An integrative genomic approach identifies p73 and p63 as activators of miR-200 microRNA family transcription

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ABSTRACT

Although microRNAs (miRNAs) are important regulators of gene expression, the transcriptional regulation of miRNAs themselves is not well understood. We employed an integrative computational pipeline to dissect the transcription factors (TFs) responsible for altered miRNA expression in ovarian carcinoma. Using experimental data and computational predictions to define miRNA promoters across the human genome, we identified TFs with binding sites significantly overrepresented among miRNA genes overexpressed in ovarian carcinoma. This pipeline nominated TFs of the p53/p63/p73 family as candidate drivers of miRNA overexpression. Analysis of data from an independent set of 253 ovarian carcinomas in The Cancer Genome Atlas showed that p73 and p63 expression is significantly correlated with expression of miRNAs whose promoters contain p53/p63/p73 family binding sites. In experimental validation of specific miRNAs predicted by the analysis to be regulated by p73 and p63, we found that p53/p63/p73 family binding sites modulate promoter activity of miRNAs of the miR-200 family, which are known regulators of cancer stem cells and epithelial-mesenchymal transitions. Furthermore, in chromatin immunoprecipitation studies both p73 and p63 directly associated with the miR-200b/a/429 promoter. This study delineates an integrative approach that can be applied to discover transcriptional regulatory mechanisms in other biological settings where analogous genomic data are available.

INTRODUCTION

Regulation of gene expression at the post-transcriptional level is governed in part by microRNAs (miRNAs), which are approximately 22 nucleotide non-protein-encoding RNAs that modulate the stability and/or translation of messenger RNAs (mRNAs) via partially complementary base-pairing interactions (1). Most microRNAs are transcribed by RNA polymerase II (2), and miRNA expression can be regulated by transcription factor (TF)

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The authors wish it to be known that, in their opinion, the first three authors should be regarded as joint First Authors.

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binding sites present in their promoters (3–7). However, for the majority of miRNAs, promoters have not been defined and the TF binding sites upstream of these miRNA loci have not been experimentally tested. Dysregulation of miRNA expression is common in human disease and contributes to pathology, since miRNAs regulate significant disease-relevant processes such as cell division, differentiation, and apoptosis (8,9). In addition, in certain cancer contexts, the pattern of miRNA expression captures important features of the developmental origin of cancers (10) and may predict the course of disease (11). However, the mechanisms underlying miRNA dysregulation are not clear, in part because the transcriptional regulation of most miRNAs is not well characterized.

In this study, we implemented an integrative computational approach to dissect the transcriptional regulation of miRNAs. We focused on the dysregulation of miRNAs in ovarian carcinoma of the serous histologic sub-type, which has a high mortality and accounts for approximately two-thirds of ovarian carcinomas. Although a subset of miRNAs dysregulated in ovarian carcinomas is associated with changes in genomic copy number and epigenetic modifications, for many miRNAs additional, unknown mechanisms appear to contribute to the re-programming of miRNA expression (12,13). We therefore sought to discover the TFs that may drive the dysregulation of miRNAs in ovarian carcinoma. We implemented a computational pipeline to annotate miRNA transcription start sites (TSS) and putative promoter regions, and then to identify the TFs with binding sites enriched in the promoters of overexpressed miRNAs in ovarian carcinoma. This approach yields putative regulatory interactions between TFs and miRNA promoters for subsequent experimental validation.

We report here that the best candidate driver of miRNA overexpression in ovarian carcinoma is the p53/p63/p73 family of TFs. Although p53 has been shown to trans-activate several miRNAs, including the miR-34 family (14–17), the transcriptional regulation of miRNA genes by p73 and p63 has not been well-described. Further analysis using data from The Cancer Genome Atlas (TCGA) suggested that, in ovarian carcinoma, p73 and p63 are primarily responsible for the altered expression of miRNAs with p53 family binding sites. We experimentally validated our approach by confirming that p73 and p63 directly regulate transcription of the miR-200 family, a novel target predicted by our analysis that is an important regulator of epithelial–mesenchymal transitions (EMTs) and of the cancer stem cell phenotype (18–22).

This study illustrates how an integrative computational analysis can identify new regulatory interactions between TFs and miRNAs. We also provide a resource by defining putative miRNA promoters and associating TF binding sites with these miRNA promoters on a genome-wide scale, and we discuss how our approach is broadly applicable to dissect TF–miRNA regulatory networks in other systems where miRNA expression data from distinct physiologic states are available.

