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# Estradiol-regulated microRNAs control estradiol response in breast cancer cells

Poornima Bhat-Nakshatri<sup>1</sup>, Guohua Wang<sup>2</sup>, Nikail R. Collins<sup>1</sup>, Michael J. Thomson<sup>3</sup>, Tim R. Geistlinger<sup>4</sup>, Jason S. Carroll<sup>4</sup>, Myles Brown<sup>4</sup>, Scott Hammond<sup>3</sup>, Edward F. Srouf<sup>2</sup>, Yunlong Liu<sup>2</sup> and Harikrishna Nakshatri<sup>1,5,\*</sup>

<sup>1</sup>Department of Surgery, <sup>2</sup>Department of Medicine, Indiana University School of Medicine, Indianapolis, IN, <sup>3</sup>Department of Cell and Developmental Biology, Lineberger Comprehensive Cancer Center, University of North Carolina, Chapel Hill, NC, <sup>4</sup>Department of Medical Oncology, Dana-Farber Medical School, Harvard Medical School, Boston, MA and <sup>5</sup>Department of Biochemistry and Molecular Biology, Indiana University School of Medicine, Indianapolis, IN, USA

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## ABSTRACT

**Estradiol (E2) regulates gene expression at the transcriptional level by functioning as a ligand for estrogen receptor alpha (ER $\alpha$ ) and estrogen receptor beta (ER $\beta$ ). E2-inducible proteins c-Myc and E2Fs are required for optimal ER $\alpha$  activity and secondary estrogen responses, respectively. We show that E2 induces 21 microRNAs and represses seven microRNAs in MCF-7 breast cancer cells; these microRNAs have the potential to control 420 E2-regulated and 757 non-E2-regulated mRNAs at the post-transcriptional level. The serine/threonine kinase, AKT, alters E2-regulated expression of microRNAs. E2 induced the expression of eight Let-7 family members, miR-98 and miR-21 microRNAs; these microRNAs reduced the levels of c-Myc and E2F2 proteins. Dicer, a ribonuclease III enzyme required for microRNA processing, is also an E2-inducible gene. Several E2-regulated microRNA genes are associated with ER $\alpha$ -binding sites or located in the intragenic region of estrogen-regulated genes. We propose that the clinical course of ER $\alpha$ -positive breast cancers is dependent on the balance between E2-regulated tumor-suppressor microRNAs and oncogenic microRNAs. Additionally, our studies reveal a negative-regulatory loop controlling E2 response through microRNAs as well as differences in E2-induced transcriptome and proteome.**

## INTRODUCTION

Estradiol (E2) controls several biological processes by functioning as a ligand for nuclear receptors estrogen receptor alpha (ER $\alpha$ ) and beta (ER $\beta$ ) (1). ERs may participate in the genomic (transcriptional) and non-genomic actions of E2 (1,2). The genomic action involves binding of ER $\alpha$  to the regulatory regions of target genes either directly or through protein–protein interaction. DNA-bound ER $\alpha$  then recruits various co-regulatory molecules to induce chromatin modifications that either increase or decrease the level of target gene transcription. Several extracellular signal activated kinases phosphorylate and modulate ER $\alpha$  activity; this may be responsible for altered E2 responses such as resistance to anti-estrogen therapy in breast cancer (2). The major kinases that modulate ER $\alpha$  activity include ERK1/2, AKT, RSK, PAK1, p38 kinase and SRC (2–4).

A significant number of studies so far have focused on E2:ER $\alpha$ -mediated transcriptome. The effect of E2 on gene expression at post-transcriptional level is yet to be elucidated. MicroRNAs are a class of evolutionarily conserved non-coding RNAs that control gene expression at the post-transcriptional level (5,6). They regulate gene expression through either direct cleavage of the target mRNAs or by translational inhibition (5,7–9). However, control of gene expression at the level of translation by microRNAs is cell cycle dependent (10). A highly conserved microRNA family is predicted to target 300 mRNA species and together affect ~30% of the protein-coding genes (11,12). Cellular stress and RNA-binding proteins, which usually bind to sequences in between microRNA-recognition sites

\*To whom correspondence should be addressed. Tel: +1 317 278 2238; Fax: +1 317 274 0396; Email: hnakshat@iupui.edu

on mRNAs, determine the target specificity of microRNAs (13,14). Additional functions of microRNAs include transcriptional activation of genes with complementary promoter sequences (15) and chromatin modification (16,17).

RNA polymerase II enzyme transcribes the majority of microRNA genes (a minority by polymerase III) to produce a primary microRNA (18,19). Approximately 50% of microRNAs are transcribed from introns of protein-coding genes while the rest are intergenic with primary transcripts as long as 4 kb (19). MicroRNA genes usually appear in polycistronic clusters and more than 50% are located in cancer associated genomic regions at fragile sites (5,20). Thus, an alteration in the expression of a single microRNA can have a profound effect on cellular physiology because of their enormous influence on the expression of multiple genes.

Three recent reports describe microRNA-expression patterns in breast cancer; two of these have evaluated the expression pattern in relation to ER $\alpha$ , ErbB2 and intrinsic subtypes (21–23). Overall, microRNA levels were lower in poorly differentiated tumors compared to well-differentiated tumors and microRNA profile rather than mRNA-expression profile correlated more accurately with cell differentiation. ER $\alpha$  positive breast cancers displayed a distinct microRNA-expression profile including elevated expression of Let-7 family members of microRNAs and miR-21 compared to ER-negative breast cancers (21). Blenkiron *et al.* described microRNA-expression profiles in relation to five intrinsic subtypes (luminal A, luminal B, normal-like, HER2+ and basal) of breast cancer (22,24,25). Luminal subtypes and basal type breast cancers display distinct microRNA-expression patterns (22). ER $\alpha$  expression and activity in breast cancer may be regulated by a Estrogen-ER $\alpha$ -microRNA-regulatory loop as mir-206 inhibits ER $\alpha$  translation by binding to the 3' UTR of ER $\alpha$  mRNA, whereas ER $\alpha$  agonists block miR-206 expression (26).

Despite enormous progress in understanding the expression pattern of microRNAs in breast cancer, regulation of their expression in different breast cancer subtypes is not known. We focused on the effect of E2 on microRNA expression in breast cancer cells with two goals: (i) to determine whether E2 regulates the expression of any of the microRNAs expressed at higher levels in luminal type A/ER $\alpha$ -positive breast cancer; and (ii) to determine whether these E2-regulated microRNAs subsequently control the expression of E2-regulated genes at the post-transcriptional level. We also examined whether extracellular signal activated kinases such as AKT modulate E2-regulated microRNA expression. Results from this study reveal a unique role for E2 in regulating the expression of luminal type A-enriched microRNAs with tumor suppressor and oncogenic functions and the ability of AKT to alter E2-regulated microRNA expression.

## MATERIALS AND METHODS

### Cell lines

MCF-7 cells containing the bicistronic vector control (MCF-7p) or the constitutively active AKT

(MCF-7AKT) were generated by retrovirus-mediated gene transfer and have been described previously (27). Virus infected cells were selected using one microgram per ml puromycin and mass culture instead of selected clones were used to avoid clonal bias. The constitutively active AKT1 construct with deletion of N-terminal pleckstrin homology domain (amino acids 4–129) but addition of myristylation signal has been described previously (3). Cells were maintained in MEM plus 10% fetal calf serum and one microgram per ml of puromycin. Cells were switched to phenol red free MEM plus 5% dextran-charcoal treated serum (CCS) for at least 4 days prior to experiments.

### MicroRNA-expression analysis

Cells were treated with the solvent ethanol or 10<sup>-8</sup> M E2 for 4 h and RNA was prepared using the mirVana<sup>TM</sup> microRNA isolation kit (Ambion Inc., Austin, TX). MicroRNA microarray and hybridization to microRNA arrays have been described previously (28,29). Assays were done in duplicate with RNA from two independent clones. Since all the probes were printed twice on the microarray in two independent blocks, four measurements were available for every microRNA under each experimental condition. Signal intensity of tRNA-Thr was used for normalization between samples and relative differences in individual microRNA expression with or without E2 treatment was then calculated. Differentially expressed microRNAs were identified using a linear mixed effect model that describes the relationship among three factors, including cell type, E2 treatment, probe blocks and interactions between cell type and E2 treatment. The linear mixed effect model can be described as:

$$y_{i,j,k,l,n} = \mu_i + \alpha_{ij} + \beta_{ik} + \gamma_{ijk} + a_{il} + b_{ln} + \varepsilon_{ijkln}$$

where  $i, j, k, l, n$  represent the the microRNA, the cell types (MCF-7p or MCF-7AKT), the conditions (with and without E2 treatment), the RNA samples and block on the array, respectively.  $\alpha_{ij}$  and  $\beta_{ik}$  represent the effect of cell types and conditions, and  $\gamma_{ijk}$  denotes the interaction of these two factors. The RNA sample and block are considered as random effects, and the RNA sample factor is nested in the cell type and condition factors.

E2-induced changes in Let-7f, miR-98 and miR-21 was verified in four independent RNA preparations by quantitative reverse transcription polymerase chain reaction (qRT-PCR) using The TaqMan<sup>®</sup> microRNA assays designed to detect and accurately quantify mature microRNAs (Applied Biosystems, Foster City, CA). Primers specific to small RNA, RNU66, were used for normalization (Applied Biosciences). Target genes of differentially expressed microRNAs were predicted using TargetScan (12).

