

The estrogen receptor- α -induced microRNA signature regulates itself and its transcriptional response

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Following estrogenic activation, the estrogen receptor- α (ER α) directly regulates the transcription of target genes via DNA binding. MicroRNAs (miRNAs) modulated by ER α have the potential to fine tune these regulatory systems and also provide an alternate mechanism that could impact on estrogen-dependent developmental and pathological systems. Through a microarray approach, we identify the subset of microRNAs (miRNAs) modulated by ER α , which include upregulation of miRNAs derived from the processing of the paralogous primary transcripts (pri-) mir-17-92 and mir-106a-363. Characterization of the mir-17-92 locus confirms that the ER α target protein c-MYC binds its promoter in an estrogen-dependent manner. We observe that levels of pri-mir-17-92 increase earlier than the mature miRNAs derived from it, implicating precursor cleavage modulation after transcription. Pri-mir-17-92 is immediately cleaved by DROSHA to pre-miR-18a, indicating that its regulation occurs during the formation of the mature molecule from the precursor. The clinical implications of this novel regulatory system were confirmed by demonstrating that pre-miR-18a was significantly upregulated in ER α -positive compared to ER α -negative breast cancers. Mechanistically, miRNAs derived from these paralogous pri-miRNAs (miR-18a, miR-19b, and miR-20b) target and downregulate ER α , while a subset of pri-miRNA-derived miRNAs inhibit protein translation of the ER α transcriptional p160 coactivator, AIB1. Therefore, different subsets of miRNAs identified act as part of a negative autoregulatory feedback loop. We propose that ER α , c-MYC, and miRNA transcriptional programs invoke a sophisticated network of interactions able to provide the wide range of coordinated cellular responses to estrogen.

AIB1 | autoregulatory feedback loop | primary transcript | processing

Upon 17- β -estradiol (E2) binding, estrogen receptors (ERs) mediate transcription by interacting directly to specific estrogen response elements (EREs) located in the promoter/enhancer region of its target genes or indirectly by tethering to nuclear proteins, such as AP1 and SP1 transcription factors (2–4). The cellular response to estrogen is highly regulated at multiple levels including transcription, RNA stability, and post-translational modifications (5–8). Following treatment with E2, ER α transcription and mRNA stability is substantially reduced within 1 h of stimulation (7). Furthermore, E2-ER α interactions accelerate receptor degradation through the ubiquitin-proteasome pathway, an effect associated with its major coactivator AIB1 (8).

MicroRNAs (miRNAs) are a class of noncoding short RNAs, 21–24 nucleotides (nt) in length, that play a role in gene regulation. They downregulate expression of their target genes by base pairing to the 3'-UTR of target messenger RNAs (mRNAs) (9). During their biogenesis most miRNAs are transcribed as part of a longer transcript named pri-miRNA (10). These molecules are processed inside the nucleus by DROSHA, producing a pre-miRNA that is a

70-nt "imperfect" stem loop RNA actively transported into the cytoplasm. In the cytoplasm the pre-miRNA is cleaved by DICER, a dual processing event that releases a small double stranded RNA, about 22 nt in length. Here, nuclear processing activity is thought to be regulated at early stages of development and in a variety of tumor cells (11–13). There is also evidence of regulation at the next step, pre-miRNA precursor processing (14, 15). After formation of the small duplex RNA, only 1 strand is loaded onto a miRNA induced silencing complex (RISC). These RISCs, guided by their miRNA, interact with the 3'-UTR or sometimes with the coding region of target mRNAs, inhibiting protein translation or degrading the mRNA target (10).

Substantial data associate changes in miRNA activity with carcinogenesis and progression (16–19). The human mir-17-92 cluster is a polycistronic gene with a chromosomal location 13q31-q32 that encodes 6 miRNAs (miR-17, miR-18a, miR-19a, miR-20a, miR-19b-1, and miR-92-1). Ancient duplications have given rise to 2 mir-17-92 paralogues in mammals: mir-106b-25 and mir-106a-363. Mir-17-92 is thought to be oncogenic in lung cancer and lymphomas (17, 20) or function as a tumor suppressor in breast cancer by downregulating AIB1 and/or cyclin D1 (21, 22). Furthermore, the genomic 13q31 area including mir-17-92 is correlated with loss of heterozygosity in breast cancer (23).

By a genome wide approach, we have elucidated the miRNAs regulated by ER α in breast cancer. Here, we show that among the few miRNAs upregulated by ER α , miR-18a encoded by the pri-mir-17-92, miR-19b encoded by both this primary transcript and its evolutionary paralogue pri-mir-106a-363, and miR-20b encoded by pri-mir-106a-363, downregulate ER α expression at the protein translational level, correlating the induction of these 2 genes during cell proliferation with a negative feedback loop. Remarkably, miR-20b also downregulates and targets the ER α coactivator AIB1. Since ER α can act as a ligand-activated oncogene, we suggest that the pri-mir-17-92 acts as a tumor suppressor in breast cancer, not only by downregulating cyclin D1 and AIB1 via the miR-17/20/106 family, but also by downregulation of ER α by miR-18, miR-19, and miR-17/20/106 members. For the first time we correlate ER α translational control by miRNAs as a further regulatory process involved in ER α transcriptional activity after ligand stimulation.

