MicroRNAs regulate methionine adenosyltransferase 1A expression in hepatocellular carcinoma

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MicroRNAs (miRNAs) and methionine adenosyltransferase 1A (MAT1A) are dysregulated in hepatocellular carcinoma (HCC), and reduced MAT1A expression correlates with worse HCC prognosis. Expression of miR-664, miR-485-3p, and miR-495, potential regulatory miRNAs of MAT1A, is increased in HCC. Knockdown of these miRNAs individually in Hep3B and HepG2 cells induced MAT1A expression, reduced growth, and increased apoptosis, while combined knockdown exerted additional effects on all parameters. Subcutaneous and intraparenchymal injection of Hep3B cells stably overexpressing each of this trio of miRNAs promoted tumorigenesis and metastasis in mice. Treatment with miRNA-664 (miR-664), miR-485-3p, and miR-495 siRNAs reduced tumor growth, invasion, and metastasis in an orthotopic liver cancer model. Blocking MAT1A induction significantly reduced the antitumorigenic effect of miR-495 siRNA, whereas maintaining MAT1A expression prevented miRNA-mediated enhancement of growth and metastasis. Knockdown of these miRNAs increased total and nuclear level of MAT1A protein, global CpG methylation, lin-28 homolog B (Caenorhabditis elegans) (LIN28B) promoter methylation, and reduced LIN28B expression. The opposite occurred with forced expression of these miRNAs. In conclusion, upregulation of miR-664, miR-485-3p, and miR-495 contributes to lower MAT1A expression in HCC, and enhanced tumorigenesis may provide potential targets for HCC therapy.

Introduction
Methionine adenosyltransferase (MAT) is an essential enzyme that is responsible for the biosynthesis of S-adenosylmethionine (SAMe), the principal biological methyl donor in mammalian cells (1). Mammals express 2 different genes, MAT1A and MAT2A, that encode for 2 homologous MAT catalytic subunits, α1 and α2, respectively (2). MAT1A is expressed mostly in adult liver and serves as a marker for normal differentiated liver, whereas MAT2A is expressed in all extrahepatic tissues and is induced during rapid liver growth and liver dedifferentiation (1). In liver, the α1 subunit forms dimer (MATIII) and tetramer (MATII) MAT isoenzymes (2). MAT1A expression is reduced in about 60% of patients with cirrhosis due to various etiologies (3) and is often silenced in human hepatocellular carcinoma (HCC) (3, 4). In addition to reduced MAT1A expression, the activity of the MATI/III isoenzymes is often reduced in patients with chronic liver disease, resulting in chronic depletion of hepatic SAMe level (5). The Mat1a-KO mouse model has illustrated the many consequences of chronic hepatic SAMe depletion, the most important of which is spontaneous development of HCC (6, 7).

The importance of hepatic SAMe depletion in HCC development is supported by several rodent models of HCC using hepatocarcinogens in which hepatic SAMe depletion develops and HCC was prevented by SAMe administration (8–10). The dominant mechanism of SAMe chemopreventive effect was thought to be from preventing hypomethylation of the promoter region of several protooncogenes, as SAMe’s chemopreventive effect was blocked by 5-azacytidine (9, 10). However, in an orthotopic liver cancer model, SAMe administration was also able to inhibit HCC establishment but was ineffective in blocking growth of already existing HCC (11). Part of this was because of a compensatory response of the liver to induce methyltransferases that removed excess SAMe to prevent its accumulation (11). A better strategy to maintain increased SAMe level in liver cancer cells is to induce the expression of MAT1A. Consistent with this, liver cancer cells with forced expression of MAT1A doubled-cellular SAMe levels grew slower in vitro and in vivo and exhibited reduced angiogenesis, ERK, and AKT activation and increased apoptosis in vivo (12). This proof of concept prompted the current work to investigate whether MAT1A expression is regulated by microRNAs (miRNAs) that are dysregulated in HCC, as miRNAs are potentially much better therapeutic targets. In the course of our investigation, we uncovered 3 miRNAs whose expression and function in HCC have not been reported to be upregulated in HCC and contribute to the downregulation of MAT1A, tumorigenicity, invasion, and metastasis.

Results
Expression of miRNAs in HCC and the effect of their knockdown on MAT1A expression in HepG2 and Hep3B hepatoma cell lines. To determine whether miRNAs can potentially regulate MAT1A expression, we used an in silico approach to search for miR-
NAs that can bind to the 3′ UTR of MAT1A. Using 3 different miRNA prediction target databases (TargetScan, mirDB, miRSVR), many miRNAs were identified (Supplemental Table 1; supplemental material available online with this article; doi:10.1172/JCI63861DS1). We focused on those with high scores (miRNA-664 [miR-664], miR-485-3p, miR-495, miR-588, miR-766) whose expression has not been reported in HCC. Figure 1A shows that miR-664, miR-485-3p, and miR-495 are induced in HCC, while miR-588 and miR-766 are downregulated. Since the dominant mechanism of miRNA regulation is that of downregulation of mRNA/protein levels (13), we focused only on the 3 that are upregulated. To see whether these miRNAs regulate MAT1A expression and to determine the level of regulation, HepG2 and Hep3B cells were treated with siRNA targeting these miRNAs. The efficiency of knockdown is shown in Figure 1B, which is comparable with 75% to 80% reduction after 24 hours. Figure 1, C and D, shows that knockdown of miR-664, miR-485-3p, and miR-495 individually raised MAT1A mRNA and protein levels comparably and combined knockdown exerted nearly an additive effect. This demonstrates that these miRNAs negatively regulate MAT1A expression and the effect is exerted at the mRNA level.

