Abstract. Background: High bone morphogenetic protein (BMP)-2 expression in lung carcinoma correlates with poor patient prognosis. The present study explored strategies to repress BMP signaling. Materials and Methods: The cytotoxicity of BMP2-knockdown, dorsomorphin derivatives, and microRNAs was tested in transformed and non-transformed lung cells. Microarray analyses of 1,145 microRNAs in A549 lung adenocarcinoma cells and two other transformed lung cell types relative to BEAS-2B bronchial epithelial cells were performed. Results: Reduced BMP2 synthesis inhibited A549 cell growth. The dorsomorphin derivative LDN-193189, but not DMH1 or DMH4, was strongly cytotoxic towards A549 cells, but not towards BEAS-2B cells. Microarray analysis revealed that 106 miRNAs were down-regulated and 69 miRNAs were up-regulated in the three transformed lines. Three down-regulated miRNAs, hsa-mir-34b, hsa-mir-34c-3p, and hsa-mir-486-3p, repressed a BMP2 reporter gene and were cytotoxic in A549 cells, but not towards BEAS-2B cells. Conclusion: The observed cytotoxicity suggests that reducing BMP signaling is a useful line of attack for therapy of lung cancer.

Lung cancer is the leading cause of cancer-related deaths in the United States and the 5-year survival rate is an abysmal 15% (1). The potent growth factor bone morphogenetic protein-2 (BMP2) is often dysregulated in non-small cell lung cancer (NSCLC). Better understanding of how such growth factors influence lung cancer is sorely needed to develop new therapies.

Normal mature lung cells do not produce BMP2, but several studies indicate that BMP2 plays a significant role in lung cancer. BMP2 RNA and protein levels are abnormally elevated in lung tumors (2-4) and in serum from patients with lung cancer (5). BMP2 activates pro-oncogenic pathways (e.g., phosphoinositide-3 kinase (PI3K)/mammalian target of rapamycin (mTOR); SMAD1, 5/ DNA-binding protein inhibitor (ID)-1) and promotes lung tumor growth in mice (6-8). The BMP2 antagonist, noggin, reduces mixed metastatic lung cancer lesions in bone (9) and inhibits the growth of transformed lung cells in monolayer and in soft agar (10). Finally, variations in BMP2 levels (11) and specific BMP2 gene polymorphisms (12) are associated with differences in patient survival and response to treatment. Therefore, blocking the pro-oncogenic signaling of BMP2 is a reasonable therapeutic strategy.

Analogs of the small organic molecule dorsomorphin have been shown to inhibit the type-I BMP receptors with different degrees of selectivity (13-15). The dorsomorphin-derived analogs DMH1 and LDN-193189 (arbitrary compound designations) target the BMP type-I receptors activin A receptor type II-like 1 (ACVRL1/ALK1; HGNC:175), activin A receptor, type I (ACVR1/ALK2; HGNC:171), bone morphogenetic protein receptor, type IA (BMPR1A/ALK3; HGNC:1076), and bone morphogenetic protein receptor, type IB (BMPR1B/ALK6; HGNC:1077). LDN-193189 also inhibits the type-I transforming growth factor, beta receptor 1, TGFBR1/ALK5; HGNC:11772), the type-2 transforming growth factor, beta receptor II (TGFBR2; HGNC:11773), and the vascular endothelial growth factor A receptor-2 (KDR/VEGFR2; HGNC:6307). DMH4 inhibits VEGFR2 (13). SB-431542 is a chemically distinct inhibitor of TGFBR1 (16).

We have previously shown that highly malignant A549 lung adenocarcinoma cells express high levels of BMP2 (10). In contrast, BMP2 is practically undetectable in immortalized but non-tumorigenic BEAS-2B bronchial epithelial cells (10). We postulated that inhibitors of BMP
signaling should differentially affect these two cell types based on their differential BMP levels. We tested this hypothesis by measuring the influence of DMH1 and LDN-193189 on cell growth and survival in culture. We also assessed the influence of microRNAs (miRNAs) that repress the expression of BMP2.

**Materials and Methods**

**Materials.** Dorsomorphin analogs DMH1, DMH4, LDN-193189, SB-14373 were provided by Charles Hong (Vanderbilt University, Nashville, TN, USA). All except SB-14373 were re-suspended in the vehicle dimethyl sulfoxide (DMSO). SB-14373 was re-suspended in phosphate buffered saline (PBS). Cell culture and molecular reagents were from Sigma–Aldrich (St. Louis, MO, USA) and Invitrogen (Carlsbad, CA, USA). Ambion Pre-miR miRNA Precursors for hsa-mir-34b (PM12727), hsa-mir-34c-3p (PM12342), and hsa-mir-486-3p (PM12986) and Negative Control #1 (AM17100) were from Applied Biosystems (Austin, TX, USA).