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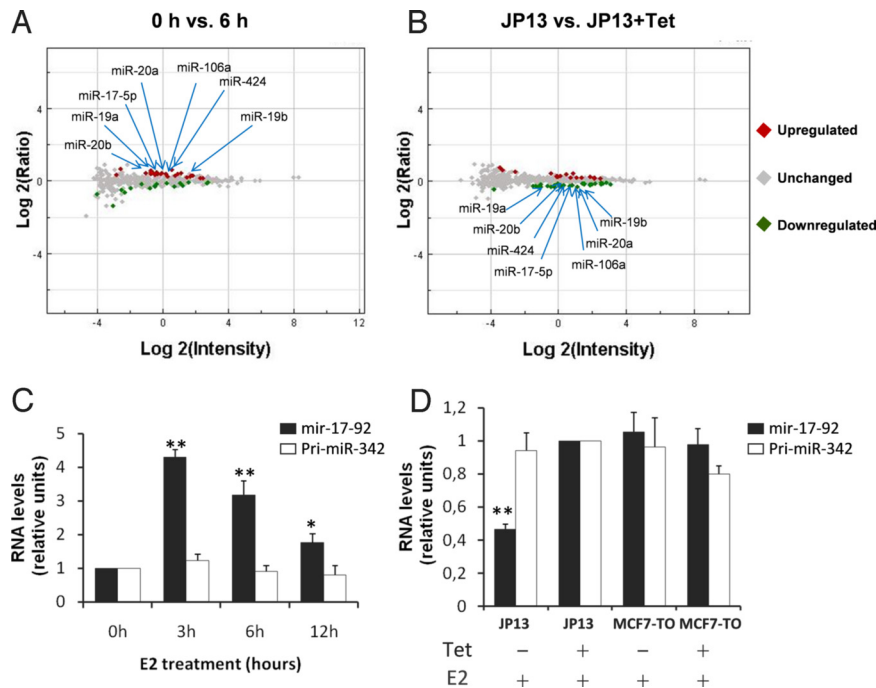
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Fig. 1. Pri-mir-17-92 is increased by E2 and decreased by overexpression of PLZF-ER α . (A) MCF-7 cell lines underwent E2 stimulation (10 nM) after 72 h of hormone deprivation. After total RNA extraction and labeling we used a microarray platform containing probes for 470 human miRNAs. After hybridization and scanning, raw data were imported into the Rosetta Resolver system for analysis. A $P < 0.01$ was used as cut-off for identification of miRNAs upregulated or downregulated between 0 h versus 6 h. (B) JP13 cells were cultured in the presence or absence of Tet for 72 h, followed by the addition of 10 nM E2 for 24 h before microarray analysis. Once again a $P < 0.01$ was used as cut-off for identification of miRNAs downregulated in JP13 + Tet versus JP13 – Tet. (C) MCF-7 cells were maintained in DMEM (minus phenol red) supplemented with 10% charcoal-dextran FBS for 3 days and then were either left untreated or treated with 10 nM E2 for the indicated time periods. After total RNA extraction, expression of pri-mir-17-92 and pri-miR-342 was analyzed by RT-qPCR using SYBR green and normalized to *GAPDH*. (D) JP13 and MCF-7-TO cells were cultured in the presence or absence of Tet for 72 h, followed by the addition of 10 nM E2 for 24 h. Once again, after total RNA extraction, expression of pri-mir-17-92 and pri-miR-342 were analyzed by RT-qPCR using SYBR green and normalized to *GAPDH*. The mean of 3 experiments each performed in triplicate are presented, error bars represent SEM. For RT-qPCR data, the asterisk indicates $P < 0.05$ in comparison to time 0 h, the double asterisk represents $P < 0.005$ in comparison to time 0 h. P values were obtained using a 2-tailed Student's t -test.



Results

ER α Induction Reveals pri-mir-17-92 Upregulation. To determine whether ER α regulates the expression of miRNAs, E2 was added to MCF-7 cells and miRNA chip hybridization was performed to elucidate early (0–3 h) and delayed (>6 h) regulation of miRNAs by ER α . As a control we used a MCF-7-TO (MCF-7-Tet-Off)-derived cell line, JP13, that conditionally overexpresses a protein composed of the zinc finger transcriptional repressor PLZF fused to ER α (PLZF-ER α), acting as a dominant-negative that inhibits expression of estrogen-regulated genes and estrogen-stimulated growth of MCF-7 cells (24). Following E2 stimulation and before microarray hybridization, we assessed the reliability of the system using quantitative real time PCR (RT-qPCR) to reveal expression levels of the ER α -regulated gene *GREB1* [supporting information (SI) Fig. S1A and ref. 25]. *GREB1* expression was reduced in PLZF-ER α cell lines treated without doxytetracycline (Tet) compared to cell lines that do not express the fusion protein (Fig. S1B). Although following array analysis we did not reveal any miRNAs with expression changes greater than 2-fold comparing 0 h to 3, 6, and 12 h ($P < 0.05$) (Table S1), we found that those miRNAs that increased following E2 in MCF-7 cells decreased in the JP13-Tet-Off system used as a control (Fig. 1A and B and Table S1). The ER α -upregulated miRNAs were generated by the processing of 3 paralogous primary miRNAs: pri-mir-17-92, pri-mir-106a-363, and pri-mir-106b-25 (Fig. 1A and B, Table S1 and Fig. S2).