Figure 2A shows the MAT1A 3′ UTR sequence containing the putative binding sites for miR-664, miR-485-3p, and miR-495. Figure 2B and C, shows that WT MAT1A 3′ UTR reduced reporter activity by more than 50% and mutation of each miRNA binding site led to slight recovery, with incremental recovery as additional sites were mutated. When miR-664, miR-485-3p, and miR-495 were all mutated, the inhibitory effect of the MAT1A 3′ UTR was completely lost.

Figure 1
miR-664, miR-485-3p, and miR-495 are induced in HCC and negatively regulate MAT1A expression in liver cancer cell lines. (A) Northern blot analysis showing expression of select miRNAs in HCC compared with adjacent nontumorous (NT) tissue. (B) Northern blot analysis confirming siRNA knockdown efficiency of miR-664, miR-485-3p, and miR-495 in HepG2 and Hep3B cells as compared with scramble siRNA (SC) control. (C and D) Northern (top) and Western (bottom) blot analyses showing the effect of siRNA knockdown of miR-664, miR-485-3p and miR-495, alone or in combination, on MAT1A expression in HepG2 (C) and Hep3B cells (D). Numbers below the blots represent densitometric values expressed as percentage of respective controls. Representative blots are shown for C and D from 3 experiments. *P < 0.01 vs. SC; †P < 0.05 vs. SC, and triple knockdown; ‡P < 0.001 vs. SC and single knockdown.
Effect of siRNA against miR-664, miR-485-3p, and miR-495 on apoptosis and cell proliferation. We previously reported that forced MAT1A expression in liver cancer cells reduced growth while inducing apoptosis (12). To see if inducing MAT1A expression by targeting miRNAs also has the same effect, we measured apoptosis and growth. Figure 3A shows that siRNA treatments (single, double, and triple knockdown) in Hep3B cells induced apoptosis only after 72 hours of transfection and that, similar to the effect on MAT1A expression, the effect on apoptosis was additive with combined knockdown. To avoid toxicity causing nonspecific effects, effect on growth was measured only after 24 hours of siRNA treatment. Figure 3B shows that single knockdown of these miRNAs reduced growth by about 20% and combined knockdown of all 3 reduced growth by about 50%. Similar findings on apoptosis and growth were observed in HepG2 cells (data not shown). To determine the contribution of MAT1A induction on the growth inhibitory effect, Hep3B cells were stably transfected with siRNAs targeting each miRNA. Control Hep3B cells were stably transfected with scramble siRNA. These cells were then transiently transfected with siRNA against MAT1A or scramble siRNA for 24 hours. Figure 3C shows that Hep3B cells with stable knockdown of each miRNA exhibited reduced cell proliferation, but this effect was significantly blunted when MAT1A was also knocked down (Figure 3C).

**Figure 2**
MAT1A 3’ UTR-driven reporter activity and the effect of mutating miRNA binding sites. (A) Diagram of MAT1A 3’ UTR fragment containing the putative binding sites for miR-664, miR-485-3p, and miR-495. Mutations created for each miRNA site are denoted in bold italics. Transient transfection assays were performed using a luciferase reporter system with WT and mutated MAT1A 3’ UTR constructs as described in Methods in (B) HepG2 and (C) Hep3B cells. *P < 0.05 vs. control; †P < 0.05 vs. MAT1A 3’ UTR; ‡P < 0.05 vs. triple miRNA siRNA knockdown. n = 3 experiments, done in triplicate.
doubled MAT1A levels. Injection of cells expressing high miRNA levels subcutaneously into the flanks of nude mice resulted in more rapid growth of tumors (Figure 4, A and C), whereas tumor growth was significantly inhibited in cells expressing siRNAs against these miRNAs (Figure 4, B and C). Manipulating the expression of miR-495 resulted in the most dramatic effect. At 8 weeks, tumors from mice injected with forced lenti–miR-664, lenti–miR-485-3p, and lenti–miR-495 showed tumor volumes that were 160%, 170%, and 360% higher as compared with empty vector (EV), respectively (Figure 4A). Tumors from mice injected with cells expressing stable knockdown of miR-664, miR-485-3p, and miR-495 siRNA or scramble siRNA (stable SC) were transiently transfected with MAT1A siRNA or SC, and BrdU incorporation and MAT1A protein levels were measured 24 hours later. *P < 0.05, **P < 0.01 vs. SC; †P < 0.05 vs. single or triple siRNA knockdown.