**MATERIALS AND METHODS**

**Clinical materials**

Normal primary human ovarian surface epithelial (HOSE) cell and serous ovarian carcinomas from which RNA was analyzed in this study have been previously described (23), and are described in Supplementary Data. HOSE specimens were obtained under a protocol that was approved by the Research Review Committee and Internal Review Board at the Fox Chase Cancer Center. All ovarian carcinoma tissue specimens were collected by the Pacific Ovarian Cancer Research Consortium Repository under a protocol approved by the Fred Hutchinson Cancer Research Center Institutional Review Board.

**Analysis of miRNA microarray data generated in the current study**

We analyzed miRNA microarray data generated on a locked nucleic acid (LNA) probe-based platform designed to profile 480 human miRNAs (Exiqon, Inc.) (23). Array quality control and data normalization was performed as previously described (23). We defined expressed miRNAs as the miRNAs expressed in at least two of the four normal or at least four of the 16 serous ovarian carcinomas. Non-specific filtering was performed as previously described (23), and genes that showed low variability across samples were removed from the analysis, which yielded 181 genes for downstream analyses. Differentially expressed miRNAs were identified by considering miRNAs to be over- or underexpressed in carcinoma compared to normal if log2 fold change was greater than 1.0 and adjusted P < 0.01.

**Identification of TFs commonly expressed in serous ovarian carcinoma**

We analyzed data from Hendrix et al. (24), where Affymetrix GeneChip Human Genome U133 Array Set HG-U133A microarrays were used to profile gene expression across a panel of 4 normal ovaries and 41 serous ovarian carcinoma tissues. The carcinoma tissues comprised the following stages: Stage IA (n = 1), Stage IC (n = 3), Stage II (n = 2), Stage IIC (n = 2), Stage III (n = 2), Stage IIIB (n = 1), IIC (n = 25), IIID (n = 1) and IV (n = 5) and the following grades: well-differentiated (n = 2), moderately differentiated (n = 5), poorly differentiated (n = 18), grade information not available (n = 16). All samples were selected to contain at least 70% malignant epithelial cells. We considered genes to be expressed if they showed expression [as determined by Affymetrix PMA (Present/Marginal/Absent) calls] in at least 2 out of 4 normal samples or 8 out of 41 carcinoma samples. We identified 83 TFs expressed in normal ovary and/or serous ovarian carcinoma tissue specimens by intersecting expressed genes with all TFs in tfbsConsFactors files downloaded from the UCSC genome browser (25,26). Out of these 83 TFs, 79 corresponded to binding sites found upstream of miRNAs overexpressed in ovarian carcinoma.
Annotation of putative miRNA promoters and their TF binding sites

miRNA coordinates were downloaded from miRBase (version 10) (27) and used to map miRNAs to their genomic locations as described in Supplementary Data. To assign TSS for each miRNA locus, we used RefSeq, AceView, expressed sequence tags (ESTs) and Eponine predictions downloaded from the UCSC genome browser (hg 18 version of the genome assembly) (26). If both 5' and 3' ESTs were available from the same clone and formed a transcript containing the miRNA, the miRNA was considered expressed by this transcript and its TSS was the 5'-end of the EST.

We examined the putative promoters of each miRNA/miRNA locus for conserved TF binding sites defined in the UCSC files tbsConsSites and tbsConsFactors (26), as described in Supplementary Data. The conservation of binding sites is based on human/mouse/rat multiple alignments and the score is computed using the Transfac Matrix Database (v7.0) (28).

Enrichment of TF binding sites in promoters of overexpressed miRNA loci

We performed a Fisher’s exact test for each of the 79 overexpressed miRNA loci to compare the number of TF binding sites in the promoters of overexpressed miRNA loci versus control miRNA loci. To assess the false discovery rate, predicted Q-values were estimated from P-values (derived from Fisher’s exact test) by using the ‘Q-value’ package in R (29).

Correlations between expression of TFs and expression of miRNAs in TCGA data set

We analyzed data from 253 serous ovarian carcinoma specimens from TCGA. Serous ovarian carcinoma tissue specimens in TCGA are generally from patients with advanced stage and high-grade disease. Clinical data on stage and grade was available on 217 of the 253 specimens we studied, and confirmed that the vast majority of these were high-grade (202 were poorly differentiated and 13 were moderately differentiated) and advanced stage (Stage IIB (n = 2), Stage IIC (n = 4), Stage IIIA (n = 2), Stage IIIB (n = 9), Stage IIIC (n = 161) and Stage IV (n = 39)]. Data on percentage malignant epithelium was available for 202 of the 253 specimens we studied: >75% malignant epithelial cells (n = 150 specimens), ≥50% and <75% (n = 43 specimens), <50% (n = 9 specimens). To derive correlations between TF and miRNA expression, we retrieved Agilent gene expression microarray and miRNA microarray data from the TCGA for 253 serous ovarian carcinomas, for which both gene expression (run on AgilentG4502A_07_3) and miRNA expression profiling data were available (http://cancergenome.nih.gov/index.asp). Agilent Level 1 microarray expression data were used for both miRNA and mRNA analyses. We used 447 miRNAs for calculating correlations; all of these miRNAs had at least one TF binding site associated with their promoters as determined by our method. All TCGA Level 1 files used to calculate correlations are listed in Supplementary Table S1. Correlation between the expression of the miRNAs and TFs was assessed by calculating Pearson correlation coefficients. Bioconductor packages ‘limma’ and ‘lim’ were used to process the raw signal data and to calculate P-values for correlation between the expression of miRNAs and TFs, respectively (30,31).

