RESEARCH ARTICLE

Genome-wide miRNA response to anacardic acid in breast cancer cells

David J. Schultz¹, Penn Muluhngwi², Negin Alizadeh-Rad², Madelyn A. Green², Eric C. Rouchka³, Sabine J. Waigel⁴, Carolyn M. Klinge²*

¹ Department of Biology, University of Louisville, Louisville, Kentucky, United States of America, ² Department of Biochemistry & Molecular Genetics, University of Louisville School of Medicine, Louisville, Kentucky, United States of America, ³ Bioinformatics and Biomedical Computing Laboratory, Department of Computer Engineering and Computer Science, Louisville, Kentucky, United States of America, ⁴ Department of Medicine, University of Louisville School of Medicine, Louisville, Kentucky, United States of America

* carolyn.klinge@louisville.edu

Abstract

MicroRNAs are biomarkers and potential therapeutic targets for breast cancer. Anacardic acid (AnAc) is a dietary phenolic lipid that inhibits both MCF-7 estrogen receptor α (ERα) positive and MDA-MB-231 triple negative breast cancer (TNBC) cell proliferation with IC₅₀ values of 13.5 and 35 μM, respectively. To identify potential mediators of AnAc action in breast cancer, we profiled the genome-wide microRNA transcriptome (microRNAome) in these two cell lines altered by the AnAc 24:1n5 congener. Whole genome expression profiling (RNA-seq) and subsequent network analysis in MetaCore Gene Ontology (GO) algorithm was used to characterize the biological pathways altered by AnAc. In MCF-7 cells, 69 AnAc-responsive miRNAs were identified, e.g., increased let-7a and reduced miR-584. Fewer, i.e., 37 AnAc-responsive miRNAs were identified in MDA-MB-231 cells, e.g., decreased miR-23b and increased miR-1257. Only two miRNAs were increased by AnAc in both cell lines: miR-612 and miR-20b; however, opposite miRNA arm preference was noted: miR-20b-3p and miR-20b-5p were upregulated in MCF-7 and MDA-MB-231, respectively. miR-20b-5p target EFNB2 transcript levels were reduced by AnAc in MDA-MB-231 cells. AnAc reduced miR-378g that targets VIM (vimentin) and VIM mRNA transcript expression was increased in AnAc-treated MCF-7 cells, suggesting a reciprocal relationship. The top three enriched GO terms for AnAc-treated MCF-7 cells were B cell receptor signaling pathway and ribosomal large subunit biogenesis and S-adenosylmethionine metabolic process for AnAc-treated MDA-MB-231 cells. The pathways modulated by these AnAc-regulated miRNAs suggest that key nodal molecules, e.g., Cyclin D1, MYC, c-FOS, PPARγ, and SIN3, are targets of AnAc activity.

Introduction

microRNAs (miRNAs) are ~ 22 nt noncoding RNAs that basepair with complementary sequences in the 3'UTR of their target mRNAs within the RNA-induced silencing complex
(RISC) resulting in translational repression and, in many cases, degradation of the target transcript [1]. The selection of the miR-5p or miR-3p arm for inclusion into the RISC complex for 3’-UTR mRNA target selection is determined by the AGO protein [2]. Each miRNA can have hundreds of gene targets resulting in coordinate regulation of cellular pathways [3]. Dysregulated miRNAs in breast cancer contribute to aberrant regulation of cell cycle, differentiation, metabolism, and cancer stem cell (CSC) survival (reviewed in [1, 4–10]).

Anacardic acid (AnAc) is a collective term for the mixture of 6-alkylbenzoic acid congeners that are produced in a number of plants [11]. AnAc has a variety of activities including inhibition of histone acetyltransferase (HAT) activity (reviewed in [12]). Previously, we reported that a specific congener AnAc 24:1n5 acts as a nuclear receptor alternate site modulator (NRAM) to inhibit breast cancer cells in an estrogen receptor (ER)-dependent manner by interfering with ER-DNA binding [13]. In addition, AnAc 24:1n5 also inhibited the growth of MDA-MB-231 triple negative breast cancer (TNBC, i.e., ERα negative, progesterone receptor negative, and ERBB2 negative) cells, albeit at a higher IC50 and through an undefined mechanism [13]. Thus, we hypothesize that additional molecular targets, including miRNAs, are affected by AnAc in breast cancer cells. High Throughput Sequencing (HTS) provides a comprehensive overview of biological processes and pathways affected by AnAc; thus, offering novel insights into potential mechanisms of action and cellular targets.

The goal of this study was to use RNA-Seq to comprehensively identify alterations in miRNAs in ERα-positive, luminal A MCF-7 and MDA-MB-231 TNBC breast cancer cell lines treated with AnAc 24:1n5. Our results identified common and divergent mRNA transcripts down- or up-regulated by AnAc. The pathways modulated by these miRNAs suggest that key nodal molecules, e.g., Cyclin D1, SMAD, SP1, MYC, c-FOS, PPARγ, BCL2, FOXO3A, MDA2, and SIN3, are targets of AnAc activity.

Materials and methods
Materials
AnAc 24:1n5 was purified to greater than 95% as previously reported [13, 14]. For our experiments, AnAc 24:1n5 (AnAc) was dissolved in ethanol (EtOH); thus, EtOH was used as a vehicle control.

Cell culture and treatments
MCF-7 and MDA-MB-231 cells were purchased from American Type Tissue Collection (ATCC, Manassas, VA). Cells were used at less than 9 passages from ATCC. MCF-7 and MDA-MB-231 cells were maintained in IMEM (Cellgro, Manassas, VA) containing 5% fetal bovine serum (FBS, Atlanta Biologicals, Lawrenceville, GA) and 1% Penicillin/Streptomycin (Cellgro). Cells were grown in phenol red-free IMEM (ThermoFisher) medium containing 5% dextran coated charcoal (DCC)-stripped FBS (hormone-depleted medium) for 48 h prior to treatment with established IC50 concentrations of AnAc 24:1n5: 13.5 μM for MCF-7 and 35.0 μM for MDA-MB-231 cells [13] for 6 h and was replicated in three separate experiments.

RNA isolation and RNA seq
RNA was isolated from MCF-7 and MDA-MB-231 breast cancer cells using the Exiqon miRCURY™ RNA Isolation kit (Woburn, MA, USA). RNA concentration was assessed using a NanoDrop spectrophotometer.
For miRNA RNA-seq

The Truseq Small RNA kit (Illumina) was used to prepare miRNA libraries from 1 μg total RNA. Each Library was individually gel purified on a Novex TBE 6% gel and resuspended in 10μL 10mM Tris-Cl, pH 8.5. Libraries were validated and quantitated by running 1μL on the Agilent Technologies 2100 Bioanalyzer DNA High Sensitivity Chip. 36-cycle single sequencing reads were generated on the Illumina NextSeq500 instrument utilizing the 500 Mid-output v2 (75 cycle) sequencing kit. The resulting samples were divided into 48 FASTQ [15] single-end raw sequencing files representing four conditions: MCF-7 control, MCF-7 treated with AnAc 24:1n5 (MCF-7 AnAc), MDA-MB-231 control, and MDA-MB-231 treated with AnAc 24:1n5 (MDA MB-231 AnAc). These raw data of our RNA-seq are available at Gene Expression Omnibus (GEO) database: accession number GSE78011.

Differential miRNA expression analysis

A total of three biological replicates for each treatment were analyzed, with four flow cell lanes per replicate. Raw sequence data files were downloaded from Illumina’s BaseSpace (https://basespace.illumina.com/) onto the KBRIN server for analysis the miRDeep2 [16] and edgeR [17]. Each of the four single-end raw. FASTQ files for each replicate (representing the four flow cells) was concatenated into one single-end. FASTQ file using the unix cat command.

Quality control (QC) of the raw sequence data was performed using FastQC (version 0.10.1) [18]. The FastQC results indicated sequence trimming was not necessary since the minimum quality value for all samples was well above Q30 (1 in 1000 error rate) (data not shown).

Given that this is a miR sequencing project, preliminary adapter trimming was performed on each of the samples using a custom file adaptersToTrim.fa which contains a subset of the Illumina TruSeq Small RNA adapter and primer sequences taken from https://support.illumina.com/content/dam/illumina-support/documents/documentation/chemistry_documentation/experiment-design/illumina-adapter-sequences_100000002694-00.pdf

Sequences were trimmed of the adapters with Trimmomatic v0.33 [19].

The trimmed sequences were directly aligned to the human hg19 reference genome assembly using the mapper.pl wrapper of the miRDeep2 package (v 0.0.7) [16]. This script used bowtie (version 1.1.1) [20], generating alignment files in arf format. The aligned sequences were then used as inputs into the miRDeep2 package and the script quantifier.pl. In addition, this script used the mirBase release 21 [21] mature miRNA and miRNA hairpin sequences downloaded from ftp://mirbase.org/pub/mirbase/CURRENT/. The result was a file containing the number of reads mapping to each of the 2,822 human (hsa) miRs for the specific sample. After quantification, the resulting counts for each miR in each sample were combined into a reads matrix. This was accomplished using a custom perl script, createReadMatrix.pl. Differentially expressed miRs were determined using edgeR [17] and a customized R script, Schultz-Klinge. miRNA.R. Using a p-value cutoff of 0.05, the number of differentially expressed miRs in each comparison is shown in Table 1.

In silico network analysis

We performed pathway and network analysis of differentially expressed genes in MetaCore™ version 6.27 (GeneGO, Thomson Reuters, New York, N.Y.). MetaCore™ is a web-based software suite for multiple applications in systems biology including RNA-seq analysis as used here. MetaCore™ analyses are based on MetaBase (http://metadatabase.org/), a 100% manually-curated integrated database of mammalian biology that contains over 6 million experimental findings on protein-protein, protein-DNA, protein-RNA, and protein-compound interactions; metabolic and signaling pathways; and other information [22].
Generation of heatmaps: Files of miRNAs significantly altered by AnAc treatment in each cell line were imported into Partek software Version 6.6 (Partek Inc., St Louis, MO.) and Partek Genomic Suite™ was used to generate heatmaps (Fig 1, S1 and S2 Figs). Each hierarchical clustering was created using Euclidean distance as similarity measure for genes and samples. We noted that one of the three MCF-7 AnAc samples appeared to behave as a hybrid between the other two AnAc treated and three control (EtOH)-treated samples (S2 Fig).

RNA isolation, RT-PCR and quantitative real-time PCR (qPCR) of miRNAs and mRNAs

Cell growth, treatment and RNA isolation and quantification/quality assessment were performed as described above. For miRNA, RNA was converted to cDNA using the Taqman® miRNA Reverse Transcription kit (PE Applied Biosystems). For mRNA, RNA was converted to cDNA using the High Capacity cDNA Reverse Transcription kit (PE Applied Biosystems). Primers for hsa-miR-268g, hsa-miR-612, hsa-miR-20b-5p, and hsa-miR-20b-3p were purchased from TaqMan (Advanced miRNA assays) and RNU48 (TaqMan) was used as the reference for normalization [23]. Primers for VIM (Vimentin) [24]: Forward, 5’-GACAATGCGTC TCTGGCACGTCTT-3’; Reverse, 5’-TCCTCCGCTCCTGCAG GTTCTT-3’; for ZFP36L1 (ZFP36 Ring Finger Protein Like 1, aka ERF1 and BRF1) [25]: Forward, 5’-AGGATGACCAC CACCCTCGTGTCT-3’; Reverse, 5’-CCC CCTGCACTGGGAGCACTA-3’, and for GAPDH [26] were purchased from IDT. qPCR was performed using ABI Viia 7 (Life Technologies) with each reaction run in triplicate. The comparative threshold cycle (Ct) method (2–ΔΔCT) was used to determine fold change relative to vehicle treated or control transfected cells [27].

Table 1. Differentially expressed miRNAs (DEmiRs). The log2-fold change with zero value in the control conditions was arbitrarily set to one and the maximum log2-fold change value and those with zero value in the treatment conditions were arbitrarily set to the minimum log2-fold change value of minus one. The number of differentially expressed genes in each comparison is shown and the number of upregulated genes indicated with the upward arrow and downregulated genes indicated by downward arrow.

| Comparison                  | Cutoff | Number of DEmiRs |
|-----------------------------|--------|------------------|
| MCF-7 AnAc vs. control      | P ≤ 0.05 | 69 (48, 21)     |
| MDA-MB-231 AnAc vs. control | P ≤ 0.05 | 37 (15, 22)     |
| All Cells AnAc vs. All Cells control | P ≤ 0.05 | 25 (13, 12)     |
| All MCF-7 vs. All MDA-MB-231 control | P ≤ 0.05 | 795 (510, 285)  |

Z All Cells is the sum of both cell lines
Y Sum of AnAc treatment and control for each cell line

https://doi.org/10.1371/journal.pone.0184471.t001

Transient transfection

MCF-7 and MDA-MB-231 cells were transiently transfected for 24 h with miR-612 mimic, miR-612 inhibitor, Anti-miR™ negative control #1, or mirVANA™ miRNA mimic negative control #1 (all from Ambion, Life Technologies, Thermo Fisher Scientific, Carlsbad, CA, USA), using Lipofectamine RNAiMAX transfection reagent (Invitrogen, Thermo Fisher Scientific) and Opti-MEM® Reduced Serum Medium (Invitrogen, Thermo Fisher Scientific). After 24 h of transfection, cells were treated with ethanol (EtOH, vehicle control) or 13.5 or 35 μM AnAc, for MCF-7 and MDA-MB-231 respectively, in phenol red-free IMEM medium containing 5% DCC-stripped FBS for 48 h prior to MTT assay (CellTiter 96, Promega, Madison, WI, USA). Two separate experiments were performed with quadruplicate wells within each experiment. For analysis of miR-612 expression in transfected cells, the medium was changed 24 h
after transfection as above, without any treatment and RNA was harvested (see above) a total of 72 h post transfection, i.e., at the same time the MTT assay was performed for qPCR of miR-612 using RNU48 as a control (see above).

Results and discussion

RNA-seq analysis of AnAc-regulated miRNAs

MCF-7 luminal A (ERα+) and MDA-MB-231 TNBC (triple negative breast cancer) cells were incubated in hormone-depleted medium for 48 h prior to a 6 h treatment with the previously established IC₅₀ concentrations of AnAc 24:1n5 for MCF-7 (13.5 μM) and MDA-MB-231 (35.0 μM) cells [13]. The 6 h time point was selected based on transcriptome studies in MCF-7 cells to identify primary gene targets [28] and because AnAc 24:1n5 has no overt effect on the viability of either MCF-7 or MDA-MB-231 at that time [13, 29]. The goal was to identify early miRNA changes in response to AnAc 24:1n5 in each cell line. For target analysis, only miRNA transcripts that showed a log2 fold-change greater than 1 (or -1 for repressed miRNAs) were included.