To confirm the change of expression detected by the microarray, we performed RT-qPCR choosing those miRNAs modulated between 1.2- to 2-fold in cells treated with E2 and those repressed similarly by PLZF-ER α in JP13 cell lines. Firstly, we examined the expression of the unprocessed pri-mir-17-92 and family members. Pri-mir-17-92 appeared upregulated within 3 h of E2 treatment reaching a 4- to 5-fold change in comparison to 0 h (Fig. 1C) defining it as a new early ER α -regulated gene. Levels of expression were significantly repressed by PLZF-ER α (Fig. 1D). Pri-mir-342, a negative control, showed no changes (Fig. 1C and D), and we obtained the same results normalizing the value of expression for *GAPDH*, for the snRNA *U6* and for the snoRNA *U47*. We excluded

from this analysis the paralogous pri-mir-106b-25 because the fold change of the miRNAs encoded by it were considered too low by our pre-established criteria (Table S1). Furthermore, levels of expression of the pri-mir-106a-363 in MCF-7 cells appeared too low to detect (we were not able to amplify it using 7 different sets of primers from 7 genomic regions). However, it is known that in the P493-6 B cell line that although there is c-MYC-regulated expression of typical miRNAs encoded by both pri-mir-17-92 and pri-mir-106a-363, it is possible to detect the pri-mir-17-92 but not pri-mir-106a-363, indicating that the latter could either be less expressed or alternatively processed more rapidly (26). Furthermore, miR-424, miR-450, and miR-542-3p located within 6 kb of the same genomic region, appeared significantly upregulated by E2 and significantly repressed by PLZF-ER α in a perfectly reciprocal manner (Table S1). We also observed a subset of miRNAs that were subtly downregulated by the E2-ER α complex from 0 to 12 h, and the majority of these belonged to the miR-181 family (Table S1).

Next, we performed RT-qPCR for miR-18a, miR-19a, miR-19b, miR-20a, miR-92 (derived from pri-mir-17-92), miR-19b, miR-20b, miR-92 (from pri-mir-106a-363), miR-424, and miR-181b; the 2 techniques showed an overall correlation (Fig. S3A–I and Table S1). Comparing the low levels of expression of the miRNAs to the higher levels of the pri-miRNA after stimulation, there was negative regulation of miRNA biogenesis, following transcriptional induction by ER α (Fig. 1C and Fig. S3).

C-MYC Directly Regulates the pri-mir-17-92 upon Estrogenic Stimula-

tion. It has already been demonstrated that pri-mir-17-92 is transcriptionally regulated by c-MYC in the P493-6 B cell line during the G1–S cellular transition phase (26). Since c-MYC mRNA is upregulated by ER α within 1 h of E2 treatment in breast cancer cells (27), c-MYC could contribute to the increased transcription of the pri-mir-17-92 upon E2 stimulation. Interestingly, we observed a half site conserved ERE 70 bp upstream of the c-MYC consensus site (E-box) of the mir-17-92 promoter (Fig. 2A and Fig. S4). Using cycloheximide (CHX), we demonstrated that new protein synthesis

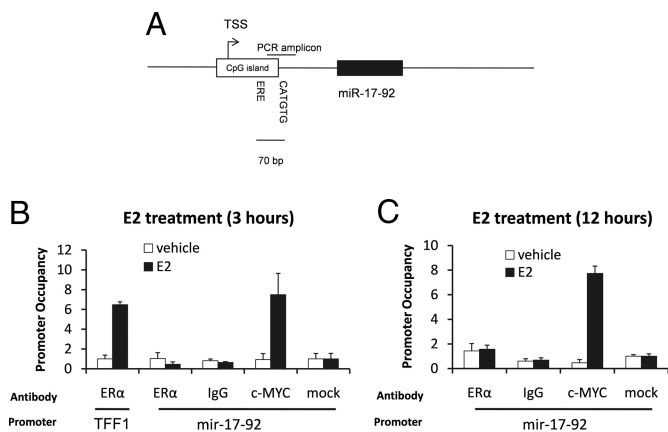


Fig. 2. c-MYC directly regulates the pri-mir-17-92 upon estrogenic stimulation. (A) Schematic representation of the mir-17-92 cluster genomic region. Both the c-MYC binding site and a putative ERE half site are indicated. (B) MCF-7 cells were maintained in estrogen-free medium for 3 days (starvation) and then either left untreated (vehicle) or treated with 10 nM E2 for 3 h after which ChIP was performed, followed by real time PCR. The c-MYC interaction site genomic region is presented. (C) After starvation, MCF-7 cells were treated with E2 for 12 h before ChIP.