### Locked nucleic acid (LNA) treatment and western blot analysis

LNA against Let-7f, miR-98 and miR-21 as well as control LNA were obtained from Exiqon, Inc (Woburn, MA). LNA (5 nM) was transfected into cells grown in 60 mm plates using Lipofectamine reagent (Invitrogen,

Carlsbad, CA). After 2 days, cells were split into two 60-mm plates. E2 or ethanol was added to cells 24 h after splitting and harvested for western analysis 24 h post E2 treatment. All experiments were done three to five times with similar results. Antibodies against AIB1 (BD biosciences, San Jose, CA), E2F1, E2F2, Maspin (Santa Cruz Biotechnology, Santa Cruz, CA), c-Myc (Upstate Biotechnology, Lake Placid, NY), EZH2 (Cell Signaling, Danvers, MA) and PDCD4 (Abcom, Cambridge, MA) were used for western analysis as per instructions of manufacturers.

#### **Total RNA isolation, northern blotting and quantitative reverse transcription polymerase chain reaction (qRT-PCR)**

Total RNA from ethanol and E2-treated cells was isolated using an RNeasy kit from Qiagen (Valencia, CA). Northern blotting was performed as described previously (3). RNA was reverse transcribed using a single-stranded cDNA synthesis kit (Invitrogen, Carlsbad, CA) and subjected to quantitative polymerase chain reaction (Q-PCR) using Syber green (Applied Biosciences, Foster City, CA). Primers used were: Dicer forward primer TCACCTGCCTCACTTGACCTGAAA; Dicer reverse CGCTTTCAAAC TGCTGCTCAT; beta-actin forward TGGATCAGCAA GCAG; and beta-actin reverse GCATTTGCGGTGGAC.

#### **Chromatin immunoprecipitation assay**

The entire ER $\alpha$  chromatin immunoprecipitation-microarray (ChIP-on-chip) data set of MCF-7p and MCF-7AKT cells as well as statistical analysis has been described (27). ER $\alpha$  ChIP assay was performed as described previously (30). ChIP DNA was subjected to Q-PCR using the primers 5'-TCTTTGACCCACTAGCCTTGACAGT-3' and 5'-TCACTTCTCTCTGGGCCACAGTTT-3' to determine E2-inducible changes in ER $\alpha$  binding to the central-binding site close to miR-21 identified in ChIP-on-chip. The chromosomal location of ER $\alpha$  binding was determined using UCSC genome browser ([www.genome.ucsc.edu](http://www.genome.ucsc.edu)). E2-inducible changes in ER $\alpha$  binding were not detected with ChIP DNA obtained by immunoprecipitation with control IgG.

#### **Transient transfection and luciferase assay**

The 3' untranslated regions of AIB1 (nucleotides 6872–7145 of NM\_181659) or c-Myc (nucleotides 1861–2360 of NM\_002467) were cloned into a cytomegalovirus (CMV) enhancer-promoter driven luciferase gene at the 3'-end of the coding sequence. MCF-7 cells maintained in MEM plus 5% CCS were transfected with the above plasmid and the internal control plasmid renilla luciferase using lipofectamine (Invitrogen). E2 was added 24 h after transfection and luciferase assay was performed 16 h after E2 addition using dual luciferase assay kit from Promega (Madison, WI).

## **RESULTS**

### **E2 regulates the expression of microRNAs**

We recently reported the ER $\alpha$ -binding pattern in MCF-7 cells with a retrovirus vector control (MCF-7p) and

MCF-7 cells overexpressing constitutively active AKT1 (MCF-7AKT) (27). We observed ER $\alpha$ -binding sites associated with 1667 and 1908 genes in one-hour E2-treated MCF-7p and MCF-7AKT cells, respectively. However, not all of the genes with ER $\alpha$ -binding sites had an altered transcript level after E2 treatment. Also, not all E2-regulated genes, as identified by microarray analysis of both cell types with or without E2 treatment for 4 h, contained ER $\alpha$ -binding sites suggesting that there are other mechanisms by which E2 regulates gene expression. In fact, among 833 E2-regulated genes in MCF-7p cells, only 299 genes contained ER $\alpha$ -binding sites (see Supplementary Data for select genes). Although secondary estrogen response through E2-regulated transcription factors such as E2F1/2 can explain the lack of correlation between ER $\alpha$  binding and target gene expression (31), control of gene expression by E2-regulated microRNAs has not been investigated in detail. To investigate this possibility, we determined microRNA-expression patterns in MCF-7p and MCF-7AKT cells with and without E2 treatment for 4 h. We observed 21 E2-inducible and 7 E2-repressible microRNAs in MCF-7p cells (statistical cutoff *P*-value <0.05 and fold change >1.5 or <0.7) (Table 1).

E2 increased the expression of eight members of the Let-7 family microRNAs ( $\geq 2.2$ -fold increase), which previously have been shown to be overexpressed in luminal type A breast cancer (21,22). Similarly, E2 increased the expression of miR-21, which is also expressed at a higher level in luminal type-A breast cancer. AKT completely changed the pattern of E2 regulation of microRNA expression. In MCF-7AKT cells, E2 increased the expression of only one microRNA but reduced the expression of 20 microRNA species. Basal expression of 11 microRNAs was lower in MCF-7AKT cells compared to MCF-7p cells; whereas, one microRNA displayed elevated expression in MCF-7AKT cells. Seven of the microRNAs that displayed lower basal expression in MCF-7AKT cells were E2-inducible in MCF-7p cells. Only three microRNAs (miR-143, miR-506 and miR-98) showed similar E2-dependent regulation in both cell types.

We performed microRNA qRT-PCR, which quantitatively measures only mature microRNAs, to confirm E2-inducible expression of Let7f, miR-21 and miR-98 (Figure 1A). Consistent with the microarray results, E2 increased the expression of these microRNAs in MCF-7p cells; none of the E2-induced changes in microRNA expression in MCF-7AKT cells was statistically significant. E2 increased the expression of all three of these microRNAs in parental MCF-7 cells that were used to generate the above cell lines (Figure 1B, top panel). E2 similarly increased Let-7f, miR-98 and miR-21 levels in T47-D cells, another ER $\alpha$ -positive cell line (Figure 1B, middle panel). Interestingly, in BT-474, an ER $\alpha$ -positive/HER2-positive cell line, E2-inducible expression of these microRNAs was modest (Figure 1B, bottom panel).

To determine the requirement of ER $\alpha$  for E2-regulated expression of the above microRNAs, we pretreated MCF-7 cells with fulvestrant for 24 h and then stimulated cells with E2 for 4 h. Fulvestrant inhibits ER $\alpha$  function through degradation. The basal levels of miR-98 and miR-21 microRNAs were elevated in fulvestrant-treated

**Table 1.** The effect of E2 on microRNA expression in MCF-7p and MCF-7AKT cells

MCF-7p cells		MCF-7AKT cells	
Name	Fold change	Name	Fold change
<b>E2-inducible microRNAs</b>			
miR-Let-7f	3.2	miR-520d	4.8
miR-Let-7a	2.9		
miR-Let-7d	2.7		
miR-Let-7c	2.6		
mir-Let-7g	2.6		
miR-203	2.4		
miR-Let-7b	2.3		
miR-Let-7e	2.2		
miR-98	2.1		
miR-21	2.0		
miR-200a	2.0		
miR-103	1.8		
miR-200c	1.8		
miR-107	1.7		
miR-17-5p	1.7		
miR-23a	1.6		
miR-200b	1.6		
miR-30c	1.6		
miR-30b	1.5		
miR-424	1.5		
miR-let7i	1.5		
<b>E2-repressible microRNAs</b>			
miR-302b*	0.4	miR-524	0.3
miR-506	0.5	miR-518d	0.3
miR-524*	0.5	miR-518e	0.3
miR-27a	0.6	miR-506	0.3
miR-27b	0.6	miR-409-5p	0.3
miR-143	0.6	miR-216	0.3
miR-9	0.7	miR-518c*	0.5
		miR-526b	0.5
		miR-34b	0.5
		miR-337	0.5
		miR-146	0.5
		miR-128b	0.5
		miR-124a	0.5
		miR-211	0.6
		miR-143	0.6
		miR-128a	0.6
		miR-126*	0.6
		miR-126	0.6
		miR-1	0.6
		miR-10b	0.7
<b>MicroRNAs that display differences in basal expression between two cell types<sup>a</sup></b>			
miR-520d	5.1	miR-200a	0.6
miR-let7g	0.5	miR-182	0.6
miR-337	0.5	miR-17-5p	0.6
miR-20b	0.5	miR-1	0.6
miR-Let-7i	0.6	miR-203	0.6
miR-98	0.6	miR-200c	0.7

Cells were treated with E2 for 4h and microRNA expression array analysis was performed. Only those microRNA whose expression differences with a *P*-value of <0.05 among untreated and E2-treated cells are presented. miR-98 was induced by E2 in both cell types (2.1-fold in MCF-7p cells, 1.4-fold in MCF-7AKT cells) whereas miR-143 (to same level) and miR-506 (0.50 in MCF-7p cells and 0.30 in MCF-7AKT cells) were repressed by E2 in both cell types.

