

MicroRNA-23a promotes neuroblastoma cell metastasis by targeting CDH1

LIN CHENG, TAO YANG, YONGQIN KUANG, BIN KONG, SIXUN YU,
HAIFENG SHU, HUTIAN ZHOU and JIANWEN GU

Department of Neurosurgery, Chengdu Military General Hospital, Chengdu, Sichuan 610083, P.R. China

Received May 13, 2013; Accepted November 18, 2013

DOI: 10.3892/ol.2014.1794

Abstract. CDH1 inactivation is important in tumor metastasis. In the present study, it was suggested that the mRNA and protein levels of CDH1 decreased in metastatic neuroblastoma (NB) tissues compared with those in primary NB tissues. The aim of the study was to explore the regulatory mechanisms of CDH1 downregulation in metastatic NB. MicroRNAs are small non-coding RNAs (~22 nt in length) that negatively regulate target mRNAs and are involved in various cancer-related processes, including metastasis. In the current study, miR-23a was shown to be upregulated in human metastatic NB tissues compared with primary NB tissues. Inhibition of miR-23a may significantly suppress NB cell migration and invasion. *In vitro* reporter assay suggested that CDH1 is a direct target gene of miR-23a. Furthermore, blocking the expression of miR-23a partly restored the expression of CDH1 in NB cells. These findings provide evidence that miR-23a is key in promoting NB cell migration and invasion through targeting CDH1, and suggest that exogenous miR-23a may have therapeutic value in treating NB metastasis.

Introduction

Characterized by spontaneous regression, maturation or aggressive progression, neuroblastoma (NB) is one of the most common forms of cancer in young children. NB typically presents with metastatic disease at diagnosis with a poor outcome (1,2). The most critical step in NB metastasis is that the cells from the primary tumor, which have enhanced capabilities of migration and invasion, invade into the surrounding normal tissues (3). To date, a set of genes associated with tumor progression in NB have been revealed (4). However, the precise mechanisms, particularly the significant molecular

pathways in the transition from primary to metastatic NB, remain to be completely understood.

The *CDH1* gene encodes the epithelial cell adhesion molecule, E-cadherin, which forms the core of the adherens junctions between adjacent epithelial cells (5). The significance of CDH1 inactivation for metastasis has been demonstrated in a variety of studies (6-11). However, few studies have analyzed the potential important roles of CDH1 in metastatic NB tissues.

Previously, a group of non-coding single-stranded RNAs, ~22 nt in length, which modulate gene expression post-transcriptionally by interacting with complementary sites within the 3' untranslated region (UTR) of target mRNA, have emerged as important regulators in cancer-related processes, including metastasis (12-15). In a previous study, Guo *et al* observed differential expression of microRNAs (miRNAs) in metastatic NB compared with primary NB using microarray analysis. In addition, the potential roles of these miRNAs in the NB metastatic process were investigated (16). However, the role of miR-23a in metastatic NB tissues compared with primary NB tissues remains unclear. Hence, we proposed that the upregulated miR-23a enhances the metastatic capability of NB cells. Exploring the miR-23a-mediated NB metastasis molecular mechanism may be useful to characterize the progression from primary to metastatic NB.

Materials and methods

Tissue samples and cell lines. In total, nine pairs of human primary and metastatic NB tissues were acquired from the Chengdu Military General Hospital (Chengdu, China). The study was approved by the ethics committee of Chengdu Military General Hospital and informed written consent was obtained from the patients. All the tissues were confirmed by pathology and immunohistochemistry, frozen in liquid nitrogen and stored at -80°C. Human NB cell lines, SK-N-SH and GI-LA-N, were cultured in Eagle's minimum essential medium (Gibco-BRL, Carlsbad, CA, USA) supplemented with 10% FBS and 100 U/ml penicillin/streptomycin. All the cells were placed in a humidified incubator with 5% CO₂ at 37°C.