**Cell culture.** A549 and non-transformed BEAS-2B cells were cultured in Dulbecco’s Modified Eagle’s Medium (DMEM, D5796, Sigma–Aldrich) supplemented with 5% fetal bovine serum and 2 mM glutamine. Cells were grown in 7% CO₂ at 37°C. For experiments to test drugs, cells were plated in 96-well plates at a density of 2.0×10⁴ cells per well. Drugs were added the following day. Plates were incubated for three days before performing the assays described below.

**RNAi-mediated BMP2 knockdown.** BMP2 (sc-270025-V; pooled viral particles with three BMP2-specific constructs encoding shRNAs) and Control (sc-108080) shRNA lentiviral particles were transduced according to the manufacturer’s instructions (Santa Cruz Biotechnology, Inc., Santa Cruz, CA, USA).

**RNA isolation.** Total RNA was isolated using Trizol reagent (Invitrogen) according to the manufacturer’s instructions.

**Reverse transcription – polymerase chain reaction (RT-PCR).** Total RNA (1 μg) was reverse-transcribed using 100 units of SuperScript™ III RT (Invitrogen) and oligonucleotide (oligo) dT or random primers according to the manufacturer’s instructions and 10% of the resulting cDNA was used for PCR. For BMP2 PCR, primers spanned the second intron (exon 2 forward primer 5'-GAGTTGAGGCTGCTACAGCAGTGT-3', exon 3 reverse primer 5'-TGCCATGGTTAGTGAGTTTGCAG-3'). The predicted size of the amplified cDNA fragment was 868 bp, whereas a product generated from genomic BMP2 gene would be 8639 bp. PCR conditions were: 95°C, 2 min; followed by 25 or 30 cycles of 94°C, 1 min; 55°C, 1 min; 72°C, 1 min; ending with 72°C, 7 min. For peptidylprolyl isomerase A (CYP4F10, cyclophilin A; HGNC:9253), the forward primer was 5'-GGCCATGGTCAACCCCACGTGGT-3' and the reverse primer was 5'-CCACTGAGCACTGGAGAGAAAGATTGGG-3'. The predicted size of the amplified cDNA fragment is 143 bp. PCR conditions were: 97°C, 5 min; followed by 29 cycles of 94°C, 30 s; 57°C, 45 s; 72°C, 1 min; ending with 72°C, 2 min.

**Enzyme linked immunosorbent assay (ELISA) for BMP2.** Conditioned medium (CM) was pre-cleared by centrifugation at 4000 × g for 2 min at 4°C to remove dead cells. Two milliliters pre-cleared CM was concentrated to about 100 μl using a Centricron Centrifugal Filter Unit with Ultracel YM-10 membrane (Millipore, Bedford, MA, USA) following the manufacturer’s instructions. The Quantikine BMP2 Immunoassay ELISA kit (R&D system, Minneapolis, MN, USA) was used to quantify the BMP2 protein level in 50 μl concentrated CM following the manufacturer’s instruction.

**Reporter gene constructs.** RLuc-empty (pSGG_3UTR_empty) and RLuc-3’UTR (pSGG_3UTR_BMP2, S208437) were described elsewhere (17). The SMAD binding element reporter was provided by Dr. Steve Harris, University of Texas Health Science Center, San Antonio, TX, USA. (18).

**Reporter gene assays.** Cells were plated and transfected with plasmid using FuGene6 Transfection Reagent (Roche, Indianapolis, IN, USA). Cells were lysed with 1×Passive Lysis Buffer (Promega, Madison, WI, USA) and luciferase activities were measured using the Luciferase Assay System (Promega). The luciferase activity of a co-transfected, constitutively expressed Renilla luciferase reporter plasmid was used to control for transfection efficiency. Protein concentration was used to control for cell density.

**Methylene blue staining.** Cells were fixed with 100% methanol. Fixative was removed and cells were stained with 1% methylene blue in 0.01 M borate buffer, pH 8.5, for 30 min (19). The stain was thoroughly washed with 0.01 M borate buffer, pH 8.5, until all excess stain was removed. Cells were photographed on a Gene Genius (Syngene, Frederick, MD, USA).

