P53-miR-191-SOX4 regulatory loop affects apoptosis in breast cancer

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ABSTRACT

miRNAs have emerged as key participants of p53 signaling pathways because they regulate or are regulated by p53. Here, we provide the first study demonstrating direct regulation of an oncogenic miRNA, miR-191-5p, by p53 and existence of a regulatory feedback loop. Using a combination of qRT-PCR, promoter-luciferase, and chromatin-immunoprecipitation assays, we show that p53 brings about down-regulation of miR-191-5p in breast cancer. miR-191-5p overexpression brought about inhibition of apoptosis in breast cancer cell lines (MCF7 and ZR-75) as demonstrated by reduction in annexin-V stained cells and caspase 3/7 activity, whereas miR-191-5p down-regulation showed the opposite. We further unveiled that *SOX4* was a direct target of miR-191-5p. *SOX4* overexpression was shown to increase p53 protein levels in MCF7 cells. miR-191-5p overexpression brought about down-regulation of *SOX4* and thus p53 levels, suggesting the existence of a regulatory feedback loop. Breast cancer treatment by doxorubicin, an anti-cancer drug, involves induction of apoptosis by p53; we thus wanted to check whether miR-191-5p affects doxorubicin sensitivity. Interestingly, Anti-miR-191 treatment significantly decreased the IC₅₀ of the doxorubicin drug and thus sensitized breast cancer cells to doxorubicin treatment by promoting apoptosis. Overall, this work highlights the importance of the p53-miR-191-SOX4 axis in the regulation of apoptosis and drug resistance in breast cancer and offers a preclinical proof-of-concept for use of an Anti-miR-191 and doxorubicin combination as a rational approach to pursue for better breast cancer treatment.

Keywords: breast cancer; apoptosis; miR-191; p53; *SOX4*

INTRODUCTION

Breast cancer is the most common invasive cancer in females that has shown improved survival owing to early detection and better therapeutic strategies. However, death due to breast cancer still shows a rising trend (Florea and Büsselberg 2013; American Cancer Society 2015). This has been attributed to multiple factors, including metastases and development of resistance to various therapies (Gonzalez-Angulo et al. 2007). Therefore, there is a need to understand breast tumor biology in its entirety and identify novel mediators involved in the pathogenesis.

Breast tumor development involves deregulated networks of specific protein coding genes, long noncoding RNAs, and miRNAs (Nana-Sinkam and Croce 2011). The key for

breast cancer treatment lies in the identification and targeting of the crucial players among these complicated and huge networks. Recently, the significant role of an miRNA, miR-191-5p, in breast tumor biology was demonstrated by multiple groups (Di Leva et al. 2013; Nagpal et al. 2013, 2015). miR-191-5p was shown to be an estrogen and hypoxia responsive miRNA that promotes cell proliferation and migration in breast cancer (Nagpal et al. 2013, 2015). The oncogenic role of miR-191-5p has also been shown in other cancers (lung cancer, colon cancer, gastric cancer) (Xi et al. 2006; Elyakim et al. 2010; Patnaik et al. 2010; He et al. 2011; Shi et al. 2011; Zhou et al. 2013; Liu et al. 2014). miR-191-5p is up-regulated in breast cancer patients with a trend of higher expression in estrogen receptor positive patients as compared to the negative ones (Di Leva et al. 2013; Nagpal et al. 2013). We became interested in studying the regulation of miR-191-5p and identified multiple p53 binding sites in its

Abbreviations: miRNAs, microRNAs; *SOX4*, SRY (sex determining region Y)-Box 4; *PLCD1*, phosphoinositide phospholipase C- Δ -1; *CDK6*, cell division protein kinase 6; *SATB1*, special AT-rich sequence binding protein1; *BDNF*, brain-derived neurotrophic factor; *CEBPB*, CCAAT/enhancer binding protein β ; *TJPI1*, tight junction protein1; *TCF7L2*, transcription factor 7 like2; ChIP, chromatin immunoprecipitation.

