miR-27a regulates cisplatin resistance and metastasis by targeting RKIP in human lung adenocarcinoma cells

Jipeng Li1, Yiping Wang1, Yulan Song2, Zhongming Fu3 and Wanjun Yu1,3*

Abstract

Background: MicroRNAs (miRNAs) have been identified as important posttranscriptional regulators involved in various biological and pathological processes of cells, but their association with tumor chemoresistance has not been fully understood.

Methods: We detected miR-27a expression in two lung adenocarcinoma cell lines, A549 and A549/CDDP, and then investigated the effects of miR-27a on the metastasis and the chemosensitivity of cancer cells, using both gain- and loss-of-function studies. The correlation between miR-27a level and chemoresistance was further investigated in clinical lung adenocarcinoma specimens.

Results: miR-27a was significantly up-regulated in cisplatin-resistant lung adenocarcinoma A549/CDDP cells compared with parental A549 cells. miR-27a regulates epithelial-mesenchymal transition (EMT) and cisplatin resistance in vitro and modulates response of lung adenocarcinoma cells to cisplatin in vivo. Further studies identified Raf Kinase Inhibitory Protein (RKIP) as a direct and functional target of miR-27a. Small interfering RNA-mediated RKIP knockdown revealed similar effects as that of ectopic miR-27a expression, while overexpression of RKIP attenuated the function of miR-27a in lung adenocarcinoma cells. Increased miR-27a expression was also detected in tumor tissues sampled from lung adenocarcinoma patients treated with cisplatin-based chemotherapy and was proved to be correlated with low expression of RKIP, decreased sensitivity to cisplatin, and poor prognosis.

Conclusion: Our results suggest that up-regulation of miR-27a could suppress RKIP expression and in turn contribute to chemoresistance of lung adenocarcinoma cells to cisplatin.

Keywords: miR-27a, RKIP, Lung adenocarcinoma, Cisplatin, Chemoresistance

Background

Lung cancer is the leading cause of cancer-related death among men and women worldwide. Approximately 70%-80% of lung cancers are non-small cell lung cancer (NSCLC), including squamous cell carcinoma, adenocarcinoma, and large cell carcinoma [1,2]. In NSCLC, the leading death cause is chemotherapy resistance and metastasis, yet the underlying mechanisms of them remain largely unclear [3-6]. Previous studies in ovarian carcinoma, breast cancer, colorectal cancer and tongue cancer have demonstrated that the drug-resistant cancer cells display features of epithelial-mesenchymal transition (EMT), characterized by the loss of the epithelial marker E-cadherin, an increase in the mesenchymal markers vimentin and N-cadherin, and an increase in the invasion and metastasis behavior [7-11]. Therefore, chemotherapy-induced EMT in tumor cells has been linked to chemotherapeutic resistance and metastasis.

MicroRNAs (miRNAs) are a class of small noncoding RNA molecules that negatively regulate the expression of target genes by either mRNA degradation or translational inhibition. miRNAs are involved in various biological and pathological processes such as differentiation, morphogenesis, and carcinogenesis [12-14]. Accumulating evidence has demonstrated that miRNAs have a key role in drug
resistance and EMT. For example, miR-141 is reported to enhance cisplatin resistance through repression of KEAP1 in ovarian cancer cells [15], miR-106a induces multidrug resistance in gastric cancer by targeting RUNX3 [16]. In contrast, MiR-200b and miR-15b reverse chemotherapy-induced EMT in human tongue cancer cells by targeting BMI1 [10]. The involvement of miRNAs in drug resistance is just beginning to emerge, and more studies are needed to identify other miRNAs, their molecular targets and the processes they affect.

In this study, we observed that miR-27a is significantly upregulated in cisplatin-resistant human lung adenocarcinoma A549/CDDP cells compared with parental A549 cells. Next we explored the roles of miR-27a and its target Raf Kinase Inhibitory Protein (RKIP) in regulating cisplatin resistance and metastasis in lung adenocarcinoma. Finally, we correlated the expression of miR-27a and RKIP with the chemotherapeutic status and prognosis of lung adenocarcinoma patients. Our results show that miR-27a has the potential as key regulatory factors for the chemotherapy resistance and metastasis of lung adenocarcinoma.