**WST-1 cell proliferation assay.** The Cell Proliferation Reagent WST-1 (Roche, Mannheim, Germany) is a colorimetric assay to quantify proliferation and cell viability after exposure to cytotoxic compounds in 96-well plates. Absorbance was measured at 450 nm using a Synergy HT, (Biotek, Winooksi, VT, USA). WST-1 Reagent was added to culture medium without cells to control for background absorbance.

**Lactate dehydrogenase (LDH) cytotoxicity assay.** The Cytotoxicity Detection kit (LDH) (Roche) quantifies cell lysis by measuring LDH released from damaged cells into the media. Absorbance was measured at 490 nm using Synergy HT (Biotek). LDH Reagent was added to culture medium without cells to control for background absorbance.

**Statistical analyses.** A Student’s t-test (two-sample assuming equal variances) was used to evaluate differences. A nominal p-value of 0.05 was considered significant.

**Results**

**RNAi-mediated inhibition of BMP2 synthesis inhibits cell growth.** BMP2, but not other BMPs, is induced in NSCLC cells and cultured transformed lung cells (2-4, 10). Previous studies used the BMP antagonist noggin to test the influence of reduced BMP2 activity on transformation, tumorigenesis, and metastasis (9, 10). Because noggin binds many members of the BMP family (20), we used RNAi-mediated inhibition of BMP2 synthesis to specifically test if reduced BMP2 levels affect the behavior of transformed lung cells. We
transduced A549 adenocarcinoma cells with BMP2 and control shRNA lentiviral particles. The lentiviral particles effectively knocked-down BMP2 mRNA levels (Figure 1A). BMP2 protein secreted into media conditioned by the A549 cells was reduced by 90% (Figure 1B).

BMP2 activates the translocation of SMAD proteins to the nucleus where they activate the expression of BMP-responsive genes. To test if BMP2 signaling was reduced, we transfected control and BMP2 knock-down A549 cells with a luciferase reporter gene bearing 12 SMAD binding elements. Luciferase activity generated by this plasmid reports the relative level of nuclear SMAD activity (18). In three independent strains of BMP2 knock-down A549 cells, SMAD-dependent luciferase activity was about half that observed in control cells (Figure 1C). This confirms that reducing the synthesis of BMP2 reduces SMAD-dependent BMP2 signaling in these cells.

Finally, we compared the effect of BMP2 knock-down on proliferation rates. The growth of A549 cells that were transduced with the BMP2 shRNA lentiviral particles was reduced compared to control cells (Figure 1B). These results are consistent with BMP2 stimulating the rapid growth of transformed lung cells.

LDN-193189 reduced cell proliferation and is cytotoxic in A549 and BEAS-2B cells. LDN-193189 inhibits type I BMP receptors ALK1, -2, -3, and -6. LDN-193189 also inhibits the type-1 TGFβ receptor ALK5, and the type-2 TGFβ receptors, as well as VEGFR2. We hypothesized that LDN-193189 should reduce growth in A549 cells that express high levels of BMP2 but not in BEAS2B cells which do not express BMP2. After equal numbers of cells were plated in 96-well plates, we added increasing concentrations of drug dissolved in DMSO or an equal volume of DMSO as a vehicle control. To