Effect of reducing the expression of miR-495, miR-485-3p, and miR-664 in an invasive liver cancer cell line in vivo. Since Hep3B cells did not metastasize to the lung, we switched to HepG2 cells, which can invade adjacent structures and metastasize to the lung in the same orthotopic liver cancer model (Figure 5B). Mice were treated with siRNA targeting miR-495, miR-485-3p, miR-664, or scramble control for up to 8 weeks. At week 8, there was a significant (35%
to 62%) reduction in tumor volume (Figure 5B) and reduced incidence of metastases to the lung, abdominal wall, pancreas, and diaphragm in mice treated with siRNAs (Figure 5B and Supplemental Table 2). Reducing miR-495 expression had the most significant inhibitory impact on tumor growth, invasion, and metastasis.

Role of MAT1A in HCC invasion and antitumorigenic effect of miR-495 siRNA. To examine the role of MAT1A directly in tumorigenesis and treatment efficacy of the most potent miRNA siRNA, miR-495 siRNA, mice were treated with siRNA against MAT1A, miR-495, and scramble control alone or together using the same invasive HepG2 cells orthotopic liver cancer model. Figure 6 shows that reducing MAT1A expression with MAT1A siRNA more than doubled the tumor volume after 8 weeks and significantly blunted the treatment efficacy of miR-495 siRNA on tumor growth. MAT1A expression also had a direct effect on invasion and metastasis, as MAT1A siRNA treatment increased...
A

- miR-495
- miR-485
- miR-664
- EV
- miR-495si
- miR-485-3psi
- miR-664si
- SC

Liver tumor:
- 1446 ± 100
- 903 ± 71
- 943 ± 68
- 677 ± 103
- 263 ± 34
- 451 ± 58
- 453 ± 25
- 728 ± 76 (mm²)

Lung metastasis:

PCNA:
- 38% ± 2.3%
- 20.6% ± 1.2%
- 19.7% ± 1.3%
- 13.8% ± 1.2%
- 6.1% ± 0.6%
- 7.1% ± 0.5%
- 6.5% ± 0.6%
- 14.5% ± 0.9%

MAI: cl

B

- SC
- miR-495si
- miR-485-3psi
- miR-664si

Liver tumor:
- 3.4 ± 0.5
- 1.3 ± 0.2
- 1.9 ± 0.3
- 2.2 ± 0.2

Tumor volume (cm³):

Lung metastasis:

Pancreas invasion:
- 2/8
- 0/8
- 0/8
- 0/8
Figure 5
Effect of varying miR-664, miR-485-3p, and miR-495 expression on tumorigenesis, invasion, and metastasis in an orthotopic liver cancer model. (A) Hep3B cells stably transfected with lentil-miR-664, miR-485, and miR-495/EV or lentiviral-siRNA against these miRNAs or SC were injected into the left hepatic lobe, and mice were sacrificed after 45 days. The top row shows H&E staining of liver tumors, and tumor volume at the site of injection are shown below for each condition (*< 0.05 vs. miR-485, miR-664 and EV; †< 0.05 vs. EV; ‡< 0.05 vs. miR-485-3psi, miR-664si, and SC; §< 0.05 vs. SC). Second row shows H&E staining of lung tissue and incidence of lung metastasis. Arrows point to tumor metastasis. Third and fourth rows show immunohistochemistry for PCNA and MAT1A protein. Numbers below PCNA represent percentage of positive cells, *< 0.01 vs. miR-485, miR-664 and EV; †< 0.05 vs. EV; ‡< 0.05 vs. miR-485-3psi, miR-664si and SC; §< 0.05 vs. SC. Original magnification, ×100 (first row); ×200 (second through fourth rows). (B) HepG2 cells capable of invasion and metastasis were injected into the left hepatic lobe as above, and lentiviral vectors containing miRNA siRNA or SC were injected into the spleen at the time of HepG2 cell injection. Two weeks later, lentiviral siRNA was injected into the tail vein, and this was repeated every 2 weeks until sacrifice at 8 weeks. H&E staining showing the effect of miR-495, miR-485-3p, and miR-664 siRNAs on tumor invasion and metastasis. Arrows point to tumor at the site of injection, and tumor volumes and invasion incidences are shown below each image. *< 0.01 vs. scrambled siRNA (SC); †< 0.05 vs. miR-485-3psi and miR-664si. Original magnification, ×100.
showed that the MAT1A 3′ UTR binds to the AU-rich RNA binding factor 1 (AUF1), which is one of the hnRNP proteins known to destabilize target mRNAs (23). HCC specimens express higher AUF1 protein levels, and knockdown of AUF1 increased MAT1A mRNA level (23). However, these mechanisms are difficult to target, since they can affect numerous other genes. This prompted us to consider the possibility that miRNAs might also regulate MAT1A expression.