is not required exclusively for pri-mir-17-92 expression (Fig. S5) and because it has been demonstrated that estrogen responsive genes can contain both ER α and c-MYC binding elements located within close proximity [13–214 bp within the promoter and regulated by both transcription factors in an E2-dependent manner (28)], we performed chromatin immunoprecipitation (ChIP) assays for both ER α and c-MYC: coprecipitated DNA was analyzed by amplifying the genomic region containing both consensus sites (Fig. 2A and Fig. S4) by real time PCR (Fig. 2B and C). Although TFF1, a known estrogen-regulated gene, is confirmed here as regulated by ER α (Fig. 2B), we observed only c-MYC interacting with the mir-17-92 promoter region analyzed (Fig. 2B and C). We demonstrated that c-MYC is recruited to the mir-17-92 promoter in breast cancer cells upon E2 stimulation.

Pri-mir-17-92 Is Negatively Regulated Following DROSHA Cleavage Prolonging miRNA Maturation over Time. Remarkably, the pri-mir-17-92 expression is striking compared to the miRNAs that are produced by its processing (miR-17, miR-18a, miR-19a, miR-19b, and miR-20a), indicative of modulation of miRNAs biogenesis at the posttranscriptional level (Figs. 1C, 3A, and Fig. S3). A primary transcript undergoes a dual processing event, the first in the nucleus by DROSHA (pre-miRNA production), the second in the cytoplasm by DICER. To define the step(s) of miRNA biogenesis in which regulation occurs, we measured levels of the pri-mir-17-92-derived pre-miR-18a after E2 treatment. DROSHA pri-mir-17-92 cleavage to pre-miR-18a was not a regulatory or “rate-limiting” step here because both were induced at similar levels (Fig. 3B). However, the primers used to amplify the pre-miR-18a also amplify pri-mir-17-92. Therefore, to establish that we could distinguish between pri- and pre-miRNA, we stimulated the cell lines with E2 and then separated the small RNA fraction from the large RNA fraction. We used the large RNA fraction to measure pri-mir-17-92 and the small RNA fraction to measure pre-miR-18a (Fig. S6). As a further control we measured the pri-mir-17-92 from the small RNA fraction without obtaining any amplification product. These data demonstrated that pri-mir-17-92 is induced by the E2-ER α complex, then it is processed by DROSHA releasing the pre-miR-18a, but the passage between pre-miR-18a and miR-18a is attenuated until at least 12 h following initial E2 stimulation. Furthermore, using RT-qPCR, we found that both miR-18a and miR-20a mature forms increase their levels of expression from 24 to 72 h

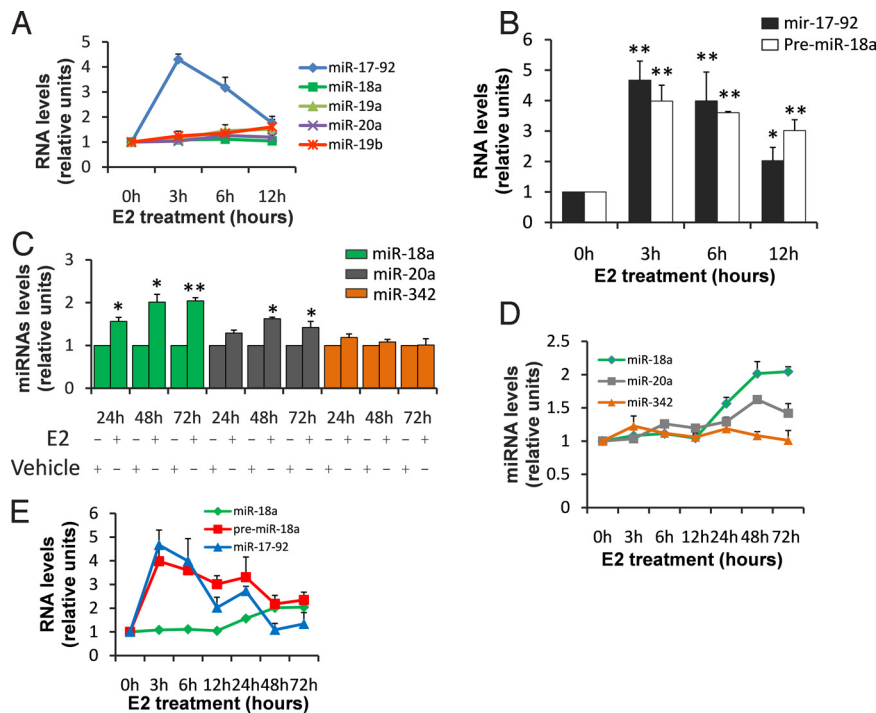
after E2 stimulation (Fig. 3C and D). Analyzing the levels of the pri-mir-17-92 and the pre-miR-18a from 0 to 72 h, we observed that pri-mir-17-92 is transcriptionally upregulated after 3 h, then DROSHA promptly processes the pri- to the pre-miR-18a, whereas the formation of the mature form from the pre-miR-18a is delayed (Fig. 3E). In addition while the miR-18a levels start to increase at 24 h, both pri-mir-17-92 and pre-miR-18a levels decline, indicative of the processing delay we observed (Fig. 3E).