Figure 1. Knocking-down bone morphogenetic protein (BMP)-2 expression and signaling inhibits A549 cell growth. A549 lung adenocarcinoma cells were transduced with control or BMP2 shRNA Lentiviral Particles. A: The average BMP2 RNA levels from semi-quantitative reverse transcription-polymerase chain reactions (RT-PCR)±standard error of the mean (SEM) (n=2 to 3). BMP2 RNA measurements were normalized to that of the constitutively-expressed cyclophilin (CYP4A) RNA. Primers specific to BMP2 and cyclophilin spanned introns within each gene to eliminate priming from genomic DNA. B: The average±SEM BMP2 protein level (n=2 to 3) from media conditioned by A549 cells transduced with control and BMP2 shRNA lentiviral particles. C: BMP2 activates the translocation of SMAD proteins to the nucleus where they activate the expression of BMP-responsive genes. Cells were transfected with a luciferase reporter gene bearing 12 SMAD binding elements (18). Luciferase activity represents the relative level of nuclear SMAD activity in three independent BMP2 knockdown lines (B1, B2, B3) relative to cells transduced with control lentiviral particles. Bar graphs show average±SEM luciferase activity (n=35). D: Inhibition of A549 cell growth. Transduced cells were plated at the same density and counted on the indicated days (n=3).
avoid any influence of a drug on plating efficiency, cells were allowed to attach prior to drug addition. Cell density was assessed after three days by staining with methylene blue. The metabolic rate of the cells in each well was determined using the Roche WST-1 cell proliferation assay. Finally, to determine if any altered cell numbers were due to decreased cell proliferation or increased cell death, cytotoxicity was assessed using the Roche LDH Cytotoxicity Detection assay. The number of cells was greatly reduced at the highest drug concentrations, as assessed by methylene blue staining (Figure 2).
2A) and the WST-1 assay (Figure 2B). Both assays indicated that LDN-193189 reduced the growth of A549 cells at lower concentrations than in BEAS-2B cells. However, LDN-193189 was particularly cytotoxic against A549 cells, as indicated by a significant increase in LDH release (Figure 2C). At the highest concentration of drug (16 μM) there was 20-fold more LDH released than in the DMSO control. In contrast, only 4-fold more LDH was released from the BEAS-2B cells. Thus, in contrast to BEAS 2B cells that do not express BMP2, LDN-193189 was selectively cytotoxic to the BMP2-expressing A549 cells.
DMH1 had little effect on cell proliferation and cytotoxicity in A549 and BEAS2B cells. DMH1 targets the BMP type-I receptors ACVRL1, ACVR1, BMPR1A, and BMPR1B more selectively (14, 21, 22). Therefore, we tested the effect of DMH1 treatment on A549 and BEAS-2B cells. In contrast to LDN-193189, DMH1 failed to significantly inhibit the growth of either cell type, as assessed by both methylene blue staining and the WST1 assay (Figure 3A, B). In addition, DMH1 was not toxic to either cell type, except at its highest concentration (16 μM, Figure 3C).

**TGFβ and VEGFR2 pathway inhibition do not account for the cytotoxic effects of LDN-193189.** Because LDN-193189 also inhibits TGFBR1, TGFβR2, and VEGFR2, we tested selective inhibitors of these signaling paths. SB-431542 inhibits TGFβ signaling, via the activin A receptor, type IB (ACVR1B; HGNC:172), TGFBR1 and the activin A receptor, type IC, (ACVR1C; HGNC:18123) but not BMP signaling (16). DMH-4 is a selective inhibitor of VEGFR2. If the cytotoxic effects of LDN-193189 are due solely to inhibition of either pathway, then one or both of these drugs would produce similar antiproliferative and cytotoxic effects. We tested this hypothesis by performing a side-by-side drug assay by treating A549 cells with increasing concentrations of DMH4, SB-431542 and LDN-193189 (Figure 4). Methylene blue staining and the WST-1 and LDH release assays demonstrated that LDN-193189 had the strongest influence on A549 cells. In contrast, SB-431542 and DMH4 had little effect on A549 cells, except at the highest concentrations used. One remaining known difference between these drugs and LDN-193189 is that LDN-193189 inhibits TGFβR2. A549 cells do not express significant levels of TGFβ. Furthermore, because TGFβ blocks A549 cell growth (9, 23, 24), inhibiting TGFβ signaling would promote, not repress, A549 cell growth.

**MicroRNAs that target BMP2 are cytotoxic.** MiRNAs are small, ~21-22-nucleotide, single-stranded non-coding RNAs that commonly down-regulate protein synthesis by messenger RNA de-stabilization and translational repression.
Figure 5. MicroRNAs that inhibit bone morphogenetic protein (BMP)-2 expression are cytotoxic. A: A549 cells were co-transfected with a constitutive 60S ribosomal protein L10 (RPL-10) promoter-driven firefly luciferase reporter gene containing the human BMP2 3'UTR (RLuc-3'UTR) or without the 3'UTR (RLuc-Empty) and lacZ control. Ninety picomoles of each Ambion Pre-miR™ miRNA Precursors alone or a pool of 30 pmol each of miR-34b, -34c, and -486-3p or the Ambion Negative Control miRNA (NC) were co-transfected as indicated. All data were normalized to that of cells transfected with the Negative Control. Relative luciferase activity was measured at 24 h (n=4-9). B: A549 cell proliferation in the presence of 320 pmol miRNA precursors was measured four days after transfection using the Roche WST-1 cell proliferation. Similar studies in non-transformed BEAS-2B cells that express high levels of these microRNAs (Table I) failed to reveal any antiproliferative effect. C: The amount of lactate dehydrogenase (LDH) released by dying A549 cells into the medium was measured four days after transfection using the Roche LDH Cytotoxicity Detection assays. The bar graph shows the relative effect of each miRNA Precursor on LDH release at 320 pmoles. The line graphs show the average LDH released after transfection with increasing amounts of miRNA precursor (n=2).
A549 and BEAStra cells are highly tumorigenic in the BEAS-2B cell line with mycoplasma (10). Despite distinct origins, mycoplasma robustly expressed in transformed BEAS-2B and A549 cells (10, 28). Bone Morphogenetic Protein (BMP) 2 is not detectable in non-transformed BEAS-2B cells, but is robustly expressed in transformed BEAS-2B and A549 cells (10, 28).