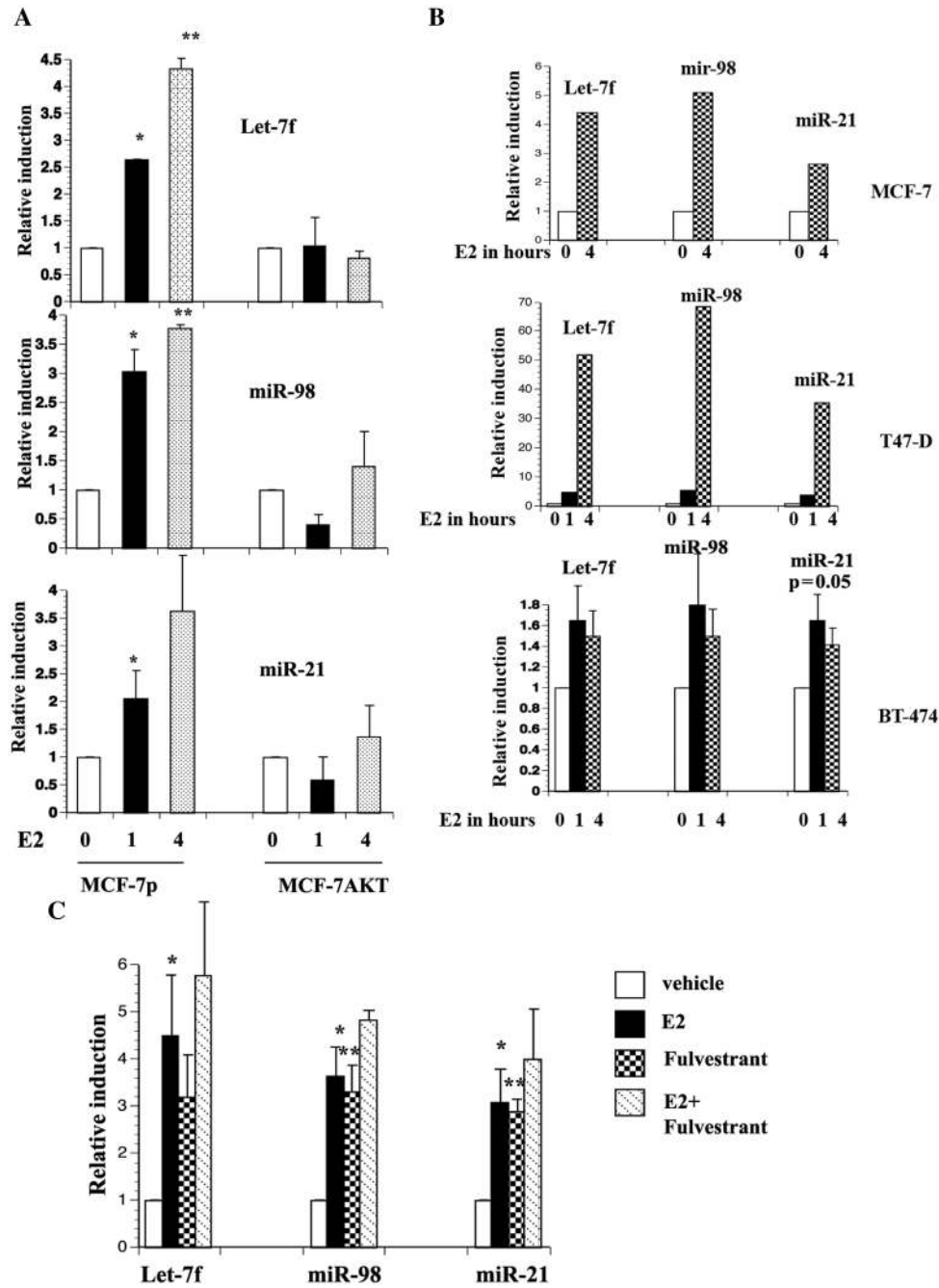
<sup>a</sup>MicroRNA with fold change >1 is expressed at higher levels in MCF-7AKT cells, whereas fold change <1 represent microRNAs expressed at higher levels in MCF-7p cells.

cells and were only marginally increased upon E2-treatment (Figure 1C). The basal level of Let-7f was marginally increased but it did not reach statistical significance. These results suggest that the unliganded ER $\alpha$  represses the expression of these microRNAs and E2 relieves this repression, which is consistent with a recent report on nuclear receptor-mediated microRNA regulation in *Caenorhabditis elegans* (32).

### Putative targets of E2-regulated microRNAs

E2-inducible expression of Let-7 family members and reduced basal and/or E2-inducible expression of this family in MCF-7AKT cells is interesting because reduced expression of this family is linked to loss of differentiation and increased self-renewal of progenitor cells (33,34). Both Let-7 family members and miR-98 have the same seed sequence and target the same mRNAs. Ras family oncogenes and HMGA2 are the well-characterized targets of Let-7 (35,36). c-Myc is also a target of Let-7 (37) and miR-21 (38). Based on TargetScan and miRGen analyses, E2F1 and E2F2, E2-inducible transcription factors involved in secondary estrogen responses (31), are predicted targets of miR-205 and Let7/miR-98, respectively. E2F1 and E2F2 are also targets of miR20 (39); mir-20b is expressed at a higher level in MCF-7p cells compared to MCF-7AKT cells (Table 1). NCOA3/AIB1 is a target of miR-17-5p (40), which in our studies is E2-inducible microRNA (Table 1). EZH2 is a multifunctional protein that integrates Wnt and E2 signaling in breast cancer cells and the abnormal function of this protein is linked to several diseases (41). Based on TargetScan analysis, EZH2 is a likely target of mir-124a and mir-506, both of which are repressed by E2. The above genes thus constituted potential targets of E2-regulated microRNAs.

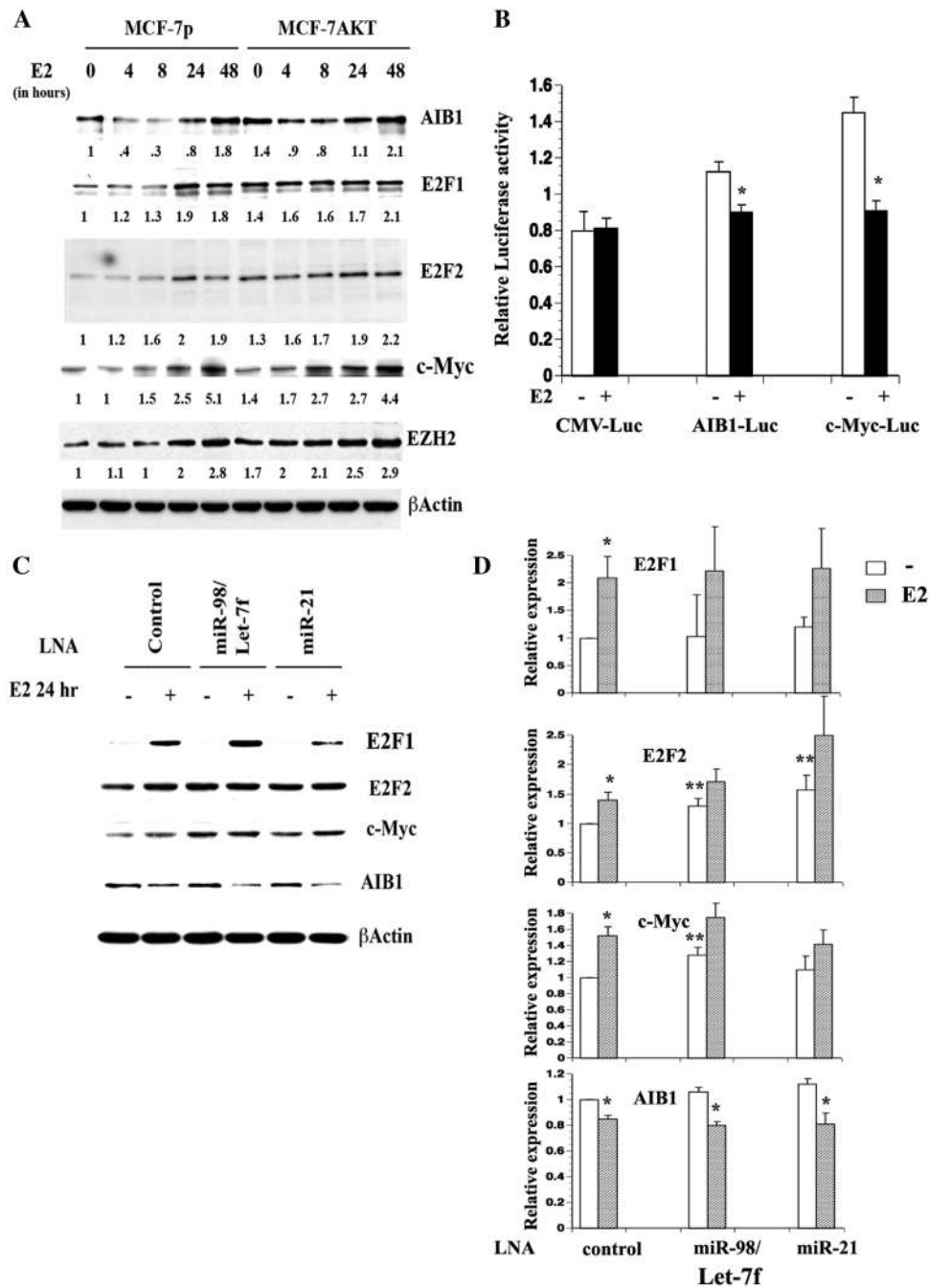
We focused our studies only to those genes that are involved in E2-signaling pathways and/or regulated by E2 at the transcriptional level and additionally controlled at the post-transcriptional level by E2-regulated microRNAs as part of an autoregulatory loop (Figure 2A). We observed repression of AIB1 and induction of EZH2 by E2 in both cell types. Additionally, while E2F1 and E2F2 displayed a delayed E2-inducible increase in protein levels in MCF-7p cells, basal levels of both of these proteins were elevated with a further E2-inducible increase in MCF-7AKT cells. Elevated basal levels of these two proteins in MCF-7AKT cells correlate with a lower level of miR20b in these cells compared to MCF-7p cells (Table 1). Previously, we reported little difference in the kinetics of E2-inducible expression of c-Myc mRNA in parental and CA-AKT overexpressing MCF-7 cells (3). However, at the protein level, E2-mediated increase in c-Myc is advanced in MCF-7AKT cells compared to MCF-7p cells (Figure 2A). None of the previous microarray profiling studies including our study observed an effect of E2 on EZH2 mRNA level and our ChIP-chip data did not identify ER $\alpha$ -binding sites associated with the *EZH2* gene. Furthermore, RT-PCR analysis has confirmed the lack of an E2 effect on the mRNA levels of EZH2 (data not shown). Thus, E2 may control translation of AIB1 and EZH2 through microRNAs.



