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High expression of CAI2, a 9p21-embedded long non-coding RNA, contributes to advanced stage neuroblastoma

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Abstract

Neuroblastoma is a pediatric cancer with significant genomic and biological heterogeneity. p16 and ARF, two important tumor suppressor genes on chromosome 9p21, are inactivated commonly in most cancers but paradoxically overexpressed in neuroblastoma. Here we report that exon γ in p16 is also part of an undescribed long non-coding RNA (lncRNA) that we have termed CAI2 (CDKN2A/ARF Intron 2 lncRNA). CAI2 is a single exon gene with a poly A signal located in but independent of the p16/ARF exon 3. CAI2 is expressed at very low levels in normal tissue but is highly expressed in most tumor cell lines with an intact 9p21 locus. Concordant expression of CAI2 with p16 and ARF in normal tissue along with the ability of CAI2 to induce p16 expression suggested that CAI2 may regulate p16 and/or ARF. In neuroblastoma cells transformed by serial passage in vitro, leading to more rapid proliferation, CAI2, p16 and ARF expression all increased dramatically. A similar relationship was also observed in primary neuroblastomas where CAI2 expression was significantly higher in advanced stage neuroblastoma, independently of MYCN amplification. Consistent with its association with high risk disease, CAI2 expression was also significantly associated with poor clinical outcomes, although this effect was reduced when adjusted for MYCN amplification. Taken together, our findings suggested that CAI2 contributes to the paradoxical overexpression of p16 in neuroblastoma, where CAI2 may offer a useful biomarker of high-risk disease.

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Conflicts of Interest: The University of California San Diego has filed a patent on CAI2 in which MBD, ALY, LB and YK are named as co-inventors.

Sequence has been submitted to Genbank with a Submission ID #1642929, accession # KF311101.

Keywords

lncRNA; CAI2; neuroblastoma; 9p21; p16

Introduction

It is becoming increasingly apparent that long non-coding RNAs (lncRNAs), which until recently were thought to have little functional significance, harbor diverse roles in regulating cell identity, differentiation, and development. They have been implicated in cellular and molecular regulation, with chromatin remodeling and gene regulation emerging as common functions (1). In addition to normal cellular regulatory roles, lncRNAs are often deregulated in a wide variety of diseases (1-4). For example, the lncRNA *CRNDE*, originally identified through its association with colorectal cancer, is elevated in multiple cancer types and is induced during neuronal differentiation (5). One of the most well characterized lncRNAs, the HOX gene antisense lncRNA HOTAIR, has been shown to serve as a tumor biomarker in various types of cancers including breast, esophageal and pancreatic, and has also been correlated with tumor progression and prognosis (6-9).

Initiating from discrete first exons separated by ~20Kb on 9p21, the transcripts of *p16* (*p16^{Ink4a}*) and *ARF* (*p14^{ARF}*) are spliced onto a common exon 2 in separate reading frames and code for two different growth regulatory proteins. In addition, a homologous third tumor suppressor gene, *p15* (*p15^{Ink4b}*), lies on 9p21 ~10Kb telomeric of *ARF*. Given the central role these proteins play in cell cycle regulation, it is not surprising to find that the inactivation of all three have been associated with the pathogenesis of many human cancers (10, 11). Additionally, genome-wide association studies (GWAS) studies also implicate this locus with coronary disease, intracranial aneurysm, and diabetes (for review see (12)), demonstrating that this chromosomal region is a major player in a wide range of diseases. 9p21 is also the home of two lncRNAs, *ANRIL* and *p15AS* (*CDKN2BAS*) (13, 14). Functionally, both act as regulators of growth. *ANRIL* has been shown to regulate its neighbor tumor suppressors *p16* and *ARF* and thereby regulate cell proliferation and senescence (15-17). Expression of exogenous *p15AS* caused *p15* silencing and increased growth through heterochromatin formation and DNA methylation (14).

Neuroblastoma is the third most common malignancy of childhood. Diagnosed in stages based on clinical and biological features, patients with a favorable (low-risk disease) diagnosis can often be cured. However, approximately 50% of patients have high-risk disease and have dismal outcomes despite multi-agent and multi-modality therapy, myeloablative therapy followed by autologous bone marrow transplantation, with more than half of the patients nevertheless relapsing and succumbing to the tumor. Recently, we have reported on the significant improvement in survival targeting the GD2 tumor associated antigen uniformly expressed by neuroblastoma (18). Nevertheless, approximately one third of the patients continue to fail, highlighting the need to identify biomarkers to better predict outcome and improve efficacy.