RNA isolation and qPCR. Total RNA was isolated using TRIzol reagent (Invitrogen Life Technologies, Carlsbad, CA, USA) according to the manufacturer's instructions. The RNA concentration and quality was measured by a NanoDrop

Correspondence to: Dr Jianwen Gu, Department of Neurosurgery, Chengdu Military General Hospital, 270 Rongdu Road, Chengdu, Sichuan 610083, P.R. China
E-mail: gujianwen2013@163.com

Key words: miRNA, miR-23a, CDH1, neuroblastoma

ND-1000 spectrophotometer (Thermo Fisher Scientific, Waltham, MA, USA). Next, 1 μ g RNA was used for reverse transcription (RT). Specific miR-23a reverse transcription (RT) primers were used for the cDNA from the miRNA RT reaction, while the common primer Oligo (dT) was used for the cDNA from total RNA, for detection of CDH1. qPCR was performed using the SYBR Green PCR master mix (Applied Biosystems, Inc., Foster City, CA, USA) according to the following conditions: 95°C for 5 min followed by 40 cycles of amplification at 95°C for 30 sec, 57°C for 30 sec and 72°C for 30 sec. U6 small nuclear B non-coding RNA was used as the internal control to normalize miR-23a expression and GAPDH was used as the internal control to normalize CDH1 expression.

Western blot analysis. The NB cells were seeded into six-well plates at a density of 3×10^4 cells/well and cells were transfected once cell density reached ~80% confluence on the second day. At 48 h following transfection, the cells were lysed using RIPA buffer for 30 min at 4°C. The protein concentration was measured by the bicinchoninic acid method using a Pierce® BCA Protein Assay kit (Thermo Fisher Scientific) and then 20 μ g protein was loaded on SDS-PAGE for analysis. The primary antibodies used were rabbit polyclonal anti-human CDH1 (1:500; Abcam, Cambridge, MA, USA) and rabbit monoclonal anti-human GAPDH (1:1,000; Abcam). The secondary antibody used was a goat anti-rabbit IgG conjugated with horseradish peroxidase (1:1,000; Abcam). The bound antibodies were detected by ECL Plus Western blotting detection system (GE Healthcare, Amersham, UK) and the chemiluminescent signals were detected by high-performance chemiluminescence film (GE Healthcare).

EGFP reporter assay. The 3'UTR of CDH1 was amplified and cloned downstream of the pcDNA3/EGFP vector. Next, the mutant 3'UTR of CDH1 (in which several nucleotides within the binding sites were deleted) was amplified using pcDNA3/EGFP-CDH1 3'UTR as the template and cloned downstream of the pcDNA3/EGFP vector. For the EGFP reporter assay, the cells were co-transfected with miR-23a mimics, ASO-miR-23a and pcDNA3/EGFP-CDH1 3'UTR or the mutant 3'UTR, together with the controls. The plasmid expressing RFP was transfected as the spike-in control. At 48 h following transfection, the cells were lysed using RIPA buffer, and EGFP and RFP intensity was measured by an F-4500 fluorescence spectrophotometer (Hitachi, Ltd., Tokyo, Japan).

Transwell migration and invasion assays. The migration and invasion assays were performed using the Transwell chamber (Millipore, Billerica, MA, USA). For the migration assay, following transfection, the cells were seeded into the upper chamber (2.5×10^4 cells/well) in 250 μ l serum-free medium, but the bottom chamber was incubated with 750 μ l medium containing 10 or 20% serum. For the invasion assay, the chamber was coated with Matrigel and the cells were seeded into the upper chamber (2.5×10^4 cells/well) in 250 μ l serum-free medium, while the bottom chamber was incubated with 750 μ l medium containing 10 or 20% serum. Following cell migration or invasion for 20 h, the cells that had not migrated

or invaded were scraped off with a cotton swab. The cells which had migrated or invaded into the chamber membrane were fixed and then stained with crystal violet for ~15 min. Finally, images of the cells were captured under a microscope (ME21 digital microscope, Olympus, Guangzhou, China) and cells were counted.

Statistics. Data are presented as the mean \pm standard deviation and the difference between groups was determined by a two-tailed Student's t-test. $P < 0.01$ or $P < 0.05$ were considered to indicate a statistically significant difference.

Results

Quantitative analysis of CDH1 expression in metastatic NB tissues. CDH1 is an important cancer metastasis-related molecule. To test the expression of CDH1 in metastatic NB tissues and matched primary NB tissues, qPCR assay was performed in nine pairs of NB tissues. It was shown that CDH1 mRNA expression levels were generally and significantly ($P < 0.01$) lower in metastatic NB tissues than in the matched primary NB tissues (Fig. 1A). In addition, western blot analysis confirmed that the CDH1 protein expression was lower in metastatic NB tissues compared with that in primary NB tissues within the representative patients 3, 6 and 7.