Differentially expressed miRNAs (DEmiRs) were identified for four pairwise comparisons (MCF-7 AnAc-treated vs. MCF-7 control; MDA-MB-231 AnAc-treated vs. MDA-MB-231 control; MCF-7 and MDA-MB-231 AnAc treated vs. MCF-7 and MDA-MB-231 control; MDA-MB-231 AnAc treated and control vs. MCF-7 AnAc treated and control) using the tuxedo suite of programs including cufflinks and cuffdiff (version 2.2.1) [30, 31]. Significant DEmiRs with fold-change and p values are listed in S1 and S2 Tables. The number of DEmiRs in each comparison is shown in Table 1. Tables 2–5 list the AnAc-regulated miRNAs in MCF-7 and MDA-MB-231 cells, their genomic location and host gene (if applicable), information about their relevance in breast or other cancers and their experimentally verified, i.e., bona
The expression of more miRNAs was significantly changed in response to AnAc in MCF-7 cells vs MDA-MB-231 cells (Figs 1 and 2). The heatmap shows that MCF-7 and MDA-MB-231 cells have different responses to AnAc with MDA-MB-231 cells showing less change in response to AnAc compared with MCF-7 cells (Fig 1). These data suggest that AnAc selectivity alters miRNA transcript expression in these two cell lines through mostly non-overlapping mechanisms.

As shown in the Venn Diagrams of Fig 2, there were no common downregulated miRNAs in AnAc-treated MCF-7 and MDA-MB-231 cells. Only two miRNA were commonly upregulated by AnAc in both MCF-7 and MDA-MB-231 cells: miR-20b and miR-612 (Fig 2, Table 2).

### Table 2. miRNAs upregulated by AnAc in both MCF-7 and MDA-MB-231 cells.

| miRNA   | Role in breast or other cancers                                                                 | Verified targets                           |
|---------|--------------------------------------------------------------------------------------------------|--------------------------------------------|
| miR-612 | Chr11, intergenic. Tumor suppressor miRNA in HCC tumors, cells and xenograft tumors (35, 36). Downregulated in colorectal cancer tumors and cells and acts as a tumor suppressor (37). | For 5p: AKT2 [37] SP1 [38]                |
| miR-20b-3p | ChrX, encoded by the miR-106a-363 cluster is intergenic (39). Oncogenic activity, i.e., stimulates soft agar colony formation in NIH-3T3 cells (39). Lower expression in taxol-resistant breast tumors and cells (40). Expression is stimulated by EGR1 [41]. | For 3p: ESR1 [42]; EPAS1 [43]; NCOA3 [40]; BRCA1, PTEN [41] |
| miR-20b-5p | MDA-MB-231                                                                                       |                                            |

For 5p: ARID4A and MYLIP [39]; HIF1A and VEGF [44]; PPARG, BAMBI, CRIM1 [45] EPHB4 and EFNB2 [46]; PTEN [47]; SOS1 and ERK2 [48].

As shown in the Venn Diagrams of Fig 2, there were no common downregulated miRNAs in AnAc-treated MCF-7 and MDA-MB-231 cells. Only two miRNA were commonly upregulated by AnAc in both MCF-7 and MDA-MB-231 cells: miR-20b and miR-612 (Fig 2, Table 2).

The common GO Processes for upregulated miR-20b and miR-612 were identified by MetaCore™ analysis and listed in Fig 2; however no matches between genes/proteins for miR-20b and miR-612 were identified in Pathway Maps by MetaCore analysis. Interestingly, AnAc increased miR-20-3p in MCF-7 and miR-20-5p in MDA-MB-231 cells. This suggests that distinct miR-20b targets would be expected to be regulated in response to AnAc upregulation of miR-20b-3p versus miR-20-5p in the two cell lines. The selection of which mature miRNA 5p or 3p arm is dominant is determined by thermodynamic and structural properties of the processed pre-miRNA-duplex AGO protein (reviewed in [32]). The functional consequences of arm selection are therefore distinct. The exact mechanism of miRNA Induced Silencing Complex (miRISC) assembly remains elusive and includes a human miRNA loading complex containing the ds-pre-miRNA, DICER1, TRBP2 and miRNA-free AGO protein as its components. [33]. Recent studies in Huh7 human hepatoma cells showed that an increase in target genes, i.e., SLC7A1 (CAT-1), increased the processing of pre-miR-122 to miR-122, implying that increases in target miRNA levels can promote miRNA biogenesis [33]. Whether this is true for other cells and miRNAs remains to be examined. The MetaCore network enrichment analysis of the miRNAs upregulated in AnAc-treated MCF-7 vs. MDA-MB-231 cells identified “Cellular response to inorganic substance” as the top GO process (S3A Fig). The network analyses for miR-20b and miR-612 are shown in S3A and S3B Fig.

There is only one previous examination of miRNAs, mRNAs, and IncRNAs in MCF-7 and MDA-MB-231 cells, but that study used a microarray expression profiling [167] rather than an...
unbiased RNA-sequencing approach. None of the AnAc-regulated miRNAs was among the miRNAs more highly expressed in MCF-7 compared with MDA-MB-231 cells [167]. In contrast, miR-4284 was more highly expressed in MDA-MB-231 cells [167] and we observed that AnAc decreased miR-4284 in MDA-MB-231 cells (Table 5). The role of miR-4284 in breast cancer is unknown and there are no validated targets of miR-4284, although microRNA.org lists 7,891 putative targets.

miRNAs downregulated by AnAc in MCF-7 cells

Twenty-one miRNAs were downregulated by AnAc in MCF-7 cells (Table 3). miRNAs are encoded within a gene (intronic or exonic) or are intergenic (reviewed in [168]). miRNAs can be regulated independently or are cotranscribed with their host gene (reviewed in [8]). To examine if the miRNA host gene was downregulated by AnAc in MCF-7 cells we searched Table 3. miRNAs downregulated by AnAc in MCF-7 cells. The genomic location of each miRNA was identified in miRAD http://bmi.ana.med.uni-muenchen.de/miriad/ [34]. Verified targets are those experimentally validated targets of the indicated miRNA as demonstrated by 3'-UTR luciferase reporter assay. Since many publications do not include whether the 5p or 3p arm of the miRNA was studied, if the sequence of the miRNA was provided, it was searched in miRBase.org to identify which arm was used in the target gene 3'-UTR luciferase reporter assay.

| miRNA   | Role in breast or other cancers                                                                 | Verified targets                      |
|---------|-------------------------------------------------------------------------------------------------|---------------------------------------|
| miR-378g| Chr1, host gene LINCO1057 [49]. Target of c-Myc [50]. High miR-378 promotes cancer stem cell (CSC) properties, increased cell survival and colony formation; acts as on oncomiR; correlates with increased SOX2 [51]. Induced during adipogenesis by increasing transactivation by C/EBPα and C/EBPβ [52]. | VIM [51] TOB2 [50] SUFU and TUSC2 [53] HDAC4 [54] |
| miR-509-1-3p, -2-3p, -3-3p | miR-509-1, -2, and -3 are in ChrX, host gene LOC107984060. Tumor suppressor miRNA [55, 56]. Inhibited by E2 in MCF-7 cells [57]. Anti-metastatic miRNA. The expression of miR-509 was reported to be attenuated in brain metastatic lesions compared to their enrichment in primary breast tumors [58]. | For 3p: RHOC [58]; For 5p: YWHAG [59] |
| miR-513b-5p | ChrX, host gene LOC107984060. Cluster with miR- 506, 507, 208, 509-1, -2, -3, 514b; Acts as a tumor suppressor in gastric cancer cells [60] | For 5p: HMGB3 [60] |
| miR-548, 548j-5p, 548l | MIR548J: Chr22: host gene HMGB1P10; MIR548L: Chr11 host gene ANKRD48. miR-548J functions as a metastasis promoter in breast cancer cells [61]. | For 3p: RHOC [58]; miR-548j-5p: TNS1 [61] |
| miR-597-3p | Chr8, host gene TNKS downregulated in colorectal cancer [63] | miR-548L: AKT [62]; miR-548j-5p: TNS1 [61] |
| miR-1238-3p | Chr19, host gene ARG4D. no publications in PubMed | For 3p: LHX2 [64] |
| miR-1915-3p | Chr10, host gene CASC10. Processing of pri-miR-1915 to pre-miR-1915 is increased by p53 [65]. | For 3p: BCL2 [66] |
| miR-3146 | Chr7, host gene TWISTNB. no publications in PubMed | miR-3146 |
| miR-4430 | Chr2 intergenic. no publications in PubMed | miR-4430 |
| miR-5002-5p | Chr3, host gene KALRN. no publications in PubMed | miR-5002-5p |
| miR-5187-5p | Chr2, host gene TOMM40L. no publications in PubMed | miR-5187-5p |
| miR-6717-5p | Chr14, host gene NDRG2. no publications in PubMed | miR-6717-5p |
| miR-6773-3p | Chr16, host gene ESRP2. no publications in PubMed | miR-6773-3p |
| miR-6804-5p | Chr19, host gene PPP6R1. no publications in PubMed | miR-6804-5p |
| miR-6814-5p | Chr21, host gene RIPK4. no publications in PubMed | miR-6814-5p |
| miR-6838-5p | Chr7, host gene PLOM. no publications in PubMed | miR-6838-5p |
| miR-6873-3p | Chr6, host gene WDR46. no publications in PubMed | miR-6873-3p |

https://doi.org/10.1371/journal.pone.0184471.t003
Table 4. miRNAs upregulated by AnAc MCF-7 cells. The genomic location of each miRNA was identified in miRAD http://bmi.ana.med.uni-muenchen.de/miRAD [34]. Verified targets are those experimentally validated targets of the indicated miRNA as demonstrated by 3'-UTR luciferase reporter assay in the cited reference. Since many publications do not include whether the 5p or 3p arm of the miRNA was studied, if the sequence of the miRNA was provided, it was searched in miRBase.org to identify which arm was used in the target gene 3'-UTR luciferase reporter assay.

| miRNA | Role in breast or other cancers | Verified targets |
|-------|---------------------------------|------------------|
| Let-7a-2-3p | Chr11; intergenic. Lower expression metastatic breast tumors [67]. Downregulated by E2 treatment in MCF-7 cells [68]. Decreased expression with breast tumor grade and upregulated KEGG pathway targets have roles in cancer-related pathways, including cycle (MCM2), Jak-STAT (SOCS1), MAPK (STMN1), PPAR signaling (ME1) [69]. Transfection of MCF-7 and MDA-MB-231 cells with let-7a mimics inhibits cell proliferation, colony formation, cell migration and invasion and HMG1A1 protein [70]. | None experimentally validated for 3p. |
| miR-378j | Chr17, host gene DDX52. no publications in PubMed | |
| miR-450a-1-3p | ChrX, intergenic, clustered with miR-424, 503, 542, 450a-2, and 450b. No publications relating to miR-450a-1 in PubMed, but miR-450a expression was higher in lymph node metastasis in breast cancer [71] and in endometrial carcinomas [72]. | None validated for 3p. For 5p: DNMT3a [73] |
| miR-520a-5p | Chr19, intergenic. miR-520a-3p inhibits proliferation by targeting HOXD8 in non-small cell lung cancer | None experimentally validated for 5p. For 5p: CCND1 and CD44 [74] |
| miR-520d-5p | Chr19, intergenic. involved in HER2-receptor-related differentiation through undefined mechanisms [75]. Overexpression by lentiviral-miR-520d infection of human HLF and Huh7 hepatoma cells converted the cells to non-tumorigenic and less differentiated normal stem cells, but no miRNA target genes were validated [76]. Acts as a tumor suppressor in colorectal cancer [77]. | For 5p: CTHRC1 [77] |
| miR-548ag-1 | Chr4, intergenic. no publications in PubMed | |
| miR-551b-5p | Chr3, intergenic. Downregulated by E2 in MCF-7 cells [57]. Down-regulated in aggressive breast tumors [78]. Upregulated in TAM-resistant MCF-7 cells [79]. Upregulated in serum samples from prostate cancer patients compared with benign prostatic hyperplasia patients [80]. Upregulated in recurrent epithelial ovarian cancer (OvCa) [81]. Upregulated in OvCa stem cells, promotes proliferation, invasion, and chemoresistance [82]. | None experimentally validated for 5p. For 5p: FOXO3 and TRIM31 [82] |
| miR-562 | Chr2, host gene DIS3L2. Upregulated in serum samples from prostate cancer patients with disseminated disease compared with benign prostatic hyperplasia patients [80]. | EYA1 [83]; IL22 [84] |
| miR-663a | Chr20, intergenic. Upregulated by E2 in ECC-1 cells [85]. Transcription increased by ZNF224 [86]. Acts as a tumor suppressor and is downregulated in gastric [87], colorectal [88], prostate [89], breast [90], pancreatic [91], non-small cell lung cancer [92]. Transcription factor Ets-2 binds the miR-663 promoter and stimulates transcription in prostate cancer cells [89]. | TP53 (P53) and CDKN1A (p21) [86] JUND [92] TGFB1 [91] HMG2A [90] |
| miR-664b-5p | ChrX, host gene DCK1. Acts as a tumor suppressor in osteosarcoma [93] and as an oncomiR in T-cell acute lymphoblastic leukemia [94] and cervical cancer [95]. | None experimentally validated for 5p. For 3p: FOXO4 [96]; MAT1A [97]; PLP2 [98]; SOX7 [93] |
| miR-921 | Chr1, host gene FAM78B. Downregulated in bladder cancer [99]. | CBRI [100] |
| miR-1229-5p | Chr 5, host gene MGAT4B. Upregulated in serum of colorectal cancer patients [101]. Overexpressed in breast cancer and correlated with poor prognosis for patients [102]. | None experimentally validated for 5p. For 5p: GSK3B, APC and ICAT [102]. |
| miR-1287-3p | Chr10, host gene PYROXD2. Downregulated in MCF-7 cells that are aromatase inhibitor resistant [103]. Hypermethylated in cervical cancer [104], downregulated in larynx carcinoma [105], anaplastic astrocytomas and/or glioblastomas [106]. | None experimentally validated for 3p. For 5p: ATF6B [107] |
| miR-1976 | Chr1, host gene RPS6KA1; Acts as a tumor suppressor in NSCLC [108]. | PLCE1 [108] |
| miR-3132 | Chr2, host gene TMEM198; no publications in PubMed | |
| miR-3195 | Chr20, intergenic; no publications in PubMed | |
| miR-3960 | Chr9, intergenic. The lncRNA HOTAIIR1 competitively binds to miR-3960 and regulates hematopoiesis [109]. | HOXA2 [110] |
| miR-4436b-1-3p | Chr2, host gene MALL. Appears to be a strong pathogenic candidate in Autism Spectrum Disorders (ASDs) [111]. | |
| miR-4436b-2-3p | Chr2, intergenic. Appears to be a strong pathogenic candidate in ASDs [111]. | |