Results
Parental A549 cells and cisplatin-resistant A549/CDDP cells differ in morphology, physiology, and miRNA expression
To better understand the biological mechanisms of chemoresistance in lung adenocarcinoma cells and search for the reversion opportunities, we made use of a cisplatin-sensitive and derived resistant lung cancer cell line pair (A549 and A549/CDDP). Compared with parental A549 cells, A549/CDDP cells displayed spindle shape and separated from one another (Figure 1A), MTT assay was then performed, and the IC50 of A549/CDDP cells increased 15.7 fold (Figure 1B). Moreover, western blotting demonstrated that the protein expression of E-cadhein decreased, while that of vimentin dramatically increased in A549/CDDP cells (Figure 1C). Additionally, invasion assay demonstrated that the invasion significantly increased in A549/CDDP as compared with A549 cells (Figure 1D).

Based on the miRNA microarray data, 13 miRNAs were found to be differentially expressed (>2-fold change) in A549/CDDP cells compared with A549 cells (Additional file 1: Table S1), among which miR-27a was the most up-regulated one (5.6-fold change). The result was validated via real-time quantitative RT-PCR (Figure 1E).

miR-27a promotes EMT and cisplatin resistance in vitro
To investigate the association of miR-27a expression with lung adenocarcinoma chemoresistance against cisplatin, A549 and A549/CDDP cells were transfected with miR-27a mimics and miR-27a inhibitors respectively. As shown in Figure 2A, miR-27a mimics increased vimentin, but suppressed E-cadherin expression in A549 cells. Invasion assay demonstrated that miR-27a mimics synergistically enhanced the invasion of A549 cells. On the other hand, silencing miR-27a expression in A549/CDDP cells using anti-sense oligonucleotides reduced vimentin and increased the expression of E-cadherin.
Furthermore, miR-27a inhibitors synergistically suppressed the invasion of A549/CDDP cells (Figure 2B). Therefore, upregulation of miR-27a may play an important role in chemotherapy-induced EMT of lung adenocarcinoma cells. Moreover, MTT assay revealed that A549 cells transfected with miR-27a mimics showed greatly decreased sensitivity to cisplatin as indicated by substantially increased IC50 values (Figure 2C). In contrast, suppression of the miR-27a level in A549/CDDP cells resulted in an enhanced sensitivity to cisplatin (Figure 2D). Similar results were obtained in H1395 and H1299 cells (Additional file 2: Figure S1).

**miR-27a regulates response of lung adenocarcinoma cells to cisplatin in vivo**

To investigate the effect of miR-27a expression on chemosensitivity of lung adenocarcinoma in vivo, A549 cells stably expressing miR-27a by lentivirus were subcutaneously inoculated into nude mice. When the average tumor size reached $= 50 \text{ mm}^3$, cisplatin was administered via intraperitoneal injection at a dose of 5 mg/kg, 1 dose every other day, with 3 doses in total. As shown in Figure 3A, the number of lung metastasis nodules was dramatically increased in miR-27a overexpression group when compared with control. Next, we engineered A549/CDDP cells to stably inhibit miR-27a with a lentivirus-mediated antagonir. The results showed that suppression of miR-27a decreased the number of lung metastases (Figure 3B). These results suggested that miR-27a could regulate the response to cisplatin in vivo.

**miR-27a directly targets RKIP**

By employing open access softwares (TargetScan and PicTarget), RKIP was chosen as a preferred candidate target gene of miR-27a because of the putative binding site within its 3'UTR (Figure 4A) and lower RKIP protein expression in A549/CDDP cells (Figure 4B). Western blot showed that overexpression of miR-27a in A549 cells significantly repressed RKIP protein expression compared to cells transfected with negative control (Figure 4C). Relatively, downregulation of miR-27a by inhibitors in A549/CDDP cells led to a moderate increase of RKIP protein level (Figure 4C). To verify whether RKIP is the direct downstream target of miR-27a, a fragment of RKIP 3'UTR containing the putative miR-27a binding site was cloned into a luciferase reporter vector. Luciferase reporter assays showed that up-regulation of miR-27a significantly decreased the relative luciferase activity of RKIP 3'UTR in A549 cells, but had no effect on the mutant of RKIP 3'UTR (Figure 4D). Taken together, these results suggest that miR-27a down-regulates RKIP expression by directly targeting its 3'UTR.
RKIP is involved in miR-27a-induced EMT and cisplatin resistance