Sequencing of TP53

All coding and flanking regulatory regions for exons 4-10 of TP53 were sequenced as previously described (32).

Cell culture

The 2008 ovarian carcinoma cell line was maintained in RPMI (Invitrogen) with 10% fetal bovine serum (Atlanta Biologicals) under 5% CO2. All transfections were performed using Lipofectamine 2000 (Invitrogen) according to the manufacturer’s instructions.

Chromatin immunoprecipitation

Chromatin immunoprecipitation was performed on 2008 ovarian carcinoma cells as previously described (33). Sheared chromatin was incubated with 12 μg of antibody overnight at 4°C and then incubated with 40 μl pre-washed Protein G agarose beads (Millipore). Antibodies used for immunoprecipitation included 4A4 (Santa Cruz) for p63, 259A (Imgenex) for p73, and normal mouse IgG (Santa Cruz) to control for non-specific binding. Quantitative PCR (qPCR) was performed as described in Supplementary Data using the primers listed in Supplementary Table S2. Triplicate immunoprecipitations were performed using each antibody, using independent batches of chromatin.

Promoter assays

Firefly luciferase promoter activity reporter plasmids were generated by PCR amplification of the region of interest from human genomic DNA (BioLine) followed by subcloning into the Mu1 and BglII sites of the pGL3-Enhancer vector (Promega), as described in Supplementary Data. Primers used are in Supplementary Table S3. To assay promoter activity, 2008 cells were plated in 96-well plates at 8 x 10^3 cells/well 24 h prior to transfection. The cells were co-transfected with 100 ng of each firefly luciferase promoter reporter vector, or pGL3-Enhancer as a control, and 25 ng of pRL-TK (Promega). Twenty-four hours after transfection, firefly and Renilla luciferase activity was measured with the Dual-Luciferase Reporter Assay System (Promega). Promoter activity was calculated as the ratio of firefly to Renilla luciferase signal.

RESULTS

Overview of computational pipeline for the identification of TFs regulating miRNA expression

We implemented a computational pipeline to study the transcriptional regulation of miRNA expression (Figure 1). Briefly, the pipeline starts with miRNA differential expression data as input and classifies miRNAs into...
two categories: miRNAs that are overexpressed in the samples of interest and control miRNAs that are expressed but not overexpressed. Putative promoters of both the overexpressed and control miRNAs are then defined via genomic mapping and annotation of TSS using a series of criteria described below. After identifying the TFs expressed in the samples based on mRNA expression data, the number of predicted binding sites for each expressed TF in the promoters of both the overexpressed miRNA and control miRNA loci is determined.

The output of the pipeline is TFs with binding sites overrepresented in the promoters of the overexpressed miRNAs, and these regulatory predictions can then be experimentally validated.

Identification of a set of miRNAs commonly differentially expressed in serous ovarian carcinoma

As input to our pipeline, we defined miRNAs that are differentially expressed in serous ovarian carcinomas compared to normal specimens. To avoid bias, we used multiple studies to identify miRNAs that are commonly altered in ovarian carcinoma. We previously profiled miRNA expression in primary serous ovarian carcinomas from 16 individuals and in cultured normal human ovarian surface epithelial (HOSE) cells from 4 other individuals using miRNA microarrays (23). We analyzed that data set here to identify miRNAs significantly dysregulated in serous ovarian carcinoma as compared to normal. We found 43 miRNAs to be overexpressed and 20 to be underexpressed in ovarian carcinoma (log2 fold change $>1.0$ and adjusted $P<0.01$) (Supplementary Table S4). We then compared the results of this analysis and six other ovarian carcinoma miRNA profiling studies (13,34–38). We found 30 miRNAs to be significantly overexpressed in two or more of the entire set of seven studies (Supplementary Table S5), which should provide a more reliable list of miRNAs commonly overexpressed in serous ovarian carcinomas. We also identified a set of 18 commonly underexpressed miRNAs (Supplementary Table S6), although these did not yield statistically significant overrepresentation of any specific TF binding sites in subsequent analysis and are therefore not discussed further.