Pri-mir-17-92 Expression Is Correlated with ER α Levels in ER α -Positive Primary Breast Cancers. To evaluate ER α modulation of the pri-mir-17-92 at the physiologic level, we examined a correlation between ER α mRNA and pri-mir-17-92, and ER α mRNA and pre-miR-18a, in breast cancer tissues by RT-qPCR. Levels of pri-mir-17-92 were correlated with ER α mRNA in tissues ($r^2 = 0.97$, $P = 0.0002$, Fig. 4A), further indicating that ER α regulates the expression of this primary miRNA. However pre-miR-18a was less correlated with ER α ($r^2 = 0.54$, $P = 0.21$, Fig. 4B). Next, we addressed whether pre-miR-18a, miR-18a, and miR-20a were differentially expressed in primary breast cancer tissues, comparing the average expression levels between ER α -positive and -negative tumors. Pre-miR-18a levels were significantly higher in ER α -positive tumors (2.52 ± 0.30) compared with negative tumors (0.90 ± 0.08 , $P = 0.006$, Fig. 4C), supporting our data. Moreover, expression levels of miR-18a showed no significant differences between the 2 groups of samples (Fig. 4D), indicating that impaired pre-miR-18a processing to miR-18a occurs in tumors.

MiR-18a, miR-20b, and miR-19b Negatively Modulate the ER α Transcriptional Activity After Estrogen Stimulation. Using the available miRNA target prediction software [TargetScan (29), Pictar (30), and Pita (31)], we observed whether ER α is a potential target of some or all of these miRNAs. Surprisingly, we found that miR-18, miR17/20/106, and miR-19 family members were predicted to target ER α . To experimentally validate this prediction, we chose miR-18a encoded by pri-mir-17-92, miR-19b encoded by both pri-mir-17-92 and the pri-mir-106a-363, and miR-20b encoded by the pri-mir-106a-363 (Fig. S2). First, we addressed whether these miRNAs influence ER α transcriptional activity. MELN cells (MCF-7 cells, stably transfected with a luciferase reporter gene under the control of an ERE using the β -globin promoter) were transfected with pre-miR-18a, pre-miR-20b, and pre-miR-negative control (pre-miR-n.c.). E2-stimulated reporter activity was significantly reduced when MELN cells were transfected with pre-miR-18a and pre-miR-20b, whereas the level of induction was not affected by pre-miR-n.c. (Fig. 5A). Remarkably, anti-miR-18a, anti-miR-20b, and anti-miR-19b molecules able to silence their miRNA function significantly increased reporter activity (Fig. 5B). The effect of miRNA silencing on luciferase reporter activity was similar to treatment with anti-miR-17-5p, previously reported to reduce the transcriptional activity of ER α by downregulating the coactivator AIB1 (21) (Fig. 5B).

Mir-17-5p, miR-106b, and miR-20a are able to negatively regulate AIB1 protein translation by a direct interaction with the 3'-UTR of AIB1 mRNA (21, 22, 32). Because we observed that miR-17/20/106 and the miR-18 family members potentially target ER α , we evaluated whether the reduction in ER α transcriptional activity induced by miR-20b overexpression was the result of the contemporary negative regulation of AIB1 and ER α and in addition, whether the reduction in ER α transcriptional activity induced by overexpression of miR-18a was the result of a reduction of ER α protein levels. To address if these miRNAs negatively regulate either ER α and/or AIB1, we overexpressed pre-miR-18a, pre-miR-19b, pre-miR-20b, and pre-miR-n.c. and measured protein levels. ER α was markedly reduced by the overexpression of all 3 premiRs analyzed in comparison to either untransfected or pre-miR-n.c.

Fig. 3. Pri-mir-17-92 is negatively regulated following DROSHA cleavage prolonging miRNA maturation over time. (A) Comparison of the levels of expression between pri-mir-17-92 (normalized to *GAPDH*) and miRNAs encoded from this cluster (normalized to *U47*). The mean of 3 experiments each performed in triplicate are presented, error bars represent SEM. (B) After starvation expression levels of both pri-mir-17-92 and pre-miR-18a has been analyzed by RT-qPCR using SYBR green and normalized to *U6* snRNA followed E2 treatment as indicated. The mean of 3 experiments each performed in triplicate are presented, error bars represent SEM. The asterisk indicates $P < 0.05$ in comparison to time 0 h, the double asterisk represents $P < 0.005$ in comparison to time 0 h. P values were obtained using a 2-tailed Student's t -test. (C) After starvation expression levels of miR-18a, miR-20a, and miR-342 were analyzed by RT-qPCR and normalized to *U47* snRNA. The mean of 3 experiments each performed in triplicate are presented, error bars represent SEM. The asterisk indicates $P < 0.05$ in comparison to vehicle treatment, the double asterisk represents $P < 0.005$ in comparison to vehicle treatment. P values were obtained using a 2-tailed Student's t -test. (D) Representation of miR-18a, miR-20a, and miR-342 levels (normalization to *U47* snoRNA) from 0 to 72 h of E2 treatment by RT-qPCR. The mean of 3 experiments each performed in triplicate are presented, error bars represent SEM. (E) Representation of the miR-18a (normalization to *U47*), pre-miR-18a and pri-mir-17-92 levels (normalization to *U6*) from 0 to 72 h of E2 treatment. The mean of 3 experiments each performed in triplicate are presented, error bars represent SEM.