While investigating the role of p16 in neuroblastoma, we observed that, in contrast to most tumors, p16 was not inactivated but paradoxically was overexpressed (19). Notably, p16

overexpression was significantly associated with advanced stage disease and a poor outcome (20). Upon continued investigation in neuroblastoma cell lines, we identified a unique transcriptional variant, p16 γ , which is formed by the splicing of the cryptic exon gamma, located in p16 intron 2, onto p16 exons 2 and 3 (21). It was our investigations into p16 γ that led to the discovery we report here that exon γ is also part of a lncRNA we call *CAI2*, a single-exon, intronlocalized, polyadenylated gene. *CAI2* is expressed at high levels in many tumor cell lines and its expression is highly correlated with the expression of *p16* and *ARF*. In neuroblastoma, *CAI2* expression was significantly associated with advanced stage disease and a poor outcome, suggesting that *CAI2* may be a novel biomarker of neuroblastoma patient risk.

Materials and Methods

DNA and RNA preparation, *RACE* and amplification

DNA was isolated using the Gentra DNA isolation kit and total RNA from cell lines extracted using Trizol reagent (Life Technologies). The 5' end of *CAI2* gene was obtained using both the GeneRacer kit (Life Technologies) and the SMART RACE cDNA amplification kit (Clontech) with DNase-treated total RNA from HeLa cells. RACE products were cloned into pCR4-TOPO vector (Life Technologies) and sequenced. For RT-PCR, DNase-treated RNA was reverse-transcribed using SuperScript III (SSIII) First-Strand Synthesis System with oligo-dT primers (Life Technologies). Quantitative RT-PCR (qRT-PCR) amplifications were performed on an ABI 7300 Real-Time PCR System using Quantitech SYBR Green Master Mix (Qiagen #204143) or Sybr Select Master Mix (Life Technologies #4472908) and compared to β -actin, or using Life Technologies Taqman assays (p16 – Hs02902543_mH, ARF – Hs99999189_m1, TH – Hs00165941_m1) and compared to GAPDH (Hs99999905_m1). Amplifications were performed in a final volume of 25 μ L with an initial incubation at 50°C for 2 min followed by one cycle of denaturation (95°C for 10 min). Sybr green amplifications utilized 40 amplification cycles of 30 sec each at 95°C, 60°C and 72°C followed by a dissociation step, while Taqman utilized 45 cycles of 95°C for 15 sec and 60°C for 1 min. Each sample was amplified in duplicate and each experiment performed at least twice. Expression was determined from the qRT-PCR data and analyzed using the 2^{-C_t} algorithm.

Primers

BS-ACT, 5'-GAAACTACCTTCAACTCCATC-3';
 BR-ACT, 5'-GTAGAAGCATTTGCGGTGGACGATGGAGGGGCC-3';
 p16S, 5'-CTGGAATTCAGCATGGAGCCGGCGGGGAG-3';
 c359R, 5'-GCGCGCAGGTACCGTGC-3';
 c257F, 5'-TCGTGCTGATGCTACTGAGG-3';
 I2796R, 5'-GAAGGCGTGAAGATGTGG-3';
 Gam517F, 5'-GTGGGTTTGTAGAAGCAGGC-3';
 ARF-SYR, 5'-CCATCATCATGACCTGGTCTTCT-3';

I1698F-Bam, 5'-CAATTAGAGCGCGGATCCAGGG-3';

3p16, 5'-CTACGAAAGCGGGGTGGGTTGT-3';

1888F-Bam, 5'-ATCGGGATCCAACCCCCTCTTCAAGCACAAT-3'

2252R-Xho, 5'-ATCGCTCGAGAAGGGCAAGATAGCTTGGGACT-3'

TH-666F, 5'-CTGACCTGGACTTGGACCAC-3'

Th-763R, 5'-GTCGCCGTGCCTGTACTG-3'

Primers (at 10 μ M unless indicated) used for qRT-PCR: *p16*: p16S-Eco & c359R (13.5 μ M each); *CAI2*: GAM517F (13.5 μ M) & I2796R (4.5 μ M); *ARF*, c257F & c359 or ARF-SYR; *β -actin*: BS-ACT and BR-ACT; TH: TH-666F and TH-763R.

Expression Constructs

Full-length *p16* (pcp16) (21) and *CAI2* (pcCAI2) expression constructs were constructed in pcDNA3.1 (Invitrogen); *CAI2* (psCAI2) was also created in pSilencer4.1-CMV vector (Ambion). *CAI2* was cloned from an amplicon generated with primers I1698 and 3p16. All constructs were sequence and expression validated, with *p16* further validated by Western blot.

For reticulocyte assay, we used full length pcp16 or *CAI2* and ORF2 (amplicon from primer set 1888F and 2252R) cloned into pCR2.1-TOPO (Invitrogen) in both the sense (*CAI2* and ORF2) and antisense (*CAI2*) directions. All constructs were sequence validated and protein production assessed with the Promega TNT Coupled Reticulocyte Lysate System using 35 S-Methionine and 35 S-Cysteine.