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promoter. Here, we report the regulation of miR-191-5p by p53 and the existence of a feedback loop wherein miR-191-5p emerges as an important regulator of p53 levels and consequently apoptosis and drug resistance in breast cancer. The work suggests the use of Anti-miR-191 therapy along with doxorubicin for effective breast cancer treatment.

RESULTS

P53 down-regulates miR-191-5p in breast cancer

First, we looked for transcription factor binding sites in the upstream region of miR-191-5p. In silico analysis of regions ~7 kb upstream of and 1 kb downstream from miR-191-5p using Promo 3.0 software showed the presence of multiple putative p53 binding sites showing <1% dissimilarity (Fig. 1A; Messegue et al. 2002; Farré et al. 2003). To evaluate the role of p53 in the regulation of miR-191-5p, we overexpressed p53 in breast cancer lines and measured p53 and miR-191-5p levels 48 h post-transcription (Fig. 1B; Supplemental Fig. S1A, B). The miR-191-5p levels were significantly reduced in p53-transfected cells relative to cells transfected with the control (Fig. 1B). To determine whether p53-mediated miR-191-5p

down-regulation is through the promoter, the regions bearing p53 binding sites (P1, four binding sites; P2, one binding site; P3, two binding sites; and P4, four binding sites) were cloned in luciferase promoter vector and scored for luciferase activity. The regions P1 and P4 showed a significant decrease in luciferase activity on p53 overexpression in MCF7 and ZR-75 cells, suggesting that p53 down-regulates miR-191-5p at the transcriptional level through the promoter (Fig. 1C). Further, in order to confirm direct binding of p53 to the predicted binding sites, chromatin immunoprecipitation (ChIP) assay was performed in MCF7 cells using p53 and IgG antibodies. We found that binding to P1 and P4 was more enriched in p53 as compared to the control (IgG) pull-down fractions (Fig. 1D,E). Based on these results, we confirm that p53 directly regulates the transcriptional expression of miR-191-5p through binding to the p53 response elements in its promoter.

miR-191-5p functions as an anti-apoptotic miRNA in breast cancer

We next wanted to check whether miR-191-5p is involved in the regulation of apoptosis in breast cancer. The PE/Annexin

FIGURE 1. Transcriptional regulation of miR-191-5p. (A) Diagram showing putative p53 binding sites with <1% dissimilarity in the upstream region of pre-miR-191. P1–P4 refers to the four regions cloned in the pGL3 luciferase vector for studying the effect of p53 overexpression in breast cancer cells. (B) Real-time PCR data showing down-regulation of miR-191-5p levels in response to p53 overexpression as compared to the control vector (pc) in breast cancer cell lines (MCF7 and ZR-75). RNU6B (U6) has been used as a control for normalization. Graph was plotted using $2^{-\Delta\Delta C_t}$ formula. (C) Luciferase activity of regions (P1–P4) was measured in MCF7 and ZR-75 48 h post-transfection with pc or p53. (D,E) p53 direct binding to the miR-191-5p upstream region was confirmed using ChIP analyses. qPCR (D) or semiquantitative (E) data showing enrichment of the p53 recognition elements (p53RE) in P1 and P4 regions on immunoprecipitation with p53 antibody as compared to the IgG control. The graphical data points represent mean $\pm$ SD of at least three independent experiments. (**) $P < 0.01$. Error bars denote $\pm$ SD.

FIGURE 2. miR-191-5p functions as an anti-apoptotic miRNA in breast cancer. FACS analysis using PE-Annexin V and 7AAD was performed in MCF7 cells (A) and ZR-75 (B) transfected with pc-191 and Anti-191 along with their controls (pc and NCtrl, respectively). (C) Graph shows percentage of Annexin V positive cells. (D) Caspase 3/7 activity in MCF7 and ZR-75 cells 48 h post-transfection with pc-191 and Anti-191 along with their controls. (E) Evaluation of apoptosis by DAPI staining in MCF7 cells in response to miR-191-5p overexpression or inhibition. (F) Graph shows percentage of apoptotic cells. The graphical data points represent mean $\pm$ SD of at least three independent experiments. (*) $P < 0.05$, (**) $P < 0.01$. Error bars denote $\pm$ SD.