In order to define a set of control miRNAs for comparison, we used data from our microarray analysis of ovarian carcinoma and normal specimens to identify 238 expressed miRNAs, which includes the 30 commonly overexpressed miRNAs defined above. We then converted this list of expressed miRNAs to a list of miRNA loci, where each locus corresponds to a single promoter, which may drive expression of multiple miRNAs. Specifically, we mapped each miRNA to its genomic position, and miRNAs that are encoded at multiple locations were assigned to each locus separately. Because many miRNAs are found in clusters in single polycistronic transcripts (39,40), we annotated all miRNAs within a genomic cluster as a single locus. After accounting for the expansion of the number of miRNA locations as a result of miRNAs encoded at multiple loci, and contraction of this number because of miRNAs in clusters with shared promoters, the 238 miRNAs corresponded to 151 loci representing a miRNA or miRNA cluster that is expressed in serous ovarian carcinomas and/or normal samples (Supplementary Table S7). We divided the 151 loci into two classes: those that encode overexpressed miRNAs and those that do not. If a single miRNA within a cluster was overexpressed, the entire miRNA locus was classified into the overexpressed class for the purpose of this analysis. From this approach, we annotated 30 loci (miRNAs or miRNA clusters) that were overexpressed in serous ovarian carcinomas, and we used...
MicroRNAs were classified into 30 loci overexpressed in ovarian carcinoma and a control group of 121 loci expressed in carcinoma and/or normal samples. The number of promoters with binding sites for a given TF was compared between the two classes. A Fisher’s exact test was performed for each of the 79 TFs that are expressed in carcinoma and/or normal samples. P-values computed by Fisher’s exact test were converted to Q-values. TFs with $P < 0.05$ are shown, while those with $Q < 0.05$ are in bold.

| TF binding site | Promoters of overexpressed miRNA | Promoters of expressed miRNAs | $P$-value | $Q$-value |
|----------------|---------------------------------|-------------------------------|-----------|-----------|
| p53/p63/p73    | 15/15                           | 17/104                        | 6.83E-05  | 0.0054    |
| EGR2           | 10/20                           | 9/12                          | 0.0006    | 0.0246    |
| SP1            | 10/20                           | 10/111                        | 0.0011    | 0.0283    |
| NFKB1          | 10/20                           | 16/105                        | 0.0145    | 0.2856    |
| RELA           | 9/21                            | 14/107                        | 0.0206    | 0.2954    |
| NR2F1          | 6/24                            | 7/114                         | 0.0233    | 0.2954    |
| EGR3           | 8/22                            | 11/110                        | 0.0262    | 0.2954    |
| PAX4           | 14/16                           | 31/90                         | 0.0429    | 0.4036    |
| XBP1           | 7/23                            | 10/111                        | 0.0460    | 0.4036    |

Table 1. TF binding sites enriched in promoters of miRNAs overexpressed in serous ovarian carcinoma

Correlation between expression of TFs and expression of miRNAs in TCGA data

To determine if expression of the candidate miRNA TFs identified correlates with expression of miRNAs with binding sites for those TFs, next we employed a larger, independent data set from TCGA, which provides comprehensive genomic profiling data from high-grade serous ovarian carcinomas. We selected samples from TCGA for which both miRNA and mRNA data were available for the same carcinoma tissue specimen. There were 253 such samples available. We calculated Pearson correlations between the expression of each of the five TFs identified above and the expression of each of 447 miRNAs in the 253 TCGA samples. We then compared correlation coefficients between the miRNAs whose promoters contain binding sites for a given TF and the remaining miRNAs lacking binding sites. We found no significant difference in the correlation of SP1 and miRNAs with or without SP1 binding sites (data not shown). However, the remaining four TFs (i.e., TP53, TP63, TP73, and EGR2) showed significantly higher expression correlations with miRNAs bearing their respective binding sites versus those without ($P < 0.01$, t-test) (Figure 2). This analysis suggests that our pipeline yields functional regulators of miRNA expression. Interestingly, we found significant negative correlations between the expression of both EGR2 probes and the expression of miRNAs with EGR2 binding sites, which suggests that EGR2 may serve as a transcriptional repressor (Figure 2D).
Expression of miRNAs with TF binding sites is correlated with TF expression in TCGA ovarian carcinoma specimens. (A–D) Data from 253 TCGA serous ovarian carcinomas were used to calculate Pearson correlations between expression of the indicated TFs and expression of 447 miRNAs. Boxplots show the correlations using mRNA expression data from two TP53 probes (A), three TP63 probes (B), four TP73 probes (C), and two EGR2 probes (D). miRNAs were classified by the presence (+) or absence (−) of the indicated TF binding site in the miRNA promoter. For each probe, t-tests were used to compare mean correlation values of the two miRNA classes, and P-values computed from the t-tests are shown. Probes: TP53-1 (NM_005427_1_1638), TP53-2 (NM_005427_2_2048), TP53-3 (A_23_P114475), TP3-1 (A_23_P114473), TP3-2 (A_23_P114472), TP3-3 (A_23_P114471), TP3-4 (NM_005427_1_1638), EGR2-1 (A_23_P46935), and EGR2-2 (A_23_P46936).  