(Continued)
| miRNA          | Role in breast or other cancers                                                                                                                                                                                                                                                                                                                                 | Verified targets                                                                 |
|---------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------|
| miR-4485-5p   | Chr11, host gene MTRNR2L8. Is transported into mitochondria and inhibits 16S rRNA processing and mitochondrial protein synthesis [112]. Acts as a tumor suppressor in MCF-7 cells in vitro and in MDA-MB-231 cells in xenograft studies in mice [112].                                                                 |                                                                                  |
| miR-4516      | Chr16, host gene PKD1. Upregulated by fine particulate matter (PM2.5) treatment of A549 NSCLC cells [113]. High expression was associated with infiltrative growth of follicular variant of papillary thyroid carcinomas [114].                                                 | STAT3 [115], RPL37 [113]                                                        |
| miR-4634      | Chr5, intergenic. One of five miRNAs in serum that detects breast cancer [116]                                                                                                                                                                                                                                                                               |                                                                                  |
| miR-4659a-3p  | Chr8, host gene AGPAT5. no publications in PubMed                                                                                                                                                                                                                                                                                                           |                                                                                  |
| miR-4661-3p   | Chr8, host gene LRRC69. miR-466i upregulates both mRNA and protein expression of IL-10 in macrophages by binding to the 3' UTR of IL10 and inhibiting RNA binding protein-induced transcript degradation [117].                                               |                                                                                  |
| miR-4675      | Chr10, intergenic. no publications in PubMed                                                                                                                                                                                                                                                                                                                |                                                                                  |
| miR-4677-3p   | Chr11, host gene STIM1. no publications in PubMed                                                                                                                                                                                                                                                                                                            |                                                                                  |
| miR-4692      | Chr11, no publications in PubMed                                                                                                                                                                                                                                                                                                                             |                                                                                  |
| miR-4695-3p   | Chr1, host gene ALDH4A1. no publications in PubMed                                                                                                                                                                                                                                                                                                          |                                                                                  |
| miR-4701-3p   | Chr12, host gene ADCY6. Downregulated in papillary thyroid carcinoma (PTC) [118].                                                                                                                                                                                                                                                                              |                                                                                  |
| miR-4741      | Chr18, host gene RBBP8. Downregulated in serum of HCC patients treated with transarterial chemoembolisation (TACE) with bad response to TACE [119].                                                                                                                                                     |                                                                                  |
| miR-4756-5p   | Chr20, host gene BCAS1. no publications in PubMed                                                                                                                                                                                                                                                                                                            |                                                                                  |
| miR-5008-3p   | Chr1, host gene WNT9A. no publications in PubMed                                                                                                                                                                                                                                                                                                             |                                                                                  |
| miR-5585-5p   | Chr1, host gene TMEM39B. no publications in PubMed                                                                                                                                                                                                                                                                                                           |                                                                                  |
| miR-6087      | ChrX, intergenic. Identified in human mesenchymal stem cells and downregulated during endothelial differentiation [120]. Uregulated in intermediate monocytes [121].                                                                                                                                   | ENG [120]                                                                        |
| miR-6126      | Chr16, host gene NAA60. Exosomal tumor suppressor is downregulated in ovarian cancer tumors and is released from ovarian cancer cells [122].                                                                                                                                                                | ITGB1 [122]                                                                     |
| miR-6131      | Chr5, host gene ROPN1L. no publications in PubMed                                                                                                                                                                                                                                                                                                             |                                                                                  |
| miR-6151-5p   | Chr19, host gene CALR. no publications in PubMed                                                                                                                                                                                                                                                                                                             |                                                                                  |
| miR-6726-5p   | Chr1, host gene ACAP3. no publications in PubMed                                                                                                                                                                                                                                                                                                             |                                                                                  |
| miR-6757-5p   | Chr12, host gene TNS2. no publications in PubMed                                                                                                                                                                                                                                                                                                             |                                                                                  |
| miR-6813-3p   | Chr20, host gene RGS19. no publications in PubMed                                                                                                                                                                                                                                                                                                             |                                                                                  |
| miR-6857-5p   | ChrX, host gene SMC1A no publications in PubMed                                                                                                                                                                                                                                                                                                              |                                                                                  |
| miR-6868-5p   | Chr17, host gene EXOC7. no publications in PubMed                                                                                                                                                                                                                                                                                                             |                                                                                  |
| miR-6874-5p   | Chr7, host gene RNF216. no publications in PubMed                                                                                                                                                                                                                                                                                                             |                                                                                  |
| miR-7151-5p   | Chr10, host gene CTNNA3. no publications in PubMed                                                                                                                                                                                                                                                                                                             |                                                                                  |
| miR-8079      | Chr13, intergenic. no publications in PubMed                                                                                                                                                                                                                                                                                                                 |                                                                                  |
| miR-8089      | Chr5, host gene BTNL9. no publications in PubMed                                                                                                                                                                                                                                                                                                             |                                                                                  |
Table 5. miRNAs downregulated by AnAc in MDA-MB-231 cells. The genomic location of each miRNA was identified in miRAD [http://bmi.ana.med.uni-muenchen.de/mirad/][34]. Verified targets are those experimentally validated targets of the indicated miRNA as demonstrated by 3'-UTR luciferase reporter assay. Since many publications do not include whether the 5p or 3p arm of the miRNA was studied, if the sequence of the miRNA was provided, it was searched in miRBase.org to identify which arm was used in the target gene 3'-UTR luciferase reporter assay.

| miRNA         | Role in breast or other cancers                                                                 | Verified targets                                                                 |
|---------------|-------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------|
| miR-23b-5p    | Chr9, host gene C9orf3. OncomiR, induced by c-Myc [123]. Lower expression in MDA-MB-231 than MCF-7 cells [124]. Stimulated by E2 in ERβ-transfected MCF-7 cells [125]. Involved in regulation of cytoskeletal remodeling and motility [126, 127]. Primary breast tumor expression of miR-23b correlates with lung metastasis [128]. Metastatic breast cancer cells in patient bone marrow had increased miR-23b [129]. Increased in MCF-7 cell derived exosomes after docosahexaenoic acid (DHA) treatment [130]. miR-23a is 2.5-fold higher in MDA-MB-231 than MCF-7 cells and downregulates CDH1 resulting in hyperactivation of Wnt/-catenin signaling, EMT, and metastasis [131].       | For 5p: PRODH [132]     |
| miR-141-3p    | Chr12, intergenic and clustered with miR-200c [133]. Both OncomiR and tumor suppressor miRNA, depending on tissue-type. Expression is repressed by ZEB1 [134], PELP1 [135], PLK1, KLF8 [136], and progesterone [137, 138] and upregulated by p53 [139]. Downregulated in metastatic breast cancer [71] and in basal-like primary tumors [140]. Expression stimulated by treatment of MDA-MB-231 cells with DNA demethylating agent 5-AZA-CdR [141]. Low circulating miR-141 was associated with lower overall survival of breast cancer patients [142, 143]. Overexpression of miR-141 stimulates brain metastasis in mouse models and high serum miR-141 levels were associated with shorter brain metastasis–free survival in human breast cancer patients [144]. miR-141 expression is higher in docetaxel-resistant breast cancer cell lines [145]. | For 3p: PGR [137]; CTNNB1 [146]; EIF4E [145]; ANP32E [140] |
| miR-499a-5p   | Chr12, host gene MYH7B. SNP rs3746444 G miR-499A>G was associated with increased breast cancer risk in Chinese population [147]. | For 5p: IFNAR1 [148]     |
| miR-664b-5p   | ChrX, host gene DKC1. No references were found in PubMed. |                                                                                   |
| miR-1247-5p   | Chr14, in the DLK1-DIO3 genomic imprinted microRNA cluster [149]. Downregulated in aromatase-resistant MCF-7 breast cancer cells [109] and lung adenocarcinomas [150]. Acts as a tumor suppressor in pancreatic cancer [151]. Silenced by DNA methylation in lung adenocarcinomas and cell lines and overexpression promotes apoptosis and inhibits cell invasion and migration [152]. Overexpressed in castration-resistant prostate cancer [153]. | For 5p: NRP1 and NRP2 [151]; SOX9 [154]; MYCBP2 [153]; MAP3K9 [155]; STMN1 [152] |
| miR-1273g-3p  | Chr1, host gene SCP2. no publications in PubMed                                                |                                                                                   |
| miR-1277-3p   | ChrX, host gene WDR44. no publications in PubMed                                              | For 3p: LPL [156]                                                             |
| miR-3611      | Chr10, host gene CUL2. no publications in PubMed                                               |                                                                                   |
| miR-3614-3p   | Chr17, host gene TRIM25. no publications in PubMed                                             |                                                                                   |

(Continued)
GSE78011. In AnAc-treated MCF-7 cells, six downregulated host genes for downregulated miRNAs were identified: MiR-548j host gene HMGB1P10; miR-597 host gene TNKS; miR-1915 host gene CASC10; miR-3146 host gene TWISTNB; miR-5187 host gene TOMM40L; and miR-6814 host gene RIPK4. Whether AnAc selectively inhibits the transcription of these genes via its p300/PCAF histone acetyltransferase (HAT) inhibitory function [169] remains to be examined. Inhibition of HAT activity would be expected to increase gene expression. Interestingly, AnAc inhibits p300/PCAF histone acetyltransferase (HAT) activity [169] and thus could coordinately downregulate this set of miRNAs and host genes by promoting a more condensed genomic state, but experimentally examining the veracity of the supposition is outside this current study and remains to be examined fully. MetaCore transcription factor (TF) network analysis identified CREB1, FosB, SOX4, TCF7L2 (TCF4), PRDM14, JunD, GATA-3, FRA-1, cFos, JunB, FOXp3, and YY1 as significantly associated with these genes. The ability of AnAc to inhibit the activity of these TFs will also need to be experimentally verified.

A decrease in a miRNA would be expected to result in an increase its target transcript expression. Validated targets of each miRNA were identified in the literature. An important note in searching the literature for miRNA targets is that often, whether the miRNA is the 3p or 5p arm is not stated. However, if the miRNA sequence is provided in a diagram along with the seed match site in a target mRNA’s 3’-UTR, the miRNA sequence can be identified as either 3p or 5p by entering the miRNA sequence in miRBase.org. Clearly, a miRNA-3p and

---

Table 5. (Continued)

| miRNA     | Role in breast or other cancers                                                                 | Verified targets          |
|-----------|--------------------------------------------------------------------------------------------------|----------------------------|
| miR-4284  | Chr7, host gene STX1A. Stimulated by treatment of primary human glioblastoma cells with a synthetic berbamine derivative [157]. Downregulated in clear cell papillary renal cell carcinoma [158]. |                                                                          |
| miR-4451  | Chr4, host gene ARHGAP24. no publications in PubMed                                             |                                                                          |
| miR-4743-5p | Chr18, host gene CTIF. no publications in PubMed                                                 |                                                                          |
| miR-5684  | Chr19, intergenic. no publications in PubMed                                                      |                                                                          |
| miR-5696  | Chr2, intergenic. no publications in PubMed                                                       |                                                                          |
| miR-6126  | Chr16, host gene NAA60. Expression is downregulated in ovarian tumors and miR-6126 acts as a tumor suppressor miRNA in ovarian cancer cells [159]. ITGB1 [159] |                                                                          |
| miR-6513-3p | Chr2, host gene PNKD. no publications in PubMed                                                  |                                                                          |
| miR-6720-5p | Chr6, host gene FOXF2. Upregulated by Alternaria spp mycotoxin alternariol (10 μM) treatment of HepG2 cells [160]. |                                                                          |
| miR-6765-3p | Chr14, host gene JAG2. no publications in PubMed                                                 |                                                                          |
| miR-6796-3p | Chr19, host gene PLD3. no publications in PubMed                                                |                                                                          |
| miR-6797-5p | Chr19, host gene RPS19. no publications in PubMed                                                |                                                                          |
| miR-6850-3p | Chr8, host gene RPL8. no publications in PubMed                                                  |                                                                          |
| miR-7109-5p | Chr22, host gene PISD. no publications in PubMed                                                |                                                                          |
miRNA-5p will have different targets, and thus potentially different cellular effects. When identified in our RNA seq study, the 3p or 5p arm is indicated. AnAc reduced miR-378g that targets VIM (vimentin) [51] and VIM mRNA transcript expression was increased in AnAc-treated MCF-7 cells (GSE78011), suggesting a reciprocal regulation. None of the other validated targets of decreased miRNAs (Table 3) were found among the upregulated mRNA transcripts identified in GSE78011. MetaCore network enrichment analysis did not match any of the downregulated miRNAs and Pathway Maps, GO processes, or Process Networks. Networks identified were 1) miR-509: positive regulation of macromolecule metabolic process; 2) miR-584: regulation of gene expression; 3) miR-509, miR584, MDM2, ERK1/2: positive regulation of gene expression (S4 Fig). Based on their CSC and tumor-promoting activities the AnAc downregulation of miR-378g, miR-548, miR-548j, miR-548l (Table 3) would be expected to contribute to the anti-proliferative activity of AnAc.

miRNAs upregulated by AnAc in MCF-7

AnAc increased the expression of 48 miRNAs in MCF-7 cells (Table 4). None of the host genes (Table 3) of intronic miRNAs was upregulated by AnAc treatment of MCF-7 cells. None of the validated targets of upregulated miRNAs in AnAc-treated MCF-7 cells (Table 4) were found among the AnAc-regulated mRNA transcripts identified in RNA seq (GSE78011). Given their roles as ‘tumor suppressor’ miRNAs in inhibiting breast and other cancer cell proliferation and activities (see Table 4), the increases in let-7a-2-3p, miR-520a-5p, miR-520d-5p, miR-551b-5p, miR-4485-5p, and miR-6126 may play roles in AnAc-mediated inhibition of breast cancer cell proliferation. miR-520a-5p and miR-520d-5p are in a cluster of miR-520 isomers (a-h) on Chr 19 that share the same seed sequence, and
thus are predicted to have common targets. miR-520f was recently reported to target ADAM9, thus inhibiting internalization of E-cadherin, and TGFBR2 that inhibits TGFβ signaling–mediated induction of ZEB1/2 and/or SNAI which thus allows CDH1 (E-cadherin) transcription, thus blocking EMT [170].

MetaCore analysis of these miRNAs identified “embryo implantation, cellular response to amino acid stimulus” as the top GO process (S5A Fig). Network analysis identified two top networks: 1) mi-1229-3p, miR-520a-5p, miR-612, miR-4516, miR-562: positive regulation of metabolic process (S5B Fig); and 2) miR 20b-3p, miR 663a, let-7a-5p, miR-1229 -3p, SMAD3: regulation of cell proliferation (S5C Fig). Network analysis of TFs associated with the 48 upregulated miRNAs identified c-Myc, N-Myc, EPAs1, E2F1, SOX2, AML1, RUNX10, NANOG, MITF, EGR1, and ZNF224 in the top ten TFs. Whether AnAc may activate these TFs to increase the transcription of the upregulated miRNAs or selectively increase miRNA stability will require further examination.

miRNAs oppositely regulated by AnAc in MCF-7 and MDA-MB-231 cells

In contrast, miR-6873 showed opposite AnAc regulation in the two cell lines: it was downregulated in MCF-7 and upregulated in MDA-MB-231 cells (Tables 2 and 5). There are no publications in PubMed on miR-6873 and miR-6873 was not listed in microRNA.org or miRTarBase. Thus, its relevance to AnAc responses in these two cell lines is unknown.

miRNAs downregulated by AnAc in MDA-MB-231 cells

Twenty-two miRNAs were downregulated by AnAc in MDA-MB-231 cells and none of these overlapped with miRNAs downregulated by AnAc in MCF-7 cells (Table 5). The chromosome location and host gene, if warranted, of each of the AnAc-downregulated miRNAs are identified in Table 5. To examine if the miRNA host gene was downregulated by AnAc in MDA-MB-231 cells, we searched GSE78011. miR-1277 host gene WDR44 was downregulated by AnAc in MDA-MB-231 cells. WDR44 encodes a protein that interacts with the small GTPase rab11 and is involved in endosome recycling [171]. There are no validated targets for miR-1277 in miRTarBase.