V staining showed that the percentage of the early apoptotic cells was lower in the cells transfected with pc-191 when compared with the control (Fig. 2A–C). In contrast, the percentage of early apoptotic cells was higher in the cells transfected with Anti-191 than in NCtrl (Fig. 2A–C). Further, to know whether the effect of miR-191-5p on apoptosis is caspase dependent, we measured caspase 3/7 activity in the transfected MCF7 and ZR-75 cells. miR-191-5p transfected cells showed a decrease in caspase 3/7 activity, whereas Anti-191 treatment increased caspase 3/7 activity (Fig. 2D). Microscopic examination of DAPI (DNA binding fluorescent dye) stained cells revealed a higher number of cells showing shrinkage and the appearance of apoptotic bodies in Anti-

191 treated cells as compared to control (Fig. 2E). In contrast, miR-191-5p overexpressing cells showed a lower number of apoptotic cells as compared to its control (Fig. 2F). Overall, these results demonstrate that miR-191-5p inhibits apoptosis in breast cancer cells.

miR-191-5p targets genes in breast cancer

Next we wanted to evaluate the mechanism of miR-191-5p-mediated inhibition of apoptosis. We used online target prediction programs (TargetScan, miRanda) to investigate the potential targets of miR-191-5p and shortlisted eight target genes (*SOX4*, *PLCD1*, *CDK6*, *SATB1*, *CEBPB*, *BDNF*,

FIGURE 3. Targets of miR-191-5p. Expression level of shortlisted target genes of miR-191-5p was determined by qRT-PCR in response to miR-191 overexpression or inhibition in MCF7 (A) and ZR-75 (B) cell lines. (C) Diagram showing miR-191-5p binding sites in the 3' UTR of target genes. 7 mer, 8 mer, and m8 refer to seed match sites. (D) Luciferase activity of the 3' UTR luciferase constructs bearing miR-191-5p binding sites in response to transfection with pc-191 or its control vector (pc) in MCF7 cells. The graphical data points represent mean $\pm$ SD of at least three independent experiments. (*) $P < 0.05$, (**) $P < 0.01$. Error bars denote $\pm$ SD.

TJP1, *TCF7L2*) that have been shown to be associated with breast cancer and also are related to regulation of apoptosis (Hoevel et al. 2004; John et al. 2005; Zahnow 2009; Hur et al. 2010; Li et al. 2010; Vanhecke et al. 2011; Chen et al. 2013; Agarwal et al. 2015; Mu et al. 2015; Tadesse et al. 2015). Interestingly, some of the targets (*SOX4*, *PLCD1*, *CDK6*, and *SATB1*) showed down-regulation; the others (*CEBPB* and *BDNF*) showed induction on pc-191 overexpression, while *TJP1* and *TCF7L2* remained unchanged (Fig. 3A,B). An opposite trend was seen on Anti-miR-191 treatment of MCF7 cells (Fig. 3A,B). To further confirm direct targeting of these transcripts by miR-191-5p, 3' UTR-luciferase reporter assay was performed for six target transcripts (*SOX4*, *PLCD1*, *CDK6*, *SATB1*, *CEBPB*, and *BDNF*) containing miR-191-5p binding sites in their 3' UTR (Fig. 3C). The 3' UTR-luciferase assay results showed that *SOX4*, *PLCD1*, *CDK6*, and *SATB1* were down-regulated by miR-191-5p, whereas *CEBPB* and *BDNF* were induced by miR-191-5p (Fig. 3D). The above results confirmed that all these genes (*SOX4*, *PLCD1*, *CDK6*, *SATB1*, *CEBPB*, and *BDNF*) are direct targets of miR-191-5p.

miR-191-5p targets *SOX4* and brings about down-regulation of p53 levels in breast cancer