**Figure 1.** (A) E2-inducible expression of Let-7f, miR98 and miR-21 in MCF-7p and MCF-7AKT cells. MCF-7p and MCF-7AKT cells were treated with E2 ( $10^{-8}$ M) for indicated time and microRNA was subjected qRT-PCR. RNU66, small RNAs encoded in the intron of RPL5 gene (chr1:93 018 360-93 018 429) was used for normalization between samples. Mean plus standard error of the mean (SEM) are shown. \* $P < 0.03$ , ethanol versus 1-h E2-treated cells; \*\* $P < 0.001$  ethanol versus 4 h E2-treated cells. (B) The effect of E2 on the expression of Let-7f, miR-98 and miR-21 in MCF-7 (top), T47-D (middle) and BT-474 (bottom) cells. (C) The effect of fulvestrant treatment on basal and E2-regulated expression of Let-7f, miR-98 and miR-21. Cells were pretreated with 100 nM fulvestrant for 24 h and then treated with E2 for 4 h. \* $P < 0.05$ , ethanol versus E2-treated; \*\* $P < 0.05$ , ethanol versus fulvestrant treated cells.

The 3' UTRs of mRNAs are considered to be the primary targets of microRNAs although recent studies indicate the coding sequences are also targeted (8,9). We cloned the predicted microRNA target sequences in the 3' UTR of AIB1 and c-Myc into a CMV-driven luciferase-reporter vector and investigated the effects of E2 on luciferase levels (Figure 2B). E2 did not have an effect on luciferase expression in cells transfected

with parental vector. In contrast, E2-reduced luciferase expression in cells transfected with vectors containing either the AIB1 or c-Myc 3' UTR. These results suggest that the 3' UTR of AIB1 and c-Myc genes contain sequences that are targeted by E2-regulated microRNAs. Note that based on CHIP-on-chip data, cloned 3'-UTR sequences of both AIB1 and c-Myc lack ER $\alpha$ -binding sites (27).



**Figure 2.** Putative targets of E2-induced microRNAs. (A) MCF-7p and MCF-7AKT cells were treated with E2 for indicated time and western blotting was performed. (B) 3'-UTR of AIB1 or c-Myc reduces E2-inducible expression of a luciferase reporter under the control of CMV enhancer-promoter. MCF-7 cells were transfected with indicated reporters along with the internal control Renilla-luciferase under TK promoter-enhancer and luciferase activity was measured in untreated and E2-treated cells. \* $P < 0.05$ , ethanol treated versus E2-treated cells. (C) LNA against Let-7f/miR-98 and miR-21 differentially affect basal and E2-inducible levels of c-Myc and E2F2 proteins in MCF-7p cells. Cells were treated with control or specific LNA and treated with E2 for 24h. Western blot analysis was done with indicated antibodies. (D) Expression levels of E2F-1, E2F-2, c-Myc and AIB1 from three to five independent experiments, as measured by densitometric scanning (Mean plus SEM), are indicated. \* $P < 0.05$ , ethanol versus E2-treated cells; \*\* $P < 0.05$ , control LNA treated versus Let-7f/miR-98 or miR-21 LNA-treated cells. Note the effects of LNA against Let-7f/miR98 and miR-21 on E2F2 and c-Myc but not on E2F1 and AIB1 protein levels.

**Let-7f/miR-98 and miR-21 may be part of negative-regulatory loop that control E2-induced levels of c-Myc, E2F1 and E2F2 proteins**

Our previous studies have shown E2-mediated increase in mRNA levels of c-Myc, E2F1 and E2F2 in MCF-7p and

MCF-7AKT cells (3,27). We used LNA-mediated knock-down of Let-7f/miR98 to determine whether reducing the levels of these microRNAs lead to changes in E2-inducible expression of c-Myc, E2F1 and E2F2 at the protein level. Neither LNA against Let-7f/miR-98 nor LNA against

miR-21 had any effect on E2-inducible expression of E2F1 (Figure 2C and D). Consistent with TargetScan prediction, LNA against Let7f/miR-98 increased E2-inducible levels of E2F2. Surprisingly, basal level of E2F2 was also increased in miR-21 LNA treated MCF-7p cells compared to control LNA-treated cells. LNA against Let-7f/miR-98 but not miR-21 increased the basal levels of c-Myc proteins. LNA against Let-7f or miR-98 individually was ineffective, possibly due to their redundant targets (data not shown). E2 mediated reduction of AIB1 protein levels was unaffected by LNA against Let-7f/miR-98 or miR-21 (Figure 2C and D). Thus, E2-induced Let7f/miR-98 controls the expression levels of expected targets (E2F2 and c-Myc), whereas miR-21 unexpectedly controls the expression of E2F2. Also, the overall protein level changes (1.5–2-fold) that we observed in cells transfected with LNA-targeting specific microRNAs are within the range predicted and verified by proteomic approaches (11,42).

#### Bioinformatics analysis of potential targets of E2-regulated microRNAs identify additional E2-responsive genes

We then investigated whether any luminal type A specific genes expressed in primary breast cancers are putative targets of E2-regulated microRNAs using available bioinformatics tools. We used TargetScan among several available search engines because this program is more robust with regards to its prediction algorithm and several of its predictions have been verified by proteomics (11,42). Indeed, major genes that define a luminal type A phenotype or are overexpressed in luminal type A breast cancers including FOXA1, GATA-3, ER $\alpha$  and MyB are putative targets of E2-regulated microRNAs (Supplementary Table S1).

We next determined whether any of the E2-regulated mRNAs are the targets of E2-regulated microRNAs. The 21 E2-induced microRNAs in MCF7p cells belong to 10 distinct microRNA families. We analyzed whether these microRNAs-target genes that are induced or repressed by E2 in MCF-7 cells. Results of microarray analysis of untreated and 4 h E2-treated MCF-7p and MCF-7AKT cells were used for this purpose (27). E2-induced microRNAs are predicted to target 94 E2-inducible, 142 E2-repressible and 541 non-E2-regulated mRNAs in MCF-7 cells. Detailed statistical analysis is presented in Table 2. Supplementary Table S2 contains name of these genes, E2-dependent changes in their mRNA levels after 4 h of E2 treatment in both cell types, and microRNAs that may target them. MicroRNAs repressed by E2,

which belongs to four families (hsa-miR-302b\* and hsa-miR-524\* are excluded from this analysis due to the lack of prediction from TargetScan), are predicted to target 67 E2-inducible, 116 repressed and 407 non-E2-regulated genes. Supplementary Table S2 contains names and E2-regulated expression of these genes. Taken together, E2-regulated microRNAs potentially target 1021 genes in MCF-7p cells. Similar analysis in MCF-7AKT cells showed that microRNAs upregulated by E2 in these cells may target 20 E2-inducible genes, 20 E2-repressed genes and 68 non-E2-regulated genes. MicroRNAs repressed by E2 in these cells may target 148 E2-inducible genes, 111 E2-repressed genes and 554 non-E2 target genes (Table 2 and Supplementary Table S3). Overall, E2-regulated microRNAs potentially control 850 genes in MCF-7AKT cells.