To further examine whether RKIP is involved in miR-27a-induced chemoresistance, we performed loss-of-function and gain-of-function analyses. Firstly, A549 cells were transfected with si-RKIP or negative control. Western blotting analysis confirmed that the expression of RKIP was suppressed (Figure 5A). As expected, RKIP knockdown significantly increased vimentin, reduced E-cadherin and decreased sensitivity to cisplatin in A549 cells (Figure 5A). Subsequently, we employed a/Xpression construct that encodes the entire RKIP coding sequence but lacks the 3′UTR. Ectopic expression of RKIP partially rescued miR-27a-mediated EMT and cisplatin resistance in miR-27a-overexpressing cells (Figure 5B). Collectively, these data suggest that miR-27a regulate chemoresistance.

Figure 3 miR-27a regulates response of lung adenocarcinoma cells to cisplatin in vivo. (A) A549 cells were performed after transduction by miR-27a-expressing or vector lentivirus. Representative images of nude mouse lungs (scale bars: 5 mm) and H&E stain of lungs (scale bars: 100 μm) are shown. (B) A549/CDDP cells were performed after transduction by miR-27a-inhibiting or antagonir-NC. Representative images of nude mouse lungs (scale bars: 5 mm) and H&E stain of lungs (scale bars: 100 μm) are shown. *P < 0.01.

Figure 4 RKIP is a direct target of miR-27a. (A) The predicted miR-27a binding site within RKIP 3′UTR and its mutated version by site mutagenesis are as shown. (B) Variable RKIP expression in A549 and A549/CDDP was obtained by western blot. (C) A549 cells were transfected with NC or miR-27a mimics, A549/CDDP cells were transfected with anti-NC or miR-27a inhibitors respectively. Western blotting was used to detect RKIP expression; β-actin was used as an internal control. (D) Luciferase assay was performed in A549 cells that were co-transfected with miRNA mimics and reporter vectors carrying RKIP 3′ UTR with wild type versus mutated miR-27a response element. Data are means of three separated experiments ± SD; *P < 0.01.
of lung adenocarcinoma cells at least in part by targeting RKIP.

**High expression of miR-27a in lung adenocarcinoma tissues is associated with decreased RKIP expression, chemotherapeutic resistance, and poor prognosis**

To better understand the association between miR-27a and RKIP expression, a total of 30 clinical tumor tissue samples were collected from patients with advanced lung adenocarcinoma and divided into “sensitive” and “insensitive” groups according to the patient’s response to cisplatin-based chemotherapy. As shown in Figure 6A, miR-27a was significantly up-regulated in the “insensitive” group tissues (n = 17) compared with that in the “sensitive” group ones (n = 13). On the contrary, RKIP mRNA expression level was significantly down-regulated in the “insensitive” group tissues (Figure 6B). The inverse correlation between miR-27a and RKIP mRNA expression was verified by linear regression analysis (r = −0.691, P < 0.01) (Figure 6C). We then analyzed the association of miR-27a expression with survival of patients. As shown in Figure 6D, Patients with high miR-27a expression showed significantly shorter overall survival than those with low miR-27a expression (P < 0.01).

**Discussion**

In the current study, we demonstrate that upregulation of miR-27a is critical for cisplatin resistance and tumor metastasis of lung adenocarcinoma cells both in vitro and in vivo, and miR-27a induces mesenchymal features and promotes tumor metastasis of chemoresistant lung adenocarcinoma via silencing RKIP. Furthermore, upregulation of miR-27a is correlated with cisplatin resistance and poor prognosis of lung adenocarcinoma patients. These findings provide new insights into the molecular functions of miR-27a as well as the role of RKIP in chemotherapeutic resistance.