miRNAs are small noncoding RNAs that regulate gene expression by targeting the 3′ UTR of mRNAs, leading to reduced protein translation and/or increased mRNA degradation in most cases (13). Dysregulation of miRNA expression plays an important role in the pathogenesis of HCC, and miRNA signatures may serve as biomarkers for HCC classification and prognostic risk stratification as well as therapy (24, 25). We used the miRWalk website (http://www.ma.uni-heidelberg.de/apps/zmf/mirwalk/index.html) to simultaneously search 8 different established miRNA prediction programs to see which miRNAs have the most positive prediction matches. Using the most popular search algorithms — miRSVR (uses miRANNA), TargetScan and mirDB — we generated a list of miRNAs that gave the best scores for MAT1A. Interestingly, while some of these known to have altered expression in HCC, the expression of most of these miRNAs in HCC is unknown. Of the 5 best-matched miRNAs whose function in HCC is unknown, 3 are induced in 4 sets of HCC samples as compared with adjacent nontumorous liver, and we focused on these for the current work, since the majority of miRNAs downregulate expression of their targets. Indeed, knockdown of miR-664, miR-485-3p, or miR-495 in both HepG2 and Hep3B cells raised MAT1A mRNA and protein levels comparably (Figure 1), which is consistent with the notion that the mechanism of these miRNAs on MAT1A is to increase its mRNA degradation. Reporter assay confirmed the presence of functional miRNA-binding sites in the MAT1A 3′ UTR (Figure 2).

Each miRNA exerted a significant influence on growth and apoptosis in vitro and tumor growth in vivo; however, although the miRNAs all have comparable effects on MAT1A expression, miR-495 exerted the most pronounced effect on tumor growth, invasion, and metastasis. This suggests miR-495 has other important targets besides MAT1A. Nevertheless, knocking down miR-664 or miR-485-3p reduced tumor growth by over 50% as well as invasion and metastasis, supporting an important role for MAT1A. Consistent with this, lowering MAT1A expression increased the tumorigenicity, invasion, and metastatic potential of HCC and blunted the therapeutic efficacy of miR-495 siRNA on these parameters.

Figure 6
Role of MAT1A in tumorigenesis and therapeutic effect of miR-495 siRNA. HepG2 cells were injected into the left hepatic lobe of male BALB/c nude mice and lentiviral vectors containing MAT1A siRNA (MAT1A si), miR-495 siRNA (miR-495 si), and scramble siRNA (SC), alone or together, were injected into the spleen at the time of HepG2 cell injection (n = 8 per group). Control group received only HepG2 cell injection. Two weeks later, lentiviral siRNAs were injected into the tail vein, and this was repeated every 2 weeks until sacrifice at 8 weeks. First row: arrows point to tumors at the site of injection, and tumor volumes are shown below. *P < 0.005 vs. SC+SC; †P < 0.005 vs. MAT1A si+SC; ‡P < 0.05 vs. miR-495 si+MAT1A si. Second and third rows show metastasis to lung and pancreas (indicated by arrows) in the various treatment groups, with the incidence shown below. Original magnification, ×200.
Furthermore, forced expression of MAT1A that cannot be inhibited by these 3 miRNAs completely eliminated the inductive effect of these miRNAs on growth and metastasis. Taken together, these results indicate that MAT1A is the key target for these miRNAs in exerting their influence on HCC tumorigenesis.

miR-485-3p has been shown to mediate topoisomerase IIA downregulation in part via altered regulation of the transcription factor nuclear factor YB and to have a role in drug responsiveness in CEM and CEM/V:1-5 cells (human leukemic lymphoblastic cells) and in Rh30 and Rh30/v1 cells (human rhabdomyosarcoma cells) (26). miR-495 has been shown to be upregulated in breast cancer stem cells, where its overexpression promoted tumorigenic activity by downregulating E-cadherin and REDD1 (regulated in development and DNA damage responses 1), an inhibitor of mTOR signaling (27). miR-495 is also thought to have a role in liver and pancreas development (28). Increase of miR-664 expression was found in atrial fibrillation patients with rheumatic heart disease (29). However, as of today, the expression and function of these miRNAs in HCC has not been reported. Recently, Koturbash et al. demonstrated that miR-29b might play a role in downregulating *Mat1a* in the preneoplastic liver tissue of rats treated with the hepatocarcinogen 2-acetylaminofluorene (30). However, miR-29a-c was found to be downregulated in HCC, and it has been shown to promote apoptosis by lowering the expression of Bcl-2 and Mcl-1 (31). Human MAT1A has not been shown to be regulated by miRNA. Our study clearly demonstrates an important role for miR-495, miR-485-3p, and miR-664 in regulating *MAT1A* expression and that also shows that downregulating their expression inhibited tumor growth, invasion, and metastasis of HCC. The next question is, how does increasing *MAT1A* expression impact on tumorigenesis, invasion and metastasis?