#### miR-21-regulatory region is associated with ER $\alpha$ -binding sites

We used our ChIP-on-chip data set (27) and other bioinformatics tools to investigate how E2 regulates the expression of microRNAs. MicroRNAs can be transcribed as independent units or as parts of intronic sequences in large transcribed genes (19). MicroRNAs that reside in the intergenic sequence are transcribed as 3–5-kb transcripts with clearly defined 5' and 3' boundaries. The promoter/enhancer regions of such microRNA genes are enriched for five transcription factor-binding sites: MSX-1, TLX2 (Hox11L1), CDC5, SRF and ZNF238 (19). Table 3 provides summary of E2-regulated microRNAs that are associated with ER $\alpha$ -binding sites in their regulatory region or located in the intronic region of known E2-regulated genes. An ER $\alpha$ -binding site was assigned to a microRNA if it is located within 20 kb of the 5' transcription start site or within 20-kb of the 3' transcription termination site (27). Additional details of ER $\alpha$ -binding site distribution around microRNAs that are regulated by E2 in MCF-7p and MCF-7AKT cells or ER $\alpha$ -binding sites along with E2-regulated expression of host genes that harbor these microRNAs are provided as supplementary files (Supplementary Table S4 and Supplementary Table S5).

We further evaluated ER $\alpha$  binding to miR-21-regulatory regions by ChIP assay. Location of the miR-21 transcribed region in relation to ER $\alpha$ -binding sites and results of the ChIP assay confirming ER $\alpha$  binding are shown in Figure 3. Unliganded ER $\alpha$  bound to three regions in MCF-7p cells (Figure 3A). E2 reduced

**Table 2.** Number of genes potentially targeted by E2-regulated microRNAs in MCF-7p and MCF-7AKT cells

	mRNAs induced by E2 (FC > 1.5)	mRNAs repressed by E2 (FC $\leq$ 1.5)	Non-E2-regulated mRNAs ( $-1.5 < \text{FC} < 1.5$ )
E2 upregulated microRNA targets in MCF-7p cells	94	142	541
E2 downregulated microRNA targets in MCF-7p cells	67	116	407
E2 upregulated microRNA targets in MCF-7AKT cells	20	20	68
E2 downregulated microRNA targets in MCF-7AKT cells	148	111	554

Cellular mRNAs are subclassified into E2-induced, E2-repressed and unaffected by E2 but expressed in MCF-7p and MCF-7AKT cells based microarray analysis of ethanol or E2 (4-h) treated cells (27). FC = Fold change.



**Table 3.** E2-regulated microRNAs associated with ER $\alpha$ -binding sites or located in the intragenic region of E2-regulated genes

MicroRNA	Fold change in microRNA expression after E2 treatment in MCF-7p cells	ER $\alpha$ -binding sites after E2 treatment	
MicroRNAs with ER $\alpha$ -binding sites			
miR-21	1.98	1	
miR-23a	1.62	4	
miR27a	0.64	4	
miR27b	0.60	5	
MicroRNA	Fold change in microRNA levels after E2 treatment of MCF-7p cells	Host gene	E2 effect on the host gene
MicroRNAs located in the intragenic regions of E2-regulated genes			
miR30c	1.58	NF-YC	-1.2
miR-9	0.654	C1orf61	1.16
Let-7c	2.6	C21orf34	-1.9
Let-7g	2.6	TMEM113	1.3

miR-27b is located within aminopeptidase gene, which is an E2-inducible gene and displayed five and eight ER $\alpha$ -binding sites in E2-treated MCF-7p and MCF-7AKT cells, respectively.

the number of binding sites to one. ChIP assay was used to confirm ER $\alpha$  binding to the region II (central region). Unliganded ER $\alpha$  binding to this region was observed in MCF-7p cells (10-fold higher levels of PCR product compared to IgG control), which was unaffected upon E2 treatment (Figure 3B). However, ER $\alpha$  binding to this region was enhanced by E2 in MCF-7AKT cells. Thus, the miR-21 gene is associated with an authentic ER $\alpha$ -binding site. The fact that E2 reduced the number of miR-21 associated ER $\alpha$ -binding sites in MCF-7p cells and that this reduction in binding correlates with induction of miR-21 suggests that unliganded ER $\alpha$  suppresses the expression of this microRNA and E2 relieves that repression. This observation is consistent with data presented in Figure 1C and recent observations in *C. elegans* (32). In summation, our studies reveal a complex mechanism of E2-mediated changes in microRNA expression and the consequences of these changes on post-transcriptional control of gene expression.

#### Dicer is an E2-inducible gene

To determine whether E2 controls microRNA processing, we searched our ChIP-on-chip datasets and microarray datasets of untreated and E2-treated MCF-7p and MCF-7AKT for ER $\alpha$  binding and E2-regulated expression of genes associated with microRNA processing (27). The only gene associated with an ER $\alpha$ -binding site is Dicer (Figure 4A) and Dicer mRNA was induced by E2 as measured by northern analysis and qRT-PCR (Figure 4B). AKT only delayed E2-mediated induction of Dicer. These results suggest that E2 regulates microRNA machinery at the level of transcription as well as processing.

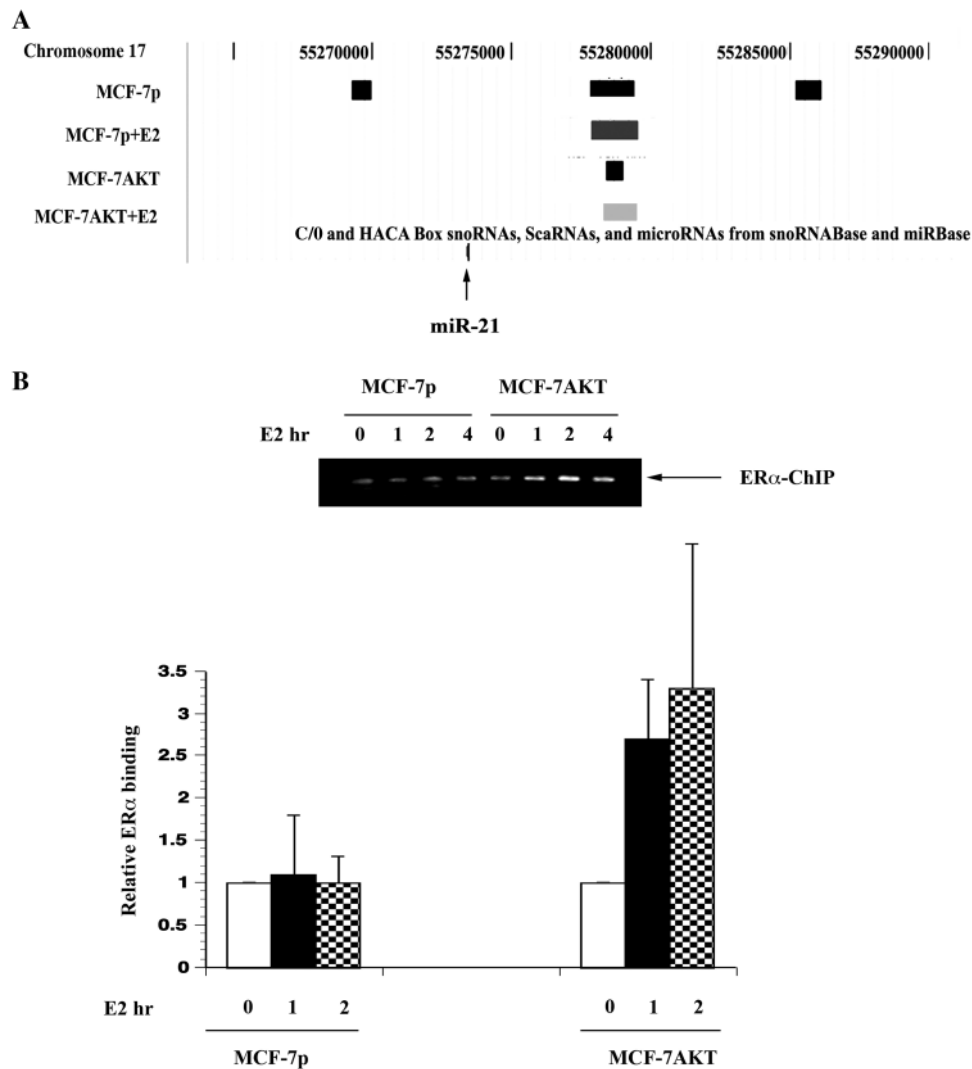
## DISCUSSION

Deregulation of microRNA expression is observed in various diseases including cancer, cardiovascular diseases and neurodegenerative disorders (43,44). While biological functions of microRNAs are being studied extensively, signaling pathways that control their expression are just beginning to be explored. Cytokines and hypoxia are known modulators of microRNA expression (45,46). In this study, we investigated the role of E2 in modulating microRNA expression in breast cancer cells. Unlike the effect of E2 on mRNA-encoding genes where the majority of them are repressed (47), 21 of the 28 E2-regulated microRNAs are induced. Our results on E2-regulated expression of miR-21 differ from a publication that appeared in this journal during revision of our manuscript (48). Wickramasinghe *et al.* (48) found lower miR-21 expression in MCF-7 but not T47-D cells treated with E2 for 6 h compared to untreated cells and this E2-mediated downregulation of miR-21 correlated with elevated expression of several miR-21 target genes. Reasons for this discrepancy are unknown. However, we note that there are several variants of MCF-7 and we have previously shown that these variants express different levels of co-activators involved in ER $\alpha$ -mediated gene expression (49). The second possibility is that E2 regulates miR-21 in a biphasic manner (induction followed by repression), which helps in fine-tuning of the E2 response. We also note that E2-mediated upregulation of miR-21 observed in our study correlates with reported higher levels of miR-21 expression in primary ER $\alpha$ + breast cancers (21).