MiR-27a is located at chromosome 19 and has been shown to be overexpressed in breast cancer, gastric cancer and cervical cancer [17-19]. In gastric cancer cells, miR-27a promotes cell growth and metastasis both in vitro and in vivo [20,21]. In addition, it could regulate endothelial differentiation of breast cancer stem like cells [22]. Moreover, miR-27a plays an important role in mediating drug resistance by targeting multiple drug-resistance related genes. MiR-27a modulated MDR1/P-glycoprotein expression in human ovarian cancer cells by targeting HIPK2 [23] and Down-regulation of miR-27a might reverse multidrug resistance of esophageal squamous cell carcinoma through regulation of MDR1 and apoptosis [24]. Despite the oncogenic role of miR-27a has been implicated by previous studies, the role of miR-27a in lung cancer chemotherapy and molecular mechanisms are not known. Here we identified RKIP as the functional target, through which miR-27a regulates metastasis and chemoresistance.

Raf Kinase Inhibitory Protein (RKIP), a member of the phosphatidylethanolamine binding protein (PEBP) family, is widely expressed in normal human tissues, highlighting its role in various physiologic processes [25], but is considered to be a metastasis suppressor in cancer, being its loss or reduced expression associated with malignancy and prognosis in many types of metastatic and aggressive cancers [26-29]. Recently, a study showed that snail, a mediator of the EMT, can inhibit RKIP transcription and
negatively correlates with RKIP levels in tumors [30]. Additionally, another study reported that RKIP inhibition in cervical cancer is associated with higher tumor aggressive behavior and resistance to cisplatin therapy [31]. Our finding showed that downregulation of RKIP induces EMT and contributes to cisplatin resistance. We further establish miR-27a-RKIP as another important pathway regulating EMT and chemoresistance of lung adenocarcinoma cells.

Chemoresistance is a major issue of treatment in the majority of human tumors, including lung cancer. Thus, detecting rationale biomarkers to predict chemotherapy sensitivity and screening for targets to overcome resistance are significant for cancer therapy [32]. A specific miRNA can affect simultaneously the expression of proteins involved in multiple cellular pathways, potentially serving as better therapeutic target or biomarker for clinical outcome than single proteins. In fact, several miRNAs, including miR-21, miR-10b and miR-125b, have been used as predictors of chemoresistance in cancers [32-34]. Herein, our observation that increased miR-27a expression is associated with chemotherapy resistance, and poor patient prognosis may provide surrogates to predict the chemotherapeutic sensitivity for lung adenocarcinoma.

**Conclusions**

In conclusion, we have reported the altered expression of miR-27a in human lung adenocarcinoma cell lines with different sensitivities to cisplatin, and have shown that miR-27a could modulate cisplatin resistance and metastasis in these cells by targeting RKIP. Therefore, targeting miR-27a-RKIP interaction may be a potential strategy for reversing chemoresistance in human lung adenocarcinoma.

**Materials and methods**

**Cell culture and transfection**

Human lung adenocarcinoma A549 cells were purchased from the American Type Culture Collection and cultured in RPMI 1640 medium containing 10% fetal bovine serum (FBS) with 100 μg/ml penicillin /streptomycin at 37°C with 5% CO₂. The cisplatin-resistant A549 cell line (A549/CDDP) was established and preserved in a 40 μmol/L final concentration of cisplatin in our laboratory.