Since MAT is responsible for SAMe synthesis and MAT/III are the products of the *MAT1A* gene (1), increasing *MAT1A* expression would increase steady state SAMe levels (3). Changes in DNA methylation, particularly hypermethylation of tumor suppressors, play a critical role in cancer pathogenesis, including HCC (32). This prompted us to examine whether *MAT1A* expression in liver cancer cells influences DNA methylation. Indeed, when *MAT1A* expression was increased by knocking down miR-664, miR-485-3p, or miR-495, nuclear SAMe levels and global DNA methylation increased, and the opposite occurred when *MAT1A* expression was reduced by forcing the expression of these miRNAs. Of the many genes deregulated in HCC, we focused on *LIN28B* overexpression in HCC and promotes transformation and invasion in HCC in part via repression of let-7 (15). The let-7 family of miRNAs regulates factors that control cell-fate decisions, including oncoproteins and cell-cycle factors (33–35). LIN28B (and LIN28) exerts a reciprocal regulation with let-7 (33). let-7 suppresses the expression of *LIN28* through let-7–binding sites in the *LIN28* 3′ UTR, while LIN28/LIN28B suppress the production of mature let-7 at multiple levels as well as enhancing let-7 degradation via 3′ terminal uridylation of let-7 precursors (33). This opposing expression pattern of *Lin28* and let-7 can be found throughout development and in oncogenesis and has been compared with a yin-yang balancing act by Ji and Wang (33). This can be illustrated by activation of *LIN28* by c-Myc and NF-κB, leading to let-7 repression and cell transformation (33). The *LIN28B* promoter region has multiple CpG sites, including 3 CpG islands (as determined by CpG island searcher: cpgislands.usc.edu), where Viswanathan et al. correlated the loss of DNA methylation of a downstream CpG island with that of its expression found in HepG2 and K562 erythromyeloblastoid leukemia cells (36). Interestingly, we found that forced miRNAs reduced *MAT1A* expression and *LIN28B* promoter methylation and increased *LIN28B* expression. This correlated with a fall in let-7a expression. Increasing *MAT1A* expression by knocking down miR-664, miR-485-3p, and miR-495 led to *LIN28B* promoter hypermethylation, reduced *LIN28B* expression, and increased let-7a expression (Figure 8D). Thus, enhancing *MAT1A* expression shifted the balance of LIN28B/let-7 toward let-7 and inhibited tumor growth, invasion, and metastasis.

Recently Reytor et al. reported finding MATI/III in the nuclei (37). The authors speculated that presence of nuclear MAT might provide a continuous source of nuclear SAMe, since SAMe is charged and whether or not it can traverse the nuclear membrane is in debate. In support of this, nuclear accumulation of the active MAT1A protein correlated with higher levels of histone H3K27 trimethylation, an epigenetic modification associated with gene repression and DNA methylation (37). Our findings are consistent with this report. Interestingly, we found that knocking down miR-664, miR-485-3p, and miR-495 increased total cellular MAT1A protein level, particularly the nuclear fraction (Figure 8E). This may explain the dramatic effect that *MAT1A* expression has on DNA methylation.

In summary, we have identified 3 miRNAs that are increased in HCC that can negatively regulate *MAT1A* expression at the mRNA level. Reducing the expression of these miRNAs raised *MAT1A* expression, which we believe is a novel strategy to shift the LIN28B/let-7 balance toward let-7. We suspect this increase in *MAT1A* expression also has an impact on the expression of many other genes involved in tumorigenesis. This will be the subject of a future investigation. Our results also help to explain why decreased *MAT1A* expression in HCC is a poor prognostic indicator (21). Few therapeutic options currently exist to treat HCC. These miRNAs are potential therapeutic targets and offer substantial promise in expanding treatment options for patients with HCC.

Methods

**Materials and reagents.** Ad-Δ32P-dCTP and γ-32P ATP (3,000 Ci/mmol) were purchased from PerkinElmer. Antibodies used for either Western blot and/or immunohistochemistry to PCNA, LIN28B, and β-actin were purchased from Cell Signaling Technology. MATa1 antibody was purchased from Novus Biologicals, whereas H3K27me3 and GFP antibody were purchased from Abcam. Lipofectamine 2000 and RNAmax were purchased from Invitrogen, whereas the MethylFlash Methylated DNA quantification kit was purchased from Epigentek. Lentivirus-MAT1A, lenti–miR-664, lenti–miR-485 (lenti–miR-485-3p is not available), lenti–miR-664 siRNA, lenti–miR-485-3p siRNA, lenti–miR-495 siRNA, and a lentiviral purification kit were purchased from SBI System Biosciences. Lenti-MAT1A siRNA was purchased from Applied Biological Material Inc. siRNA to hsa–miR-664 (AGGGTGGGATAATTGGAAT), hsa–miR-485-3p (AGAGGAGGCGGTG-TATGAC), and hsa–miR-495 (5′-AGAAGTGACCATGTGTTTGT-3′) were purchased from Exiqon. pMir-Target vector for MAT1A 3′ UTR clone was purchased from OriGene Technologies. GFP expression after injection of lenti-miRNAs or lenti-siRNAs was visualized on paraffin sections by immunohistochemistry with mouse anti-GFP antibody (1:200; Clontech, BD Biosciences) using the ABC method (Vector Laboratories). All other reagents were of analytical grade and obtained from commercial sources.