Downregulation of a miRNA would be expected to increase the expression of its targets; hence, we searched our data of mRNAs upregulated by AnAc in MDA-MB-231 cells (550 genes, GSE78011) for the validated targets in Table 5, but none were reciprocally upregulated. This may be because the miRNA and mRNA for RNA seq were extracted at the same time, i.e., after 6 h of AnAc treatment, or that these mRNAs are not expressed or targeted in MDA-MB-231 cells. Given their roles as putative oncomiRs the downregulation miR-23b and miR-1247 may play a role in the anti-proliferative activity of AnAc in MDA-MB-231 cells.

Analysis of the data identified ZFP36L1 as a putative target of miR-3614 in MDA-MB-231 cells. Interestingly, AnAc downregulated miR-3614 and upregulated ZFP36L1 transcript expression in MDA-MB-231 cells, suggesting an inverse correlation. ZFP36L1 has been identified as a cancer gene due to mutations in breast cancer and acts in a recessive manner [172]. ZFP36L1 is a member of the TFP family of tandem zinc finger proteins that bind AU-rich elements (AURE) in the 3'-end of target gene transcripts and promote target degradation, e.g. STARD1 [173], VEGFA [174], NR4A2 [175], BCL2 [176], LDLR [177], STAT5B [178], and CDK6 [179]. Of these genes, only VEGFA and LDLR were identified as differentially expressed genes in AnAc-treated cells. LDLR was downregulated whereas VEGFA was upregulated in AnAc-treated MDA-MB-231 cells. Interestingly, medroxyprogesterone acetate (MPA, a synthetic progestin), but not E2, upregulates ZFP36L1 transcription in MCF-7 cells [25].
MetaCore analysis of the AnAc-downregulated miRNAs in MDA-MB-231 cells identified one canonical pathway map: "Development: miRNA-dependent regulation of EMT" and the 10 GO processes in S6A Fig. Network analysis identified two top networks: 1) miR-23b-3p, miR-499, miR-499-3p, miR-499-5p, c-Fos (S6B Fig), and miR-141, miR-141-3p, miR-1247-5p, PPAR-gamma, BMI-1 (S6C Fig).

miRNAs upregulated by AnAc in MDA-MB-231 cells

Fourteen miRNAs were increased by AnAc-treatment of MDA-MB-231 cells (Table 6). We have described miR-20b-5p and miR-612 upregulation in the context of similar results in AnAc-treated MCF-7 cells (Table 2, Fig 2, S2 Fig). The chromosome location and host gene, if warranted, of each of the AnAc-upregulated miRNAs are identified in Table 6. Interestingly, most of the downregulated miRNAs were intergenic. miR-1298 is in encoded in HTR2C, but HTR2C was not among the AnAc-regulated genes in MDA-MB-231 cells in GSE78011. An increase in a miRNA would be expected to result in a decrease of its target transcript. miR-20b-5p target EFNB2 (ephrin B2) expression was downregulated in AnAc-treated MDA-MB-231 cells, but none of the validated targets of the upregulated miRNAs (Table 6) were found among the AnAc-downregulated mRNA transcripts identified in RNA seq (GSE78011). Given their roles as 'tumor suppressor' miRNAs (see Table 6), the increases in miR-29b, miR-612, and miR-1298 may contribute to the antiproliferative activity of AnAc in MDA-MB-231 cells.

MetaCore analysis of these upregulated miRNAs identified "cellular response to inorganic substance" as the top GO process (S7A Fig). MetaCore analysis identified two networks: 1)

Table 6. miRNAs upregulated by AnAc in MDA-MB-231 cells. The genomic location of each miRNA was identified in miRAD http://bmi.ana.med.uni-muenchen.de/mirrad/ [34]. Verified targets are those experimentally validated targets of the indicated miRNA as demonstrated by 3'-UTR luciferase reporter assay. Since many publications do not include whether the 5p or 3p arm of the miRNA was studied, if the sequence of the miRNA was provided, it was searched in miRBase.org to identify which arm was used in the target gene 3'-UTR luciferase reporter assay.

| miRNA       | Role in breast or other cancers                                                                 | Verified targets                      |
|-------------|-------------------------------------------------------------------------------------------------|---------------------------------------|
| miR-378f    | Chr1, intergenic. Downregulated by E6/E7 silencing in HeLa cells [161].                        |                                       |
| miR-1257    | Chr20, intergenic. Downregulated in dedifferentiated liposarcoma [162].                        |                                       |
| miR-1298-5p | ChrX, host gene HTR2C clustered with miR-764, miR1912, miR1264, miR-1911, and miR-448. Downregulated in neuroglioma [163]. Identified as an inhibitor the growth of KRAS-driven colon cancer cells both in vitro and in vivo [164]. | For 5p: GJA1 [165], PTK2 and LAMB3 [164] |
| miR-1304-5p | Chr11, intergenic. Downregulated in NSCLC cells [166].                                          |                                       |
| miR-3116-1  | Chr1, host gene PATJ. no publications in PubMed                                               |                                       |
| miR-3139    | Chr4, host gene GAB1. no publications in PubMed                                               |                                       |
| miR-3159    | Chr11, intergenic. no publications in PubMed                                                   |                                       |
| miR-3936    | Chr5, intergenic. no publications in PubMed                                                    |                                       |
| miR-4473    | Chr9, host gene MLLT3. no publications in PubMed                                              |                                       |
| miR-6794-5p | Chr19, host gene MAST1. no publications in PubMed                                             |                                       |
| miR-6873-3p | Chr6, host gene WDR46. no publications in PubMed                                               |                                       |
| miR-7113-5p | Chr11, host gene NDUFS8. no publications in PubMed                                             |                                       |

https://doi.org/10.1371/journal.pone.0184471.t006
miR-1257, Bcl-2, PAX6, FOXO3A, and FOXP3; and 2) miR-20b-5p, PPARγ, MDA2, p57, and Sin3A (S7B and S7C Fig).

qPCR validation of select AnAc-mediated changes in miRNAs

We selected miR-612, increased by AnAc in both MCF-7 and MDA-MB-231 cells (Table 2); miR-20b-3p and miR-29-5p, upregulated by AnAc in MCF-7 and MDA-MB-231, respectively (Table 2), and miR-378g that was downregulated by AnAc in MCF-7 cells for validation. miR-378g was selected because miR-378g targets VIM [51] and VIM mRNA transcript expression was increased in AnAc-treated MCF-7 cells (GSE78011), suggesting a reciprocal regulation. Cells were grown in hormone-depleted medium for 48 h prior to 6 h treatment with 13.5 or 35 μM AnAc. As anticipated, AnAc increased miR-612 in both cell lines (Fig 3A). Also as anticipated, AnAc increased miR-20b-3p in MCF-7 cells. We did not detect the anticipated decrease in miR-378g in AnAc-treated MCF-7 cells; however, AnAc reduced miR-378g in MDA-MB-231 cells. We did not detect miR-20b-5p in MDA-MB-231 cells (CT values were undetermined). CT values show that miR-20b-3p is the dominant arm of miR-20b expressed in both cell lines (Fig 3B).

Effect of altered miR-612 on cell viability

Since AnAc increased miR-612 in both MCF-7 and MDA-MB-231 cells (Table 2, Fig 2) and miR-612 has reported tumor suppressor activity in HCC [35, 36] and colorectal cancers [37] (Table 4), we examined how altering miR-612 levels affected cell viability of MCF-7 and MDA-MB-231 cells and their responses to AnAc. Alterations in miR-612 levels in each cell line in response to transfection of miR-612 mimic and anti-miR-612 were demonstrated (Fig 4A). As expected, AnAc inhibited cell viability in both cell lines (Fig 4B). Transfection with miR-612 mimic inhibited cell viability in each cell line with a larger effect in MCF-7 than MDA-MB-231 cells. Transfection with a miR-612 inhibitor had no effect in MCF-7 cells, but inhibited the viability of MDA-MB-231 cells ~ 20%. Notably, the miR-612 inhibitor abrogated the anti-proliferative activity of AnAc in MCF-7 cells and reduced AnAc’s anti-proliferative activity in MDA-MB-231 cells. These results are consistent with a model in which the increase in miR-612 in AnAc-treated MCF-7 and MDA-MB-231 cells plays a role in the anti-proliferative activity of AnAc (Fig 4C).

Fig 3. qPCR analysis of select AnAc-regulated miRNA expression. MCF-7 and MDA-MB-231 cells were grown in hormone-depleted medium for 48 h prior to 6 h treatment with 13.5 or 35 μM AnAc. A, qPCR using TaqMan assays for miR-378g, miR-612, miR-20b-5p, and miR-20b-3p was performed using U48 as normalizer. B, CT values for miR-20b-5p and miR-20b-3p expression. miR-20b-5p was not detected in MDA-MB-231 (CT values undetermined). For both A and B: Values are the mean ± SEM of triplicates in one experiment for MCF-7 cells and are the mean ± SEM of two independent experiments for MDA-MB-231 cells.
We selected VIM, a target of miR-378g downregulated by AnAc in MCF-7 cells, and ZFP36L, a target of miR-3614 downregulated by AnAc in MDA-MB-231 cells for validation by qPCR. As anticipated from the decrease in miR-378g in RNA seq data (Table 3), we detected a slight increase in VIM transcript expression in MCF-7 as well as an increase in VIM in MDA-MB-231 cells (Fig 5). However, because qPCR indicated an increase in miR-378g levels in AnAc-treated MCF-7 cells (Fig 5), it is possible that VIM is upregulated by AnAc by mechanisms unrelated to miR-378g. In addition, miRNA and mRNA were extracted at the same time, i.e., after 6 h of AnAc treatment, and it may be that changes in VIM mRNA levels require a longer time to be degraded after miR-378g targeting. Transcript levels of ZFP36L were increased in AnAc-treated MDA-MB-231 cells (Fig 5), corresponding with the observed downregulation of miR-3614 (Table 5). These data confirm the reciprocal expression of these mRNA transcripts detected in RNA seq and their target miRNAs in the respective AnAc-treated cell line.

Pathways affected by DEGs and DEmiRs in AnAc-treated MCF-7 cells

MetaCore analysis of DEGs from both mRNA and miRNA data sets of AnAc-treated MCF-7 cells identified NETosis in SLE as the top pathway. The release of neutrophil extracellular traps (NETs) by dying cells (NETosis) was first described as the release of nuclear chromatin, nuclear histones and many granular antimicrobial proteins from neutrophils as one of the first lines of defense against pathogens (reviewed in [180]). The top GO processes were chromatin silencing, negative regulation of gene expression (epigenetic, nucleosome assembly, chromatin assembly, and nucleosome organization. The three gene networks identified were 1): PDEGF, PDE6G, APOBEC3H, GGTF II beta, CDIP, p53; 2) miR-499, BMCC1, Histone H1, miR-20b, miR-23b; 3) UCHL1, Protein C, PDK4, EGR1, miR-1298 5p. Network #2 processes include
anoikis, negative regulation of fat cell proliferation, regulation of DNA metabolic processes, which reflect the antiproliferative, pro-apoptotic, and NRAM activity of AnAc detected previously in MCF-7 cells [13].

Pathways affected by DEGs and DEmiRs in AnAc-treated MDA-MB-231 cells

MetaCore analysis of DEGs from both mRNA and miRNA data sets of AnAc-treated MDA-MB-231 cells identified “Immune response, IL-3 signaling via JAK/STAT, p38, JNK, and NFkB” as the top pathway. The top GO processes were “Positive regulation of biological process; cellular response to oxygen-containing compound, positive regulation of cellular process, response to oxygen-containing compound, regulation of developmental process, and response to lipid”. The three gene networks identified were Network #1: Axin, Frizzled, cMyc, WNT, PI3K reg classIA: canonical Wnt signaling pathway, beta-catenin destruction complex disassembly, regulation of cell proliferation, cell surface receptor signaling pathway involved in cell-cell signaling, cell-cell signaling by wnt. Network #2: C/EBPbeta, SOS, NGFR, H-Ras, NGF: positive regulation of cellular metabolic process, positive regulation of MAPK cascade, positive regulation of metabolic process, positive regulation of macromolecule metabolic process, and positive regulation of intracellular signal transduction. Network #3: GALNT4, Keratin80, BCMP101, HEXIM1, PNRC1: translational elongation, translation, amide biosynthetic process, peptide biosynthetic process, peptide metabolic process.

Fig 5. qPCR analysis of mRNA targets of AnAc-downregulated miRNAs. MCF-7 and MDA-MB-231 cells were grown in hormone-depleted medium for 48 h prior to 6 h treatment with 13.5 or 35 μM AnAc. qPCR was performed using GAPDH as normalizer. Values are the mean ± SEM of triplicates in one experiment for MCF-7 cells and are the mean ± SEM of two independent experiments for MDA-MB-231 cells.

https://doi.org/10.1371/journal.pone.0184471.g005
Conclusions

In summary, we describe the first comprehensive assessment of miRNA expression in response to anacardic acid in ERα+, luminal A MCF-7 and MDA-MB-231 TNBC breast cancer cells. The pathways modulated by these miRNAs suggest that key nodal molecules, e.g., Cyclin D1, SMAD, SP1, MYC, c-FOS, PPARγ, BCL2, FOXO3A, MDA2, and SIN3, are targets of AnAc activity. In agreement with the pathway analysis, we previously reported that AnAc reduced CCND1 transcript expression in MCF-7 and MDA-MB-231 cells [13]. The roles of the other proteins and pathways in AnAc responses remains to be investigated.

Supporting information

S1 Fig. Heat map of miRNAs significantly altered in AnAc-treated MCF-7 cells. miRNAs significantly affected by AnAc were analyzed using Partek Genomic Suite™ to generate the heat map.

S2 Fig. Heat map of miRNAs significantly altered in AnAc-treated MDA-MB-231 cells. miRNAs significantly affected by AnAc were analyzed using Partek Genomic Suite™ to generate the heat map.

S3 Fig. MetaCore analysis of upregulated miRNAs in AnAc-treated MCF-7 and MDA-MB-231 cells. A) Gene Ontology (GO) processes. The hatched bars are common whereas orange indicates MCF-7 cells. MetaCore Analyze Networks algorithm identified B) miR-20b-5p, Cyclin D1, DEC1 (Stra13), SMAD4 network: circadian regulation of gene expression (41.2%), negative regulation of nucleobase containing compound metabolic process (82.4%), negative regulation of cellular biosynthetic process (82.4%), rhythmic process (58.8%), negative regulation of nitrogen compound metabolic process (82.4%). C) miR-612, SP1, MyCH, gamma-ENaC, DR5 network: muscle filament sliding (36.4%), actin-myosin filament sliding (36.4%), actin filament-based movement (43.2%), muscle contraction (50.0%), actin-mediated cell contraction (36.4%)

S4 Fig. MetaCore analysis of downregulated miRNAs in AnAc-treated cells. MetaCore Analyze Networks algorithm identified A) miR509: B) miR-584, C/EBPbeta, HOX10A; 3) miR-509, miR-584, MDM2, ERK1/2.