miR-27a mimics and negative control mimics (NC), miR-27a inhibitors (anti-miR-27a) and negative control inhibitors (anti-NC) and RKIP siRNAs were synthesized by GenePharma Company (Shanghai, China). Transfection was performed with Lipofectamine 2000 (Invitrogen, CA, USA).
A total of 30 lung adenocarcinoma tissues were collected from patients with advanced lung adenocarcinoma who received chemotherapy at Yinzhou People’s Hospital (Ningbo, China) between January 2009 and March 2010. Informed consent was obtained from all subjects and this study was approved by the Clinical Research Ethics Committee of Yinzhou People’s Hospital. Patients met all of the following criteria: primary lung adenocarcinoma; histological diagnosis of lung adenocarcinoma with at least 1 measurable lesion; clinical stage IIIIB-IV; first-line chemotherapy either with cisplatin 100 mg/m^2 and docetaxel 75 mg/m^2 or cisplatin 100 mg/m^2 and gemcitabine 1000 mg/m^2 administered every 3 weeks for a maximum of 5 cycles. Samples were divided into “sensitive” (complete response or partial response) and “insensitive” (stable disease or progressive disease) groups according to the patient’s responses assessed via medical image analysis and detection of serum tumor markers after 4 or 5 cycles of cisplatin-based chemotherapy.

**RNA extraction and qRT–PCR**

Total RNA was extracted from the cultured cells and the lung adenocarcinoma tissue specimens using Trizol Reagent (Invitrogen, CA, USA) according to the manufacturer’s protocol. The expression level of mature miR-27a was measured by TaqMan miRNA assays (Applied Biosystems, CA, USA) according to the provided protocol, U6 snRNA levels were used for normalization. RKIP expression was measured by SYBR green qPCR assay (Takara, Dalian, China) and GAPDH was used as an endogenous control.

**Western blot analysis**

Protein extracts were prepared by a modified RIPA buffer with 0.5% sodium dodecyl sulfate (SDS) in the presence of proteinase inhibitor cocktail (Complete mini, Roche, Indianapolis, IN, USA). Polycrylamide gel electrophoresis, tank-based transfer to Immobilon Hybond-C membranes (Amersham Biosciences) and immunodetection were performed with standard techniques. Antibodies to RKIP (catalog no. sc-28837), E-cadherin (catalog no. sc-8426), vimentin (catalog no. sc-32322) and β-actin (catalog no. sc-1616) were purchased from Santa Cruz Biotechnology, Inc. (Santa Cruz, CA). Signals were visualized with SuperSignal® West Pico chemiluminescent substrate (Pierce, Rockford, Ill, USA) by exposure to films.

**Cell viability assay**

Cells were seeded into 96-well plates (2 × 10^3 cells/well) directly or 24 hours after transfection and allowed to attach overnight. Freshly prepared cisplatin was then added with different final concentrations. Forty-eight hours later, cell viability was assessed via 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl-tetrazolium bromide (MTT) assay as described previously [35].

**Transwell invasion assay**

2 × 10^5 cells were added into the upper chamber of the insert precoated with Matrigel (ECM gel, Sigma-Aldrich, St. Louis, MO). Cells were plated in medium without serum, and medium containing 10% fetal bovine serum in the lower chamber served as chemoattractant. After several hours of incubation, the cells that did not invade through the pores were carefully wiped out with cotton wool, and the filters were fixed by treatment with 95% ethanol for 30 minutes and stained with 0.2% crystal violet solution for 30 minutes. Invasive cells adhering to the undersurface of the filter were counted (five fields/chamber; 0.24 mm^2/field) using an inverted microscope, and each experiment was repeated three times.

**Plasmid construction and luciferase reporter assay**

Wild-type 3’ untranslated region (3’ UTR) of RKIP containing predicted miR-27a target sites were amplified by PCR from A549 cell genomic DNA. Primers used: Forward: GAT CTG CAG GGG TTA GCT TGG GGA C; Reverse: GAT CAT ATG AGA GTG ACA TAC TGA TGC CTA C. Mutant 3’UTRs were generated by overlap-extension PCR method. Both wild-type and mutant 3’UTR fragments were subcloned into the pGL3-control vector (Promega, Madison, WI) immediately downstream of the stop codon of the luciferase gene. DNA fragment coding RKIP protein was amplified by PCR from A549 cell cDNA, and cloned into pCMV-Myc expression vector (Clonetech, Mountain View, CA). Primers used: Forward: GCT GAA TTC ATG CCG GTG GAC CTC AGC AAG T; Reverse: CTG CTC GAG CTA CTT CCC AGA CAG CTG CTC G. For luciferase assay, the reporter plasmid was cotransfected with a control Renilla luciferase vector into A549 cells in the presence of either miR-27a or NC. After 48 h, cells were harvested, and the luciferase activity was measured using the Dual-Luciferase Reporter Assay System (Promega, Madison, WI, USA).