#### Possible mechanisms of E2-regulated expression of microRNAs

Based on the results of our ChIP-on-chip study and microarray expression analysis (27), at least three distinct mechanisms appear to be involved in E2-mediated upregulation of microRNAs. First is the direct binding of ER $\alpha$  to the regulatory regions of microRNAs. The regulatory regions of miR-21 and miR-23a contain ER $\alpha$ -binding sites. With respect to miR-21, unliganded ER $\alpha$  binds to three regions in MCF-7p cells but only one region in MCF-7AKT cells. E2 either enhances or decreases interaction of ER $\alpha$  to the region shared by both cell types, depending on AKT activity. How this unusual interaction pattern of ER $\alpha$  to the miR-21-regulatory region impacts the expression of this microRNA is unknown.

The second mechanism of E2-regulated microRNA expression may involve E2-inducible expression of mRNA-encoding genes that harbor microRNA genes in their intronic regions. NF-YC and C21orf34 are repressed by E2 (5) yet microRNAs within their introns are induced by E2. It remains to be determined whether these microRNAs are transcribed from an intronic promoter independent of the host gene. The third possibility is that E2 regulates the expression of transcription factors that control the expression of microRNAs. In this respect, TRANSFAC database analysis revealed selective enrichment of binding sites for MRF-2, PBX-1, RUNX1, RUNX2 and GATA-3-transcription factors in



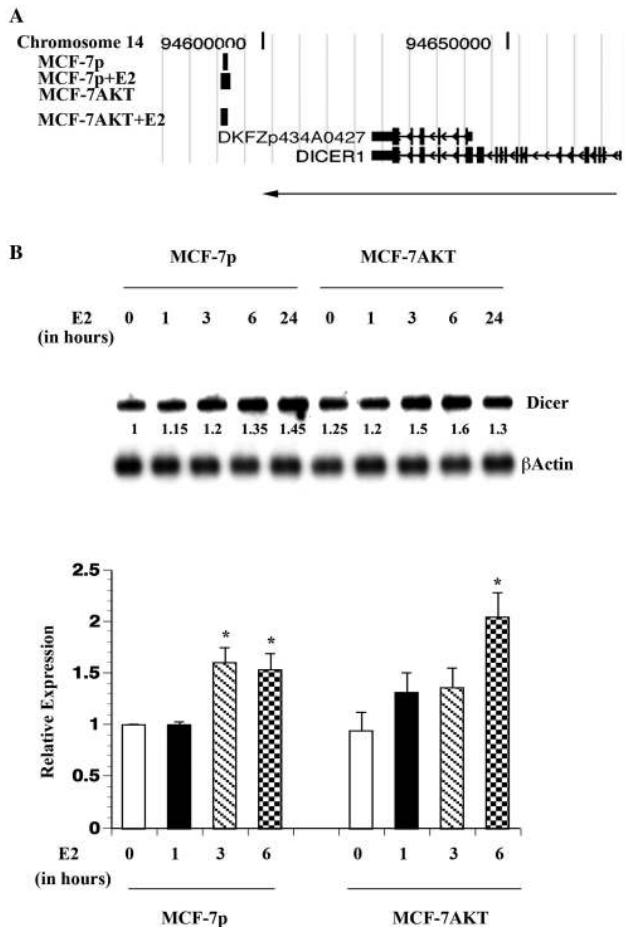
**Figure 3.** The regulatory regions of miR-21 contain ER $\alpha$ -binding sites. (A) ER $\alpha$  DNA-binding patterns to the genomic region harboring miR-21 gene on chromosome 17. Black bars represent ER $\alpha$ -binding sites in two cell types observed under ethanol and E2-treated condition. (B) ChIP analysis of ER $\alpha$  binding to the genomic region (central bar) shown in A. ChIP DNA obtained with ER $\alpha$  antibody was subjected to semi-quantitative PCR (top) or Q-PCR (bottom) with specific primers and relative enrichment of ER $\alpha$  binding is shown after normalizing ER $\alpha$  binding in ethanol MCF-7p or MCF-7AKT cells to one unit. The level of ER $\alpha$  binding in E2-treated MCF-7AKT cells was significantly higher than in untreated cells.

microRNA genes regulated by E2. Microarray and subsequent validation by qRT-PCR revealed MSX1, RUNX1 and RUNX2 as E2-inducible genes and MRF-2 and PBX-1 as E2-repressible genes in MCF-7 cells (data not shown). Previous studies have shown E2-regulated expression of GATA-3 in breast cancer cells (50). Which of these transcription factors contribute to E2-dependent microRNA expression remains to be addressed.

The fact that eight members of the Let-7 family are induced by E2 is interesting and suggests that a common pathway is involved in the expression and/or processing of Let-7 family microRNAs. This pathway is under the control of E2. In this respect, LIN28 has been shown to suppress the processing of all Let-7 family microRNAs without interfering with processing of other microRNAs (51). However, neither LIN28a nor LIN28b has ER $\alpha$ -binding sites and our microarray studies and RT-PCR

analysis indicate lack of E2-regulated LIN28 expression. We cannot rule out a post-translational activation mechanism of LIN28 by E2-regulated signaling pathways, which may induce enhanced processing of Let-7 family members. Among the several genes studied that are known to be involved in the processing of microRNAs, only Dicer is associated with ER $\alpha$ -binding sites and is induced by E2 (Figure 4). Oncomine database analysis revealed overexpression of Dicer in ER $\alpha$ -positive breast cancers compared to ER $\alpha$ -negative breast cancers, which is consistent with our results of E2-regulated expression of this gene. Thus, E2 not only regulates the expression of specific microRNAs, but also may have global effects on microRNA-regulated gene expression by altering their rate of processing.

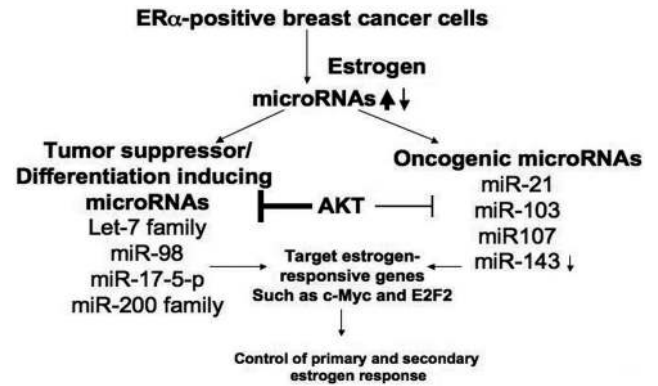
Among the microRNAs repressed by E2, only miR-27a and miR-27b contain ER $\alpha$ -binding sites. E2 may indirectly



**Figure 4.** Dicer is an E2-inducible gene. (A) ERα DNA-binding patterns to the genomic region harboring Dicer. Black bars represent ERα-binding sites in two cell types observed under ethanol-treated or E2-treated condition. (B) E2-inducible expression of Dicer in MCF-7p and MCF-7AKT cells was measured by northern blotting (top panel). The same blot was reprobed with 36B4 to ensure integrity of RNA. qRT-PCR confirming E2-inducible expression of Dicer is also shown (mean plus SEM, bottom panel). \**P* < 0.04 ethanol-treated versus E2-treated cells.

repress the expression of other microRNAs through a transcriptional repressor. In this respect, c-Myc has been shown to be a global repressor of microRNA expression (52). c-Myc is an E2-inducible gene, which is further negatively regulated by E2-induced Let-7 (Figure 2). Let-7 directly represses c-Myc translation and also destabilizes c-Myc transcript levels by inhibiting translation of IMP-1, a RNA-binding protein that stabilizes c-Myc transcripts (37,53). Thus, pathways that disrupt the finely balanced positive and negative effects of E2 on c-Myc expression may play a significant role in breast cancer progression.

The global effect of AKT on E2-regulated microRNA expression is puzzling and may involve AKT-dependent changes in the activity of transcription factors. The failure of E2 to induce the expression of miR-200a, miR-17-5p, miR-200a, miR-200c and miR-203 in MCF-7AKT cells could be due to AKT-mediated downregulation of the basal expression of these microRNAs and the effect of E2 on them being below the detection level in our assay



**Figure 5.** A model depicting the possible effects of E2-regulated microRNAs on E2 response and differentiated phenotype of ERα-positive breast cancer cells.

(Table 1). miR-520d appears to be an exception to the general phenomena of AKT-dependent decrease in microRNA expression as AKT increased the basal levels of this microRNA by ~5-fold and this microRNA was further induced by E2 in AKT overexpressing cells. miR-520d may be E2-inducible in parental cells also but its expression is below the level for detection. The significance of AKT-mediated induction of miR-520d could not be predicted as targets of this microRNA are yet to be identified.

**Potential role of E2-regulated microRNAs in breast cancer**

The majority of E2-regulated microRNAs appear to have a tumor suppressor role by participating in a negative feedback loop to restrict E2 action. For example, E2-inducible expression of Let-7 family members may limit the expression of Ras and c-Myc oncogene levels and promote differentiation of cancer cells. Control of c-Myc levels by Let-7 is particularly relevant to E2 signaling because c-Myc cooperates with ERα in the expression of select target genes and Let-7 may play a role in attenuating such cooperative gene expression (54). E2-induced microRNAs may control secondary estrogen responses by limiting the expression levels of E2F family members (31). Additionally, E2-induced microRNA miR-17-5p may control ERα primary responses by reducing the levels of the AIB1 coactivator (40). Figure 5 schematically represents how E2-regulated microRNAs may control E2 responses by targeting c-Myc and E2F2.