miR-23a is a candidate regulator of CDH1 in NB. miRNAs function as tumor suppressors or oncogenes, through the direct regulation of associated oncogenes or tumor suppressor genes (14,17,18). The oncogene miRNAs (onco-miRs), such as miR-23a, are usually upregulated in tumors and may cause the loss of expression of tumor suppressors and contribute to the tumorigenesis of cancer, including metastasis. The loss of CDH1 expression in metastatic NB tissues implies that there may be important miRNAs involved in the regulation of CDH1 in NB metastasis. Greater specificity in miRNA predictions may be attained by the consensus of multiple algorithms. Therefore, three programs (TargetScan, microRNA.org and miRDB) were used for the prediction of which miRNAs target the 3'UTR of CDH1 and regulate its expression. Table I shows the 20 miRNAs that were identified to target the 3'UTR of CDH1. In addition, Table II shows the 34 downregulated miRNAs collected from metastatic NB tissues to compare with primary NB tissues, according to the previous study by Guo *et al* using the microarray analysis (16). Based on these two tables, miR-23a (Fig. 2) was identified as a candidate for directly targeting CDH1 whose mRNA 3'UTR contains a putative binding site. This analysis is consistent with the model in which tumor onco-miRs were found to promote tumor development by targeting and negatively regulating tumor suppressors (13).

miRNA-23a is upregulated in metastatic NB tissues and regulates NB cell migration and invasion. miR-23a has been previously reported to play a role as an onco-miR in several types of cancer, and the microarray analysis identified that miR-23a is upregulated in metastatic NB tissues compared with primary NB tissues. Next, qPCR assay was used to detect the expression of miR-23a in nine pairs of NB samples (metastatic and matched primary NB tissues). It was

Table I. Predicted miRNAs targeting CDH1 by three algorithms.

Gene name	miRNA	microRNA.org	miRDB	TargetScan
CDH1	hsa-miR-1248	+	+	+
CDH1	hsa-miR-219-2-3p	+	+	+
CDH1	hsa-miR-504	+	+	+
CDH1	hsa-miR-340	+	+	+
CDH1	hsa-miR-580	+	+	+
CDH1	hsa-miR-296-3p	+	+	+
CDH1	hsa-miR-1299	+	+	+
CDH1	hsa-miR-1296	+	+	+
CDH1	hsa-miR-515-5p	+	+	+
CDH1	hsa-miR-1323	+	+	+
CDH1	hsa-miR-1264	+	+	+
CDH1	hsa-miR-548o	+	+	+
CDH1	hsa-miR-619	+	+	+
CDH1	hsa-miR-544	+	+	+
CDH1	hsa-miR-339-5p	+	+	+
CDH1	hsa-miR-23b	+	+	+
CDH1	hsa-miR-23a	+	+	+
CDH1	hsa-miR-1207-5p	+	+	+
CDH1	hsa-miR-219-5p	+	+	+
CDH1	hsa-miR-338-3p	+	+	+

miRNA, microRNA.

Table II. Differential expression of miRNAs in NB between metastatic and primary tumors (increased expression, >2-fold).

miRNA	Fold changes (metastatic/primary tumors)	miRNA	Fold changes (metastatic/primary tumors)
hsa-miR-342-3p	6.2879	hsa-miR-335	2.5920
hsa-miR-92b	11.8088	hsa-miR-483-5p	3.2567
hsa-miR-130b	2.5771	hsa-miR-23a	2.5497
hsa-miR-129-3p	3.0205	hsa-miR-29a	2.4728
hsa-miR-483-3p	9.9583	hsa-miR-423-3p	2.0051
hsa-miR-345	3.1005	hsa-miR-128a	2.3610
hsa-miR-24	5.0395	hsa-miR-130a	3.1655
hsa-miR-486-5p	2.4739	hsa-miR-214	2.8887
hsa-miR-99b	3.2086	hsa-miR-15a	2.5875
hsa-miR-145	4.9442	hsa-miR-191	2.3789
hsa-miR-375	3.5245	hsa-miR-100	2.2647
hsa-miR-92a	5.3560	hsa-miR-10b	3.1618
hsa-miR-361-5p	4.0653	hsa-miR-29b	2.4477
hsa-miR-484	2.9674	hsa-miR-30c	2.5702
hsa-miR-339-5p	3.8349	hsa-miR-324-5p	2.2561
hsa-miR-148b	4.3914	hsa-miR-602	4.4085
hsa-miR-140-3p	3.7209	hsa-miR-107	2.1387

