to p53; promotes stress-induced autophagy</td>
</tr>
<tr>
<td>MRE11A</td>
<td>stress response; DNA repair</td>
<td>activates ATM - p53 signaling and homologous recombination or non-homologous end joining (NHEJ) to repair DNA</td>
</tr>
<tr>
<td>TDG</td>
<td>stress response; DNA repair</td>
<td>mediates base excision repair of DNA</td>
</tr>
</tbody>
</table>

**Legend**

No. of Screen/Validation assays that each target passed:

- 3 GOF/LOF, pull-down & luciferase
- 2 GOF/LOF, & pull-down or luciferase
- 1 GOF/LOF
- 0 predicted target; validated in other species
potential tumor suppressor by suppressing proliferation in cell-culture models [19–23]. Our identification of 20 direct targets of \textit{miR-125b} in the p53 network reconciles these findings because \textit{miR-125b} modulates the expression of both apoptosis regulators and cell-cycle regulators. Although \textit{miR-125b}’s suppression of p53 itself is not conserved in mice, \textit{miR-125b}’s anti-apoptotic role – through suppression of multiple pro-apoptosis regulators in the p53 network – appears to be conserved in vertebrates. \textit{miR-125b}’s ability to fine-tune the subtle balance of apoptosis vs. cell-cycle regulators and thus buffer the p53 network dosage in different contexts, could explain why \textit{miR-125b} dysregulation can lead to either tumor suppression or oncogenesis depending on the context. It is possible that this buffering feature of \textit{miR-125b} represents a general principle of miRNA regulation of gene networks.

Figure 6. Models of \textit{miR-125b} regulation of p53 networks in humans, mice, and zebrafish. (A) Human p53 network. (B) Mouse p53 network. (C) Zebrafish p53 network. Models were constructed by Ingenuity Pathway Analysis. Red: predicted targets validated by 3 assays; Orange: predicted targets validated by 2 assays; Yellow: predicted targets validated by 1 assay; Pink: predicted targets not validated by any assay, but validated by 3 assays in another species. (D) Incoherent feedforward loop (FFL) motifs characterize \textit{miR-125b} regulation of p53 network genes that mediate apoptosis or cell cycle arrest. doi:10.1371/journal.pgen.1002242.g006
Materials and Methods

Prediction of miR-125b targets in the p53 network

A list of p53-associated genes was compiled from the p53 Knowledgebase website [30] and from the Ingenuity Pathway Analysis™ database [29]. The targets of miR-125b in human and mouse were predicted by TargetScan [31]. The targets of miR-125b in zebrafish were predicted by MicroCosm [32]. The human homologues of mouse and zebrafish targets were identified by the DAVID gene ID conversion tool.

Cell culture and transfection

Human lung fibroblast cells, human neuroblastoma SH-SY5Y cells, mouse neuroblastoma Neuro-2A cells, mouse fibroblast Swiss-3T3 cells and human HEK-293T cells were maintained in DMEM media, supplemented with 10% fetal bovine serum and 1% penicillin-streptomycin (Invitrogen). Neuro-2A cells, 3T3 cells, SH-SY5Y cells and human lung fibroblast cells were transfected in suspension with 5x10⁵ cells per well in 6-well plates using lipofectamin-2000 (Invitrogen). After 48 hours, the cell extract was obtained; Firefly and Renilla luciferase activities were measured with the Dual-Luciferase reporter system (Promega) according to the manufacturer’s instructions.

Western blot assay

Cells were lysed in RIPA buffer (Pierce). Proteins were separated by a 10% polyacrylamide gel and transferred to a methanol-activated PVDF membrane (GE Healthcare). The membrane was blocked for one hour in PBS containing 7.5% milk and subsequently probed with primary antibodies (Santa Cruz) overnight at 4°C. After 1-hour incubation with goat-antirabbit HRF-conjugated secondary antibody (Santa Cruz), the protein level was detected with luminol reagent (Santa Cruz).

Statistical analysis

Two-tail T-tests were used to determine the significance of differences between the treated samples and the controls where values were obtained from luciferase reporter assay or qRT-PCR. The tests were performed using Microsoft Excel where the test type is always set to two-sample equal variance.

Supporting Information

Figure S1 Mature miR-125b levels before and after overexpression or knockdown. (A) The level of miR-125b in human SH-SY5Y cells one day after a transfection with mock (lipofectamin-2000 only), negative control duplex (NC-DP) or miR-125b duplex (125b-DP). (B) The level of miR-125b in mouse N2A cells one day after a transfection with mock, NC-DP or 125b-DP. (C) The level of miR-125a and miR-125b in human lung fibroblasts one day after a transfection with mock, negative control antisense (NC-AS) or miR-125a antisense and miR-125b antisense cotransfection (125ab-AS). (D) The level of miR-125a and miR-125b in mouse SWISS-3T3 fibroblasts one day after a transfection with mock, NC-AS or (125ab-AS). In all panels, the levels of miR-125a and miR-125b were quantified by real-time PCR, and presented as log₂ (fold change) ± s.e.m. (n≥3) relative to the levels of RNU6B loading control.

Table S1 Genes in p53 network with predicted miR-125b binding sites. *Hsa: Homo sapiens, humans. †Mmu: Mus musculus, mouse. ‡Dre: Danio rerio, zebrafish. *Non-official but common gene name that is used in this paper.

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References
15. Sonoki T, Iwanaga E, Mitsuya H, Asou N (2005) Insertion of microRNA-125b-1 into the human lung fibroblasts and SH-SY5Y cells; Wa Xian and Frank McKeon for providing Swiss-3T3 cells; Lim Qing En and Too Heng Phon for providing Neuro-2A cells; Michael Kharas for help with flow cytometry.

Author Contributions
Conceived and designed the experiments: MTNL NS-C HFL BL. Performed the experiments: MTNL NS-C SLK LC JT CT. Analyzed the data: MTNL NS-C. Contributed reagents/materials/analysis tools: VK EO AL HY. Wrote the paper: NS-C MTNL HFL BL JL.