All three members of the p53 family featured probes that showed positive, significantly higher correlations with miRNAs bearing p53 family binding sites than those without (Figure 2A–C). Three out of seven TP63 probes and one out of three TP53 probes showed lower correlations that were not statistically significant (data not shown), which may be related to sequence-specific variation in probe performance. Of the three members of the p53 family, TP73 expression showed the strongest positive correlations with miRNAs bearing p53 family binding sites (Figure 2C). Taken together, these results suggest that members of the p53 family, and in particular p73, may drive the overexpression of a panel of miRNAs in ovarian carcinoma.

Overexpression of miRNAs with p53 binding sites in their promoters is not associated with the mutational status of TP53 in serous ovarian carcinomas

While expression of p53 mRNA appeared to correlate with miRNA expression, regulation of p53 occurs primarily at the protein level. Therefore, we sought to determine whether changes in the activity of the p53 protein may affect the expression of miRNAs bearing p53 family binding sites. Mutation is the leading cause of p53 dysfunction in serous ovarian carcinoma, as 60–70% of serous ovarian cancers are known to have TP53 mutations (48,49), which in general ablate the ability of p53 to transactivate its target genes. However, because mutations to TP53 can result in a dominant negative protein that alters the activity of the other p53 family members, we examined whether TP53 mutational status affects expression of miRNAs overexpressed in ovarian carcinoma. Somatic mutation data for ovarian carcinomas in TCGA were not available. We therefore determined the p53 mutation status of the 16 ovarian carcinoma specimens that we previously used for identifying differentially expressed miRNAs. We sequenced all coding and flanking regulatory regions for exons 4-10 of TP53, which include mutation hotspots and also coincide with the most highly conserved region of the gene (50). We found non-synonymous TP53 mutations accompanied by loss of the wild-type allele in 9 out of our 16 ovarian carcinoma specimens (Supplementary Table S10). Comparing the expression of miRNAs in the TP53 mutant versus wild-type specimens demonstrated no significant difference in the expression of the 17 overexpressed miRNAs that are encoded within the 15 overexpressed loci with p53 binding sites (Supplementary Figure S1). While these results do not exclude p53 as a possible driver of...
Correlations between TP73 or TP63 and miRNA expression in TCGA ovarian carcinomas

| miRNA     | Correlation | p-value |
|-----------|-------------|---------|
| TP73      |             |         |
| miR-107   | 0.389       | 1.39E-10|
| miR-200a  | 0.389       | 1.47E-10|
| miR-200c  | 0.383       | 2.77E-10|
| miR-23a   | 0.373       | 8.27E-10|
| miR-24    | 0.371       | 1.17E-09|
| miR-200c  | 0.364       | 2.36E-09|
| miR-141   | 0.355       | 6.12E-09|
| miR-30c   | 0.348       | 1.30E-08|
| miR-29b   | 0.346       | 1.57E-08|
| miR-429   | 0.337       | 4.03E-08|
| miR-103   | 0.332       | 6.62E-08|
| miR-200b  | 0.318       | 2.31E-07|
| miR-26-5p | 0.303       | 9.31E-07|
| miR-30e   | 0.303       | 9.41E-07|
| miR-212   | 0.302       | 9.77E-07|
| miR-93    | 0.301       | 1.06E-06|
| miR-7     | 0.295       | 1.85E-06|
| miR-106b  | 0.291       | 2.46E-06|
| let-7c    | 0.289       | 2.88E-06|
| miR-20a   | 0.284       | 4.62E-06|
| miR-130a  | 0.278       | 7.03E-06|
| miR-34a   | 0.276       | 8.11E-06|
| miR-27a   | 0.274       | 9.62E-06|
| miR-135b  | 0.271       | 1.25E-05|
| miR-219-5p| 0.266       | 1.77E-05|
| miR-19a   | 0.259       | 3.02E-05|
| TP63      |             |         |
| miR-200a  | 0.322       | 1.64E-07|
| miR-29c   | 0.320       | 1.91E-07|
| miR-30e   | 0.316       | 2.94E-07|
| miR-30c   | 0.311       | 4.50E-07|
| miR-107   | 0.310       | 5.06E-07|
| miR-429   | 0.294       | 1.99E-06|
| miR-141   | 0.293       | 2.09E-06|
| miR-24    | 0.287       | 3.50E-06|
| miR-103   | 0.283       | 4.66E-06|
| miR-29b   | 0.278       | 7.10E-06|
| miR-200c  | 0.275       | 9.19E-06|
| miR-200b  | 0.263       | 2.20E-05|
| miR-93    | 0.257       | 3.57E-05|