Taken from (16). miRNA, microRNA; NB, neuroblastoma.

shown that miR-23a expression levels were generally higher in metastatic NB tissues compared with primary NB tissues (Fig. 3A). Transwell migration (without Matrigel) and inva-

sion (with Matrigel) assays were used to evaluate the effect of miR-23a on NB cell migration and invasion. As Fig. 3C shows, blocking the expression of miR-23a with miR-23a ASO

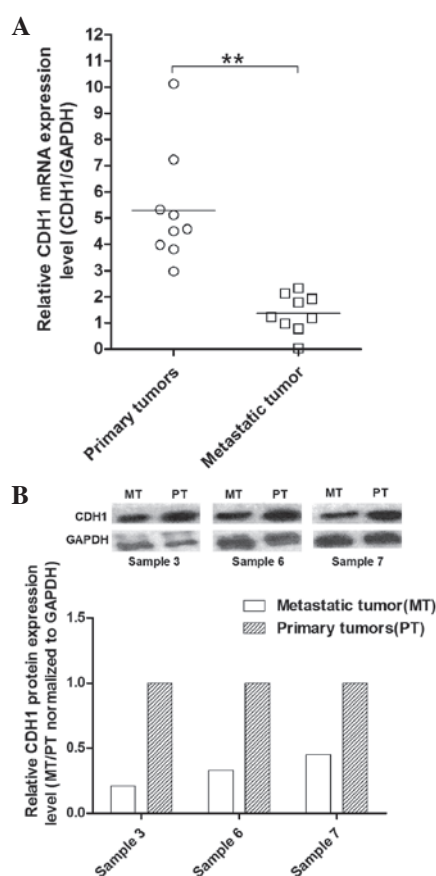


Figure 1. CDH1 is downregulated in metastatic NB tissues compared with primary NB tissues. The expression levels of CDH1 (A) mRNA and (B) protein were analyzed by real-time polymerase chain reaction and western blot analysis, respectively and were normalized to GAPDH in metastatic and primary NB tissues. The representative results of patients 3, 6 and 7 are shown. ** $P < 0.01$. NB, neuroblastoma.

significantly decreased the number of invaded SK-N-SH and GI-LA-N cells compared with the ASO control group. This implies that the upregulated miR-23a promotes the invasion of NB cells. Consistent with the invasion assay results, blockage of miR-23a significantly inhibited the migration of SK-N-SH and GI-LA-N cells compared with the control group (Fig. 3D). Thus, it was concluded that miR-23a is overexpressed in metastatic NB tissues and inhibition of miR-23a suppresses the migration and invasion of NB cells.

miR-23a directly targets CDH1 expression in NB cells. To verify that the regulation of CDH1 expression by miR-23a is direct, an EGFP reporter system was used in SK-N-SH cells. The alignment of miR-23a with the CDH1 3'UTR insert is illustrated in Fig. 4A. In addition, a mutated 3'UTR was constructed by deleting 4 nt in the seed sequence of miR-23a (asterisk marked in Fig. 4A). The EGFP reporter analysis showed that with the CDH1-3'UTR insert, the EGFP intensity was significantly lower than that in the CDH1-3'UTR-mutant group when treated with control oligo. When miR-23a was blocked by miR-23a ASO, the EGFP intensity levels were significantly higher than those in the control group. However, the EGFP intensity with the mutated 3'UTR was not affected by miR-23a (Fig. 4B). Next, it was investigated whether the endogenous CDH1 is regulated by miR-23a in NB cells. qPCR

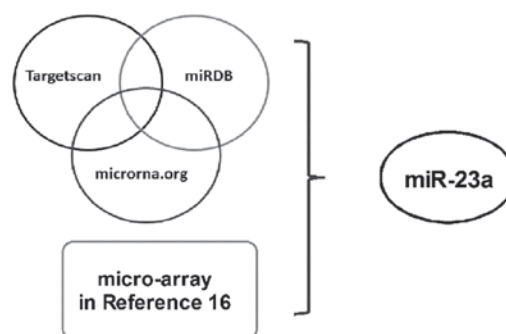


Figure 2. miR-23a is a candidate regulator of CDH1 in NB. miR-23a simultaneously appeared among the 20 miRNAs shown in Table I (miRNAs that may potentially target the 3' untranslated region of CDH1, predicted by TargetScan, microRNA.org and miRDB) and 34 miRNAs shown in Table II, which were upregulated in metastatic NB tissues compared with primary NB tissues. NB, neuroblastoma.

assay revealed that the CDH1 mRNA levels were inversely correlated with the miR-23a levels (Fig. 4C). The trend of CDH1 protein levels was similar to that of the mRNA levels, according to the western blot analysis (Fig. 4D). Therefore, it was concluded that CDH1 is a direct target of miR-23a in SK-N-SH cells, and that miR-23a regulates the expression of CDH1 at the mRNA and protein levels.