Table I. Relative abundances of miR-34b, -34c, -486 in non-transformed BEAS-2B cells, in two mycoplasma-transformed BEAS-2B lines (BEAStra1, BEAStra2), and in A549 cells by Illumina v2 MiRNA Expression Profiling Assay (www.illumina.com/documents/products/workflows/workflow_microrna_assay). Bone Morphogenetic Protein (BMP) 2 is not detectable in non-transformed BEAS-2B cells, but is robustly expressed in transformed BEAS-2B and A549 cells (10, 28).

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(25-27). Using computational methods, we predicted that over 40 microRNAs may target 178 sites within the BMP2 3′ untranslated region (UTR) (http://www.targetscan.org and http://pictar.bio.nyu.edu). To identify miRNAs that regulate BMP2 expression specifically in lung tumors, we used the Illumina v2 MiRNA Expression Profiling Assay (http://www.illumina.com/technology/miRNA_assay.ilmn; platform #GPL8179) to compare the abundance of miRNAs from BEAS-2B bronchial epithelial cells and three lines of neoplastic lung cells [A549, BEAStra1, and BEAStra2 (10, 28)]. A549 cells were derived from a human alveolar cell carcinoma (29). In contrast, BEAStra cells were transformed by infecting the immortalized, but non-tumorigenic, BEAS-2B cell line with mycoplasma (10). Despite distinct origins, A549 and BEAStra cells are highly tumorigenic in immunocompromised mice. By examining the relative abundances of 1,145 miRNAs, we discovered that 106 miRNAs were down-regulated and 69 miRNAs were up-regulated in all three transformed lines (GEO accession #GSE557487). The abundance of several miRNAs predicted to interact with the BMP2 transcript inversely correlated with BMP2 secretion. That is, several miRNAs were expressed at high levels in normal BEAS-2B cells which do not synthesize any BMP2 but were significantly repressed in all three transformed lung cell lines which do synthesize BMP2 (10, 28). Such miRNAs may repress BMP2 synthesis in normal lung cells.

We analyzed three of the BMP2-targeting miRNAs repressed in malignant lung cells (Table I), previously shown to have interesting attributes relevant to lung cancer. For example, miR-34b and -34c are induced by p53 and are expressed at abnormally low levels in lung tumors (30-32). The level of miR-486 in sputum and serum of patients with lung adenocarcinoma differed from that of healthy subjects and predicted patient survival (33, 34). We tested if these miRNAs repress BMP2 in A549 adenocarcinoma cells, by co-transfecting Ambion Pre-miRTM miRNA Precursor Molecules and a reporter gene driven by a constitutive ribosomal protein promoter and bearing the human BMP2 3′UTR (Figure 5A). These small, chemically-modified double-stranded RNA molecules mimic endogenous mature miRNAs. Mimics of miR-34b, -34c, and -486 activity significantly reduced the expression of the BMP2 reporter gene bearing the BMP2 3′UTR (RLuc-3′UTR), but not the promoter-only construct (RLuc-empty). This demonstrates that the computer-predicted target sites in the BMP2 3′UTR are functional. Ninety picomoles of each miRNA alone or a pool consisting of 30 pmol of each miRNA inhibited reporter gene expression to comparable levels.

Because BMP2 promotes A549 cell growth and survival (Figure 1), (4, 6-8, 10), we then tested if miRNAs that target the BMP2 mRNA alter A549 cell proliferation. A549 cells were transfected with Ambion mimics of miR-34b, -34c, and -486 or the Ambion Negative Control mimic. Four days after transfection, the WST-1 colorimetric assay and protein concentration were used to compare the number of proliferative cells in each situation. All three experimental miRNAs strongly reduced the number of A549 cells (Figure 5B). In contrast, growth of non-transformed BEAS-2B cells was not affected by these miRNAs (Figure 5B). Because the reduced number of cells may have resulted from decreased growth rate or increased cytotoxicity, we also assessed the amount of LDH released by dying cells into the media. The level of LDH inversely correlated with the decreased number of A549 cells following transfection with miR-34b, -34c, and -486 (compare Figure 4B and C).