Tissue, Cell Lines, Cell Culture, Cell Growth and Transfections

Normal human tissue RNA was from Clontech. FS15 are low passage cultures of foreskin fibroblasts generously provided by Dr. Bruce Barshop (UCSD). The source of the cell lines and culture conditions has been documented previously (19, 22). Cells were transfected at 70-90% confluence using Lipofectamine 2000 reagent (Invitrogen) or with the NEON electroporation apparatus (Invitrogen). Post-transfection incubations proceeded for 24h to 72h, after which cells were observed, counted, assayed and/or harvested as appropriate. Primary sample and cell line RNA was isolated using Trizol (Invitrogen). All RNA samples were subjected to DNase treatment and repurification using the Zymo Research RNA Clean & Concentrator kit.

Primary Tumor Samples

A total of 62 samples of primary neuroblastoma tumors with a tumor cell content of 80% were obtained from the Children's Oncology Group. The patients had an average age of diagnosis of 1.6 years (range 0 -12.2 years), with an average follow-up of those who have had no event (relapse or death) of 3.8 years (range 0.8 -9.5 years). Samples distributed as follows: 10 stage 4S, 11 stage 1, 12 stage 2, 9 stage 3 and 20 stage 4. The clinical stages were classified as favorable stage/low risk (stages 1, 2, and 4S), unfavorable stage/moderate

risk (stage 3) or unfavorable stage/high risk (stage 4) based on COG staging criteria. Tumor samples were collected after informed consent and used under an UCSD approved IRB.

NMB7 cells

For the serial passage experiments, a fresh thaw of low passage NMB7 was plated in complete media and, at the passage indicated, cells were released from the plate with trypsin, and a portion of the cells lysed in Trizol for RNA extraction while a separate portion was reseeded for continued growth. The serial passage experiments were done twice, while RNA extractions from low and high passage cultures were done multiple times. For treatment comparisons, NMB7 cells at passage 5 or passage 16 were plated in 6 well plates at 125,000 cells in 10% FBS supplemented RPMI with antibiotics. After an attachment period, the media was removed and replaced with fresh media either without serum, with serum plus 0.5% DMSO or with serum plus 5 μ M RA (Sigma R2625 with 0.5% DMSO final concentration) and cells allowed to grow for an additional 24h or 72h, after which they were evaluated and, in some experiments, lysed in Trizol. Cells were photographed in independent experiments either directly or after H&E staining at 40X under phase-contrast. For growth experiments, 50,000 low or high passage cells were plated, allowed 24h to adhere, then observed and counted at 24h intervals up to 96h. Doubling time (DT) was calculated as $DT = h * \ln(2) / \ln(c2/c1)$, where h is time in hours and C1 and C2 are the beginning and ending number of cells present.

Statistical analysis

A Spearman's Rank Correlation (SRC) was used to analyze assess correlations between in expression data and a continuous outcome, a Kruskal-Wallis test used to test the association between CAI2 expression and a categorical variable such as stage, and a Wilcoxon Rank sum test used to test the association between CAI2 expression and a binary variable such as MYCN status. A logrank test was used to assess the association a survival outcome and a binary predictor, and a Cox model was used to assess these associations adjusted for other covariates such as age at diagnosis. Anova was used to compare expression across multiple parameter or time points, with individual analyses compared by t-test. P values less than 0.05 were considered statistically significant.

Results

I. Discovery of CAI2 (CDKN2A /ARF/ Intron 2 non-coding RNA) by RACE

While exploring *p16* γ expression, we observed that the gamma exon was present in many more samples than in which *p16* γ could be detected. Speculating that the gamma exon may be part of another alternatively spliced transcript of *p16*, we employed RACE (Rapid Amplification of cDNA Ends) to explore potential novel 5' ends using DNase-treated total RNA from HeLa cells. The 5' RACE amplification product using the Invitrogen GeneRacer kit resulted in a sharp band and a uniform smear of increasing molecular weight (Fig 1A). Sequence analysis of the extracted, purified and cloned RACE products revealed the sharp RACE band encompassed not only exon γ , but extended upstream to 1256 bases from the 3' end of *p16* exon 2 (Fig 1B). Sequence analysis of the smear yielded identical sequence except contained a concatemer of the RNA oligonucleotide adapter on the 5' end. A second

RACE analysis using the DNA-oligo adaptor method (Clontech) also yielded a single amplicon, which subcloning and sequence characterization revealed a 5' end that was within 30 bases of the transcript characterized using the GeneRacer kit and a 3' end that terminated 226 bases after the p16 stop codon (Fig 1B). Amplifications with sense primers to exon 2, exon 1α or exon 1β yielded amplicons with p16 3'-UTR primers but not with *RACE*-region antisense primers. On the other hand, *RACE*-region sense primers easily amplified with the 3'-UTR primers (data not shown). Furthermore, transcript amplification could only be demonstrated from RNA subjected to the RT reaction (Fig 1E). Taken together, this data demonstrates that this intronless transcript is not DNA contamination and is independent of p16 and ARF.