E2-regulated miR-21 may have dual roles in cancer. It may protect cancer cells against cell death and promote metastasis: this effect of miR-21 is likely due to translational repression of pro-apoptotic PDCD4 and metastasis suppressor Maspin (55). In this respect, overexpression of miR-21 in MCF-7 cells leads to enhanced tumor growth in nude mice (56). On the other hand, miR-21 may reduce self-renewal of cancer cells with stem-cell-like properties by inhibiting the translation of stem cell renewal genes Oct-4, c-Myc, Nanog and Sox2 (38). For unknown reasons, we did not observe an effect of LNA against miR-21 on PDCD4 expression in E2-treated MCF-7 cells and these cells do not express Maspin (data not

shown). Our results differ from that of Wickramasinghe *et al.* (48) with respect to PDCD4 possibly reflecting differences in the variants of MCF-7 cells used in the two studies. In our MCF-7 cells, PDCD4 is an E2-inducible gene and is associated with ER $\alpha$ -binding sites. Additionally, based on TargetScan prediction, PDCD4 is a target of E2-inducible miR-200a (Table 1 and Supplementary Table S1). Therefore, these multiple levels of E2 action may have obscured the effects of miR-21 LNA on PDCD4 protein levels in E2-treated cells.

AKT is a major signaling molecule in breast cancer. MMTV-AKT1 transgenic mice develop ER $\alpha$ -positive breast cancer when exposed to chemical carcinogens suggesting a significant crosstalk between AKT and ER $\alpha$  in breast cancer (57). The extent to which this crosstalk relies on modulation of microRNA expression is not known and may potentially provide important clues into the initiation of ER $\alpha$ -positive breast cancers. AKT decreased E2-inducible expression of both tumor suppressor and oncogenic microRNAs. E2-dependent reduction of miR-126 in only AKT overexpressing cells is relevant to breast cancer as this microRNA is considered both a tumor suppressor and metastasis inhibitor (58).

In summary, we show the effects of E2 on modulating the microRNA-expression pattern as well as the potential for extracellular signal activated kinases to modulate this action of E2. With respect to breast cancer, our observations may have implications on responses to endocrine therapy as signaling pathways that modulate ER $\alpha$  function may change the balance between tumor suppressor and oncogenic microRNAs induced by E2 and thus alter the response to anti-estrogen therapy.

## SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online.

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## REFERENCES

- Jordan, V.C. and O'Malley, B.W. (2007) Selective estrogen-receptor modulators and antihormonal resistance in breast cancer. *J. Clin. Oncol.*, **25**, 5815–5824.
- Ali, S. and Coombes, R.C. (2002) Endocrine-responsive breast cancer and strategies for combating resistance. *Nat. Rev. Cancer*, **2**, 101–112.
- Campbell, R.A., Bhat-Nakshatri, P., Patel, N.M., Constantinidou, D., Ali, S. and Nakshatri, H. (2001) Phosphatidylinositol 3-kinase/AKT-mediated activation of estrogen receptor  $\alpha$ : a new model for anti-estrogen resistance. *J. Biol. Chem.*, **276**, 9817–9824.
- Gururaj, A.E., Rayala, S.K., Vadlamudi, R.K. and Kumar, R. (2006) Novel mechanisms of resistance to endocrine therapy: genomic and nongenomic considerations. *Clin. Cancer Res.*, **12**, 1001s–1007s.
- Calin, G.A. and Croce, C.M. (2007) Chromosomal rearrangements and microRNAs: a new cancer link with clinical implications. *J. Clin. Invest.*, **117**, 2059–2066.
- Hammond, S.M. (2007) MicroRNAs as tumor suppressors. *Nat. Genet.*, **39**, 582–583.
- Chendrimada, T.P., Finn, K.J., Ji, X., Baillat, D., Gregory, R.I., Liebhaber, S.A., Pasquinelli, A.E. and Shiekhattar, R. (2007) MicroRNA silencing through RISC recruitment of eIF6. *Nature*, **447**, 823–828.
- Tay, Y., Zhang, J., Thomson, A.M., Lim, B. and Rigoutsos, I. (2008) MicroRNAs to Nanog, Oct4 and Sox2 coding regions modulate embryonic stem cell differentiation. *Nature*, **455**, 1124–1128.
- Forman, J.J., Legesse-Miller, A. and Coller, H.A. (2008) A search for conserved sequences in coding regions reveals that the let-7 microRNA targets Dicer within its coding sequence. *Proc. Natl Acad. Sci. USA*, **105**, 14879–14884.
- Vasudevan, S., Tong, Y. and Steitz, J.A. (2007) Switching from repression to activation: microRNAs can up-regulate translation. *Science*, **318**, 1931–1934.
- Bartel, D.P. (2009) MicroRNAs: target recognition and regulatory functions. *Cell*, **136**, 215–233.
- Lewis, B.P., Burge, C.B. and Bartel, D.P. (2005) Conserved seed pairing, often flanked by adenosines, indicates that thousands of human genes are microRNA targets. *Cell*, **120**, 15–20.
- Kedde, M., Strasser, M.J., Boldajipour, B., Vrielink, J.A., Slanchev, K., le Sage, C., Nagel, R., Voorhoeve, P.M., van Duijse, J., Orom, U.A. *et al.* (2007) RNA-binding protein Dnd1 inhibits microRNA access to target mRNA. *Cell*, **131**, 1273–1286.
- Bhattacharyya, S.N., Habermacher, R., Martine, U., Closs, E.I. and Filipowicz, W. (2006) Relief of microRNA-mediated translational repression in human cells subjected to stress. *Cell*, **125**, 1111–1124.
- Place, R.F., Li, L.C., Pookot, D., Noonan, E.J. and Dahiya, R. (2008) MicroRNA-373 induces expression of genes with complementary promoter sequences. *Proc. Natl Acad. Sci. USA*, **105**, 1608–1613.
- Amaral, P.P., Dinger, M.E., Mercer, T.R. and Mattick, J.S. (2008) The eukaryotic genome as an RNA machine. *Science*, **319**, 1787–1789.
- Lippman, Z. and Martienssen, R. (2004) The role of RNA interference in heterochromatic silencing. *Nature*, **431**, 364–370.
- Borchert, G.M., Lanier, W. and Davidson, B.L. (2006) RNA polymerase III transcribes human microRNAs. *Nat. Struct. Mol. Biol.*, **13**, 1097–1101.
- Saini, H.K., Griffiths-Jones, S. and Enright, A.J. (2007) Genomic analysis of human microRNA transcripts. *Proc. Natl Acad. Sci. USA*, **104**, 17719–17724.
- Calin, G.A., Sevignani, C., Dumitru, C.D., Hyslop, T., Noch, E., Yendamuri, S., Shimizu, M., Rattan, S., Bullrich, F., Negrini, M. *et al.* (2004) Human microRNA genes are frequently located at fragile sites and genomic regions involved in cancers. *Proc. Natl Acad. Sci. USA*, **101**, 2999–3004.
- Mattie, M.D., Benz, C.C., Bowers, J., Sensinger, K., Wong, L., Scott, G.K., Fedele, V., Ginzinger, D., Getts, R. and Haqq, C. (2006) Optimized high-throughput microRNA expression profiling provides novel biomarker assessment of clinical prostate and breast cancer biopsies. *Mol. Cancer*, **5**, 24.
- Blenkiron, C., Goldstein, L.D., Thorne, N.P., Spiteri, I., Chin, S.F., Dunning, M.J., Barbosa-Morais, N.L., Teschendorff, A.E., Green, A.R., Ellis, I.O. *et al.* (2007) MicroRNA expression profiling