S5 Fig. MetaCore analysis of upregulated miRNAs in AnAc-treated MCF-7 cells. A) Gene Ontology (GO) processes. MetaCore Analyze Networks algorithm identified B) miR 1229 3p, miR 520a 5p, miR 612, miR 4516, miR 562: positive regulation of metabolic process (60.5%), negative regulation of apoptotic process (37.2%), negative regulation of programmed cell death (37.2%), negative regulation of cell death (37.2%), viral process (34.9%); C) miR 20b 5p, miR 663a, miR let 7a 5p, miR 1229 3p, SMAD3: regulation of cell proliferation (43.5%), response to growth factor stimulus (43.5%), response to growth factor (43.5%), positive regulation of macromolecule metabolic process (71.7%), response to lipid (52.2%)

S6 Fig. MetaCore analysis of downregulated miRNAs in AnAc-treated MDA-MB-231 cells. A) Gene Ontology (GO) processes. MetaCore Analyze Networks algorithm identified B) miR-23b-3p, miR-499, miR-499-3p, miR-499-5p, c-Fos: response to drug (37.8%), response to abiotic stimulus (48.9%), response to mechanical stimulus (28.9%), cellular response to hormone
stimulus (37.8%), response to inorganic substance (37.8%). C) miR-141, miR-141-3p, miR-1247-5p, PPAR-gamma, BMI-1: positive regulation of transcription from RNA polymerase II promoter (76.6%), regulation of transcription from RNA polymerase II promoter (85.1%), positive regulation of nucleic acid-templated transcription (76.6%), positive regulation of transcription, DNA-templated (76.6%), negative regulation of RNA metabolic process (74.5%).

S7 Fig. MetaCore analysis of upregulated miRNAs in AnAc-treated MDA-MB-231 cells. A) Gene Ontology (GO) processes. MetaCore Analyze Networks algorithm identified B) miR-1257, Bcl-2, PAX6, FOXO3A, and FOXP3; and C) miR-20b-5p, PPARγ, MDA2, p57, Sin3.

S1 Table. miRNAs regulated by AnAc in MCF-7 cells. Cells were grown in phenol red-free IMEM (ThermoFisher) medium containing 5% dextran coated charcoal (DCC)-stripped FBS (hormone-depleted medium) for 48 h prior to treatment with established IC50 concentrations of AnAc 24:1n5: 13.5 μM for MCF-7 cells [13] for 6 h and was replicated in three separate experiments. Differentially expressed miRNAs (DEmiRs) were identified for pairwise comparisons (MCF-7 AnAc-treated vs. MCF-7 control using the tuxedo suite of programs including cufflinks and cuffdiff (version 2.2.1) Significant DEmiRs with fold-change and p values are listed. These raw data of our RNA-seq are available at Gene Expression Omnibus (GEO) database: accession number GSE78011.

S2 Table. miRNAs regulated by AnAc in MDA-MB-231 cells. Cells were grown in phenol red-free IMEM (ThermoFisher) medium containing 5% dextran coated charcoal (DCC)-stripped FBS (hormone-depleted medium) for 48 h prior to treatment with established IC50 concentrations of AnAc 24:1n5: 35.0 μM for MDA-MB-231 cells [13] for 6 h and was replicated in three separate experiments. Differentially expressed miRNAs (DEmiRs) were identified for pairwise comparisons (MDA-MB-231 AnAc-treated vs. MDA-MB-231 control using the tuxedo suite of programs including cufflinks and cuffdiff (version 2.2.1) Significant DEmiRs with fold-change and p values are listed. These raw data of our RNA-seq are available at Gene Expression Omnibus (GEO) database: accession number GSE78011.

Acknowledgments

We thank Brandie N. Radde for performing the cell treatments and isolating the RNA for RNA seq. This research was funded by a grant from the University of Louisville Center for Genetics and Molecular Medicine (CGeMM) Next Generation Pilot Grant to D.J.S.; a grant from the University of Louisville School of Medicine to C.M.K., and by a University of Louisville Executive Vice President for Research and Innovation grant to C.M.K. Bioinformatics support for this work by E.C.R. and S.J.W. was provided by National Institutes of Health grants P20GM103436 (Kentucky IDeA Networks of Biomedical Research Excellence, Nigel Cooper, PI).

Author Contributions

Conceptualization: David J. Schultz, Carolyn M. Klinge.

Data curation: Eric C. Rouchka, Sabine J. Waigel.

Formal analysis: Eric C. Rouchka, Carolyn M. Klinge.
**Investigation:** Penn Muluhngwi, Negin Alizadeh-Rad, Carolyn M. Klinge.

**Methodology:** Penn Muluhngwi, Negin Alizadeh-Rad, Madelyn A. Green, Carolyn M. Klinge.

**Project administration:** Carolyn M. Klinge.

**Resources:** David J. Schultz.

**Supervision:** Carolyn M. Klinge.

**Writing – original draft:** Carolyn M. Klinge.

**Writing – review & editing:** David J. Schultz, Carolyn M. Klinge.

**References**

1. Iorio MV, Croce CM. microRNA involvement in human cancer. Carcinogenesis. 2012; 33(6):1126–33. https://doi.org/10.1093/carcin/bgs140 PMID: 22491715

2. Meijer HA, Smith EM, Bushell M. Regulation of miRNA strand selection: follow the leader? Biochem Soc Trans. 2014; 42(4):1135–40. Epub 2014/08/12. https://doi.org/10.1042/BST20140142 PMID: 25110015

3. Pasquinelli AE. MicroRNAs and their targets: recognition, regulation and an emerging reciprocal relationship. Nat Rev Genet. 2012; 13(4):271–82. https://doi.org/10.1038/nrg3162 PMID: 22411466

4. McGuire A, Brown JA, Kerin MJ. Metastatic breast cancer: the potential of miRNA for diagnosis and treatment monitoring. Cancer Metastasis Rev. 2015; 34(1):145–55. Epub 2015/02/28. https://doi.org/10.1007/s10555-015-9551-7 PMID: 25721950

5. Zhang H, Li Y, Lai M. The microRNA network and tumor metastasis. Oncogene. 2010; 29(7):937–48. https://doi.org/10.1038/onc.2009.406 PMID: 19935707

6. Zhang W, Liu J, Wang G. The role of microRNAs in human breast cancer progression. Tumour Biol. 2014; 35(7):6235–44. Epub 2014/06/19. https://doi.org/10.1007/s13277-014-2202-8 PMID: 24938874

7. Hayes EL, Lewis-Wambi JS. Mechanisms of endocrine resistance in breast cancer: an overview of the proposed roles of noncoding RNA. Breast Cancer Res. 2015; 17(1):542. Epub 2015/03/18.

8. Klinge CM. miRNAs regulated by estrogens, tamoxifen, and endocrine disruptors and their downstream gene targets. Mol Cell Endocrinol. 2015; 418:273–97. https://doi.org/10.1016/j.mce.2015.01.035 PMID: 25659536

9. van Schooneveld E, Wildiers H, Vergote I, Vermeulen PB, Dirix LY, Van Laere SJ. Dysregulation of microRNAs in breast cancer and their potential role as prognostic and predictive biomarkers in patient management. Breast Cancer Res. 2015; 17:21. Epub 2015/04/08. https://doi.org/10.1186/s13058-015-0526-y PMID: 25849621

10. Pinweha P, Rattanapornsompong K, Charoensawan V, Jitrapakdee S. MicroRNAs and oncogenic transcriptional regulatory networks controlling metabolic reprogramming in cancers. Computational and structural biotechnology journal. 2016; 14:223–33. Epub 2016/07/01. https://doi.org/10.1016/j.csbj.2016.05.005 PMID: 27358718

11. Schultz DJ, Wickramasinghe NS, Klinge CM. Biosynthesis and bioactivity of anacardic acid. In: Romeo J, editor. Recent Advances in Phytochemistry. 402006. p. 131–56.

12. Hemshekar M, Sebabast Santhosh M, Kemparaju K, Girish KS. Emerging Roles of Anacardic Acid and Its Derivatives: A Pharmacological Overview. Basic Clin Pharmacol Toxicol. 2012; 110(2):122–32. https://doi.org/10.1111/j.1742-7843.2011.00833.x PMID: 22103711

13. Schultz DJ, Wickramasinghe NS, Ivanova MM, Isaacs SM, Dougherty SM, Imbert-Fernandez Y, et al. Anacardic acid inhibits estrogen receptor alpha-DNA binding and reduces target gene transcription and breast cancer cell proliferation Mol Cancer Ther. 2010; 9(3):594–605. Epub Epub 2010 Mar 2.; https://doi.org/10.1158/1535-7186.MCT-09-0976 PMID: 20197399

14. Schultz DJ, Olsen C, Cobbs GA, Stolowich NJ, Parrott MM. Bioactivity of anacardic acid against Colorado potato beetle (Leptinotarsa decemlineata) larvae. J Agric Food Chem. 2006; 54(20):7522–9. https://doi.org/10.1021/jf061418u PMID: 17002417

15. Cock PJ, Fields CJ, Goto N, Heuer ML, Rice PM. The Sanger FASTQ file format for sequences with quality scores, and the Solexa/Illumina FASTQ variants. Nucleic Acids Res. 2010; 38(6):1767–71. Epub 2009/12/18. https://doi.org/10.1093/nar/gkp1137 PMID: 20015970;
16. Friedlander MR, Mackowiak SD, Li N, Chen W, Rajewsky N. miRDeep2 accurately identifies known and hundreds of novel microRNA genes in seven animal clades. Nucleic Acids Res. 2012; 40(1):37–52. Epub 2011/09/14. https://doi.org/10.1093/nar/gkr688 PMID: 21911355;

17. Robinson MD, McCarthy DJ, Smyth GK. edgeR: a Bioconductor package for differential expression analysis of digital gene expression data. Bioinformatics. 2010; 26(1):139–40. Epub 2009/11/17. https://doi.org/10.1093/bioinformatics/btp616 PMID: 19910308;

18. Andrews S. FastQC: A Quality Control Tool for High Throughput Sequence Data 2014. http://bioinformatics.babraham.ac.uk/projects/fastqc/.

19. Bolger AM, Lohse M, Usadel B. Trimmomatic: a flexible trimmer for Illumina sequence data. Bioinformatics. 2014; 30(15):2114–20. Epub 2014/04/04. https://doi.org/10.1093/bioinformatics/btu170 PMID: 24695404;

20. Langmead B, Salzberg SL. Fast gapped-read alignment with Bowtie 2. Nat Methods. 2012; 9(4):357–9. Epub 2012/03/06. https://doi.org/10.1038/nmeth.1923 PMID: 22388286;

21. Griffiths-Jones S, Saini HK, van Dongen S, Enright AJ. miRBase: tools for microRNA genomics. Nucleic Acids Res. 2008; 36(suppl_1):D154–8. https://doi.org/10.1093/nar/gkm952 PMID: 17991681;

22. Bolser DM, Chibon PY, Palopoli N, Gong S, Jacob D, Del Angel VD, et al. MetaBase— the wiki-data-base of biological databases. Nucleic Acids Res. 2012; 40(Database issue):D1250–4. Epub 2011/12/06. https://doi.org/10.1093/nar/gkr1099 PMID: 22139927;

23. Pennanen PT, Sarvilinna NS, Purmonen SR, Ylikomi TJ. Changes in protein tyrosine phosphatase type IVA member 1 and zinc finger protein 36 C3H type-like 1 expression demonstrate altered estrogen and progesterin effect in medroxyprogesterone acetate-resistant and estrogen-independent breast cancer cell models. Steroids. 2009; 74(4–5):404–9. https://doi.org/10.1016/j.steroids.2008.12.005 PMID: 19146866;

24. Schmittgen TD, Livak KJ. Analyzing real-time PCR data by the comparative C(T) method. Nat Protoc. 2008; 3(6):1101–8. Epub 2008/06/13. PMID: 18546601.

25. Hah N, Kraus WL. Hormone-regulated transcriptomes: Lessons learned from estrogen signaling pathways in breast cancer cells. Mol Cell Endocrinol. 2014; 382(1):56–67. https://doi.org/10.1016/j.mce.2013.10.010 PMID: 24141032;

26. Smirnoff LM, Appana SN, Kingley CM. COUT-TFII inhibits NFkappaB activation in endocrine-resistant breast cancer cells. Mol Cell Endocrinol. 2014; 382(1):358–67. https://doi.org/10.1016/j.mce.2013.10.010 PMID: 24141032;

27. Schmittgen TD, Livak KJ. Analyzing real-time PCR data by the comparative C(T) method. Nat Protoc. 2008; 3(6):1101–8. Epub 2008/06/13. PMID: 18546601.