Correlation coefficients for expression of miRNAs with TP73 expression and TP63 expression across 253 TCGA ovarian carcinomas were calculated. MicroRNAs listed showed positive correlation coefficients >0.25, P < 0.05 and contain p53/p63/p73 binding sites in their promoters. MicroRNAs are ranked in order of increasing P-value (unadjusted). The miR-200 family miRNAs, indicated in bold, had significant positive correlations with TP73 and TP63 expression. Correlation coefficients shown were calculated using representative probes for TP73 (A_23_P74081) and TP63 (A_32_P114473).

differential miRNA expression, this analysis suggests that p53 is not the primary regulator of these miRNAs in ovarian carcinoma.

p73 and p63 as putative positive regulators of miRNA expression in ovarian cancer

Our miRNA expression correlation analyses from TCGA data suggested that p73, and potentially p63, may drive the overexpression of miRNAs in ovarian carcinoma (Figures 2B and C). To identify specific miRNAs that may be regulated by p73 or p63 for further study, we used the P-values from the Pearson correlations to rank the miRNAs that were overexpressed in ovarian carcinoma and contained p53 family binding sites. We found multiple miRNAs with significant positive correlation with TP73 or TP63 expression (Table 2). A complete list of miRNAs showing positive correlations (Pearson correlation coefficient >0.25 and P < 0.05) with all p73 and p63 probes is provided in Supplementary Tables S11 and S12. To provide a biologically relevant reference for interpreting these correlation values, we also determined the correlations of 88 host gene-intronic miRNA pairs using the same data set of 253 TCGA samples. Intrinsic miRNAs (i.e., lying in host genes on the same strand) are generally co-transcribed with their host gene and therefore show strong correlations in expression with their host gene. Among miRNA-host gene pairs analyzed by the same criteria (correlation coefficients >0.25 and P < 0.05), host gene-intronic miRNA pairs showed an average correlation of 0.37 (+0.16). The correlations between the expression of p73 and p63 and miRNAs in Table 2 range between 0.25 and 0.39, and thus reflect a similar magnitude of correlation coefficients as observed with the host gene-intronic miRNA correlations.

p73 and p63 directly regulate miR-200 transcription

To experimentally validate our computational predictions, we examined the transcriptional regulation of the miR-200 family of miRNAs. Expression of all five members of the miR-200 family (miR-200a, miR-200b, miR-200c, miR-141 and miR-429) was positively correlated with p73 and p63 (all with P < 2.2 × 10^{-5}), with correlation coefficients ranging from +0.318 to +0.389 for p73, and from +0.263 to +0.322 for p63 (Table 2). Our previous work has implicated the miR-200 family in ovarian carcinoma pathogenesis (51), and the miR-200 family has a well-established role in governing EMTs and repressing tumor-initiating cells (i.e. cancer stem cells) (18–20,22) in a variety of other cancer contexts.

The miR-200 family miRNAs are transcribed from two chromosomal clusters: miR-200b/a/429 from chromosome 1, and miR-200c/141 from chromosome 12. The promoters of both miR-200 clusters contain p53 family binding sites. We first tested whether the predicted p53 family binding sites in the miRNA promoters modulate miR-200 transcription by employing a luciferase reporter system. We cloned a 2-kb fragment of the miR-200b/a/429 promoter, which contains one predicted p53 family binding site, upstream of the luciferase gene in the pG3L-Enhancer vector. Similarly, we also produced a miR-200c/141 promoter construct by cloning a 1.5-kb region of the miR-200c/141 promoter containing two predicted p53 family binding sites into the vector. We then generated mutant versions of each promoter construct by mutating the consensus nucleotides in each predicted p53 family binding site, with the miR-200c/141 mutant promoter featuring mutations at both of the two predicted binding sites (Figure 3A). We transfected the wild-type and mutated constructs into 2008 ovarian carcinoma cells. Transfection with the wild-type promoter constructs led to increased luciferase activity as compared to the empty vector.
vector lacking a promoter, indicating that the minimal miR-200 promoters were functional (data not shown). To determine if the p53 family binding sites contribute to promoter activity, we compared the luciferase activity of the wild-type versus mutated promoters (Figure 3B). We found that mutation of the p53 binding sites of the miR-200b/a/429 and miR-200c/141 promoters each resulted in a 40% decrease in luciferase activity ($P < 0.003$, t-test), consistent with the notion that members of the p53 family directly regulate miR-200 transcription.