Pan et al. (2009) demonstrated that *SOX4* regulates p53 protein stability in colon cancer. Therefore, we focused on *SOX4* to explore the miR-191-5p-mediated effects on an apoptotic pathway. For this, we first confirmed the direct effect of miR-191-5p on *SOX4* levels by performing luciferase assay with wild-type or mutated miR-191-5p binding site in the 3' UTR of *SOX4* (Fig. 4A). We found a significant decrease in the luciferase activity of wild-type *SOX4* 3'UTR but not the mutated one (Fig. 4B). Next, we measured *SOX4* protein levels in response to miR-191-5p modulation. We found that miR-191-5p overexpression decreased the *SOX4* protein levels, whereas Anti-191 treatment showed the opposite trend (Fig. 4C). To evaluate the effect of *SOX4* on p53 protein stability in breast cancer, MCF7 cells were transfected with *SOX4* or control plasmid, and 48 h post-transfection, the transcript as well as the protein level of p53 were measured (Fig. 4D,E; Supplemental Fig. S1C). We found that while the p53 transcript remained unaffected, the p53 protein levels were increased on *SOX4* overexpression. Therefore,

FIGURE 4. miR-191-5p down-regulates SOX4 and p53 levels. (A) Diagram showing wild-type/mutated miR-191-5p binding site in the *SOX4* 3' UTR. (B) Luciferase activity of the *SOX4* wild-type/mutated miR-191-5p binding site bearing constructs was measured in response to pc or p53 overexpression. (C) *SOX4* levels were measured at the protein level in MCF7 cells expressing pc-191 or Anti-191 along with their controls. The results show that miR-191-5p overexpression down-regulates *SOX4* at the protein level. β -Actin was used for normalization of Western blotting data. (D,E) To evaluate the effect of *SOX4* on p53 protein stability in breast cancer, MCF7 cells were transfected with *SOX4* or control vector (pc), and 48 h post-transfection, p53 transcript as well as protein levels were measured by qRT-PCR and Western blotting. The real-time graph was plotted using $2^{-\Delta\Delta C_t}$ formula. (F,G) The transcript and protein levels of p53 and p21 were evaluated by qRT-PCR and Western blotting in response to miR-191-5p overexpression or inhibition in MCF7 cells. The real-time graph was plotted using $2^{-\Delta\Delta C_t}$ formula. (H) p53 protein level was determined by Western blotting in MCF7 cells transfected with pc-191 alone or in combination with *SOX4*. The graphical data points represent mean $\pm$ SD of at least three independent experiments. (*) $P < 0.05$, (**) $P < 0.01$. Error bars denote $\pm$ SD.

the regulation of p53 by *SOX4* is more likely at the post-transcriptional level. This hinted at the existence of a feedback loop between miR-191-5p and p53, and we hypothesized that miR-191-5p by down-regulation of *SOX4* may affect p53 levels. We found that overexpression of miR-191-5p in MCF7 cells resulted in a decrease of p53 protein levels with a minimal effect on transcript levels. A similar trend was seen on the levels of a well-known p53 target gene, *p21* on miR-191-5p overexpression/inhibition (Fig. 4F,G). Replenishing *SOX4* in miR-191-5p overexpressing cells brought back the p53 protein levels when compared to the control (Fig. 4H). Collectively, all these results demonstrated that miR-191-5p targets *SOX4* and brings about down-regulation of p53 levels in breast cancer.

Anti-miR-191 treatment sensitizes breast cancer cells to doxorubicin-mediated apoptotic death

Apoptosis is a common and preferred target of several breast cancer treatment strategies. Since miR-191-5p is shown here

to be anti-apoptotic miRNA, we wanted to evaluate the effect of using anti-miR-191 oligos (Anti-191) either alone or in combination with the chemotherapeutic drug doxorubicin for breast cancer treatment. Interestingly, we found that the IC₅₀ value of doxorubicin was significantly decreased with the combination treatment of Anti-191 and doxorubicin when compared with the controls (NCtrl + Dox and Dox free) and displayed higher sensitivity to the treatment (Fig. 5A; Supplemental Fig. S2). This result suggested that inhibition of miR-191-5p sensitized breast cancer cells to chemotherapy. Similar results were obtained on using Anti-191 along with mitomycin C and paclitaxel, other chemotherapeutic drugs used for breast cancer treatment (Supplemental Fig. S3). To investigate whether the combination of Anti-191 with doxorubicin affects apoptosis in breast cancer cells, FACS analysis was done. The combination of Anti-191 with doxorubicin clearly increases the percentage of apoptotic cells (69.15%) as compared to its control (Fig. 5B,C). This was further confirmed by caspase 3/7 assay and DAPI-based microscopic analyses (Fig. 5D,E). These findings