transfected cells although the reduction with pre-miR-19b was less pronounced (Fig. 5 *C* and *D*). Furthermore, miR-20b downregulated AIB1 because transfection of pre-miR-20b into MCF7 cells reduced AIB1 protein levels (Fig. 5 *C* and *E*). On the other hand, the transfection of anti-miR-20b increased AIB1 (a dose-response was also observed here; Fig. 5*F*). Because a reduction in either *ER α* or *AIB1* mRNA levels after transfection of precursors was not observed, it appears likely that this regulation occurs at the protein translation step (Fig. S7 *A* and *B*).

To confirm whether these miRNAs directly target *ER α* , we inserted into the luciferase reporter vector 4 fragments of the 3'-UTR of *ER α* : the full length (FL, containing all of the putative miRNAs interaction sites), the first half part of the 3'-UTR (ERup for miR-17/20/106 and miR-18), a fragment containing just the putative miR-18 family interaction sites (ER1), and finally a fragment corresponding to the second half section, containing the miR-19 family interaction sites (ER2) (Fig. 6*A*). Transfection of miR-18a, miR-19b, and miR-20b, derived from pri-mir-17-92 and pri-mir-106a-363, were used to investigate direct interactions with the 3'-UTR of *ER α* constructs and we demonstrated that these miRNAs profoundly downregulate luciferase activity for the constructs containing miRNA interaction sites, but not for the ones in which these sites are absent (Fig. 6*B*). This indicates direct targeting of *ER α* by a number of miRNAs derived from these paralogous primary miRNAs. We did not observe any downregulation of luciferase reporter activity upon miR-17-5p overexpression, according to a recent report (33).

Discussion

In this study we were able to classify miRNAs upregulated by estrogen as the members encoded by the paralogous transcripts pri-mir-17-92 and pri-mir-106a-363. For individual miRNAs small changes were observed, but as multiple mature molecules derived from these primary transcripts target *ER α* and/or AIB1 this increases both the overall level of the miRNAs regulating these 2 proteins after E2 induction and the

effects of silencing; it is known that multiple molecules affecting a single target increase their inhibitory effect (34).

Changes in pri-mir-17-92 were significantly greater than the miRNAs derived from it, implicating inhibition during miRNA biogenesis: DROSHA cleavage of pri- to pre-miRNAs occurred rapidly, indicating that this step is not rate limiting (Fig. S8). Such regulation however has been described regarding let-7 family members during stem cell differentiation: LIN-28 is able to interact with pri-let-7 and/or pre-let-7 impairing its processing (12, 15). It has also been reported that c-MYC downregulates let-7 maturation increasing the transcription of LIN-28b in P493-6 B cell lines (35). Furthermore, it has been shown that the RNA binding protein KSRP, interacting with DICER, promotes the biogenesis of a subset of miRNAs comprising miR-20a and miR-106a in both HeLa and NIH 3T3 cells (36). Because we did not observe any estrogen-mediated upregulation of LIN-28 and/or any expression of LIN-28b, or any estrogen-mediated downregulation of DICER and/or KSRP in our models, this indicates that these factors are not responsible. Our data indicate that pri-mir-17-92 (not only let-7), is regulated after induction. Additionally, many expression studies note discordance in the levels of mature miRNAs derived from polycistronic precursors. Although regulation of pri-mir-17-92-derived microRNAs could not be explained by the candidate factors we tested, the apparent prevalence of regulated miRNA maturation strongly suggests involvement of additional RNA binding proteins in this process.

The importance of miRNA activity in breast cancer biology is also highlighted by the finding that a number of miRNAs show a differential expression between *ER α* positive and *ER α* negative breast cancers (37, 38). We demonstrated that pri-mir-17-92 expression is highly correlated with the level of *ER α* in breast cancers, and that pre-miR-18a derived from DROSHA-pri-mir-17-92 cleavage is also significantly more expressed in *ER α* -positive compared to *ER α* -negative tumors. This indicates that a specific increase of this pri-miRNA also occurs in physiologic conditions. It is interesting that miR-18a produced by pri-mir-17-92 is not expressed