II. CAI2 is a non-conserved non-coding RNA

A bioinformatic analysis of this transcript revealed 5 open reading frames (ORF) with a length of 40 amino acids or greater (Fig 1C). The largest, ORF2, predicts a putative protein of 107 residues that a cross-species comparative analysis revealed only weak homologies (57%) that were limited to upper primates, consistent with the presence of an Alu element in this region. A comprehensive comparative analysis of the entire *CAI2* nucleotide sequence also revealed no significant homologies with the exception of the expected p16 exon 3 and the Alu region of ORF2. We nevertheless assessed possible translation using the TNT Reticulocyte Lysate System (Promega Corp). However, no evidence of a protein was observed from the transcript oriented in either direction nor ORF2 in the sense direction (Fig 1D). These experiments revealed the gamma exon is part of an uninterrupted 1643 base non-coding transcript we call *CAI2* (*CDKN2A/ARF* Intron 2 lncRNA) that is distinct from *p16*, *ARF* and *p16γ* (Fig 1B). The complete sequence is presented in Supplemental Table S1.

III. Expression of CAI2 in relation to the p16 and ARF

We compared the expression profile of *CAI2* in 46 tumor cell lines and 5 non-tumor samples, with representative amplifications shown in Fig 1F and summarized in Supplemental Table S2. The expression profiles of *p16*, *ARF* and *CAI2* were similar in most cell lines, where 20 of 26 cell lines with an intact 9p21 locus expressed all three genes robustly. Of the remaining six, all expressed *ARF* high and *p16* low while expression of *CAI2* was mixed. In contrast to tumor cell lines, none of the five non-tumor samples appreciably expressed *CAI2*. Considering whether this was a general characteristic of non-neoplastic cells, we undertook a quantitative assessment of *CAI2* expression in 25 normal samples. When each gene was normalized and graphed individually against expression in fibroblasts, most tissues exhibited similar levels of expression (>0.5 to <5.0 fold), with expression much lower (<0.5 of control) in just six and very high in two (spinal cord and adrenal). In contrast, p16 and ARF expression were much lower than fibroblasts in most tissues (Supplemental Figure S1). However, normalizing each gene to itself hides fact that *CAI2* expression is much lower than that of p16 and ARF in many tissues, including fibroblasts. As illustrated in Figure 2A, when all three genes were normalized to *CAI2* in fibroblasts, the low expression of *CAI2* in comparison to p16 and ARF was almost ubiquitous with the exception of spinal cord, which expressed *CAI2* ~200-fold higher than normal fibroblasts, and adrenal gland at ~16-fold higher. In contrast, other tissues tested expressed *CAI2* at levels similar to or much lower than fibroblasts, with the lowest levels

(~0.1-fold) seen in fetal tissue, kidney and heart. *P16* and *ARF* are independent genes with independent promoters that share correlated expression profiles. Consistent with this, a Spearman Rank Correlation (SRC) analysis of *ARF* and *p16* expression showed a significant association (SRC = 0.68, $p < 0.001$), with an even stronger correlation of *CAI2* with both genes (*p16* SRC = 0.73, $p < 0.0001$; *ARF* SRC = 0.89, $p < 0.0001$) (Figure 2B). We hypothesized that *CAI2* could be a regulator of p16 and/or ARF. No neuroblastoma cell line was low for all three genes, though NMB7, IMR32 and PCL1643 were low for both *CAI2* and p16 (Supplemental Table S2). Using the NMB7 cell line, which unlike the other two cell lines grow rapidly, as dispersed cell populations and are easily transfected (Fig 2C), we overexpressed *CAI2* and observed a small induction of both *p16* and *ARF* expression (2-7 fold), consistent with this hypothesis, though the lack of an influence on cell growth (Fig 2D) suggests that the levels of induction may be too small to be functionally significant.