FIGURE 5. The effects of miR-191-5p silencing on breast cancer chemotherapy. (A) MCF7 cells were treated with different concentrations of doxorubicin (free), or doxorubicin along with control oligo (NCtrl), or Anti-191. Subsequently, cell viability was evaluated by MTT assay. (B) FACS analysis using PE-Annexin V and 7AAD was performed in MCF7 cells treated with Anti-191 and doxorubicin. (C) Graph showing percentage of annexin V positive cells. (D) Caspase 3/7 activity was measured in MCF7 cells treated with Anti-191 and doxorubicin. (E) Microscopic images of DAPI-stained MCF7 cells treated with Anti-191 or its control and doxorubicin. (F) Graph shows percentage of apoptotic cells. The graphical data points represent mean $\pm$ SD of at least three independent experiments. (*) $P < 0.05$ and (**) $P < 0.01$. Error bars denote $\pm$ SD.

together demonstrate that inhibition of miR-191-5p promotes doxorubicin-induced apoptosis in breast cancer cells.

DISCUSSION

Apoptosis-mediated cancer cell death remains central to most treatment strategies. Thus, identification of miRNAs involved in the regulation of apoptosis holds immense importance. Here, we identify miR-191-5p as a negative regulator of apoptosis in breast cancer. We looked for miR-191-5p target genes to study its mechanism of action. We identified *BDNF*, *CDK6*, *CEBPB*, *SATB1*, *SOX4*, and *PLCD1* as direct miR-191-5p targets. We have previously reported *SATB1*, *CDK6*, and *BDNF* as miR-191-5p targets but have included these here for their reported link with breast cancer apoptosis. miR-191-5p regulation of their levels has also been independently confirmed here in other breast cancer cell lines ZR-75

and T47D (Fig. 3B; Supplemental Fig. S4). *BDNF* has been shown to inhibit apoptosis in breast cancer cell lines MDA-MB-231 and MCF7 (Yang et al. 2012). *SATB1* has also been shown to inhibit apoptosis and promote chemoresistance in MCF7 cells (Li et al. 2010). Similarly, high *CDK6* protects cells from fulvestrant-mediated apoptosis in breast cancer (Alves et al. 2016). *PLCD1* has been reported to function as a tumor suppressor in breast cancer that inhibits breast tumor cell formation in vivo by inducing apoptosis (Mu et al. 2015). *CEBPB* has been shown to promote tumorigenesis and its high levels are seen in breast cancer patients (Grimm and Rosen 2003). *SOX4* has also been shown to affect apoptosis in breast cancer (Song et al. 2015). Overall, aberrations in miR-191-5p levels significantly affect the levels of proteins known to affect apoptosis.

We particularly focused on *SOX4* because it has previously been shown to regulate the levels of p53, the master regulator