28. Hah N, Kraus WL. Hormone-regulated transcriptomes: Lessons learned from estrogen signaling pathways in breast cancer cells. Mol Cell Endocrinol. 2014; 382(1):562–64. https://doi.org/10.1016/j.mce.2013.06.021 PMID: 23819798;

29. Radde BN, Alizadeh-Rad N, Price SM, Schultz DJ, Kingley CM. Anacardic Acid, Salicylic Acid, and Oleic Acid Differentially Alter Cellular Energetic Function in Breast Cancer Cells. J Cell Biochem. 2016. Epub 2016/03/19. https://doi.org/10.1002/jcb.25544 PMID: 26990649;

30. Trapnell C, Hendrickson DG, Sauvageau M, Goff L, Rinn JL, Pachter L. Differential analysis of gene regulation at transcript resolution with RNA-seq. Nat Biotechnol. 2013; 31(1):46–53. Epub 2012/12/12. https://doi.org/10.1038/nbt.2450 PMID: 23222703;

31. Trapnell C, Roberts A, Goff L, Pertea G, Kim D, Kelley DR, et al. Differential gene and transcript expression analysis of RNA-seq experiments with TopHat and Cufflinks. Nat Protocols. 2012; 7(3):562–78. https://doi.org/10.1038/nprot.2012.016 PMID: 22383036;

32. Griffiths-Jones S, Hui JH, Marco A, Ronshaugen M. MicroRNA evolution by arm switching. EMBO Rep. 2011; 12(2):172–7. Epub 2011/01/08. https://doi.org/10.1038/embor.2010.191 PMID: 21212805;

33. Bose M, Bhattacharyya SN. Target-dependent biogenesis of cognate microRNAs in human cells. Nature Communications. 2016; 7:12200. https://doi.org/10.1038/ncomms12200 PMID: 27448149;

34. Hinske LC, Franca GS, Torres HAM, Ohara DT, Lopes-Ramos CM, Heyn J, et al. miR3AD—integrating microRNA inter- and intragenic data. Database. 2014; 2014. https://doi.org/10.1093/database/bau099 PMID: 25288656;

35. Tang J, Tao Z-H, Wen D, Wan J-L, Liu D-L, Zhang S, et al. miR-612 suppresses the stemness of liver cancer via Wnt/β-catenin signaling. Biochem Biophys Res Commun. 2014; 447(1):210–5. https://doi.org/10.1016/j.bbrc.2014.03.135 PMID: 24704424
36. Tao ZH, Wan JL, Zeng LY, Xie L, Sun HC, Qin LX, et al. miR-612 suppresses the invasive-metastatic cascade in hepatocellular carcinoma. J Exp Med. 2013; 210(4):789–803. Epub 2013/03/13. https://doi.org/10.1084/jem.20120153 PMID: 23478189;

37. Sheng L, He P, Yang X, Zhou M, Feng Q. miR-612 negatively regulates colorectal cancer growth and metastasis by targeting AKT2. Cell Death Dis. 2015; 6:e1808. https://doi.org/10.1038/cddis.2015.184 PMID: 26158514;

38. Liu Y, Liu DL, Dong LL, Wen D, Shi DM, Zhou J, et al. miR-612 suppresses stem cell-like property of hepatocellular carcinoma cells by modulating Sp1/Nanog signaling. Cell Death Dis. 2016; 7(9):e2377. Epub 2016/09/30. https://doi.org/10.1038/cddis.2016.282 PMID: 27685621;

39. Landais S, Landry S, Legault P, Rassart E. Oncogenic potential of the miR-106-363 cluster and its implication in human T-cell leukemia. Cancer Res. 2007; 67(12):5699–707. Epub 2007/06/19. https://doi.org/10.1158/0008-5472.CAN-06-4478 PMID: 17575136;

40. Ao X, Nie P, Wu B, Xu W, Zhang T, Wang S, et al. Decreased expression of microRNA-17 and microRNA-20b promotes breast cancer resistance to taxol therapy by upregulation of NCOA3. Cell Death Dis. 2016; 7:e2463. https://doi.org/10.1038/cddis.2016.367 PMID: 27831559;

41. Li D, Ilnytskyi Y, Kovalchuk A, Khachigian LM, Bronson RT, Wang B, et al. Crucial role for early growth response-1 in the transcriptional regulation of miR-20b in breast cancer. Oncotarget. 2013; 4(9):1373–87. Epub 2013/08/16. https://doi.org/10.18632/oncotarget.1116 PMID: 23945289;

42. Castellano L, Giamas G, Jacob J, Coombes RC, Lucchesi W, Thiruchelvam P, et al. The estrogen receptor-alpha induced microRNA signature regulates itself and its transcriptional response. Proc Natl Acad Sci USA. 2009; 106(37):15732–7. https://doi.org/10.1073/pnas.0906971106 PMID: 19706389;

43. Giraud-Triboult K, Rochon-Beaucourt C, Nissan X, Champon B, Aubert S, Pietu G. Combined miRNA and microRNA profiling reveals that miR-148a and miR-20b control human mesenchymal stem cell phenotype via EPAS1. Physiol Genomics. 2011; 43(2):77–86. https://doi.org/10.1152/physiolgenomics.00077.2010 PMID: 21081659;

44. Xue TM, Tao LD, Zhang M, Zhang J, Liu X, Chen GF, et al. Clinicopathological Significance of MicroRNA-20b Expression in Hepatocellular Carcinoma and Regulation of HIF-1α and VEGF Effect on Cell Biological Behaviour. Dis Markers. 2015; 2015:325176. Epub 2015/11/28. https://doi.org/10.1155/2015/325176 PMID: 26612965;

45. He J, Zhang JF, Yi C, Lv Q, Xie WD, Li JN, et al. miRNA-mediated functional changes through co-regulation function related genes. PLoS One. 2010; 5(10):e13558. Epub 2010/11/03. https://doi.org/10.1371/journal.pone.0013558 PMID: 21042576;

46. Wang W, Feng L, Zhang H, Hachy S, Satohisa S, Laurent LC, et al. Preeclampsia Up-Regulates Angiogenesis-Associated MicroRNA (i.e., miR-17, -20a, and -20b) That Target Ephrin-B2 and EPHB4 in Human Placenta. The Journal of Clinical Endocrinology & Metabolism. 2012; 97(6):E1051–E9. https://doi.org/10.1210/jc.2011-3131 PMID: 22438230;

47. Wang B, Yang J, Xiao B. MicroRNA-20b (miR-20b) Promotes the Proliferation, Migration, Invasion, and Tumorigenicity in Esophageal Cancer Cells via the Regulation of Phosphatase and Tensin Homologue Expression. PLoS One. 2016; 11(10):e0164105. Epub 2016/10/05. https://doi.org/10.1371/journal.pone.0164105 PMID: 27701465;

48. Hong S, Yu S, Li J, Yin Y, Liu Y, Zhang Q, et al. MiR-20b Displays Tumor-Suppressor Functions in Papillary Thyroid Carcinoma by Regulating the MAPK/ERK Signaling Pathway. Thyroid. 2016; 26(12):1733–43. Epub 2016/11/03. https://doi.org/10.1089/thy.2015.0578 PMID: 27717302;

49. Krist B, Florczyk U, Pietraszek-Gremplewicz K, Jozkowicz A, Dulak J. The Role of miR-378a in Metabolism, Angiogenesis, and Muscle Biology. International journal of endocrinology. 2015; 2015:281756. Epub 2016/02/04. https://doi.org/10.1155/2015/281756 PMID: 26839547;

50. Feng M, Li Z, Auu M, Wong CH, Yang X, Yu Q. Myc/miR-378/TB2/bicclin D1 functional module regulates oncogenic transformation. Oncogene. 2011; 30(19):2242–51. https://doi.org/10.1038/onc.2010.602 PMID: 21242960;

51. Deng Z, Du WW, Fang L, Shan SW, Qian J, Lin J, et al. The Intermediate Filament Vimentin Mediates MicroRNA miR-378 Function in Cellular Self-renewal by Regulating the Expression of the Sox2 Transcription Factor. J Biol Chem. 2013; 288(1):319–31. https://doi.org/10.1074/jbc.M112.418830 PMID: 23135265;

52. Gerin I, Bommer GT, McCoIn CS, Sousa KM, Krishnan V, MacDougald OA. Roles for microRNA-378/378* in adipocyte gene expression and lipogenesis. Am J Physiol Endocrinol Metab. 2010; 299(2):E198–206. https://doi.org/10.1152/ajpendo.00179.2010 PMID: 20484008;

53. Lee DY, Deng Z, Wang C-H, Yang BB. MicroRNA-378 promotes cell survival, tumor growth, and angiogenesis by targeting Smad2 and Fas-1 expression. Proceedings of the National Academy of Sciences. 2007; 104(51):20350–5. https://doi.org/10.1073/pnas.0706901104 PMID: 18077375
54. Wei X, Li H, Zhang B, Li C, Dong D, Lan X, et al. miR-378a-3p promotes differentiation and inhibits proliferation of myoblasts by targeting HDAC4 in skeletal muscle development. RNA Biology. 2016; 13 (12):1300–9. https://doi.org/10.1080/15476286.2016.1239008 PMID: 27661135.

55. Yoon S, Han E, Choi YC, Kee H, Jeong Y, Yoon J, et al. Inhibition of cell proliferation and migration by miR-509-3p that targets CDK2, Rac1, and PIK3CA. Mol Cells. 2014; 37(4):314–21. Epub 2014/05/08. https://doi.org/10.14348/molcells.2014.2360 PMID: 24802056;

56. Zhang G, Liu Z, Han Y, Wang X, Yang Z. Overexpression of miR-509 Increases Apoptosis and Inhibits Invasion via Suppression of Tumor Necrosis Factor-alpha in Triple-Negative Breast Cancer HS78T Cells. Oncol Res. 2016; 24(4):233–8. Epub 2016/09/24. https://doi.org/10.3727/09650416X1464870144777 PMID: 27866833.

57. Shi X, Zhan L, Xiao C, Lei Z, Yang H, Wang L, et al. miR-1238 inhibits cell proliferation by targeting Bcl-2 to modulate multidrug resistance by targeting high mobility group-box 3 protein in gastric cancer. Tumour Biol. 2014; 35(11):11081–9. Epub 2014/08/07. https://doi.org/10.1007/s13277-014-2405-z PMID: 25095978.

58. Wang P, Deng Y, Fu X. MiR-509-5p suppresses the proliferation, migration, and invasion of non-small cell lung cancer by targeting YWHAG. Biochem Biophys Res Commun. 2017; 482(4):935–41. https://doi.org/10.1016/j.bbrc.2016.11.136 PMID: 27894843.

59. Chen X, Zhao G, Wang F, Gao F, Luo H, Wang Y, et al. Upregulation of miR-513b inhibits cell proliferation, migration, and promotes apoptosis by targeting high mobility group-box 3 protein in gastric cancer. Tumour Biol. 2014; 35(11):11081–9. Epub 2014/08/07. https://doi.org/10.1007/s13277-014-2405-z PMID: 25095978.

60. Wang P, Deng Y, Fu X. MiR-509-5p suppresses the proliferation, migration, and invasion of non-small cell lung cancer by targeting YWHAG. Biochem Biophys Res Commun. 2017; 482(4):935–41. https://doi.org/10.1016/j.bbrc.2016.11.136 PMID: 27894843.

61. Chen X, Zhao G, Wang F, Gao F, Luo H, Wang Y, et al. Upregulation of miR-513b inhibits cell proliferation, migration, and promotes apoptosis by targeting high mobility group-box 3 protein in gastric cancer. Tumour Biol. 2014; 35(11):11081–9. Epub 2014/08/07. https://doi.org/10.1007/s13277-014-2405-z PMID: 25095978.

62. Chen WC, Lin MS, Ye YL, Gao HJ, Song ZY, Shen XY. microRNA expression pattern and its alteration following celecoxib intervention in human colorectal cancer. Experimental and therapeutic medicine. 2012; 3(6):1039–48. Epub 2012/09/13. https://doi.org/10.3892/etm.2012.531 PMID: 22970014;

63. Xu K, Liang X, Cui D, Wu Y, Shi W, Liu J. miR-509-3p that targets CDK2, Rac1, and PIK3CA. Mol Cells. 2014; 37(4):314–21. Epub 2014/05/08. https://doi.org/10.14348/molcells.2014.2360 PMID: 24802056;

64. Nakazawa K, Dashzeveg N, Yoshida K. Tumor suppressor p53 induces miR-1915 processing to inhibit Bcl-2 in the apoptotic response to DNA damage. FEBS J. 2014; 281(13):2937–44. Epub 2014/05/13. https://doi.org/10.1111/febs.12831 PMID: 24814047.

65. Xu K, Liang X, Cui D, Wu Y, Shi W, Liu J. miR-1915 inhibits Bcl-2 to modulate multidrug resistance by increasing drug-sensitivity in human colorectal carcinoma cells. Mol Carcinog. 2013; 52(1):70–8. Epub 2012/11/29. https://doi.org/10.1002/mc.21832 PMID: 22121083;

66. Iorio MV, Ferracin M, Liu CG, Veronese A, Spizzo R, Sabbioni S. MicroRNA gene expression deregulation in human breast cancer. Cancer Res. 2005; 65:7065–70. https://doi.org/10.1158/0008-5472.CAN-05-1783 PMID: 16103053.

67. Wei X, Li H, Zhang B, Li C, Dong D, Lan X, et al. Widespread Estrogen-Dependent Repression of microRNAs Involved in Breast Tumor Cell Growth. Cancer Res. 2009; 69 (21):8332–40. https://doi.org/10.1158/0008-5472.CAN-09-2206 PMID: 19826037.

68. Oztemur Y, Bekmez T, Aydos A, Yulug IG, Bozkurt B, Dedeoglu BG. A ranking-based meta-analysis reveals let-7 family as a meta-signature for grade classification in breast cancer. PLoS One. 2015; 10 (5):e0126837. Epub 2015/05/16. https://doi.org/10.1371/journal.pone.0126837 PMID: 25978727;

69. Liu K, Zhang C, Li T, Ding Y, Yu T, Zhou F, et al. Let-7a inhibits growth and migration of breast cancer cells by targeting HMGA1. Int J Oncol. 2015; 46(6):2526–34. Epub 2015/04/08. https://doi.org/10.3892/ijo.2015.2949 PMID: 25846193.

70. Baffa R, Fassan M, Volinia S, O’Hara B, Liu CG, Palazzo JP, et al. MicroRNA expression profiling of human metastatic cancers identifies cancer gene targets. J Pathol. 2009; 219(2):214–21. Epub 2009/07/14. https://doi.org/10.1002/path.2586 PMID: 19583777.

71. Castilla MA, Moreno-Bueno G, Romero-Pérez L, De Vijver KV, Biscuola M, López-García MA, et al. Micro-RNA signature of the epithelial–mesenchymal transition in endometrial carcinomas. The Journal of Pathology. 2011; 223(1):72–80. https://doi.org/10.1002/path.2802 PMID: 21125666.
73. Weng Z, Wang D, Zhao W, Song M, You F, Yang L, et al. microRNA-450a targets DNA methyltransferase 3a in hepatocellular carcinoma. Experimental and therapeutic medicine. 2011; 2(5):951–5. Epub 2012/09/15. https://doi.org/10.3892/etm.2011.288 PMID: 22977604;

74. Li J, Wei J, Mei Z, Yin Y, Li Y, Lu M, et al. Suppressing role of miR-520a-3p in breast cancer through CCND1 and CD44. American journal of translational research. 2017; 9(1):146–54. Epub 2017/01/27. PMID: 28123641;

75. Lowery A, Miller N, Devaney A, McNeill R, Davoren P, Lemetre C, et al. MicroRNA signatures predict oestrogen receptor, progesterone receptor and HER2/neu receptor status in breast cancer. Breast Cancer Research. 2009; 11(3):R27. https://doi.org/10.1186/bcr2257 PMID: 19432961;

76. Tsuno S, Wang X, Shomori K, Hasegawa J, Miura N. Hsa-miR-520d induces hepatoma cells to form normal liver tissues via a stemness-mediated process. Scientific reports. 2014; 4:3852. https://doi.org/10.1038/srep03852 PMID: 24458129

77. Yan L, Yu J, Tan F, Ye GT, Shen ZY, Liu H, et al. SP1-mediated microRNA-520d-5p suppresses tumor growth and metastasis in colorectal cancer by targeting CTHRC1. American journal of cancer research. 2015; 5(4):1447–59. Epub 2015/06/24. PMID: 26101709;

78. Cimino D, De Pittà C, Orso F, Zampini M, Casara S, Penna E, et al. miR148b is a major coordinator of breast cancer progression in a relapse-associated miRNA signature by targeting ITGA5, ROCK1, PIK3CA, NRAS, and CSF1. The FASEB Journal. 2013; 27(3):1223–35. https://doi.org/10.1096/fj.12-214692 PMID: 23233531

79. Ward A, Balwierz A, Zhang JD, Kublbeck M, Pawliuk R, Hielscher T, et al. Re-expression of microRNA-375 reverses both tamoxifen resistance and accompanying EMT in breast cancer. Oncogene. 2013; 32(9):1173–82. https://doi.org/10.1038/onc.2012.128 PMID: 22508479

80. Haldrup C, Kosaka N, Ochiya T, Borre M, Hoyer S, Orntoft TF, et al. Profiling of circulating microRNAs for prostate cancer biomarker discovery. Drug delivery and translational research. 2014; 4(1):19–30. Epub 2015/03/20. https://doi.org/10.1007/s13346-013-0169-4 PMID: 25786615.