**Animal studies**

Five-week-old female BALB/c nude mice were purchased from the Animal Center of Zhejiang University (Hangzhou, China). For in vivo chemosensitivity and metastasis assays, A549 cells (infected with either the miR-27a-overexpressing lentivirus or the mock lentivirus) and A549/CDDP cells (infected with either the miR-27a-knockdown lentivirus-mediated antagonir or the antagonir-NC) were subcutaneously inoculated into...
nude mice (six per group, 1 × 10⁶ cells for each mouse). Tumor growth was examined every other day, and tumor volumes were calculated using the equation V = A × B²/2 (mm³), where A is the largest diameter and B is the perpendicular diameter. When the average tumor size reached ≈ 50 mm³, cisplatin was administered via intraperitoneal injection at a dose of 5 mg/kg, 1 dose every other day, with 3 doses in total. After 2 weeks, all mice were sacrificed. Transplanted tumors were excised, and tumor tissues were used to perform hematoxylin & eosin (H&E) staining. All research involving animal complied with protocols approved by the Zhejiang medical experimental animal care commission.

Statistical analysis
Statistical analyses were performed using SPSS 16.0 software (SPSS Inc.). All data from 3 independent experiments were expressed as mean ± SD. Differences were assessed by two-tailed Student's t test. P < 0.05 was considered statistically significant.

Additional files

Additional file 1: Table S1. miRNAs differentially expressed in the A549/DDP and A549 cell lines.

Additional file 2: Figure S1. miR-27a regulates EMT and cisplatin resistance in H1395 and H1299 cells. (A) The expression of miR-27a was examined by qRT-PCR. U6 small nuclear RNA was used as an internal control. (B) Western blotting was used to detect E-cadherin and vimentin expression. β-actin was used as an internal control. (C and D) MTT assay was used to measure cisplatin sensitivity. Data are means of three separated experiments ± SD; *P<0.05.

Competing interests
The authors declare that they have no competing interests.

Authors' contributions
JL designed and performed experiments, wrote the manuscript. YS and YW performed experiments. ZF gave suggestion on study design, discussed and interpreted the data. WY designed and supervised study, discussed and interpreted the data. All authors read and approved the final manuscript.

Acknowledgements
This work was supported by Zhejiang Provincial Natural Science Foundation of China (Grant No. LY14H160002), Zhejiang Provincial Medicine and Health Science Research Foundation of China (Grant No. 2014KYB248), Ningbo Municipal Medical Science and Technique Foundation (Grant No. 2013A330) and Yinzhou Science and Technology bureau (2011–111, 2013–107).

Author details
1Department of Clinical Laboratory, Yinzhou People's Hospital, Ningbo 315040, China. 2Department of Nephrology, Yinzhou People's Hospital, Ningbo 315040, China. 3Department of Respiratory and Critical Care Medicine, Yinzhou People's Hospital, 251 East Baizhang Road, Ningbo 315040, China.