We next sought to determine which p53 family members are physically associated with the miR-200 promoters in ovarian carcinoma cells. We performed chromatin immunoprecipitations on the 2008 cell line using antibodies directed against p53, p63, or p73, as well as normal IgG as a control for non-specific binding. We did not find enrichment of p53 at several known targets, including p21 and MDM2, nor at the miR-200 promoter regions, which is consistent with western blots revealing that 2008 cells do not express stable p53 protein (data not shown). In the p73 and p63 immunoprecipitations, however, we found a strong enrichment of the miR-200b/a/429 p53 family binding site relative to the IgG control (Figure 3C and D). The levels of enrichment were similar to those seen at the p21 promoter, a known transcriptional target of the entire p53 family (46,52). As a further negative control for non-specific enrichment, we assayed the immunoprecipitates for a region 1-kb 3′ to the miRNAs at the miR–200b/a/429 locus. No enrichment was found at this control region, indicating that we were specifically enriching the promoter of the miR-200b/a/429 locus. When the locus on chromosome 12 corresponding to the miR-200c/141 promoter was analyzed in the p73 and p63 chromatin immunoprecipitates, no enrichment was observed compared to the IgG control (data not shown). Although such negative results are difficult to interpret, this observation may be a false negative due to the limitations of the sensitivity of chromatin immunoprecipitation, such as the limited set of promoter regions assayed by qPCR as well as potentially limited antibody accessibility related to chromatin structure or interference from transcriptional co-factors. As an additional negative control, we assayed the promoter of miR-29a/b−1, which is overexpressed in ovarian carcinomas yet in our computational analysis was not found to have p53 family TF binding sites. Our
containing ovarian carcinoma, with 10 overexpressed miRNA loci binding sites upstream of miRNA loci overexpressed in ovarian carcinoma. We also observed an overrepresentation of conserved TF binding sites for the p53/p63/p73 family as significant activators of miRNA transcriptional regulation mechanisms have been investigated for specific miRNAs in the past, genome-scale studies have been limited, partially due to the lack of well-annotated miRNA primary transcripts and their corresponding transcriptional start sites. MicroRNAs are excised through a series of processing steps from primary transcripts, which are highly variable in length and can be several kilobases long; identifying the TSS for miRNAs at a genome scale is therefore not trivial. Although experimental methods using chromatin signatures to identify miRNA promoters have been useful in identifying approximate locations of miRNA TSS in some cases (53,54), such approaches have their limitations because miRNA expression is highly tissue specific and therefore experimental data applies to a relatively limited subset of miRNAs in specific cell line or tissue contexts. Our genome-wide approach relies heavily on transcript information derived from ESTs, both from EST clusters (AceView) and from ESTs derived from a common clone, to provide transcript-based evidence for defining the TSS of intergenic miRNAs. AceView is one of the most comprehensive assemblies of human transcriptome data, drawing cDNA sequences from dbEST, Genbank and RefSeq and representing a broad panel of tissue and cell type contexts. There are currently more than 8 million ESTs in dbEST and notably approximately 2 million 5' capped ESTs in Genbank submitted by Kimura et al. (55).

An important aspect of our study, therefore, is the genome-wide definition of presumptive miRNA TSS and promoters using such data, which provides a resource for further investigations of miRNA transcriptional regulation. The key biological finding of this work is the identification of the p63 and p73 TFs as significant activators of miRNA overexpression in ovarian carcinoma. We found conserved TF binding sites for the p53/p63/p73 family upstream of 17 miRNA loci overexpressed in serous ovarian carcinoma, suggesting p63 and/or p73 may coordinate regulate an ensemble of miRNAs that could significantly modify the cellular gene expression landscape via repression of a cohort of miRNA targets. We also observed an overrepresentation of conserved EGR2 binding sites upstream of miRNA loci overexpressed in ovarian carcinoma, with 10 overexpressed miRNA loci containing EGR2 binding sites. In the TCGA ovarian carcinoma data set, EGR2 expression was significantly negatively correlated with the expression of miRNAs containing EGR2 binding sites. EGR2 has been suggested to be a tumor suppressor, based on the fact that it is frequently underexpressed in human cancers (including ovarian) and cancer cell lines (56,57). We hypothesize that the 10 miRNA loci mentioned above are normally repressed by EGR2, but that decreased EGR2 expression in ovarian carcinoma leads to their overexpression.