81. Chong GO, Jeon HS, Han HS, Son JW, Lee YH, Hong DG, et al. Differential MicroRNA Expression Profiles in Primary and Recurrent Epithelial Ovarian Cancer. Anticancer Res. 2015; 35(5):2611–7. Epub 2015/05/13. PMID: 25964536.

82. Wei Z, Liu Y, Wang Y, Zhang Y, Luo Q, Man X, et al. Downregulation of Foxo3 and TRIM31 by miR-148b in side population promotes cell proliferation, invasion, and drug resistance of ovarian cancer. Medical oncology (Northwood, London, England). 2016; 33(11):126. Epub 2016/10/16. https://doi.org/10.1007/s12032-016-0942-9 PMID: 27743201;

83. Ward A, Balwierz A, Zhang JD, Kublbeck M, Pawliuk R, Hielscher T, et al. Downregulation of Foxo3 and TRIM31 by miR-663a in breast cancer patients with triple-negative breast cancer. Scientific reports. 2016; 7(21):31177–90. Epub 2016/04/23. https://doi.org/10.18632/oncotarget.8870 PMID: 27105517;

84. Pan J, Hu H, Zhou Z, Sun L, Peng L, Yu L, et al. Tumor-suppressive miR-663 gene induces mitotic catastrophe growth arrest in human gastric cancer cells. Oncol Rep. 2010; 24(1):105–12. Epub 2010/06/02. PMID: 20514450.

85. Till E, Michaille J-J, Adair B, Alder H, Limagne E, Taccioli C, et al. Resveratrol decreases the levels of miR-155 by upregulating miR-663, a microRNA targeting JunB and JunD. Carcinogenesis. 2010; 31(9):1561–6. https://doi.org/10.1093/carcin/bgp143 PMID: 20622002

86. Jiao L, Deng Z, Xu C, Yu Y, Li Y, Yang C, et al. miR-663 induces castration-resistant prostate cancer transformation and predicts clinical recurrence. J Cell Physiol. 2014; 229(7):834–44. Epub 2013/11/19. https://doi.org/10.1002/jcp.24510 PMID: 24243035.

87. Huang W, Li J, Guo X, Zhao Y, Yuan X. miR-663a inhibits hepatocellular carcinoma cell proliferation and invasion by targeting HMGAA2. Biomed Pharmacother. 2016; 81:431–8. https://doi.org/10.1016/j.biopha.2016.04.034 PMID: 27216623

88. Mody HR, Hung SW, AlSaggar M, Griffin J, Govindarajan R. Inhibition of S-Adenosylmethionine-Dependent Methyltransferase Attenuates TGFbeta1-Induced EMT and Metastasis in Pancreatic
Anacardic acid and miRNAs

PLOS ONE | https://doi.org/10.1371/journal.pone.0184471 September 8, 2017 25 / 29

92. Zhang Y, Xu X, Zhang M, Wang X, Bai X, Li H, et al. MicroRNA-663a is downregulated in non-small cell lung cancer and inhibits proliferation and invasion by targeting JunD. BMC Cancer. 2016; 16:315. Epub 2016/05/18. https://doi.org/10.1186/s12885-016-2350-x PMID: 27184257;

93. Bao Y, Chen B, Wu Q, Hu K, Xi X, Zhu W, et al. Overexpression of miR-664 is associated with enhanced osteosarcoma cell migration and invasion ability via targeting SOX7. Clinical and experimental medicine. 2017; 17(1):51–8. Epub 2015/10/31. https://doi.org/10.1007/s10238-015-0398-6 PMID: 26518513.

94. Zhu H, Miao M-h, Ji X-q, Xue J, Shao X-j. miR-664 negatively regulates PLP2 and promotes cell proliferation and invasion in T-cell acute lymphoblastic leukaemia. Biochem Biophys Res Commun. 2015; 459(2):340–5. https://doi.org/10.1016/j.bbrc.2015.02.116 PMID: 25735976

95. Yang Y, Liu H, Wang X, Chen L. Up-regulation of microRNA-664 inhibits cell growth and increases cis-platin sensitivity in cervical cancer. International journal of clinical and experimental medicine. 2015; 8(10):18123–9. Epub 2016/01/16. PMID: 26770409;

96. Chen B, Bao Y, Chen X, Yi J, Liu S, Fang Z, et al. Mir-664 promotes osteosarcoma cells proliferation via downregulating of FOXO4. Biomed Pharmacother. 2015; 75:1–7. https://doi.org/10.1016/j.biopha.2015.08.012 PMID: 26463624

97. Yang H, Cho ME, Li TW, Peng H, Ko KS, Mato JM, et al. MicroRNAs regulate methionine adenosyltransferase 1A expression in hepatocellular carcinoma. J Clin Invest. 2013; 123(1):285–98. Epub 2012/12/18. https://doi.org/10.1172/JCI63861 PMID: 23241961;

98. Ding Z, Jian S, Peng X, Liu Y, Wang J, Zheng L, et al. Loss of MiR-664 Expression Enhances Cutaneous Malignant Melanoma Proliferation by Upregulating PLP2. Medicine (Baltimore). 2015; 94(33):e1327. Epub 2015/08/20. https://doi.org/10.1097/md.0000000000001327 PMID: 26287415;

99. Pignot G, Cizeron-Clairac G, Vacher S, Susini A, Tozlu S, Vieilledont A, et al. microRNA expression profile in a large series of bladder tumors: identification of a 3-miRNA signature associated with aggressiveness of muscle-invasive bladder cancer. Int J Cancer. 2013; 132(11):2479–91. Epub 2012/11/22. https://doi.org/10.1002/ijc.27949 PMID: 23169479.

100. Kalabus JL, Cheng Q, Blanco JG. MicroRNAs differentially regulate carbonyl reductase 1 (CBR1) gene expression dependent on the allele status of the common polymorphic variant rs9024. PLoS One. 2012; 7(1):e48622. Epub 2012/11/08. https://doi.org/10.1371/journal.pone.0048622 PMID: 23133646;

101. Ogata-Kawata H, Izumiya M, Kurioka D, Honma Y, Yamada Y, Furuta K, et al. Circulating exosomal microRNAs as biomarkers of colon cancer. PLoS One. 2014; 9(4):e92921. Epub 2014/04/08. https://doi.org/10.1371/journal.pone.0092921 PMID: 24703249;

102. Tan Z, Zheng H, Liu X, Zhang W, Zhu J, Wu G, et al. MicroRNA-1229 overexpression promotes cell proliferation and tumorigenicity and activates Wnt/beta-catenin signaling in breast cancer. Oncotarget. 2016; 7(17):24076–87. Epub 2016/03/19. https://doi.org/10.18632/oncotarget.11919 PMID: 26992223;

103. Vilquin P, Donini CF, Villedieu M, Grisard E, Corbo L, Bachelot T, et al. MicroRNA-125b upregulation confers aromatase inhibitor resistance and is a novel marker of poor prognosis in breast cancer. Breast Cancer Res. 2015; 17(1):13. Epub 2015/01/31. https://doi.org/10.1186/s13058-015-0515-1 PMID: 25639043;

104. Yao T, Rao Q, Liu L, Zheng C, Xie Q, Liang J, et al. Exploration of tumor-suppressive microRNAs silenced by DNA hypermethylation in cervical cancer. Virology journal. 2013; 10:175. Epub 2013/06/05. https://doi.org/10.1186/1743-422X-10-175 PMID: 23732000;

105. Wang Y, Chen M, Tao Z, Hua Q, Chen S, Xiao B. Identification of predictive biomarkers for early diagnosis of larynx carcinoma based on microRNA expression data. Cancer Genetics. 2013; 206(9–10):340–6. https://doi.org/10.1016/j.cancergen.2013.09.005 PMID: 24238754;

106. Wolter M, Werner T, Malzkorn B, Reifenberger G. Role of microRNAs Located on Chromosome Arm 10q in Malignant Gliomas. Brain Pathol. 2016; 26(3):344–58. Epub 2015/08/01. https://doi.org/10.1111/bpa.12294 PMID: 26223576.

107. Pieper LA, Strotbek M, Wengert T, Olayioye MA, Haussur A. ATP6beta-based fine-tuning of the unfolded protein response enhances therapeutic antibody productivity of Chinese Hamster Ovary cells. Biotechnol Bioeng. 2017. Epub 2017/02/07. https://doi.org/10.1002/bit.26263 PMID: 28165157.

108. Chen G, Hu J, Huang Z, Yang L, Chen M. MicroRNA-1976 functions as a tumor suppressor and serves as a prognostic indicator in non-small cell lung cancer by directly targeting PLCE1. Biochem Biophys Res Commun. 2016; 473(4):1144–51. Epub 2016/04/12. https://doi.org/10.1016/j.bbrc.2016.04.030 PMID: 27063799.

109. Xin J, Li J, Feng Y, Wang L, Zhang Y, Yang R. Downregulation of long noncoding RNA HOTAIRM1 promotes monocyte/dendritic cell differentiation through competitively binding to endogenous miR-
110. Hu R, Liu W, Li H, Yang L, Chen C, Xia ZY, et al. A Runx2/miR-3960/miR-2861 regulatory feedback loop during mouse osteoblast differentiation. J Biol Chem. 2011; 286(14):12328–39. Epub 2011/02/18. https://doi.org/10.1074/jbc.M110.176099 PMID: 21324897;

111. Marrale M, Albanese NN, Cali F, Romano V. Assessing the impact of copy number variants on miRNA genes in autism by Monte Carlo simulation. PLoS One. 2014; 9(3):e90947. Epub 2014/03/29. https://doi.org/10.1371/journal.pone.0090947 PMID: 24667286;

112. Sripada L, Singh K, Lipatova AV, Singh A, Prajapati P, Tomar D, et al. hsa-miR-4485 regulates mitochondrial functions and inhibits the tumorigenicity of breast cancer cells. Journal of molecular medicine (Berlin, Germany). 2017. Epub 2017/02/22. https://doi.org/10.1007/s00109-017-1517-5 PMID: 28220193.

113. Li X, Lv Y, Hao J, Sun H, Gao N, Zhang C, et al. Role of microRNA-4516 involved autophagy associated with exposure to fine particulate matter. Oncotarget. 2016; 7(29):45385–97. Epub 2016/06/23. https://doi.org/10.18632/oncotarget.9976 PMID: 27329587;

114. Borrelli N, Denaro M, Ugolini C, Poma AM, Miccoli M, Vitti P, et al. miRNA expression profiling of 'non-invasive follicular thyroid neoplasms with papillary-like nuclear features' compared with adenomas and infiltrative follicular variants of papillary thyroid carcinomas. Mod Pathol. 2017; 30(1):39–51. Epub 2016/09/03. https://doi.org/10.1038/modpathol.2016.157 PMID: 27586203.

115. Chowdhari S, Saini N. hsa-miR-4516 mediated downregulation of STAT3/CDK6/UBE2N plays a role in PUVA induced apoptosis in keratinocytes. J Cell Physiol. 2014; 229(11):1630–8. Epub 2014/03/13. https://doi.org/10.1002/jcp.24608 PMID: 24610393.

116. Shimomura A, Shiino S, Kawauchi J, Takizawa S, Sakamoto H, Matsu zaki J, et al. Novel combination of serum microRNA for detecting breast cancer in the early stage. Cancer Sci. 2016; 107(3):326–34. Epub 2016/01/11. https://doi.org/10.1111/cas.12880 PMID: 26749252;

117. Ma F, Liu X, Li D, Wang P, Li N, Lu L, et al. MicroRNA-466l Upregulates IL-10 Expression in TLR-Trig-gered Macrophages by Antagonizing RNA-Binding Protein Tristetraproli n-Media ted IL-10 mRNA Deg-radation. The Journal of Immunology. 2010; 184(11):6053. https://doi.org/10.4049/jimmunol.0902308 PMID: 20410487.

118. Yu S, Liu X, Zhang Y, Li J, Chen S, Zheng H, et al. Circulating microRNA124-3p, microRNA9-3p and microRNA196b-5p may be potential signatures for differential diagnosis of thyroid nodules. Oncotarget. 2016; 7(51):84165–77. Epub 2016/10/06. https://doi.org/10.18632/oncotarget.12389 PMID: 27798935.

119. Liu J, Yan J, Zhou C, Ma Q, Jin Q, Yang Z. miR-1285-3p acts as a potential tumor suppressor miRNA via downregulating JUN expression in hepatocellular carcinoma. Tumour Biol. 2015; 36(1):219–25. Epub 2014/09/19. https://doi.org/10.1007/s13277-014-2622-5 PMID: 22142236;

120. Yoo JK, Kim J, Choi SJ, Noh HM, Kwon YD, Yoo H, et al. Discovery and character ization of novel microRNAs during endothelial differentiat ion of human embryoni c stem cells. Stem cells and develop -ment. 2012; 21(11):2049–57. Epub 2011/12/07. https://doi.org/10.1089/scd.2011.0500 PMID: 21422366.

121. Zawada AM, Zhang L, Emrich IE, Rogacev KS, Krezdorn N, Rotter B, et al. MicroRNA profiling of human intermediate monocytes. Immunobiology. 2017; 222(3):587–96. https://doi.org/10.1016/j.imbilo.2016.11.006 PMID: 27876379.

122. Kanlikilicer P, Rashed MH, Bayraktar R, Mitra R, Ivan C, Aslan B, et al. Ubiquitous Release of Exosomal Tumor Suppressor miR-6126 from Ovarian Cancer Cells. Cancer Res. 2016; 76(24):7194–207. Epub 2016/10/16. https://doi.org/10.1158/0008-5472.CAN-16-0714 PMID: 27742688.

123. Kim JW, Mori S, Nevins JR. Myc-induced microRNAs integrate Myc-mediated cell proliferation and cell fate. Cancer Res. 2010; 70(12):4820–8. Epub 2010/06/03. https://doi.org/10.1158/0008-5472. CAN-10-0659 PMID: 20516112;

124. Wang C, Bian Z, Wei D, Zhang JG. miR-29b regulates migration of human breast cancer cells. Mol Cell Biochem. 2011; 352(1–2):197–207. Epub 2011/03/02. https://doi.org/10.1007/s11010-011-0755-z PMID: 21359530.

125. Paris O, Ferraro L, Grober OMV, Ravo M, De Filippo MR, Giurato G, et al. Direct regulation of microRNA biogenesis and expression by estrogen receptor beta in hormone-responsive breast cancer. Oncogene. 2012; 31(38):4196–206. https://doi.org/10.1038/onc.2011.583 PMID: 22231442.