IV. *CAI2* and neuroblastoma cell line transformation

The ability to be induced to differentiate is a characteristic of many neuroblastoma cell lines. We observed that NMB7 cells, with serial passage and without additional treatment, morphologically change from cells with a rounded, trapezoidal morphology (Fig 3A) to a thinner, more elongated phenotype with long processes (Fig 3B). Tyrosine Hydroxylase (TH) expression is known to increase as neuroblastoma cells differentiate (23). However, with increasing passage number, *TH* expression *decreased* by more than 100-fold (Fig 3C, $p < 0.001$, Anova), suggesting that the morphological change did not represent a normal differentiated phenotype. The NMB7 cell line at low passage number is also one of the few neuroblastoma cell lines that does not appreciably express *CAI2* (Supplemental Table S2). However, with serial passage, *CAI2*, *p16* and *ARF* expression each exhibited greater than 100-fold increases in expression (Fig 3C, $p < 0.001$ for *CAI2* and *ARF*, $p < 0.01$ for *p16*, Anova). And counter to the growth suppressor effects expected of p16 and ARF, high passage cells proliferated more rapidly than low passage cells, with a doubling time of 22h vs. 29h (Supplemental Figure S2).

Retinoic acid (RA) can induce differentiation and death in neuroblastoma cell lines. Treatment of low passage NMB7 cells with 5 μ M RA resulted in a visible loss of viability within 24h, with more than 95% of the cells having detached by 72h, while high passage NMB7 cells showed no effect from the RA treatment (Supplemental Table S3). The 24h RA treatment had no significant effect on TH, ARF or p16 gene expression in the high or low passage cell lines, though *CAI2* expression decreased in the low passage cell line (0.39-fold, $p = 0.02$, t-test, Fig 3D). Serum deprivation can also induce neuroblastoma cell line differentiation. Serum starved low passage NMB7 showed visible morphological signs of stress with cells looking “stringy” by 24h, rounded by 48h (Supplemental Figure S3) and detached by 72h; *TH* expression increased moderately within 24h of serum starvation (8.8-fold, $p = 0.05$, t-test, Fig 3D), an indication differentiation was beginning. On the other hand, there were no significant changes in the expression of *CAI2*, *p16* or *ARF* ($p > 0.05$, Anova). In contrast high passage NMB7 cells showed no negative effects from serum starvation, with no visible morphological changes or changes in the (already elevated) expressions of *CAI2*, p16 or ARF (Fig 3D and Supplemental Figure S3).

V. Enhanced mRNA levels of CAI2 in advanced stage neuroblastoma

As most neuroblastoma cell lines are derived from advanced stage tumors with high risk disease (24), and many of the neuroblastoma cell lines highly expressed *CAI2* (Supplemental Table S2), we considered whether this represented a unique feature of cell lines or reflected a characteristic of advanced stage tumor. An examination using RNA from primary neuroblastoma tumors revealed that the 33 prognostically favorable tumors (stages 4s, 1 and 2) expressed low levels of *CAI2* that were similar to that found in most normal tissues, though an occasional very high expressor was noted (Figure 4A). In comparison, the 29 high risk/stage 4 neuroblastoma expressed significantly higher levels of *CAI2* than favorable outcome group (Figure 4A, $p=0.005$, Kruskal-Wallis test comparing stage 4s, 1 and 2 vs. stage 3 vs. stage 4). Interestingly, a significant trend could be established when comparing lowest stage to highest stage neuroblastoma ($\text{SRC} = 0.33$, $p=0.01$), though expression was not significantly associated with *MYCN* amplification status (Wilcoxon Rank sum test, $p=0.18$, $n=62$ patients at all stages; $p=0.88$, $n=29$ stage 3 and 4 patients only). *CAI2* expression was also significantly associated with age at diagnosis ($\text{SRC}=0.39$, $p<0.002$, Figure 4B). Age over the age of 1 year is a negative prognostic factor in neuroblastoma, and most of the patients with high levels of *CAI2* delineated to the high risk category. Thus we were concerned that an association with age could be an artifact of prognostic features. However, when we limited the analysis to only high risk stage 3 and 4 patients, *CAI2* expression remained significantly associated with age of diagnosis ($\text{SRC}=0.38$, $p<0.05$, Figure 4C).

We next considered survival outcomes, with the three patients who had no follow up data censored at day 0, and dichotomized *CAI2* expression at 35.9, which was the median of *CAI2* values from the stage 3 and 4 patients, resulting in a “high *CAI2* expression” population that was comprised of 4 out of 33 stages 1, 2, and 4s patients, 2 out of 9 stage 3 patients, and 13 out of 20 stage 4 patients (Figure 4A). Among all patients, higher *CAI2* expression was significantly associated with shorter event-free survival (hazard ratio (HR) for higher vs. lower *CAI2* expression = 2.88, 95% confidence interval (CI) = (1.12, 7.45), logrank test $p=0.022$, Figure 5A) and showed a significant association with overall survival (HR = 3.27, 95% CI = (1.1, 9.77), logrank test $p=0.025$, Figure 5B). However, when we limited our analysis to only stage 3 and 4 patients, the association of *CAI2* expression with either EFS or OS was no longer apparent (HR=1.84 and $p=0.26$ for EFS, Figure 5C, and HR=1.37 and $p=0.58$ for OS, Figure 5D). Thus *CAI2* expression, in this limited data set, was a prognostic factor associated with a poor outcome, but by itself did not predict survival.