**Source of normal and cancerous liver tissues.** Normal and cancerous liver tissues were obtained as described (38).
**Figure 7**

MAT1A is the key mediator of miR-664, miR-485, and miR-495 on tumor growth and metastasis. (A) MAT1A Northern and Western blots of Hep3B cells stably transfected with lentiviral miR-664, lentiviral miR-485, and lentiviral miR-485/EV singly or all 3 miRNAs together (miRS), lenti-siRNA against miRNAs (alone or together, miRSi), or SC. *P* < 0.05 vs. respective controls; †*P* < 0.05 vs. individual miRNA or miRNAs. (B) MAT1A Western blots of Hep3B cells stably transfected with lentiviral MAT1A with or without 3′ UTR, †*P* < 0.05 vs. EV. (C) BrDU measurement in Hep3B cells stably expressing MAT1A with or without 3′ UTR, then transiently transfected with lentiviral vector containing all 3 miRNAs, or EV for 24 hours and expressed as percentage of control (EV+EV). Results are mean ± SEM from 3 experiments done in triplicate. †*P* < 0.01 vs. EV+EV; †*P* < 0.05 vs. EV+MAT1A no 3′ UTR; †*P* < 0.05 vs. EV+miRNAs and MAT1A with 3′ UTR+miRNAs; †∗*P* < 0.05 vs. EV+MAT1A with or without 3′ UTR. (D). Hep3B cells stably transfected with MAT1A without 3′ UTR or EV were injected into the left hepatic lobe and treated with lentiviral vector expressing all miRNAs, siRNA against all 3 miRNAs (miRNA siRNAs), or EV. Tumor volumes at the site of injection 45 days later are shown below for each condition. †*P* < 0.05 vs. EV+EV; †∗*P* < 0.05 vs. EV+miRNAs. MAT1A protein levels and incidence of lung metastasis are shown below. †*P* < 0.05 vs. EV+EV. Original magnification, ×200. Numbers below all blots refer to densitometric values expressed as percentage of respective controls.

**Cell lines and stable transfection of lenti-miRNAs and lenti-siRNAs.** 293T, Hep3B, and HepG2 cell lines were obtained from Cell Culture Center of the USC Research Center for Liver Diseases and cultured in DMEM supplemented with 10% fetal bovine serum.

To establish stable expression of miRNA or its siRNA, 10^6 Hep3B cells were seeded in a 24-well plate 1 day prior to infection. To generate cells stably expressing miRNAs or siRNAs, Hep3B cells were transfected with lentiviral miR-664, lenti-miR-485, lenti-miR-495, lenti-EV, lenti-miR-664 siRNA, lenti-miR-485-3p siRNA, lenti-miR-495 siRNA, and lenti-siRNA vector for 3 hours by using Lipofectamine 2000 (Invitrogen). Following selection with puromycin (Invitrogen), stable clonal cell lines were established and examined for the expression of miRNA or siRNA and GFP expression by Northern analysis.

In separate experiments, stable cell lines expressing siRNA against miR-495, miR-485-3p, or miR-664 were transiently transfected with scramble siRNA or siRNA against MAT1A (S90009387; QIAGEN) in Hep3B for 24 hours using RNAiMax (Invitrogen). MAT1A expression, apoptosis, and BrdU incorporation were measured as described below.

**Construction of vectors and stable cell lines expressing multiple miRNAs or their siRNAs and MAT1A with or without its 3′ UTR.** Single miRNA expression vectors for premiR-485 (PMIRH485PA-1), premiR-495 (PMIRH495PA-1), and premiR-664 (PMIRH664PA-1) were obtained from SBI System Biosciences. The multiple miRNA expression vector was constructed by sequentially cloning the premiR-485 and premiR-664 insert into the premiR-495 lentiviral expression vector BamHI and EcoRI sites, respectively.

Oligonucleotides used to construct the lenti-vector containing siRNAs targeting miR-664, miR-485-3p, and miR-495 are shown in Supplemental Table 4. miR-664 siRNA oligonucleotide was inserted into the BamHI site of pGreenPuro (SBI System Biosciences). CMV promoter was amplified by PCR using the primers 5′-TGTATTTAATGATATCAACGTACCCG-3′ (forward primer) and 5′-GATCTGACGGTTACATAAACCAG-3′ (reverse primer) from pGreenPuro and cloned into PCR 2.1 vector by TA cloning (Invitrogen). miR-485-3p siRNA oligonucleotide was inserted at the EcoRV, followed by miR-495 inserted at the SacI and SpeI sites. A second CMV promoter was cloned into the HindIII and KpnI sites. The fragment containing the 2 CMV promoters miR-485-3p and miR-495 was subcloned into the miR-664 containing pGreenPuro EcoRI site. Each shRNA and its CMV promoter was confirmed by sequencing.

MAT1A expression vector cloning was described previously (12). MAT1A 3′ UTR fragment was subcloned into pcDH-CMV-MCS-EF1-copGFP vector (SBI System Biosciences) EcoRI site to create MAT1A without 3′ UTR. Lenti-reporter-luciferase vector containing only the MAT1A 3′ UTR (ABM) was excised and subcloned into the BamHI site of pcDH-CMV-MCS-EF1-copGFP vector to generate MAT1A with 3′ UTR. Stable Hep3B cell lines expressing these vectors were established as described above.