CDH1 suppresses NB cell migration and invasion in vitro. Previous studies have shown that the downregulation of CDH1 is an important feature of a number of transformed cells (19), and that CDH1 functions as a tumor suppressor. Accordingly, the present study investigated whether CDH1 affects NB cell metastasis. Overexpression plasmids (pCMV6/CDH1) were constructed that ectopically overexpress the CDH1 protein in NB cells. Subsequently, the effect of CDH1 on cell metastasis was evaluated by Transwell assay. The results showed that the invasion rate was reduced by ~50 and ~60% with the overexpression of CDH1 in SK-N-SH and GI-LA-N cells, respectively, compared with that of the control group (Fig. 5A). In addition, the migration rate of the SK-N-SH and GI-LA-N cells transfected with pCMV6/CDH1 was markedly lower than that of the control group (Fig. 5B). These results provided further evidence that CDH1 is a tumor suppressor in NB metastasis. The tumor suppressor role of CDH1 in NB metastasis may explain the manner in which the upregulation of miR-23a promotes NB cell migration and invasion and contribute to the transformation of primary to metastatic NB (Fig. 5C).

Discussion

As in the case of various types of cancer metastasis, the transformation of primary to metastatic NB is a complex multistep process including cell adhesion, migration, angiogenesis, immune escape and homing to target organs. The increased NB cell migration and invasion capabilities are essential features of the metastatic transformation process. Identifying the molecules and pathways that control NB cell migration and invasion is critical to understanding NB metastasis.

Accumulating previous evidence has suggested that genetic or epigenetic alterations in CDH1 or alterations in their protein