of apoptosis (Hur et al. 2010; Zhou et al. 2015). There have been conflicting reports on the effect of SOX4 on p53 levels. While some studies demonstrate that SOX4 brings about the degradation of p53 levels, others show that it rather promotes p53 stabilization (Pan et al. 2009). Here, we found that SOX4 overexpression in MCF7 cells brought about a significant increase in p53 protein levels. This is in line with Pan et al. (2009), who reported that SOX4 is a novel mediator for p53 stabilization and activation in response to DNA damage (Pan et al. 2009). In our study, results revealed that miR-191-5p-mediated targeted down-regulation of SOX4 brings about down-regulation of p53, which further leads to inhibition of apoptosis in breast cancer. In turn, silencing of miR-191-5p by using Anti-191 brings about up-regulation of SOX4 and p53, which triggers apoptosis in breast cancer. This suggests that SOX4 may contribute to p53 protein stabilization in breast cancer cells, and by targeting SOX4, miR-191-5p brings about down-regulation of p53 in breast cancer cells. Several other miRNAs too have been shown to be associated with apoptosis and p53 pathways in breast cancer (Feng et al. 2011; Sharma et al. 2016). While miR-504 directly targets p53, other miRNAs were shown to indirectly affect p53 signaling by targeting upstream regulators or downstream mediators of p53 to affect apoptosis in breast cancer (Feng et al. 2011; Sharma et al. 2016). p53 has also been shown to regulate several miRNAs (miR-143, miR-145, miR-15, miR-16, miR-26a, and miR-34) involved in apoptosis in breast cancer (Feng et al. 2011). We found that p53 brings about down-regulation of miR-191-5p levels through binding to specific p53 binding sites in the promoter of miR-191-5p. Thus there exists an interesting p53-miR-191-SOX4 feedback loop that affects apoptosis in breast cancer. The role of miR-191-5p in regulation of apoptosis has also been demonstrated in hepatic and gastric cancer, but the mechanism was not elucidated (Elyakim et al. 2010; Shi et al. 2011). miR-191-5p has also been shown to target *MDM4* in ovarian cancer and thus affect p53 levels (Wynendaele et al. 2010). Whether *MDM4* is a target of miR-191-5p in breast cancer remains to be evaluated. miR-191-5p has been shown to be regulated by other transcription factors such as estrogen receptor and hypoxia-inducible factor-1 (Di Leva et al. 2013; Nagpal et al. 2013, 2015). The mutual regulation between p53 and ER- α or p53 and HIF-1 or HIF-1 and ER- α has been demonstrated by various groups in breast cancer (Berger et al. 2013; Obacz et al. 2013). However, what effect this cross-talk may have on miR-191-5p levels in tumors and consequently its impact on tumor aggressiveness and treatment outcome needs to be thoroughly investigated.

Doxorubicin is a commonly used effective drug for the treatment of breast cancer patients. However, patients developing resistance to this drug is common, representing a major hindrance to successful treatment (Smith et al. 2006; Raguz and Yagüe 2008). Here, we found that pretreatment of breast cancer cells with Anti-miR-191 sensitizes the cells to doxorubicin treatment and decreases its IC₅₀ significantly.

One of the mechanisms by which Anti-miR-191 treatment promotes doxorubicin sensitivity is by increasing p53 levels, leading to increased apoptosis in the cells, as demonstrated here. However, whether Anti-miR-191 also affects doxorubicin-mediated disruption of topoisomerase II-mediated DNA repair and generation of free radicals remains to be studied (Thorn et al. 2011). Latorre et al. (2012) previously demonstrated that HuR, an RNA binding protein, promotes doxorubicin sensitivity in breast cancer cells. HuR has also been shown to be a miR-191-5p target in breast cancer (Nagpal et al. 2015). Thus, miR-191-5p-mediated down-regulation of HuR can also be one of the mechanisms of doxorubicin resistance in breast cancer cells. Other miRNAs, such as miR-21, miR-34a, miR-181a, miR-17/20, and miR-218, also have been shown to affect doxorubicin resistance in breast cancer. The target genes of these miRNAs are involved in cell signaling or apoptotic pathways (miR-21-*PTEN*, miR-34a-*NOTCH1*, miR-181a-*BCL2*, miR-17/20-*AKT1*, and miR-218-*SURVIVIN*) (Wang et al. 2011; Li et al. 2012; Zhu et al. 2013; Yu et al. 2014; Hu et al. 2015). Considering that resistance to hormone therapy or chemotherapy drugs apart from doxorubicin is also partly due to inhibition of p53-mediated apoptosis, the effect of Anti-miR-191 on sensitizing cells to other therapies needs to be checked.

In summary, our findings add new insight into the role of miR-191-5p in the regulation of apoptosis in breast cancer. We also show the existence of a regulatory negative feedback loop between miR-191-5p and p53 in breast cancer. Considering previous reports of miR-191-5p as part of hypoxia and estrogen receptor signaling, it emerges as a critical mediator of multiple oncogenic and tumor suppressor pathways that are known to be deregulated in breast cancer. Thus, Anti-miR-191 treatment may lead to simultaneous inhibition of HIF and ER signaling and promotion of p53 signaling, leading to a multipronged attack on breast cancer. This, combined with the fact that Anti-miR-191 also promotes sensitivity toward doxorubicin treatment, makes it an attractive target for breast cancer treatment.