It is important to note that both p63 and p73 can be expressed as multiple isoforms that were not distinguishable on the expression platforms involved in this study. The isoforms of p63 and p73 can be divided into two classes: transactivating TAp63/TAp73 isoforms and the inhibitory ΔNp63/ΔNp73 isoforms. In ovarian carcinoma, multiple inhibitory isoforms of p73 and p63 are overexpressed and can be associated with poor prognosis (58,59). In contrast, transactivating TAp73 increases the response rate to chemotherapy in BRCA-1-associated ovarian carcinoma (60). We hypothesize that p73 and p63 may drive the expression of multiple miRNAs that contribute to tumor aggressiveness and/or treatment response. For instance, low levels of miR-200 family miRNA expression are associated with poor treatment response in ovarian carcinoma (61); thus, TAp73-mediated activation of miR-200 could serve as a mechanism for increased sensitivity to chemotherapy. Future studies will be required to determine the role of specific isoforms of p73 and/or p63 in determining miRNA transcription in ovarian carcinoma.

In experimental validation of the pipeline predictions, we focused on the miR-200 family because of its important roles in tumor development and progression. Members of the miR-200 family inhibit EMT and suppress tumor invasion by directly repressing the TFs Zeb1 and Zeb2 (18–20). Additionally, the miR-200 family has been shown to repress Bmi1 and Suz12, two essential components of polycomb repressor complexes that are responsible for the maintenance of tumor-initiating cells (21,22). Previous work on miR-200 regulation has largely focused on mechanisms of repression, including transcriptional inhibition and epigenetic modifications (18,62,63); mechanisms of transcriptional activation of this miRNA family are not well-understood. Our results with p63 and p73 shed light onto positive transcriptional regulation of the miR-200 family miRNAs, which has potential therapeutic value in the development of approaches aimed at modifying miR-200 family miRNA expression for the treatment of diverse forms of human cancer.

Although our work focused on miR-200 transcriptional regulation by p73 and p63, recent studies have identified different mechanisms by which the p53 family can modulate miR-200 expression. p63 can serve as a transcriptional regulator of Dicer to modulate miRNA maturation, including processing of the miR-200 family (64). Given our observation of p63 binding to the miR-200b/a/429 promoter, these studies suggest that p63 can act both directly and indirectly to alter miRNA expression, although the mechanism may be cellular context-dependent. Additionally, a recent report found that in mammary epithelial cells, p53 serves as a transcriptional activator of miR-200c, but not of the miR-200b/a/429
cluster (17). In ovarian carcinomas, TP53 is commonly mutated and thus unlikely to drive the overexpression of miR-200 that we detected in our samples; however, TP53 mutations can frequently yield dominant negative proteins that interfere with the actions of p63 and p73 through promoter competition and through the formation of inactive heterotetrameric complexes (65). Although we did not find a relationship between the TP53 mutational status of ovarian carcinomas and their expression of miRNAs with p53 family binding sites, we cannot exclude the role of p53 in the transcriptional regulation of the miR-200 family. In contrast, our results indicate that p63 and p73 are involved in overexpression of this miRNA family in ovarian carcinoma. Taken together with the report of p53-driven miR-200c expression in mammary epithelial cells, our study suggests that cellular context determines the relative importance of different p53 family members on the transcriptional activation of the miR-200 family.

Through our integrative analysis, we identified transcriptional mechanisms that govern miRNA expression in ovarian carcinoma and also found novel activators of miRNAs implicated in tumorigenesis. Our results indicate that p73 and potentially also p63 contribute to broad dysregulation of miRNA expression in ovarian carcinoma. Although we have directed our analysis here toward ovarian cancer, our approach can be broadly applied to other cancer types (e.g., TCGA data sets are expected to expand to include nearly 25 cancer types), or any biological state(s) in which mRNA and differential miRNA expression data are available (e.g., expression profiles across stages of development or across different cell differentiation states). Because different cancer and tissue types show unique mRNA and miRNA expression patterns, utilizing our computational pipeline with other systems may reveal distinct TFs that regulate miRNA activity in varied biological contexts.

SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online.

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E.C.K., K.G. and J.D.A. wrote the manuscript. K.G. designed and implemented the computational pipeline and TCGA data analyses, D.S. performed the Affymetrix mRNA microarray analysis and R.G. and W.L.R. supervised parts of the computational analyses. E.C.K., J.D.A., Y.C. and R.K.P. performed the experimental validation. K.W. and E.M.S. sequenced and E.M.S. edited the manuscript. C.W.D., A.K.G., K.C.O. and N.D.U. helped procure and/or process specimens, and C.W.D. helped design the study. M.T. designed and supervised the study and helped to write the manuscript. All authors read and approved the final manuscript. The content of this publication is solely the responsibility of the authors and does not necessarily reflect the views or policies of the NIH or Department of Health and Human Services or other funders, nor does mention of trade names, commercial products or organizations imply endorsement by the U.S. Government.

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