Received: 15 May 2014 Accepted: 12 August 2014 Published: 16 August 2014

References
1. Jemal A, Thun MJ, Ries LA, Howe HL, Weir HK, Center MM, Ward E, Wu XC, Ehliman C, Anderson R, Ayeni UA, Kohler B, Edwards BK: Annual report to the nation on the status of cancer, 1975–2005, featuring trends in lung cancer, tobacco use, and tobacco control. J Natl Cancer Inst 2008, 100:1672–1694.
2. Jemal A, Siegel R, Xu J, Ward E: Cancer statistics, 2010. CA Cancer J Clin 2010, 60:277–300.
3. Gupta GP, Massague J: Cancer metastasis: building a framework. Cell 2006, 127:679–696.
4. Goldman B: Multidrug resistance: can new drugs help chemotherapy score against cancer? J Natl Cancer Inst 2003, 95:255–257.
5. Gao D, Vahdat LT, Wong S, Chang JC, Mittal V: Microenvironmental regulation of epithelial-mesenchymal transitions in cancer. Cancer Res 2012, 72:4883–4889.
6. Sato M, Shames DS, Gazdar AF, Minna JD: A translational view of the molecular pathogenesis of lung cancer. J Thorac Oncol 2007, 2:327–343.
7. Kajiyama H, Shibata K, Terauchi M, Yamasita M, Ino K, Nawa A, Kikka F: Chemoresistance to paclitaxel induces epithelialmesenchymal transition and enhances metastatic potential for epithelial ovarian carcinoma cells. Int J Oncol 2007, 31:277–283.
8. Hiscox S, Jiang W, Obermeier K, Taylor K, Morgan L, Burmi R, Barrow D, Nicholson RI: Tamoxifen resistance in MCF7 cells promotes EMT-like behaviour and involves modulation of beta-catenin phosphorylation. Int J Cancer 2006, 118:290–301.
9. Yang AD, Fan F, Camp ER, Van Buren G, Liu W, Somico R, Gray M, Cheng H, Hoff PM, Ellis LW: Chronic oxaliplatin resistance induces epithelial-to-mesenchymal transition in colorectal cancer cells. Clin Cancer Res 2006, 12:4147–4153.
10. Sun L, Yao Y, Liu B, Lin L, Yang M, Zhang W, Chen W, Pan C, Liu G, Song E, Li J: miR-200b and miR-15b regulate chemotherapeutic-induced epithelial-mesenchymal transition in human tongue cancer cells by targeting BMI1. Oncogene 2012, 31:432–445.
11. Singh A, Sett Row J: EMT, cancer stem cells and drug resistance: an emerging axis of evil in the war on cancer. Oncogene 2010, 29:3747–3751.
12. Godeladze JO, Djouad F, Brondello JM, Duroux-Richard J, Apparailly F, Jorgensen C: Concerted stimuli regulating osteo-chondral differentiation from stem cells: phenotype acquisition regulated by microRNAs. Acta Pharmacol Sin 2009, 30:1389–1394.
13. Wu F, Yang Z, Li G: Role of specific microRNAs for endothelial function and angiogenesis. Biochem Biophys Res Commun 2009, 386:549–555.
14. Skaffrinozko KO, Prestegarden L, Mickle DR, Lorens JB: MicroRNAs in tumorigenesis. Curr Pharm Biotechnol 2007, 8:320–325.
15. Van Jaarsveld MT, Helleman J, Boersma AW, Van Kuijk PF, Van Lichen WF, Despierie E, Vergote I, Mathijssen RH, Bens EM, Verweij J, Pothof J, Wiemer EA: miR-141 regulates KEAP1 and modulates cisplatin sensitivity in ovarian cancer cells. Oncogene 2013, 32:4284–4293.
16. Zhang Y, Lu Q, Cai X: MicroRNA-106a induces multidrug resistance in gastric cancer by targeting RUNX3. FEBS Lett 2013, 587:3069–3075.
17. Mertens-Talcott SU, Chintaphaladi S, Li X, Safe S: The oncogenic microRNA-27a targets genes that regulate specificity protein transcription factors and the G2-M checkpoint in MDA-MB-231 breast cancer cells. Cancer Res 2007, 67:1101–1111.
18. Huang D, Wang H, Liu R, Li H, Ge S, Bai M, Deng T, Yao G, Ba Y: miR-27a is a biomarker for predicting chemosensitivity and prognosis in metastatic or recurrent gastric cancer. J Cell Biochem 2014, 15:549–556.
19. Wang X, Tang S, Le SY, Lu R, Rader JS, Meyers C, Zheng ZM: Aberrant expression of oncogenic and tumor-suppressive microRNAs in cervical cancer is required for cancer cell growth. PLoS One 2008, 3:e2557.
20. Zhao X, Yang L, Hu J: Down-regulation of miR-27a might inhibit proliferation and drug resistance of gastric cancer cells. J Exp Clin Cancer Res 2011, 30:55.
21. Zhang Z, Liu S, Shi R, Zhao G: miR-27 promotes human gastric cancer cell metastasis by inducing epithelial-to-mesenchymal transition. Cancer Genet 2011, 204:486–491.
22. Tang W, Yu F, Yao H, Cui X, Jiao Y, Lin L, Chen J, Yin D, Song E, Liu Q: miR-27a regulates endothelial differentiation of breast cancer stem like cells. Oncogene 2014, 33:2629–2638.
23. Li Z, Hu S, Wang J, Cai J, Xiao L, Yu L, Wang Z: miR-27a modulates MDRI/ Pglycoprotein expression by targeting HIPK2 in human ovarian cancer cells. Gynecol Oncol 2010, 119:125–130.
24. Zhang H, Li M, Han Y, Hong L, Gong T, Sun L, Zheng X: Down-regulation of miR-27a might reverse multidrug resistance of esophageal squamous cell carcinoma. Dig Dis Sci 2010, 55:2545–2551.
25. Granovsky AE, Rosner MR: Raf kinase inhibitory protein: a signal transduction modulator and metastasis suppressor. Cell Res 2008, 18:452–457.