MATERIALS AND METHODS

Cell culture

The breast cancer cell lines (MCF7, ZR-75, and T47D) were obtained from Cell Repository of NCCS, Pune. All the cells were maintained in RPMI 1640 (GIBCO) medium. The media was supplemented with 100 U/mL penicillin, 100 μ g/mL streptomycin, and 10% fetal bovine serum and incubated at 37°C in a 5% CO₂ incubator.

Transfections

Cells were seeded (5×10^5 cells per well) in six-well plates. Of note, 2.5 μ g of miR-191-5p plasmid (pc-191) or its control plasmid pcdna3.1 (pc) and 30nM anti-miR-191 (Anti-191) or the negative control (NCtrl) (Sigma-Aldrich) was transfected into six-well plates using lipofectamine 2000 (Invitrogen). After 5 h, media was

changed, and after 48 h post-transfection the samples were assayed for transcript levels through quantitative reverse-transcription-PCR (qRT-PCR) or used for other cellular assays. Each experiment was repeated three times.

qRT-PCR

Total RNA was isolated using an RNA Extraction Kit (Fermentas) and reverse transcribed using a cDNA Synthesis Kit (Bio-Rad). The cDNA formed was then further amplified for the predicted genes with respective primers sets using SsoFast EvaGreen Master Mix (Bio-Rad). For miR191-5p detection, a set of stem-loop primers were used (see Supplemental Fig. S5 for a list of primers). Glyceraldehyde 3-phosphate dehydrogenase (*GAPDH*) and RNU6B (*U6*) were used as controls for the normalization of the data. The expression fold change values were determined by $2^{-\Delta\Delta C_t}$ formula.

Target prediction

The probable targets for miR-191-5p were selected by established target prediction programs. Two target prediction programs, TargetScan and miRanda, were used to select the miR-191-5p targets. The candidate targets were selected on the basis of published literature demonstrating their association with apoptosis.

Construction of *SOX4* overexpression plasmid: pWPXL-*SOX4* was a gift from Bob Weinberg (Addgene plasmid # 36984). To create the pc-*SOX4* plasmid, a pWPXL-*SOX4* plasmid was digested with *NdeI* (Fermentas) and treated with Vent Polymerase (NEB) at 72°C for 15 min, then the plasmid was purified and digested with *BamHI*. Upon digestion with *BamHI*, the *SOX4* insert was released from the pWPXL plasmid vector and ligated into the pcDNA3.1 plasmid (digested with *BamHI* and *EcoRV* restriction enzymes [Fermentas]).

Construction of 3' untranslated region- or promoter-luciferase constructs

To determine whether miR-191-5p down-regulates its target transcripts through its direct binding to the 3' untranslated region (UTR), the region containing the miR-191-5p binding site in the 3'UTRs of the genes (*SOX4*, *PLCD1*, *CDK6*, *SATB1*, *CEBPB*, and *BDNF*) was amplified (using *Taq* polymerase; Fermentas) and cloned in the luciferase reporter vectors (pmiR-Report) downstream from a firefly luciferase gene (Supplemental Fig. S5). To further confirm whether the predicted target is the functional target of miR-191-5p, the miR-191-5p binding site in the 3'UTR of the gene (*SOX4*) was mutated through site-directed mutagenesis using inverse PCR and luciferase activity was observed (Supplemental Fig. S5). For promoter analysis, we amplified the miR-191-5p promoter fragments predicted to encompass p53 binding sites. The p53 binding sites upstream of miR-191-5p were cloned upstream of a luciferase gene in PGL3-tk-luciferase vector (Promega). All the clones were confirmed with PCR, restriction digestion, and sequence analysis.

Apoptosis assay

MCF7 cells were transiently transfected with pc-191 or anti-miR-191 (Anti-191) or respective control oligos (pc and NCtrl), and

48 h post-transfection apoptosis was assayed by fluorescence-activated cell sorting (BD Bioscience) using a PE Annexin V Staining Kit (BD Bioscience). The data obtained were analyzed using Cyflogic software. The experiment was repeated three times.