MYCN amplification is a very strong prognostic factor in neuroblastoma. Therefore, we wanted to determine if *CAI2* expression added prognostic information over and above *MYCN* amplification status, using Cox proportional hazards regression models which contained both factors. As expected, *MYCN* amplification status was a strong negative prognostic factor for both EFS and OS, and remained so even when *CAI2* expression was added to the model. On the other hand, the association previously observed between higher *CAI2* expression and outcome was attenuated in the model containing both factors in the entire population for both EFS (adjusted HR for higher vs. lower *CAI2* expression = 2.19,

95% CI = (0.81, 5.94), $p=0.12$) and OS (adjusted HR = 2.39, 95% CI = (0.76, 7.52), $p=0.14$). Similarly, when we limited the analysis to the stage 3 and 4 population only in the model containing both factors, the association of *CAI2* expression with either EFS or OS remained unapparent (HR=1.67, 95% CI = (0.56, 4.97), $p=0.36$ for EFS, or HR=1.25, 95% CI = (0.40, 3.88), $p=0.70$ for OS).

Discussion

The hallmark of neuroblastoma is its clinical and biological heterogeneity, with treatment based on subgroups defined by age at diagnosis, extent of disease and tumor biology. Clinically, risk stratification has been very successful. However, it remains true that some patients with identical risk factors who receive the same treatment can have considerably different outcomes. For example, though immunotherapy with ch14.18 has resulted in significant decreases in relapse and a significant improvement in cure rate, almost one-third of the children continue to fail therapy and succumb to the disease (18). The challenge, therefore, is to refine risk stratification through the identification of biomarkers and novel biological targets.

The over- and under-expression of ncRNAs have been associated with neuroblastoma prognosis. We and others have shown that microRNA (miRNA) expression profiles can differentially identify high versus low risk neuroblastoma (25, 26). The lncRNA *lncRAN* is overexpressed and prognostically significant in neuroblastoma and bladder cancer (27, 28). In a bladder cancer cell line, overexpression of *lncRAN* enhanced the proliferation, migration, and invasion of the cells, while its suppression enhanced chemosensitivity (28). Expression of the ncRNA *NDM29* correlated with growth rate of neuroblastoma and other tumor cell lines in an inverse fashion, with low expression correlating with a faster growth while overexpression in the SKNB2 neuroblastoma cell line led to their differentiation and an increased chemosensitivity (29). The diverse family of genes known as transcribed ultra-conserved regions (T-UCRs) are another class lncRNAs of that have been implicated in neuroblastoma through their association with MYCN amplification, outcome of high risk/stage 4 patients and RA response (30-32). Many of the characteristics of *lncRAN*, *NDM29* and the T-UCRs are similar to what we find with *CAI2* in neuroblastoma. In NMB7, cellular “transformation” is accompanied by not only an increase in *CAI2* expression, but a loss of sensitivity to RA. High passage NMB7 cells also express significantly higher levels of *CAI2* and proliferate faster than low passage NMB7 cells. *CAI2* was also expressed highly in most tumor cell lines but low in normal cells, suggesting a possible correlation with growth rate.

9p21 is a chromosomal region implicated in many different cancers, with deletion/mutation/epigenetic inactivation of p16 and ARF almost universal events in cancer. However, neuroblastoma has been a glaring exception. In fact, it has been more than a decade since we first reported on the paradoxical overexpression of *p16* in advanced stage neuroblastoma and its association with poor outcome (20). As ncRNAs play regulatory roles, aberrant expression of these molecules may contribute to the paradoxical p16 expression. We considered mir-24, a regulator of p16 (33). However, we and others have not observed this microRNA to be differentially expressed in neuroblastoma (25, 26). ANRIL has also been shown to negatively regulate p16 expression (15, 16), though the expression status of

ANRIL in neuroblastoma remains unknown. With the discovery that CAI2 can modulate *p16* and *ARF* expression *in vitro*, there is now an alternative pathway of p16 regulation to consider. A role in neuroblastoma is also supported by our findings that CAI2 is overexpressed in advanced stage neuroblastoma and is an independent prognostic factor.