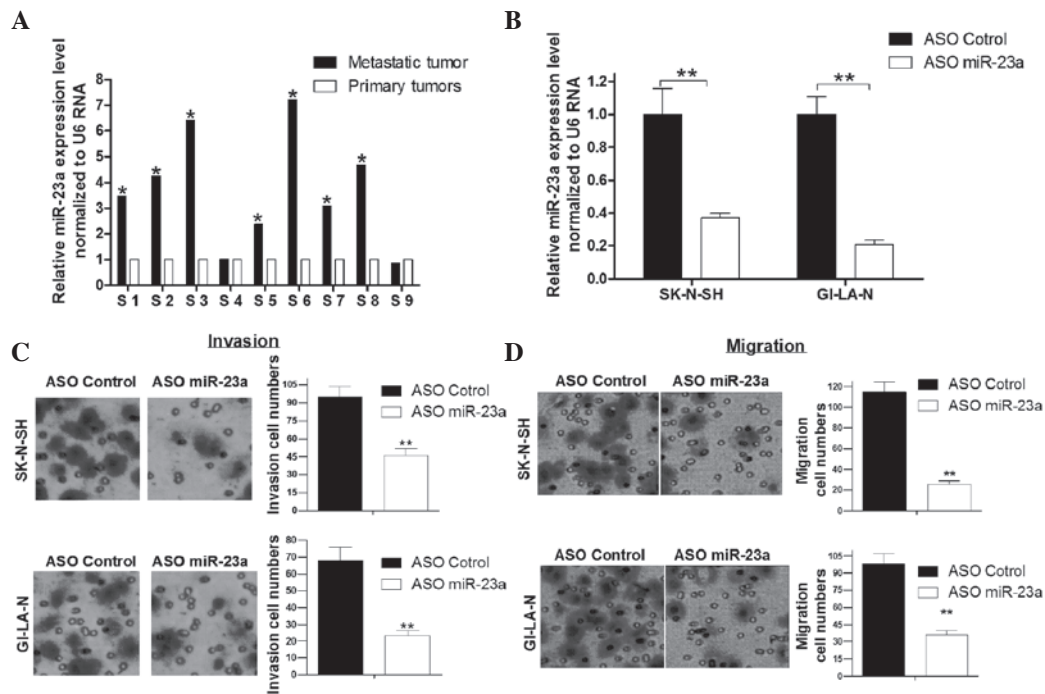


Figure 3. miR-23a is upregulated in metastatic NB tissues and suppresses the NB cell metastasis. (A) qPCR was performed to test the expression of miR-23a in the nine pairs of metastatic and primary NB tissues and was normalized to U6. (B) The SK-N-SH and GI-LA-N cells were transfected with ASO miR-23a, together with the controls. The miR-23a expression levels were measured by qPCR and normalized to U6. (C) Invasion and (D) migration assays were performed to detect the effects of miR-23a on the invasion and migration capabilities of SK-N-SH and GI-LA-N cells. The NB cells were seeded onto Transwell chambers for migration (without Matrigel) and invasion (with Matrigel). These assays were performed three times for each group; the number of migratory or invasive cells per field (graphs) and the invasive and migratory SK-N-SH and GI-LA-N cells stained with crystal violet (images) are presented. ** $P < 0.01$. NB, neuroblastoma; qPCR, real-time polymerase chain reaction; UTR, untranslated region; miRNA, microRNA.

expression, often result in tissue disorder, cellular dedifferentiation, increased invasiveness of tumor cells and ultimately metastasis (19-21). It has been previously demonstrated that the downregulation of CDH1 leads to loss of cell polarity and derangement of normal tissue architecture (22,23). In various types of cancer, CDH1-mediated cell-cell adhesion is lost concomitantly with acquisition of an invasive phenotype, high tumor grade and low patient survival rates (24-29). Based on the important role of CDH1 in cancer tumorigenesis, the investigation of the regulatory mechanisms of CDH1 in the process of primary to metastatic NB transition may highlight new prognostic markers and therapy targets for the metastasis of NB in the future. To date, somatic mutations, chromosomal deletions, proteolytic cleavage and silencing of the CDH1 promoter have been reported to cause the loss or reduction of CDH1 expression (30-32). In a previous study by Hoebeeck *et al*, the authors tested 42 primary NB tumors and the frequencies of methylation were 8% with CDH1 (19). This implies that there may be potential other mechanisms that are involved in the downregulation of CDH1 in the NB metastatic process. In the present study, nine pairs of metastatic NB tissues and matched primary NB tissues were tested, and the mRNA and protein levels of CDH1 were found to be significantly decreased in the NB metastatic transformation process (Fig. 1).

Previously, miRNAs have been identified as important regulators of gene expression. Several miRNAs have been found to exhibit prometastatic (miR-10b, -21 and -373/520c) or antimetastatic (miR-34b/c, -126, -148a, -206 and -335) activity (33-35). In a previous study, miRNA microarray

assay was performed to screen out the miRNAs that were differentially expressed between metastatic and primary NB tissues (16). In addition, a previous study has shown that miR-9, which is directly bound and upregulated by MYC and MYCN in breast cancer cells, directly targets CDH1, leading to increased cell motility and invasiveness (26). miR-10b has been found to modulate breast cancer metastasis by targeting CDH1 and may be a useful biomarker of advanced progression and metastasis of breast cancer (36). miR-25 has also been demonstrated to promote esophageal squamous cell carcinoma cell migration and invasion by directly targeting CDH1 (37). Recently, Cao *et al* (38) found that miR-23a regulates TGF- β -induced epithelial-mesenchymal transition by targeting E-cadherin (CDH1) in lung cancer cells. However, few studies have considered the manner in which the miRNAs contribute to the transformation of primary to metastatic NB through the CDH1-induced pathway. Therefore, the current study analyzed the microarray results (Table II) and miRNA target prediction algorithms (Table I). miR-23a was identified as a candidate molecule in the regulation of NB metastasis by the CDH1 pathway (Fig. 2).

miR-23a belongs to the miR-23a/24/27a cluster, which is located in chromosome 19p13.12 and may be induced by TGF- β . In several types of human cancer, this gene cluster functions as an oncogenic miRNA and has been previously reported to be upregulated in types of human cancer. As one of the most famous members of the miRNA-cluster, miR-23a has been shown to promote the growth of gastric adenocarcinoma cells by targeting the interleukin-6 receptor (39,40).

Acknowledgements

The current study was supported by the National Natural Science Foundation of China (grant nos. 81071037 and 81271395).