**DNA constructs and dual luciferase assay.** A partial forward and reverse MAT1A 3′ UTR fragment from +1822 bp to +1961 bp and mutant fragments containing Sgf1 and MluI linkers were synthesized. Two base pair mutants were performed for miR-664 (AAATAAT→AAATAAT, where italics represent the targeted nucleotides within the miRNA sequence for mutation), miR-485-3p (TGTTATGA→TGTTGCA), and miR-495 (TTGTGT→TTATT) putative target site(s) in MAT1A 3′ UTR (Figure 2A). The annealed fragments and pmirTarget vector were digested with Sgf1 and MluI. WT and mutant MAT1A 3′ UTR were cloned into the pmirTarget vector containing a luciferase reporter ( OriGene Technologies). Hep3B cells were plated in 24-well plates the day before transfection. The WT, mutated 3′ UTR of MAT1A pmirTarget vector, or pmirTarget EV (200 ng) and a control Renilla luciferase expression vector (2.5 ng) were cotransfected into Hep3B cells with Superfect (QIAGEN) following the manufacturer’s instructions. Luciferase assays were performed 24 hours later using the Dual Luciferase Reporter Assay System (Promega) as directed by the manufacturer’s suggested protocol. Firefly luciferase activity was normalized to Renilla luciferase activity.

**Xenograft model.** Sixty-four 4-week-old male BALB/c nude mice from Jackson ImmunoResearch Laboratories Inc. were divided equally into 8 groups (n = 8 per group) and given the following Hep3B stable cell line injection: group 1, lenti-EV; group 2, lenti-miR-664; group 3, lenti-miR-485; group 4, lenti-miR-495; group 5, lenti-miRNA scramble siRNA; group 6, lenti-miR-664 siRNA; group 7, lenti-miR-485-3p siRNA; group 8, lenti-miR-495 siRNA. Hep3B cells (1 × 10^7) in 100 μL PBS were injected subcutaneously into the right flank of each nude mouse. From week 3 on, xenograft tumor size was measured by calipers. The tumor volume was calculated according to the formula: π/6 (length x width^2) (39). Animals were sacrificed at week 8. Parts of the tumor tissues were used for RNA and protein analysis; the rest were fixed in 4% formalin for histology and immunohistochemistry.

**Orthotopic liver cancer model using Hep3B cells expressing varying levels of miRNAs and MAT1A.** Hep3B cells stably transfected with lentiviral vectors expressing miRNAs or siRNAs against the miRNAs (1.5 × 10^6 cells/50 μl) were slowly injected into the left hepatic lobe of 4-week-old male BALB/c nude mice (n = 8 per group). Animal groups and number were the same as described for the xenograft model. The tumor size in liver tissues was measured as above at day 45 (pilot experiment showed 50% mice died at day 56 in the miR-495 group, but all mice in different groups survived at day 45) and the tumor volume was calculated. Lung, liver, and adjacent tissues were harvested for DNA, RNA, and protein assays as well as standard pathologic studies as described for the xenograft model.

In separate experiments, 4-week-old male BALB/c nude mice (n = 8 per group) were injected in the left hepatic lobe as above with Hep3B cells (1.5 × 10^6 cells/50 μl) and stably transfected with MAT1A expression vector that does not have the 3′ UTR (MAT1A-no 3′ UTR) or EV. Concurrently, mice were also injected into the spleen with lentiviral vector that expresses either all 3 miRNAs (miRNAs), siRNAs against all 3 miRNAs (miRNAsi), or EV. The packaging was done using the Trans Lentiviral pGIPz Packaging system (TLP4614; Open Biosystems). Viral harvesting was done as described in the Open Biosystems protocol. A total of 1 × 10^9 Hep3B cells were infected at a multiplicity of 20 PFU/cell for 24 hours. 2 × 10^9 transducing units (final
Figure 8
Possible mechanism of miR-664, miR-485-3p, miR-495, and MAT1A involvement in tumorgenesis, invasion and metastasis. (A) 5-mC levels in tumors derived from Hep3B cells stably expressing lower (with siRNA or si) or higher levels of miRNAs. *P < 0.05; **P < 0.01; ***P < 0.001 vs. SC; *P < 0.05 vs. EV; n = 8 per condition. (B) Effect of varying miRNA expression on nuclear H3K27me3 levels in tumors (Western blots) and SAIme levels. Results are mean ± SEM from 8 per condition. *P < 0.05; **P < 0.01 vs. respective controls. (C) Diagram shows HpaII and MspI sites between PvuII and NdeI in human LIN28B promoter. Black squares, CCGG sites; TSS, transcriptional start site. Numbers are relative to TSS. Southern blot analysis of LIN28B promoter region between −1576 and +2432 (right). DNA samples from liver tumor derived from stably transfected Hep3B containing miR-664, miR-485, and miR-495 and their srRNAs were digested as indicated. MspI digestion results in a band size of 1369 bp as control for HpaII digestion. (D) Effect of overexpressing miR-664, miR-485, and miR-495 and their srRNAs on MAT1A promoter and LIN28B protein expression (top) and let-7a miRNA expression (bottom). Numbers below the blots are densitometric values expressed as percentage of respective controls. *P < 0.01 vs. EV; *P < 0.05 vs. miR-495; 1P < 0.01 vs. SC; 5P < 0.05 vs. miR-495si. (E) Increased nuclear localization of MAT1A protein is seen after knockdown of miR-664, miR-485-3p, and miR-495. Original magnification, ×630 (oil immersion).