Discussion

We demonstrated that an inhibitor of BMP signaling (LDN-193189) reduces the growth of A549 lung cancer cells, but not immortalized, non-transformed BEAS-2B cells. Thus, reducing BMP signaling is a valid therapeutic strategy for lung cancer. Secondly, we provide a global profile of miRNAs expressed in three transformed lung cancer lines relative to BEAS-2B cells. Finally, we demonstrate that restoring the level of three specific miRNAs whose expression is reduced in A549 cells (miR-34b, -34c, and -486) is cytotoxic. Understanding these inhibitory molecules may lead to strategies that pharmacologically re-awaken BMP2-repressive mechanisms in patients with lung cancer. Pursuing all therapeutic strategies is essential because existing lung cancer treatments are woefully inadequate.

An important aspect of drug development is the demonstration that normal cells are unaffected relative to tumor cells. We found that LDN-193189, an inhibitor of BMP signaling is more cytotoxic to A549 lung adenocarcinoma cells than to non-transformed BEAS-2B cells. While this work was in progress, LDN-193189 was reported to inhibit lung cancer cell growth and increase cell death, as assessed by different
methods (35). Our study complements the published study by testing cytotoxicity in non-transformed cells that do not express BMP2 (10). The induction of cell death, rather than inhibition of proliferation accounted for most of the cytotoxicity. In contrast to LDN-193189, more selective inhibitors of type I BMP, TGFβ, and VEGF receptor (VEGFR) exhibited little toxicity to either cell type. The limited cytotoxicity of DMH1, SB-431542, and DMH4 suggests that activating the receptors specific to these compounds is not sufficient to mediate the toxicity of LDN-193189. The ability of LDN-193189 to inhibit several compounds is not sufficient to mediate the toxicity of LDN-193189. The ability of LDN-193189 to inhibit several pathways simultaneously may contribute to its effect on lung cancer cells. The precise mechanism by which LDN-193189 kills cells remains to be determined.

A sequence in the BMP2 3′UTR is 70% identical between mammals and fishes. We discovered that the function of this region in non-transformed lung cells differs from that in malignant cells (28). This sequence, which is targeted by the miRNAs tested here, is a potent repressor of gene expression in non-transformed BEAS-2B cells. In contrast, the sequence strongly stimulates BMP2 expression in A549 adenocarcinoma and mycoplasma-transformed BEAS-2B lung cells (28). Thus the ultra-conserved sequence functions as a post-transcriptional cis-regulatory switch. Three BMP2-targeting miRNAs are less abundant in strains of lung cells transformed by highly distinct means (Table I). The transformation-associated reduction in the abundance of miR-34b, -34c, and -486 miRNAs in transformed lung cells would impair the repression mediated by this switch and contribute to the elevated BMP2 levels observed in lung tumors (2-4, 10, 28).

The transcription of the bicistronic primary transcript bearing miR-34b and -34c is directly induced by the p53 tumor suppressor, and both miRNAs also inhibit proliferation in several cancer cell types (30-32). The list of experimentally-verified miR-34b and -34c targets is highly weighted to genes that regulate cell behavior (36). We now demonstrate that BMP2, a growth factor that strongly influences proliferation, apoptosis, and differentiation, is also a target of miR-34b and -34c. MicroRNA-34b and -34c are most highly expressed in lung tissues (30). Inactivation of the MIR-34b/c gene and the subsequently dysregulated expression of BMP2 and other influential genes in lung cells would profoundly influence the onset and progression of lung tumors.

Down-regulation of miR-486 also is associated with many types of human cancer, including human lung adenocarcinomas (37). A reduced level of miR-486 in sputum and serum is associated with lung cancer in diagnosis (33) and prognosis (34). Furthermore, the MIR-486 gene resides within a region frequently deleted in cancer tissue (Chr 8p11) (38). We have now demonstrated that miR-486 has a repressive interaction with the BMP2 3′UTR and is cytotoxic to lung adenocarcinoma cells (Figure 5). The abnormally low abundance of tumor suppressive miRNAs such as miR-34b, -34c, and -486 may contribute to neoplastic cell behavior. In miRNA replacement therapy, miRNA mimics are used to restore these miRNAs, and most importantly their tumor-suppressor function, to normal levels. Restoring the natural level of these miRNAs may repress BMP2 synthesis and ameliorate the neoplastic behavior of lung cancer cells.

Acknowledgements

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References