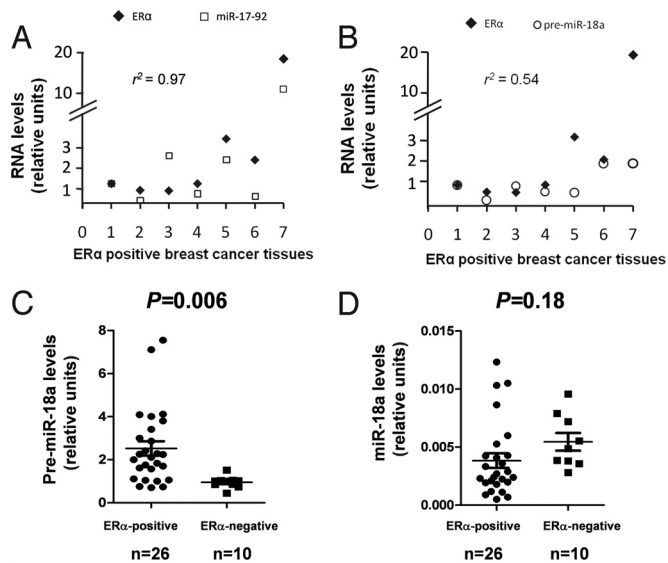


Fig. 4. ER α modulates pri-mir-17-92 in breast cancer tissues. (A) Expression levels of ER α and pri-mir-17-92 (Pearson correlation 0.97) or (B) pre-miR-18a (Pearson correlation 0.54) was measured by RT-qPCR in ER α -positive breast cancers. (C) RT-qPCR showed that expression levels of pre-miR-18a are significantly higher in ER α -positive than in ER α -negative tumors (unpaired, 2-tailed Student's *t*-test *P* = 0.006). Error bars represent SEM. (D) RT-qPCR showed that expression levels of miR-18a are not different between ER α -positive and ER α -negative tumors (unpaired, 2-tailed Student's *t*-test *P* = 0.18). Error bars represent SEM.

preferentially in ER α -positive tumors. This further suggests that ER α -positive tumors escape the inhibitory targeting of ER α caused by miRNAs by in turn downregulating DICER processing of those miRNAs during tumor progression. Here we demonstrate that the factors implicated in attenuation of miRNA processing are also active in cancer tissues themselves.

The modulation of the pri-mir-17-92 by ER α appears mediated by the c-MYC oncogene by its direct interaction with the mir-17-92 promoter. It has been reported that c-MYC directly downregulates the expression of a set of miRNAs in B cells (39). Because we have not observed any reduction of those after estrogenic stimulation, we

conclude that the upregulation of pri-mir-17-92 through ER α -c-MYC is specific to breast cells.

By forming a complex with several coactivators or corepressors, ER α transcriptionally modulates several genes implicated in cell proliferation and apoptosis such as BCL2, c-MYC, and cyclin D1. AIB1, SRC1, and TIF2 belong to the same family of coactivators that interact and collaborate with ER α in the transcriptional regulation of target genes (40). MiR-17-5p and miR-20a encoded by pri-mir-17-92, and the homologue miR-106b, downregulate the translation of AIB1 (21, 22, 32). Because following E2-mediated upregulation: (i) miR-18a, miR-19b, and miR-20b downregulate ER α and (ii) miR-20a, miR-17-5p, miR-106a, and miR-20b downregulate AIB1, we conclude that both primary transcripts are implicated in the regulation of ER α transcriptional activity upon estrogenic stimulation. Several studies have indicated that after estrogenic induction, both ER α and AIB1 are rapidly downregulated. This attenuation occurs at transcriptional, posttranscriptional, and posttranslational levels (5–8). We propose here the translational regulation by miRNAs as a further step of ER α transcriptional activity attenuation after estradiol-mediated ER α activation. Interestingly, this regulation occurs especially at a later time and in a negative feedback loop because DICER pri-mir-17-92 processing appeared inhibited after early ER α upregulation (Fig. S8).

Methods

MiRNA Microarray. Isolated RNA was labeled using the Agilent labeling kit following the manufacturer's instruction (Agilent Technologies). The Agilent human (V1) miRNA microarray platform, containing probes for 470 human (and 64 viral miRNAs from the Sanger database v9.1), was used to perform miRNA expression profiling.

RT-qPCR Assay. For RT-qPCR assays, cDNA was synthesized from 1 μ g of purified Dnase-treated RNA by the SuperScript III First-Strand cDNA synthesis system (Invitrogen); RT-qPCR was performed on a 7900HT Thermocycler using the Power SYBR green PCR master mix (both from Applied Biosystems). For detection of mature miRNAs, the TaqMan MicroRNA assay kit (Applied Biosystems) was used. Sequences of primers used are provided in Table S2.