References

1. Maris JM, Hogarty MD, Bagatell R and Cohn SL: Neuroblastoma. *Lancet* 369: 2106-2120, 2007.
2. Maris JM: Recent advances in neuroblastoma. *N Engl J Med* 362: 2202-2211, 2010.
3. Steeg PS: Tumor metastasis: mechanistic insights and clinical challenges. *Nat Med* 12: 895-904, 2006.
4. Nevo I, Oberthuer A, Botzer E, *et al*: Gene-expression-based analysis of local and metastatic neuroblastoma variants reveals a set of genes associated with tumor progression in neuroblastoma patients. *Int J Cancer* 126: 1570-1581, 2010.
5. Gumbiner BM: Regulation of cadherin-mediated adhesion in morphogenesis. *Nat Rev Mol Cell Biol* 6: 622-634, 2005.
6. Ceteci F, Ceteci S, Karreman C, *et al*: Disruption of tumor cell adhesion promotes angiogenic switch and progression to micro-metastasis in RAF-driven murine lung cancer. *Cancer Cell* 12: 145-159, 2007.
7. Derksen PW, Liu X, Saridin F, *et al*: Somatic inactivation of E-cadherin and p53 in mice leads to metastatic lobular mammary carcinoma through induction of anoikis resistance and angiogenesis. *Cancer Cell* 10: 437-449, 2006.
8. Onder TT, Gupta PB, Mani SA, Yang J, Lander ES and Weinberg RA: Loss of E-cadherin promotes metastasis via multiple downstream transcriptional pathways. *Cancer Res* 68: 3645-3654, 2008.
9. Vleminckx K, Vakaet L, Jr., Mareel M, Fiers W and van Roy F: Genetic manipulation of E-cadherin expression by epithelial tumor cells reveals an invasion suppressor role. *Cell* 66: 107-119, 1991.
10. Frixen UH, Behrens J, Sachs M, *et al*: E-cadherin-mediated cell-cell adhesion prevents invasiveness of human carcinoma cells. *J Cell Biol* 113: 173-185, 1991.
11. Perl AK, Wilgenbus P, Dahl U, Semb H and Christofori G: A causal role for E-cadherin in the transition from adenoma to carcinoma. *Nature* 392: 190-193, 1998.
12. Bartel DP: MicroRNAs: genomics, biogenesis, mechanism, and function. *Cell* 116: 281-297, 2004.
13. Gabriely G, Wurdinger T, Kesari S, *et al*: MicroRNA 21 promotes glioma invasion by targeting matrix metalloproteinase regulators. *Mol Cell Biol* 28: 5369-5380, 2008.
14. O'Donnell KA, Wentzel EA, Zeller KI, Dang CV and Mendell JT: c-Myc-regulated microRNAs modulate E2F1 expression. *Nature* 435: 839-843, 2005.
15. Pillai RS, Bhattacharyya SN and Filipowicz W: Repression of protein synthesis by miRNAs: how many mechanisms? *Trends Cell Biol* 17: 118-126, 2007.
16. Guo J, Dong Q, Fang Z, *et al*: Identification of miRNAs that are associated with tumor metastasis in neuroblastoma. *Cancer Biol Ther* 9: 446-452, 2010.
17. Johnson SM, Grosshans H, Shingara J, *et al*: RAS is regulated by the let-7 microRNA family. *Cell* 120: 635-647, 2005.
18. Cimmino A, Calin GA, Fabbri M, *et al*: miR-15 and miR-16 induce apoptosis by targeting BCL2. *Proc Natl Acad Sci USA* 102: 13944-13949, 2005.
19. Hoebeek J, Michels E, Pattyn F, *et al*: Aberrant methylation of candidate tumor suppressor genes in neuroblastoma. *Cancer Lett* 273: 336-346, 2009.
20. Paredes J, Figueiredo J, Albergaria A, *et al*: Epithelial E- and P-cadherins: role and clinical significance in cancer. *Biochim Biophys Acta* 1826: 297-311, 2012.
21. Jiang T, Jiang H, Su XM, *et al*: Expressions of E-cadherin and alpha-catenin in benign, malignant and metastatic prostate tumors. *Zhonghua Nan Ke Xue* 18: 499-503, 2012 (In Chinese).
22. Larue L, Ohsugi M, Hirchenhain J and Kemler R: E-cadherin null mutant embryos fail to form a trophectoderm epithelium. *Proc Natl Acad Sci USA* 91: 8263-8267, 1994.