- of human breast cancer identifies new markers of tumor subtype. *Genome Biol.*, **8**, R214.
23. Volinia,S., Calin,G.A., Liu,C.G., Ambs,S., Cimmino,A., Petrocca,F., Visone,R., Iorio,M., Roldo,C., Ferracin,M. *et al.* (2006) A microRNA expression signature of human solid tumors defines cancer gene targets. *Proc. Natl Acad. Sci. USA*, **103**, 2257–2261.
  24. Perou,C.M., Sorlie,T., Eisen,M.B., van de Rijn,M., Jeffrey,S.S., Rees,C.A., Pollack,J.R., Ross,D.T., Johnsen,H., Akslen,L.A. *et al.* (2000) Molecular portraits of human breast tumours. *Nature*, **406**, 747–752.
  25. Sorlie,T., Perou,C.M., Tibshirani,R., Aas,T., Geisler,S., Johnsen,H., Hastie,T., Eisen,M.B., van de Rijn,M., Jeffrey,S.S. *et al.* (2001) Gene expression patterns of breast carcinomas distinguish tumor subclasses with clinical implications. *Proc. Natl Acad. Sci. USA*, **98**, 10869–10874.
  26. Adams,B.D., Furneaux,H. and White,B.A. (2007) The micro-ribonucleic acid (miRNA) miR-206 targets the human estrogen receptor-alpha (ERalpha) and represses ERalpha messenger RNA and protein expression in breast cancer cell lines. *Mol. Endocrinol.*, **21**, 1132–1147.
  27. Bhat-Nakshatri,P., Wang,G., Appaiah,H., Luktuke,N., Carroll,J.S., Geistlinger,T.R., Brown,M., Badve,S., Liu,Y. and Nakshatri,H. (2008) AKT Alters genome-wide estrogen receptor alpha binding and impacts estrogen signaling in breast cancer. *Mol. Cell Biol.*, **28**, 7487–7503.
  28. Thomson,J.M., Parker,J.S. and Hammond,S.M. (2007) Microarray analysis of miRNA gene expression. *Methods Enzymol.*, **427**, 107–122.
  29. Thomson,J.M., Parker,J., Perou,C.M. and Hammond,S.M. (2004) A custom microarray platform for analysis of microRNA gene expression. *Nat. Methods*, **1**, 47–53.
  30. Carroll,J.S., Liu,X.S., Brodsky,A.S., Li,W., Meyer,C.A., Szary,A.J., Eeckhoutte,J., Shao,W., Hestermann,E.V., Geistlinger,T.R. *et al.* (2005) Chromosome-wide mapping of estrogen receptor binding reveals long-range regulation requiring the forkhead protein FoxA1. *Cell*, **122**, 33–43.
  31. Bourdeau,V., Deschenes,J., Laperriere,D., Aid,M., White,J.H. and Mader,S. (2008) Mechanisms of primary and secondary estrogen target gene regulation in breast cancer cells. *Nucleic Acids Res.*, **36**, 76–93.
  32. Bethke,A., Fielenbach,N., Wang,Z., Mangelsdorf,D.J. and Antebi,A. (2009) Nuclear hormone receptor regulation of microRNAs controls developmental progression. *Science*, **324**, 95–98.
  33. Johnson,C.D., Esquela-Kerscher,A., Stefani,G., Byrom,M., Kelnar,K., Ovcharenko,D., Wilson,M., Wang,X., Shelton,J., Shingara,J. *et al.* (2007) The let-7 microRNA represses cell proliferation pathways in human cells. *Cancer Res.*, **67**, 7713–7722.
  34. Ibarra,I., Erlich,Y., Muthuswamy,S.K., Sachidanandam,R. and Hannon,G.J. (2007) A role for microRNAs in maintenance of mouse mammary epithelial progenitor cells. *Genes Dev.*, **21**, 3238–3243.
  35. Mayr,C., Hemann,M.T. and Bartel,D.P. (2007) Disrupting the pairing between let-7 and Hmga2 enhances oncogenic transformation. *Science*, **315**, 1576–1579.
  36. Johnson,S.M., Grosshans,H., Shingara,J., Byrom,M., Jarvis,R., Cheng,A., Labourier,E., Reinert,K.L., Brown,D. and Slack,F.J. (2005) RAS is regulated by the let-7 microRNA family. *Cell*, **120**, 635–647.
  37. Sampson,V.B., Rong,N.H., Han,J., Yang,Q., Aris,V., Soteropoulos,P., Petrelli,N.J., Dunn,S.P. and Krueger,L.J. (2007) MicroRNA let-7a down-regulates MYC and reverts MYC-induced growth in Burkitt lymphoma cells. *Cancer Res.*, **67**, 9762–9770.
  38. Singh,S.K., Kagalwala,M.N., Parker-Thornburg,J., Adams,H. and Majumder,S. (2008) REST maintains self-renewal and pluripotency of embryonic stem cells. *Nature*, **453**, 223–227.
  39. Sylvestre,Y., De Guire,V., Querido,E., Mukhopadhyay,U.K., Bourdeau,V., Major,F., Ferbeyre,G. and Chartrand,P. (2007) An E2F/miR-20a autoregulatory feedback loop. *J. Biol. Chem.*, **282**, 2135–2143.
  40. Hossain,A., Kuo,M.T. and Saunders,G.F. (2006) Mir-17-5p regulates breast cancer cell proliferation by inhibiting translation of AIB1 mRNA. *Mol. Cell Biol.*, **26**, 8191–8201.
  41. Shi,B., Liang,J., Yang,X., Wang,Y., Zhao,Y., Wu,H., Sun,L., Zhang,Y., Chen,Y., Li,R. *et al.* (2007) Integration of estrogen and Wnt signaling circuits by the polycomb group protein EZH2 in breast cancer cells. *Mol. Cell Biol.*, **27**, 5105–5119.
  42. Baek,D., Villen,J., Shin,C., Camargo,F.D., Gygi,S.P. and Bartel,D.P. (2008) The impact of microRNAs on protein output. *Nature*, **455**, 64–71.
  43. Couzin,J. (2008) MicroRNAs make big impression in disease after disease. *Science*, **319**, 1782–1784.
  44. Wang,W.X., Rajeev,B.W., Stromberg,A.J., Ren,N., Tang,G., Huang,Q., Rigoutsos,I. and Nelson,P.T. (2008) The expression of microRNA miR-107 decreases early in Alzheimer's disease and may accelerate disease progression through regulation of beta-site amyloid precursor protein-cleaving enzyme 1. *J. Neurosci.*, **28**, 1213–1223.
  45. Kulshreshtha,R., Ferracin,M., Wojcik,S.E., Garzon,R., Alder,H., Agosto-Perez,F.J., Davuluri,R., Liu,C.G., Croce,C.M., Negrini,M. *et al.* (2007) A microRNA signature of hypoxia. *Mol. Cell Biol.*, **27**, 1859–1867.
  46. Taganov,K.D., Boldin,M.P., Chang,K.J. and Baltimore,D. (2006) NF-kappaB-dependent induction of microRNA miR-146, an inhibitor targeted to signaling proteins of innate immune responses. *Proc. Natl Acad. Sci. USA*, **103**, 12481–12486.
  47. Frasier,J., Danes,J.M., Komm,B., Chang,K.C., Lyttle,C.R. and Katzenellenbogen,B.S. (2003) Profiling of estrogen up- and down-regulated gene expression in human breast cancer cells: insights into gene networks and pathways underlying estrogenic control of proliferation and cell phenotype. *Endocrinology*, **144**, 4562–4574.
  48. Wickramasinghe,N.S., Manavalan,T.T., Dougherty,S.M., Riggs,K.A., Li,Y. and Klinge,C.M. (2009) Estradiol downregulates miR-21 expression and increases miR-21 target gene expression in MCF-7 breast cancer cells. *Nucleic Acids Res.*, **37**, 2584–2595.
  49. Kishimoto,H., Wang,Z., Bhat-Nakshatri,P., Chang,D., Clarke,R. and Nakshatri,H. (2005) The p160 family co-activators regulate breast cancer cell proliferation and invasion through autocrine/paracrine activity of SDF-1{alpha}/CXCL12. *Carcinogenesis*, **26**, 1706–1715.
  50. Eeckhoutte,J., Keeton,E.K., Lupien,M., Krum,S.A., Carroll,J.S. and Brown,M. (2007) Positive cross-regulatory loop ties GATA-3 to estrogen receptor alpha expression in breast cancer. *Cancer Res.*, **67**, 6477–6483.
  51. Viswanathan,S.R., Daley,G.Q. and Gregory,R.I. (2008) Selective blockade of microRNA processing by Lin28. *Science*, **320**, 97–100.
  52. Chang,T.C., Yu,D., Lee,Y.S., Wentzel,E.A., Arking,D.E., West,K.M., Dang,C.V., Thomas-Tikhonenko,A. and Mendell,J.T. (2008) Widespread microRNA repression by Myc contributes to tumorigenesis. *Nat. Genet.*, **40**, 43–50.
  53. Boyerinas,B., Park,S.M., Shomron,N., Hedegaard,M.M., Vinther,J., Andersen,J.S., Feig,C., Xu,J., Burge,C.B. and Peter,M.E. (2008) Identification of let-7-regulated oncofetal genes. *Cancer Res.*, **68**, 2587–2591.
  54. Cheng,A.S., Jin,V.X., Fan,M., Smith,L.T., Liyanarachchi,S., Yan,P.S., Leu,Y.W., Chan,M.W., Plass,C., Nephew,K.P. *et al.* (2006) Combinatorial analysis of transcription factor partners reveals recruitment of c-MYC to estrogen receptor-alpha responsive promoters. *Mol. Cell*, **21**, 393–404.
  55. Zhu,S., Wu,H., Wu,F., Nie,D., Sheng,S. and Mo,Y.Y. (2008) MicroRNA-21 targets tumor suppressor genes in invasion and metastasis. *Cell Res.*, **18**, 350–359.
  56. Si,M.L., Zhu,S., Wu,H., Lu,Z., Wu,F. and Mo,Y.Y. (2007) miR-21-mediated tumor growth. *Oncogene*, **26**, 2799–2803.
  57. Blanco-Aparicio,C., Perez-Gallego,L., Pequeno,B., Leal,J.F., Renner,O. and Carnero,A. (2007) Mice expressing myrAKT1 in the mammary gland develop carcinogen-induced ER-positive mammary tumors that mimic human breast cancer. *Carcinogenesis*, **28**, 584–594.
  58. Tavazoie,S.F., Alarcon,C., Oskarsson,T., Padua,D., Wang,Q., Bos,P.D., Gerald,W.L. and Massague,J. (2008) Endogenous human microRNAs that suppress breast cancer metastasis. *Nature*, **451**, 147–152.