26. Escara-Wilke J, Yeung K, Keller ET: Raf kinase inhibitor protein (RKIP) in cancer. Cancer Metastasis Rev 2012, 31:515–520.

27. Das SK, Bhutia SK, Sokhi UK, Azab B, Su ZZ, Boukerche H, Anwar T, moen EL, Chatterjee D, Pellecchia M, Sarkar D, Fiser PB: Raf kinase inhibitor RKIP inhibits MDA-9/syntenin-mediated metastasis in melanoma. Cancer Res 2012, 72:6217–6226.

28. Zhang XM, Gu H, Yan L, Zhang GY: RKIP inhibits the malignant phenotypes of gastric cancer cells. Neoplasma 2013, 60:196–202.

29. Koelzer VH, Karamitopoulou E, Dawson H, Kondi-Pafiti A, Zlobec I, Lugli A: Geographic analysis of RKIP expression and its clinical relevance in colorectal cancer. Br J Cancer 2013, 108:2088–2096.

30. Beach S, Tang H, Park S, Dhillon AS, Keller ET, Kolch W, Yeung KC: Snail is a repressor of RKIP transcription in metastatic prostate cancer cells. Oncogene 2008, 27:3243–3248.

31. Martinho O, Pinto F, Granja S, Miranda-Goncalves V, Moreira MA, Ribeiro LF, di Loreto C, Rosner MR, Longatto-Filho A, Reis RM: RKIP inhibition in cervical cancer is associated with higher tumor aggressive behavior and resistance to cisplatin therapy. PLoS One 2013, 8(3):e59104.

32. Hwang JH, Voortman J, Giovannetti E, Steinberg SM, Leon LG, Kim YT, Funel N, Park JK, Kim MA, Kang GH, Del Chiaro M, Peters GJ, Giaccone G: Identification of microRNA-21 as a biomarker for chemoresistance and clinical outcome following adjuvant therapy in resectable pancreatic cancer. PLoS One 2010, 5:e10630.

33. Nishida N, Yamashita S, Mimori K, Sudo T, Tanaka F, Shibata K, Yamamoto H, Ishii H, Doki Y, Mori M: MicroRNA-10b is a prognostic indicator in colorectal cancer and confers resistance to the chemotherapeutic agent 5-fluorouracil in colorectal cancer cells. Ann Surg Oncol 2012, 19:3065–3071.

34. Wang H, Tan G, Dong L, Cheng L, Li K, Wang Z, Luo H: Circulating miR-125b as a marker predicting chemoresistance in breast cancer. PLoS One 2012, 7:e34210.

35. Tsang WP, Kwok TT: The miR-18a* microRNA functions as a potential tumor suppressor by targeting on K-Ras. Carcinogenesis 2009, 30:953–959.

doi:10.1186/1476-4598-13-193
Cite this article as: Li et al.: miR-27a regulates cisplatin resistance and metastasis by targeting RKIP in human lung adenocarcinoma cells. Molecular Cancer 2014 13:193.