23. Karayiannakis AJ, Syrigos KN, Savva A, Polychronidis A, Karatzas G and Simopoulos C: Serum E-cadherin concentrations and their response during laparoscopic and open cholecystectomy. *Surg Endosc* 16: 1551-1554, 2002.
24. Alexiou D, Karayiannakis AJ, Syrigos KN, *et al*: Serum levels of E-selectin, ICAM-1 and VCAM-1 in colorectal cancer patients: correlations with clinicopathological features, patient survival and tumour surgery. *Eur J Cancer* 37: 2392-2397, 2001.
25. Cavallaro U and Christofori G: Cell adhesion and signalling by cadherins and Ig-CAMs in cancer. *Nat Rev Cancer* 4: 118-132, 2004.
26. Karatzas G, Karayiannakis AJ, Syrigos KN, *et al*: E-cadherin expression correlates with tumor differentiation in colorectal cancer. *Hepatogastroenterology* 46: 232-235, 1999.
27. Karayiannakis AJ, Syrigos KN, Chatzigianni E, *et al*: Aberrant E-cadherin expression associated with loss of differentiation and advanced stage in human pancreatic cancer. *Anticancer Res* 18: 4177-4180, 1998.
28. Karayiannakis AJ, Syrigos KN, Chatzigianni E, Papanikolaou S and Karatzas G: E-cadherin expression as a differentiation marker in gastric cancer. *Hepatogastroenterology* 45: 2437-2442, 1998.
29. Syrigos KN, Krausz T, Waxman J, *et al*: E-cadherin expression in bladder cancer using formalin-fixed, paraffin-embedded tissues: correlation with histopathological grade, tumour stage and survival. *Int J Cancer* 64: 367-370, 1995.
30. Bex G, Becker KF, Höfler H and van Roy F: Mutations of the human E-cadherin (CDH1) gene. *Hum Mutat* 12: 226-237, 1998.
31. Maretzky T, Reiss K, Ludwig A, *et al*: ADAM10 mediates E-cadherin shedding and regulates epithelial cell-cell adhesion, migration, and beta-catenin translocation. *Proc Natl Acad Sci USA* 102: 9182-9187, 2005.
32. Strathdee G: Epigenetic versus genetic alterations in the inactivation of E-cadherin. *Semin Cancer Biol* 12: 373-379, 2002.
33. Voorhoeve PM, le Sage C, Schrier M, *et al*: A genetic screen implicates miRNA-372 and miRNA-373 as oncogenes in testicular germ cell tumors. *Cell* 124: 1169-1181, 2006.
34. Zhu S, Si ML, Wu H and Mo YY: MicroRNA-21 targets the tumor suppressor gene tropomyosin 1 (TPM1). *J Biol Chem* 282: 14328-14336, 2007.
35. Ma L, Teruya-Feldstein J and Weinberg RA: Tumour invasion and metastasis initiated by microRNA-10b in breast cancer. *Nature* 449: 682-688, 2007.
36. Liu Y, Zhao J, Zhang PY, *et al*: MicroRNA-10b targets E-cadherin and modulates breast cancer metastasis. *Med Sci Monit* 18: BR299-308, 2012.
37. Xu X, Chen Z, Zhao X, *et al*: MicroRNA-25 promotes cell migration and invasion in esophageal squamous cell carcinoma. *Biochem Biophys Res Commun* 421: 640-645, 2012.
38. Cao M, Seike M, Soeno C, *et al*: MiR-23a regulates TGF- β -induced epithelial-mesenchymal transition by targeting E-cadherin in lung cancer cells. *Int J Oncol* 41: 869-875, 2012.
39. Zhu LH, Liu T, Tang H, *et al*: MicroRNA-23a promotes the growth of gastric adenocarcinoma cell line MGC803 and downregulates interleukin-6 receptor. *FEBS J* 277: 3726-3734, 2010.
40. Huang S, He X, Ding J, *et al*: Upregulation of miR-23a approximately 27a approximately 24 decreases transforming growth factor-beta-induced tumor-suppressive activities in human hepatocellular carcinoma cells. *Int J Cancer* 123: 972-978, 2008.
41. Jahid S, Sun J, Edwards RA, *et al*: miR-23a promotes the transition from indolent to invasive colorectal cancer. *Cancer Discov* 2: 540-553, 2012.
42. Wang Z, Wei W and Sarkar FH: miR-23a, a critical regulator of 'migR'ation and metastasis in colorectal cancer. *Cancer Discov* 2: 489-491, 2012.