Despite strong associations, we did not see an effect of CAI2 on NMB7 cell growth, either negative or positive, after transient transfections, and the effect on p16 was only minimal. It is possible that the conditions we are evaluating the effect of CAI2 on growth and p16 expression are inappropriate. For example, if CAI2 optimally influences p16 expression under conditions of stress or DNA damage response, p16 levels may not appreciably change when CAI2 is overexpressed in normal, proliferating cell lines. This would be similar to the actions performed by ANRIL, which suppresses p16 and ARF expression in response to DNA damage (16). The observation that the highest levels of CAI2 are seen in spinal cord, a conduit of stress response, supports this idea. Alternatively, if CAI2 is a transcript “stabilizer” like the lncRNA BASE1-AS (34), then overexpression might again only exhibit a minimal effect. Several lncRNAs have been shown to transcriptionally regulate indirectly through epigenetic actions. For example, the recently reported lncRNA ecCEBPA is a regulator of DNA methylation (35) while HOTAIR and ANRIL bind and modulate the Polycomb-repressive complex (36), which is required for epigenetic silencing during development and cancer. However, we showed many years ago that p16 is not methylated in neuroblastoma (19, 22), perhaps precluding the observation of a significant influence of CAI2 on p16 expression.

The fact that CAI2 has only a marginal impact on its nearest neighbor genes while seeming to have a considerable impact on tumor cell growth and survival, as evidenced by its preferential expression in advanced stage neuroblastoma and association with outcome, may suggest that CAI2 acts in a more global manner and the regulation of p16 and/or ARF are not its primary functions. The NMB7 cells during “transformation” undergo a significant change in morphology, growth, and drug sensitivity consistent with advanced stages of neuroblastoma. The inappropriate overexpression of CAI2 may at least facilitate such a “reprogramming”. The possibility of a more global function would be consistent with the histone modification actions of HOTAIR and ANRIL (37). Studies have shown lncRNAs play roles in multiple biological processes, including dosage compensation, genomic imprinting, chromatin remodeling, alternative splicing and nuclear organization (38). A recent study has also shown actions on proliferation, apoptosis, metastasis and autophagy (39). Exploring possible binding partners of CAI2 will give insight into pathways associated with the gene and clues to its function, as will its overexpression and/or silencing in moderate p16/ARF/CAI2 expressing model systems.

In summary, it is clearer than ever that the 9p21 locus, which is undoubtedly important in cancer, represents a complicated locus. It harbors three tumor suppressor genes *p16*, *ARF*, *p15*, the alternative spliced *p16γ* (21) and *p10* (40), the contiguously transcribed *p12* (41), mutation generated hybrid proteins such as chimeric ARF (chARF) and p16-Alternate Carboxy Terminus (p16-ACT) (42), the metabolic gene MTAP and its variants (43) and the interferon (IFN) gene cluster. Thus alterations to this region have a profound impact on cell growth, metabolic activity and signaling. It also contains the lncRNAs *ANRIL*, *p15AS* and

now *CAI2*. As *CAI2* is embedded in intron 2 of the *p16/ARF* gene, alterations to this locus will undoubtedly also influence *CAI2* as well. *CAI2* was significantly associated with high risk neuroblastoma, a population known to be the least likely to respond to chemotherapy, and a population where the many of patients will relapse and die (18)). Thus *CAI2* could play an additional role as a biomarker of advanced stage neuroblastoma, but it is premature to implicate it in response to chemotherapy. Nevertheless, it is clear now that lncRNAs may act not only as biomarkers and/or be prognostically significant, but that their modulation may influence growth and response to therapy. Therefore, one must now also consider the role of *CAI2* when considering how alterations at this locus influence cell behavior. As such, the role *CAI2* plays in diseases where the inactivation p16 and ARF, such as coronary artery disease, are not consistent with the known cell cycle regulatory functions of the proteins, should be considered. Based on this data and the central role this locus plays in multiple human diseases, we believe that *CAI2* must be considered in any future analyses of the genes of this locus on tumorigenesis specifically, and 9p21-associated disease pathogenesis in general.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Abbreviations

lncRNA	long non-coding RNA
ncRNA	non-coding RNA
CAI2	CDKN2A/ARF Intron 2 non-coding RNA

MNC	Peripheral blood mononuclear cells
RACE	Raid Amplification of cDNA Ends
ORF	Open Reading Frame
AA	Amino Acids
chARF	chimeric ARF
p16-ACT	p16-Alternate Carboxy Terminus
RT	reverse transcriptase
qRT-PCR	quantitative RT-PCR
SRC	Spearman-Rank correlation
HR	hazard ratio
CI	confidence interval
EFS	event-free survival
OS	Overall survival