Volume 0.1 ml were injected into the tail veins of mice. In order to maintain a high level of miRNA knockdown, repeated tail-vein injections were done at week 2 and 4. Tumor volumes in the liver and presence or absence of lung metastasis were documented at day 45 as above. Orthotopic liver cancer model using HepG2 cells and treatment with siRNAs against miR-495, miR-485-3p, miR-664, or MAT1A. HepG2 cells have the ability to invade and metastasize (40). To test the effect of knocking down miRNAs, HepG2 cells (1.5 × 10⁶ cells/50 μl) were injected into the left hepatic lobe of 4-week-old male BALB/c nude mice following spleen injection of miRNA siRNA. Then tail-vein injection was done every 2 weeks. Animal groups (n = 8 per group) were as follows: group 1, lenti-scramble siRNA; group 2, lenti-miR-495 siRNA; group 3, lenti-485 siRNA; and group 4, lenti-664 siRNA. Mice were sacrificed at day 56. Lenticival packaging and harvesting were as described above except HeG2 cells were used and tail vein injection was repeated at week 6. Immunohistochemistry and Western blot were done to assess transduction efficiency using GFP.

To examine the role of MAT1A expression on tumorigenicy and the therapeutic effect of miR-495 siRNA, in separate experiments, mice were injected with HepG2 cells directly into the left lobe as above and treated with lenti-MAT1A siRNA and lenti-miR-495 siRNA alone or in combination as described above. Animal groups (n = 8 per group) were as follows: group 1, lenti-scramble vector and lenti-scramble siRNA; group 2, HepG2 injection; group 3, lenti-MAT1A siRNA+lenti-scramble siRNA; group 4, lenti-miR-495 siRNA+lenti-MAT1A siRNA; and group 5, lenti-miR-495 siRNA+lenti-scramble siRNA. Mice were sacrificed at day 56, liver tumor volume at the site of original injection was measured and tissues (lung, liver, pancreas) were harvested for pathological exam as described above.

Global DNA and LIN28B promoter methylation assay. Levels of 5-methylcytosine (5-mC) in hepatic tumors derived from Hep3B cells stably expressing miR-664, miR-485, and miR-495 or their siRNAs were measured by the MethylFlash Methylated DNA Quantification Kit.

DNA samples of tumor tissues from mice injected with Hep3B cells expressing forced miR-664, miR-485, and miR-495 and their respective siRNAs were extracted and digested by MspI, HpaII, PvuII, and NdeI. Southern blot was done as described previously (41).

Northern and Western blot analysis. Northern blotting probes for miR-664, miR485-3p, miR-495, and let-7a were purchased from EXIQON. Total cellular RNA was extracted by using TRizol reagent (Invitrogen) according to the manufacturer’s instructions. Equal amounts of total RNA (15 μg) were denatured, fractionated by electrophoresis on a 15% polyacrylamide–8 M urea gel, electroblotted, and cross-linked onto a nylon membrane. Northern blot analysis was performed as described using Ultrahyb-Oligo (Ambion). As a control for normalization of RNA expression levels, blots were hybridized with an oligonucleotide probe complementary to the U6 RNA (5′-GACAGGCGCCATGCTTCTCTGATCG-3′).

Xenograft and liver tissues isolated from the different treatment groups were subjected to Western blot analysis. Fifteen micrograms of total protein extract were resolved on 12.5% SDS-polyacrylamide gels. Membranes were probed with antibodies to LIN28B, H3K27me3, and MAT1A. To ensure equal loading, membranes were stripped and reprobed with anti-β-actin antibodies. Semi-quantitative analysis was performed for both Northern and Western blots using Quantity One (Bio-Rad).

Histology and immunohistochemistry. Sections from xenograft, liver, lung, and pancreas were fixed with formalin for 4 hours, embedded in paraffin, sectioned, and stained with H&E, as previously described (12). Staining and counting of PCNA were performed according to the manufacturer’s suggested protocol (Invitrogen), whereas MAT1A antibody was diluted to 1:200. Immunohistochemical staining of MAT1A was performed with the Vector ABC Kit according to the manufacturer’s method. For quantifying immunohistochemical staining, a total of 5 fields at x100 magnification were randomly selected (minimum of 1000 cells total), and positive nuclei or cells were counted and expressed as a percentage of the total using MetaMorph imaging software. Control with no antibody showed no staining. Apoptosis and BrdU incorporation. Apoptosis was measured as described (12). BrdU incorporation was measured with BrdU Detection Kit according to the manufacturer’s protocol (BD Biosciences — Pharmingen).

Nuclear SAIme levels. SAIme levels were measured as described (11) in purified nuclear fractions from tumors expressing varying levels of miRNAs or their siRNAs.

Statistics. Data are given as mean ± SEM. Statistical analysis was performed using ANOVA followed by Fisher’s test for multiple comparisons. Significance was defined as P < 0.05.

Study approval. The study protocol conformed to the ethical guidelines of the 1975 Declaration of Helsinki as reflected in a priori approval by Keck School of Medicine University of University of Southern California Health Science Institutional Review Board (Los Angeles, California, USA). All procedure protocols, use, and the care of the animals were reviewed and approved by the Institutional Animal Care and Use Committee at UCLA.

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