126. Pellegrino L, Stebbing J, Braga VM, Frampton AE, Jacob J, Buluwela L, et al. miR-23b regulates cytoskeletal remodeling, motility and metastasis by directly targeting multiple transcripts. Nucleic Acids Res. 2013; 41(10):5400–12. Epub 2013/04/13. https://doi.org/10.1093/nar/gkt245 PMID: 23580553;
127. Pellegrino L, Krell J, Roca-Alonso L, Stebbing J, Castellano L. MicroRNA-23b regulates cellular architecture and impairs motogenic and invasive phenotypes during cancer progression. Bioarchitecture. 2013; 3(4):119–24. Epub 2013/09/05. https://doi.org/10.4161/bio.26134 PMID: 24002530;

128. Li P, Xu T, Zhou X, Liao L, Pang G, Luo W, et al. Downregulation of miRNA-141 in breast cancer cells is associated with cell migration and invasion: involvement of ANP32E targeting. Cancer medicine. 2015; 2(8):2135–44. Epub 2015/04/18. https://doi.org/10.1002/cam4.1024 PMID: 28220627;

129. Ell B, Qiu Q, Wei Y, Mercatali L, Ibrahim T, Amadori D, et al. The microRNA-23b/27b/141 cluster targets proline oxidase, a novel tumor suppressor protein in renal cancer. Oncogene. 2010; 29(35):4914–24. https://doi.org/10.1038/onc.2010.237 PMID: 20562915.

130. Uhlmann S, Zhang JD, Schwager A, Mannsperger H, Riazalhosseini Y, Burmester S, et al. miR-200bc/429 cluster targets PLC[gamma]1 and differentially regulates proliferation and EGF-driven invasion than miR-200a/141 in breast cancer. Oncogene. 2010; 29(30):4257–66. https://doi.org/10.1038/onc.2010.201 PMID: 20514023.

131. Ma F, Li W, Liu C, Li W, Yu H, Lei B, et al. MiR-23a promotes TGF-beta1-induced EMT and tumor metastasis in breast cancer cells by directly targeting CDH1 and activating Wnt/beta-catenin signaling. Oncotarget. 2017. Epub 2017/06/18.

132. Liu W, Zabirnyk O, Wang H, Shiao YH, Nickerson ML, Khalil S, et al. miR-23b/27b/24 cluster targets ZEB1 and members of the miR-200 family promotes EMT and invasion in cancer cells. EMBO Rep. 2008; 9(6):582–8. https://doi.org/10.1038/embr.2008.74 PMID: 18483486.

133. Roy SS, Gonugunta VK, Bandypadhyay A, Rao MK, Goodall GJ, Sun L, et al. Significance of PELP1/HDAC2/miR-200 regulatory network in EMT and metastasis of breast cancer. Oncogene. 2014; 33(28):3707–16. https://doi.org/10.1038/onc.2013.332 PMID: 23975430.

134. Li T, Lu H, Mukherjee D, Lahini SK, Chen C, Yu L, et al. Identification of epidermal growth factor receptor and its inhibitory microRNA141 as novel targets of Kruppel-like factor 8 in breast cancer. Oncotarget. 2015; 6(25):21428–42. Epub 2015/05/31. https://doi.org/10.18632/oncotarget.4077 PMID: 26025929.

135. Finlay-Schultz J, Wellner U, Schmalhofer O, Vincan E, Spaderna S, et al. A reciprocal repression between ZEB1 and members of the miR-200 family promotes EMT and invasion in cancer cells. J Biol Chem. 2015; 290(5):3033–44. Epub 2014/12/17. https://doi.org/10.1074/jbc.M114.615179 PMID: 25505268.

136. Kim T, Veronesi A, Pichiorri F, Lee TJ, Jeon Y-J, Volinia S, et al. p53 regulates epithelial–mesenchymal transition through microRNAs targeting ZEB1 and ZEB2. The Journal of Experimental Medicine. 2011; 208(5):875–83. https://doi.org/10.1084/jem.20110235 PMID: 21518799.

137. Dimri M, Cho JH, Kang M, Dimri GP. PLK1 inhibition down-regulates polycomb group protein BMI1 via modulation of the miR-200c/141 cluster. J Biol Chem. 2015; 290(5):3033–44. Epub 2014/12/17. https://doi.org/10.1074/jbc.M114.615179 PMID: 25505268.

138. Vrba L, Jensen TJ, Garbe JC, Heimark RL, Cress AE, Dickinson S, et al. Role for DNA Methylation in the Regulation of miR-200c and miR-141 Expression in Normal and Cancer Cells. PLoS ONE. 2010; 5(1):e8697. https://doi.org/10.1371/journal.pone.0008697 PMID: 20841174.

139. Antolin S, Calvo L, Blanco-Calvo M, Santiago MP, Lorenzo-Patino MJ, Haz-Conde M, et al. Circulating miR-200c and miR-141 and outcomes in patients with breast cancer. BMC Cancer. 2015; 15:297. Epub 2015/04/18. https://doi.org/10.1186/s12885-015-1238-5 PMID: 25885099.

140. Madhavan D, Peng C, Wallwiener M, Zucknick M, Nees J, Schott S, et al. Circulating miRNAs with prognostic value in metastatic breast cancer and for early detection of metastasis. Carcinogenesis. 2016; 37(5):461–70. https://doi.org/10.1093/carcin/bgw008 PMID: 26785733.

141. Debeb BG, Lacerda L, Anfossi S, Diagaradjane P, Chu K, Bambhroliya A, et al. miR-141-Mediated Regulation of Brain Metastasis From Breast Cancer. J Natl Cancer Inst. 2016; 108(8). Epub 2016/04/15. https://doi.org/10.1093/jnci/djw026 PMID: 27075851;
145. Yao YS, Qiu WS, Yao RY, Zhang Q, Zhuang LK, Zhou F, et al. miR-141 confers docetaxel chemoresistance of breast cancer cells via regulation of EIF4E expression. Oncol Rep. 2015; 33(5):2504–12. Epub 2015/03/31. https://doi.org/10.3892/or.2015.3866 PMID: 25813250.

146. Abedi N, Mohammadi-Yeganeh S, Koochaki A, Karami F, Paryan M. miR-141 as potential suppressor of beta-catenin in breast cancer. Tumour Biol. 2015; 36(12):9895–901. Epub 2015/07/15. https://doi.org/10.1007/s13277-015-3737-y PMID: 26164002.

147. Xu Y, Gu L, Pan Y, Li R, Gao T, Song G, et al. Different effects of three polymorphisms in MicroRNAs on cancer risk in Asian population: evidence from published literatures. PLoS One. 2013; 8(6):e65123. Epub 2013/06/12. https://doi.org/10.1371/journal.pone.0065123 PMID: 23750236.

148. Jarrett A, McFarland AP, Horner SM, Kell A, Schwerk J, Hong M, et al. Hepatitis-C-virus-induced microRNAs dampen interferon-mediated antiviral signaling. Nat Med. 2016; 22(12):1475–81. https://doi.org/10.1038/nm.4211 PMID: 27841874.

149. Benetatos L, Hatzimichael E, Londin E, Vartholomatos G, Loher P, Rigoutsos I, et al. The microRNAs within the DLK1-DIO3 genomic region: involvement in disease pathogenesis. Cell Mol Life Sci. 2013; 70(5):795–814. https://doi.org/10.1007/s00018-012-1080-8 PMID: 22825660.

150. Cinegaglia NC, Andrade SCS, Tokar T, Pinheiro M, Severino FE, Oliveira RA, et al. Integrative transcriptome analysis identifies deregulated microRNA-transcription factor networks in lung adenocarcinoma. Oncotarget. 2016; 7(20).

151. Shi S, Lu Y, Qin Y, Li W, Cheng H, Xu Y, et al. miR-1247 is correlated with prognosis of pancreatic cancer and inhibits cell proliferation by targeting neuropilins. Curr Mol Med. 2014; 14(3):316–27. Epub 2014/03/05. PMID: 24588767.

152. Zhang J, Fu J, Pan Y, Zhang X, Shen L. Silencing of miR-1247 by DNA methylation promoted non-small-cell lung cancer cell invasion and migration by effects of STMN1. OncoTargets and Therapy. 2016; 9:7297–307. Epub 2016/12/13. https://doi.org/10.2147/OTT.S11291 PMID: 22825660.

153. Scaravilli M, Porkka KP, Brofeldt A, Annala M, Tammela TL, Jenster GW, et al. MiR-1247-5p is overexpressed in castration resistant prostate cancer and targets MYCBP2. Prostate. 2015; 75(8):798–805. Epub 2015/03/04. https://doi.org/10.1002/pros.22961 PMID: 25731699.

154. Martinez-Sanchez A, Murphy CL. miR-1247 Functions by Targeting Cartilage Transcription Factor SOX9. J Biol Chem. 2013; 288(43):30802–14. https://doi.org/10.1074/jbc.M113.496729 PMID: 24014021.

155. Zhao F, Lv J, Gan H, Li Y, Wang R, Zhang H, et al. MiRNA profile of osteosarcoma with CD117 and stro-1 expression: miR-1247 functions as an onco-miRNA by targeting MAP3K9. Int J Clin Exp Pathol. 2015; 8(2):1451–8. Epub 2015/05/15. PMID: 25973030.

156. Caussy C, Charrière S, Meirhaегhe A, Dallongeville J, Lefai E, Rome S, et al. Multiple microRNA regulation of lipoprotein lipase gene abolished by 3'UTR polymorphisms in a triglyceride-lowering haplotype harboring p.Ser474Ter. Atherosclerosis. 2016; 246:280–6. https://doi.org/10.1016/j.atherosclerosis.2016.01.010 PMID: 26820803.

157. Yang F, Nam S, Brown CE, Zhao R, Starr R, Ma Y, et al. A novel berbamine derivative inhibits cell viability and induces apoptosis in cancer stem-like cells of human glioblastoma, via up-regulation of miRNA-4284 and JNK/AP-1 signaling. PLoS One. 2014; 9(4):e94443. Epub 2014/04/16. https://doi.org/10.1371/journal.pone.0094443 PMID: 24732116.

158. Munari E, Marchionni L, Chitre A, Hayashi M, Martignoni G, Brunelli M, et al. Clear cell papillary renal cell carcinoma: micro-RNA expression profiling and comparison with clear cell renal cell carcinoma and papillary renal cell carcinoma. Hum Pathol. 2014; 45(6):1130–8. http://dx.doi.org/10.1016/j.humpath.2014.01.013 PMID: 24703100.

159. Hisaoa M, Matsumoto M, Aberrant Calreticulin Expression Is Involved in the Dedifferentiation of Dedifferentiated Liposarcoma. The American Journal of Pathology. 2012; 180(5):2076–83. https://doi.org/10.1016/j.ajpath.2012.01.042 PMID: 22429966.
163. Yang H, Wang Y. Five miRNAs considered as molecular targets for predicting neuroglioma. Tumour Biol. 2016; 37(1):1051–9. Epub 2015/08/14. https://doi.org/10.1007/s13277-015-3898-9 PMID: 26269115.

164. Zhou Y, Dang J, Chang KY, Yau E, Aza-Blanc P, Moscat J, et al. miR-1298 Inhibits Mutant KRAS-Driven Tumor Growth by Repressing FAK and LAMB3. Cancer Res. 2016; 76(19):5777–87. Epub 2016/10/05. https://doi.org/10.1158/0008-5472.CAN-15-2936 PMID: 27698189;

165. Hu W, Wang M, Yin H, Yao C, He Q, Yin L, et al. MicroRNA-1298 is regulated by DNA methylation and affects vascular smooth muscle cell function by targeting connexin 43. Cardiovasc Res. 2015; 107(4):534–45. Epub 2015/05/31. https://doi.org/10.1093/cvr/cvv160 PMID: 26025955.

166. Othman N, In LL, Harikrishna JA, Hasima N. Bcl-xL silencing induces alterations in hsa-miR-608 expression and subsequent cell death in A549 and SK-LU1 human lung adenocarcinoma cells. PLoS One. 2013; 8(12):e81735. Epub 2013/12/18. https://doi.org/10.1371/journal.pone.0081735 PMID: 24339958;

167. Wu Q, Guo L, Jiang F, Li L, Li Z, Chen F. Analysis of the miRNA-mRNA-lncRNA networks in ER+ and ER- breast cancer cell lines. J Cell Mol Med. 2015; 19(12):2877–87. Epub 2015/09/30. https://doi.org/10.1111/jcmm.2015.19.issue-12 PMID: 26416600;

168. Saini HK, Griffiths-Jones S, Enright AJ. Genomic analysis of human microRNA transcripts. Proceedings of the National Academy of Sciences. 2007; 104(45):17719–24. https://doi.org/10.1073/pnas.0703890104 PMID: 17965236.

169. Gerhauser C. Cancer Chemoprevention and Nutri-Epigenetics: State of the Art and Future Challenges. In: Pezzuto JM, Suh N, editors. Natural Products in Cancer Prevention and Therapy. Topics in Current Chemistry. 329: Springer Berlin Heidelberg; 2013. p. 73–132.

170. van Kampen JGM, van Hooij O, Jansen CF, Smit FP, van Noort PI, Schultz I, et al. miRNA-520f Reverses Epithelial-to-Mesenchymal Transition by Targeting ADAM9 and TGFBR2. Cancer Res. 2017; 77(8):2008–17. Epub 2017/02/18. https://doi.org/10.1158/0008-5472.CAN-16-2609 PMID: 28209612;

171. Mammoto A, Sasaki T, Kim Y, Takai Y. Physical and functional interaction of rabphilin-11 with mammalian Sec13 protein. Implication in vesicle trafficking. J Biol Chem. 2000; 275(18):16715–70. Epub 2000/04/05. PMID: 10747849.

172. Nik-Zainal S, Davies H, Staaf J, Ramakrishna M, Glodzik D, Zou X, et al. Landscape of somatic mutations in 560 breast cancer whole-genome sequences. Nature. 2016; 534(7605):47–54. https://doi.org/10.1038/nature17676 PMID: 27135926.

173. Duan H, Cherradi N, Feige J-J, Jefcoate C. cAMP-Dependent Posttranscriptional Regulation of Steroidogenic Acute Regulatory (STAR) Protein by the Zinc Finger Protein ZFP36L1/TIS11b. Mol Endocrinol. 2009; 23(4):497–509. https://doi.org/10.1210/me.2008-0296 PMID: 19179481;

174. Vignudelli T, Selmi T, Martello A, Parenti S, Grande A, Gemelli C, et al. ZFP36L1 Negatively Regulates Erythroid Differentiation of CD34+ Hematopoietic Stem Cells by Interfering with the Stat5b Pathway. Mol Biol Cell. 2010; 21(19):3340–51. https://doi.org/10.1091/mbc.E10-01-0040 PMID: 20702587.

175. Adachi S, Homoto M, Tanaka R, Hori Y, Murakami H, Suga H, et al. ZFP36L1 and ZFP36L2 control LDLR mRNA stability via the ERK-RSK pathway. Nucleic Acids Res. 2014; 42(15):10037–49. Epub 2014/08/12. https://doi.org/10.1093/nar/gku652 PMID: 25106868;

176. Remijisen Q, Kuipers TW, Wirawan E, Lippens S, Vandenabeele P, Vanden Berghe T. Dying for a cause: NETosis, mechanisms behind an antimicrobial cell death modality. Cell Death Differ. 2011; 18(4):581–8. https://doi.org/10.1038/cdd.2011.1 PMID: 21293492