Fig 1. The discovery of *CAI2* as a tumor expressed ncRNA

(A) The 5' and 3' RACE smear (RS) and RACE band (RB) cut out of an agarose gel and cloned are boxed. A 100bp molecular weight ladder is included in the 3' RACE figure; (B) The fragment of 9p21 containing the genomic orientation of ARF, p16, p16γ and *CAI2* exons and transcripts are illustrated, with the RACE primers and results illustrated. ARF exon 1 (X1β), p16 exon 1 (X1α), p16/ARF exon 2 (X2) and 3 coding (X3) and exon γ (Xγ) are shown; (C) The ORFs greater than 40 AA are illustrated relative to the entire *CAI2* transcript. The many ORFs smaller than 40 AA are unlabeled and shaded in grey; (D) *CAI2*, ORF2 and p16 were assessed for protein production using the Promega TNT Coupled Reticulocyte Lysate Systems. The luciferase control yielded a sharp band of the correct molecular weight (lane 1). p16 yielded a faint band at the correct molecular weight of 16 kDa (lane 2), consistent with the paucity of methionine (N=5) and cysteine (N=1) residues. Neither ORF2 (lane 3) nor *CAI2* in either the sense (lane 4) or antisense (lane 5) orientations yielded any evidence of a protein; (E) *CAI2* was detected only in reverse-transcribed RNA. And though *CAI2* sequence could be amplified from genomic DNA, amplification was fully abolished by the same DNase treatment the RNA samples were subjected to; (F) Representative non-quantitative amplifications demonstrate robust expression of *CAI2* in most tumor cell lines (HL60, NB20 and HeLa), though some tumor cell lines (PCL1643) and most normal samples (MNC) express *CAI2* only weakly; Molt4 are deleted (summarized in Supplemental Table 2).

Fig 2. CAI2, p16 and ARF expression in normal tissue and after CAI2 transfection

(A) A quantitative PCR analysis of *p16*, *ARF* and *CAI2* in normal human tissues. SJ-SA-1 (strongly expresses *p16*, *ARF* and *CAI2*) and Be2C/ADR5 (deleted for all three genes) cell lines are included as controls. All expressions are compared to *CAI2* in fibroblasts whose expression is set as 1 for comparison, demonstrating *CAI2* expression is very low in most tissues compared to *p16* and *ARF*. (B) A Spearman-Rank Correlation (SRC) analysis of *CAI2*, *p16* and *ARF* expression in primary tissue demonstrates the association each gene has with the other; (C) NMB7 cells were transfected with psCAI2 and expression of *CAI2*, *p16* and *ARF* after 48h were compared to cells transfected with a vector containing a scrambled sequence; (D) The same cultures of transfected cells as in Figure 2C were counted prior to lysing and RNA extraction. The percentage of viable cells after transfection with the control scrambled vector (psSc2) or the *CAI2* expression vector (psCAI2) versus lipid only is shown.

Fig 3. NMB7 cells morphologically change and p16, ARF and CAI2 expression increase with serial passage

(A) NMB7 cells at pre-transformation (shown at passage 10) have a rounded, trapezoid-like morphology; (B) After prolonged serial passage (shown at passage 22), cells morphologically change into an elongated structure with long protrusions; (C) RNA was extracted from NMB7 during serial passage (P) and *CAI2*, *p16*, *ARF* and *TH* expression determined by qRT-PCR; (D) Low and high passage (pass) NMB7 cells were cultured for 24h in the presence of complete media (untreated), complete media + 5 μ M RA, or media without sera (0% FBS) for 24h before being visually assessed and expression determined. Parallel cultures were also continued for 72h, but loss of viability in the treated low passage cells precluded harvesting of RNA. See also Supplemental Figure S3 and Supplemental Table 3.

Fig 4. CAI2 expression is associated with advanced stage neuroblastoma and age at diagnosis (A) CAI2 expression is significantly associated with high risk/advanced stage neuroblastoma ($P=0.005$, Kruskal-Wallis test, stage 4S, 1 and 2 vs. stage 3 vs. stage 4). MYCN amplified samples are shown as hollow symbols and were not significantly associated with CAI2 expression. For illustration purposes, the single highest CAI2 expression data point in a stage 4 tumor sample, with a relative expression value of 771, is not shown. A Spearman-Rank Correlation (SRC) analysis of CAI2 expression vs. stage illustrates the significant trend toward higher CAI2 expression in advanced stage neuroblastoma vs. low stage (B) CAI2 expression is significantly associated with age at diagnosis (dx) for all patients ($p<0.002$); and (C) for patients diagnosed with advanced stage disease (stage 3 and 4) only ($p<0.05$).

Fig 5. CAI2 expression is an independent prognostic risk factor

CAI2 expression is plotted versus event-free survival (A) or overall survival (B) in all neuroblastoma patients where it is significantly associated with survival. It is also plotted versus event-free survival (C) and overall survival (D) in advanced stage 3 and 4 neuroblastoma patients only where despite its association with advanced stage disease, did not independently associate with outcome in this patient population. Single variable logrank analyses are shown.