Western blot

MCF7 cells were transfected with miR-191-5p or *SOX4* along with their controls and lysed using protein lysis buffer (Tris-50 mM, EDTA-5 mM, NaCl-150 mM and 1% Triton X-100 with 1 mM β -mercaptoethanol and protease inhibitor cocktail) 48 h post-transfection. The protein concentration was determined by using Bradford reagent (Sigma-Aldrich). An equal amount of protein lysates was separated with 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis and transferred to nitrocellulose membrane (Bio-Rad). The membrane was then probed with a specific primary antibody at a dilution of 1:1000 (p53; Santa Cruz Biotechnology) or 1:3000 (β -actin; Santa Cruz Biotechnology) or 1:1000 (*SOX4*; Santa Cruz Biotechnology) followed by washing and incubation with the respective secondary antibody (p53 and β -actin-anti-mouse, horseradish peroxidase-linked, Santa Cruz Biotechnology; *SOX4*-anti-goat horseradish peroxidase-linked). The specific protein band was visualized by autoradiography using an ECL kit (Bio-Rad).

Chromatin immunoprecipitation

MCF7 cells were cultured in 100-mm culture dishes at 90%–100% confluency; then cells were harvested and crosslinked with formaldehyde, washed with phosphate-buffered saline, and resuspended in the RIPA buffer. Sonication was performed to obtain DNA fragment of 100- to 500-bp fragments (sonicated lysate was confirmed for fragment size through 2% agarose gel). For chromatin immunoprecipitation (CHIP), specific antibodies against p53 (Santa Cruz Biotechnology), salmon sperm DNA (Sigma-Aldrich), and protein A-sepharose beads (30 μ L) were added to the chromatin extract and incubated overnight, washed and eluted with 0.5% w/v sodium dodecyl sulfate solution. Decrosslinking was done at 65°C for 4–5 h and DNA was then purified with a PCR Purification Kit (HiMedia). Rabbit IgG was used as control antibody, whereas the chromatin extract without any antibody/beads treatment was used as positive control. For DNA sequence-specific quantification, qRT-PCR was done with an equal amount of chromatin extract using sequence-specific primers (Supplemental Fig. S5). The experiment was repeated twice.

Cell viability assay

Cell viability was determined by 3-(4,5-dimethylthiazole-2-yl)-2,5-diphenyl tetrazolium bromide (MTT). MCF7 and ZR-75 cells were seeded in a 96-well plate containing 100 μ L of culture medium, and then MTT reagent was added at different time points. Subsequently, the plate was incubated for 2 h at 37°C in a humidified incubator containing 5% CO₂ and quantified at 595 nm. The experiments were repeated three times.

Caspase-Glo 3/7 assay

Caspase-Glo 3/7 activity was determined by a Caspase-Glo 3/7 Assay Kit (Promega). Cells were seeded in a 96-well plate and transfections

were done with miR-191-5p or its control. Forty-eight hours post-transfection, 50 μ L of Caspase-Glo 3/7 reagent was added to the medium and the mixture was incubated at room temperature for 2 h, after which the luminescence was analyzed in a luminometer (Berthold).

DAPI staining

MCF7 cells were seeded in a 12-well plate and transfections were done with miR-191-5p or its control. Forty-eight hours post-transfection, cells were fixed with 4% paraformaldehyde for 20 min at room temperature. Fixed cells were washed with 1 $\times$ PBS and permeabilized with 0.2% Triton X-100 for 10 min and again washed with PBS. Cells were stained with DAPI (1:1000, Sigma-Aldrich) and incubated in the dark for 20 min at room temperature. Cells were washed with PBS and analyzed under a fluorescent microscope.

SUPPLEMENTAL MATERIAL

Supplemental material is available for this article.

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Author contributions: R.K. conceptualized and supervised the whole study. P.C.G. supervised and helped with the analyses of the FACS data. S.S. performed all the experiments. N.N. made some constructs for the luciferase assays.

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