ChIP. Cross-linked chromatin was prepared from MCF-7 cells as described previously with minor modifications (43). Aliquots of 20 μ g were incubated

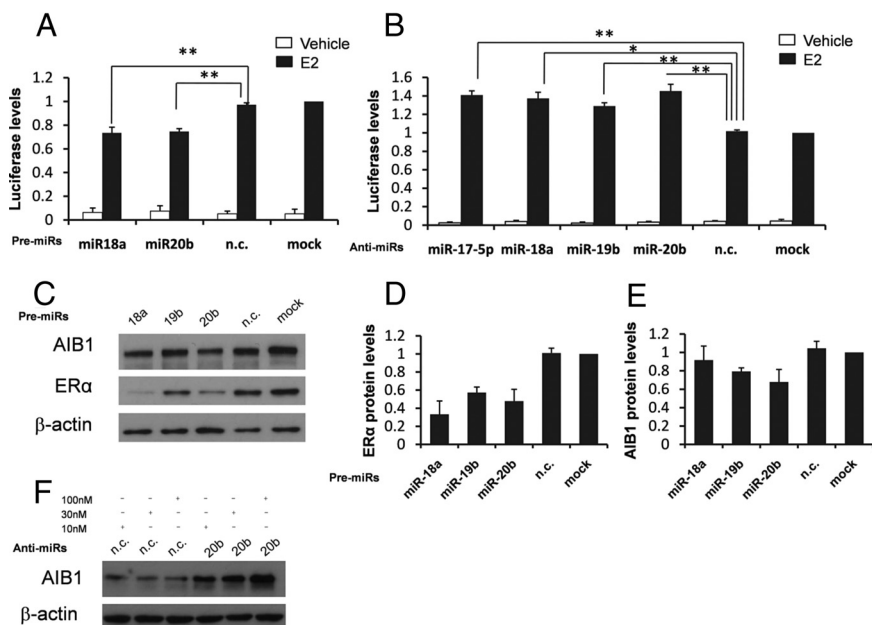


Fig. 5. MiR-18a, miR-19b, and miR-20b suppress ER α -mediated signaling. (A) Luciferase activity in MELN cells untransfected or transiently transfected for 48 h with pre-miR-18a, pre-miR-20b, and pre-miR-n.c. in the absence or presence of 10 nM of E2 for 24 h. (B) Luciferase activity in MELN cells transiently transfected for 48 h with anti-miR-17-5p, anti-miR-18a, anti-miR-19b, anti-miR-20b, anti-miR-n.c. or untransfected in the absence or presence of 10 nM of E2 for 24 h. (C) Western blot showing ER α , AIB1, and β -actin in MCF-7 cells untransfected or transiently transfected with pre-miR-18a, pre-miR-19b, pre-miR-20b, and pre-miR-n.c. (D) Densitometric analysis of ER α Western blot shown in C normalized to β -actin. (E) Densitometric analysis of AIB1 Western blot (shown in C normalized to β -actin). The mean of 3 independent experiments are presented, error bars represent SEM. (F) Western blots showing AIB1 and β -actin in MCF-7 cells transfected with anti-miR-n.c. and anti-miR-20b at 10, 30, and 100 nM concentrations. One representative experiment from 3 independent experiments is shown.

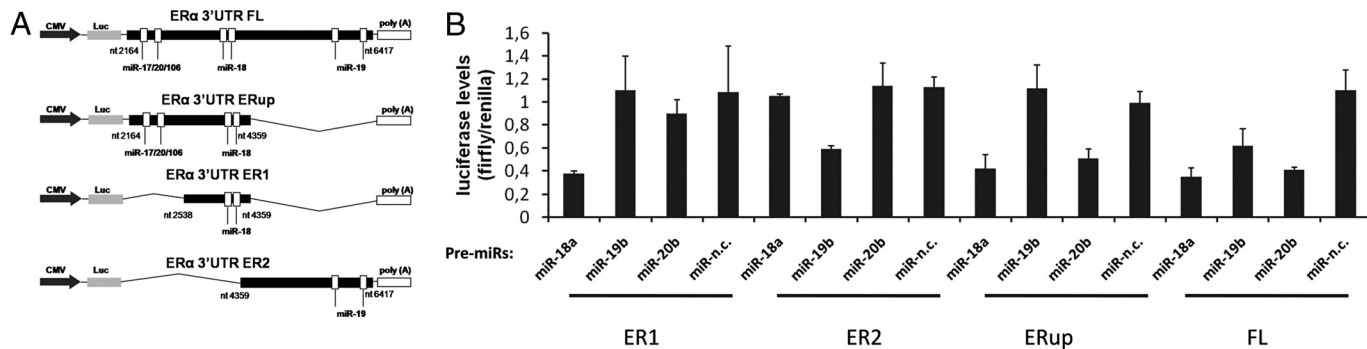


Fig. 6. ER α is directly regulated by miRNA-3'-UTR interaction. (A) Representation of the 4 different lengths of ER α 3'-UTR cloned in the pMIR-REPORT luciferase vector and miRNAs interaction sites. (B) Luciferase activity from cells cotransfected with pre-miR-18a, pre-miR-19b, pre-miR-20b, pre-miR-n.c., and different lengths of DNA fragments corresponding to the ER α 3'-UTR. Firefly luciferase was normalized for transfection levels to Renilla luciferase as indicated in experimental procedures.

overnight with 2 μ g of c-Myc (sc-764) and ER α (sc-543) antibodies (Santa Cruz Biotechnology) or without (mock controls) in a total volume of 1 mL and immunoprecipitated. Triplicate samples of 5 μ L of immunoprecipitated genomic DNA were amplified by real time PCR. Values are expressed as fold of enrichment with respect to input DNA. Primer sequences used in this assay are listed in Table S2.

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