Elevated microRNA-141-3p in placenta of non-diabetic macrosomia regulate trophoblast proliferation

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Abstract

Background: Several studies have reported microRNAs (miRNAs) could regulate the placental development, though the role and mechanism of miRNAs in the development of non-diabetic macrosomia (NDFMS) remains unclear.

Methods: To identify the aberrantly expressed key miRNAs in placenta of NDFMS, we employed a strategy consisting of initial screening with miRNA microarray and further validation with quantitative RT-PCR assay (qRT-PCR). In vitro cellular model and a mouse pregnancy model were used to delineate the functional effects of key miRNA on proliferation, invasion, and migration.

Findings: miR-141-3p was identified as the key miRNA with expression level significantly higher in placentas of NDFMS compared with those from normal controls. Overexpressed miR-141-3p in HTR-8/SVneo cells contributed to increased cell proliferation, invasion, and migration. miR-141-3p inhibition in HTR-8/SVneo cells resulted in decreased cell proliferation and invasion. Significantly increased infant birth weight was observed in late pregnancy of C57BL/6j mice treated with miR-141-3p agomir. However, no significant difference was found in early pregnancy of C57BL/6j mice treated with miR-141-3p agomir.

Interpretation: miR-141-3p could stimulate placental cell proliferation to participate in the occurrence and development of NDFMS.

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1. Introduction

Macrosomia is a common perinatal complication of pregnancy defined as infant’s birth weight of 4000 g or greater at term. There has been an increased incidence of macrosomia in recent decades. Though various interventions for the management of diabetic macrosomia, there is no clear consensus on antepartum prediction and management of non-diabetic macrosomia (NDFMS) and its accurate diagnosis is only made retrospectively [1]. It is known that fetal macrosomia increases risk of adverse outcomes in mother and long-term health problems in the infant [2,3]. However, the potential causes and mechanisms of NDFMS need further studies.

Neonatal birth weight is associated with a variety of factors, including genetic factors, gestational nutrition, endocrine, and placental function, etc. [4–6]. The placenta acts as a bridge between mother and fetus, and can also protect the fetus from the maternal immune system attack and secrete pregnancy-related hormones and growth factors [7]. Therefore, placental development and abnormal function may have an important impact on the growth and development of the fetus [8–10].
Research in context

Evidence before this study

Fetal macrosomia is associated with an increased risk of adverse outcomes for mother and long-term health problems of the infant. Previous studies confirmed a variety of miRNAs with abnormal expression in placental tissues are related to fetal growth and development. However, miRNAs expression pattern in placenta of non-diabetic macrosomia (NDFMS) and the underlying mechanism is still unclear.

Added value of this study

miR-141-3p was identified as the key miRNA, which expression level was significantly higher in placentas of NDFMS compared with those from normal controls. Overexpressed miR-141-3p in HTR-8/SVneo cells resulted in increased cell proliferation, invasion, and migration. miR-141-3p inhibition in HTR-8/SVneo cells resulted in decreased cell proliferation and invasion. Significantly increased of infant birth weight was observed in late pregnancy of C57BL/6J mice treated with miR-141-3p agomir. miR-141-3p could stimulate placental cell proliferation to participate in the occurrence and development of NDFMS.

Implication of all the available evidence

These findings reveal an important role of miR-141-3p in development of NDFMS, which may provide new therapeutic approaches for NDFMS.

Previous studies reported that genes dysregulation in placentas played an important role in the pathology of fetal growth restriction and macrosomia. Sabri et al. found 338 and 41 genes were significantly dysregulated in the growth-restricted and macrosomic placentas through gene expression microarray analysis [11]. Yuan et al. reported that Lep-regulated in the growth-restricted and macrosomic placentas through macrosomia. Sabri et al. found 338 and 41 genes were significantly dysregulated in the growth-restricted and macrosomic placentas through gene expression microarray analysis [11]. Yuan et al. reported that Lep-regulated in the growth-restricted and macrosomic placentas through macrosomia. Sabri et al. found 338 and 41 genes were significantly dysregulated in the growth-restricted and macrosomic placentas through gene expression microarray analysis [11]. Yuan et al. reported that Lep-regulated in the growth-restricted and macrosomic placentas through macrosomia. Sabri et al. found 338 and 41 genes were significantly dysregulated in the growth-restricted and macrosomic placentas through gene expression microarray analysis [11]. Yuan et al. reported that Lep-regulated in the growth-restricted and macrosomic placentas through macrosomia. Sabri et al. found 338 and 41 genes were significantly dysregulated in the growth-restricted and macrosomic placentas through gene expression microarray analysis [11]. Yuan et al. reported that Lep-regulated in the growth-restricted and macrosomic placentas through macrosomia. Sabri et al. found 338 and 41 genes were significantly dysregulated in the growth-restricted and macrosomic placentas through gene expression microarray analysis [11]. Yuan et al. reported that Lep-regulated in the growth-restricted and macrosomic placentas through macrosomia. Sabri et al. found 338 and 41 genes were significantly dysregulated in the growth-restricted and macrosomic placentas through gene expression microarray analysis [11]. Yuan et al. reported that Lep-regulated in the growth-restricted and macrosomic placentas through macrosomia. Sabri et al. found 338 and 41 genes were significantly dysregulated in the growth-restricted and macrosomic placentas through gene expression microarray analysis [11]. Yuan et al. reported that Lep-regulated in the growth-restricted and macrosomic placentas through macrosomia. Sabri et al. found 338 and 41 genes were significantly dysregulated in the growth-restricted and macrosomic placentas through gene expression microarray analysis [11]. Yuan et al. reported that Lep-regulated in the growth-restricted and macrosomic placentas through macrosomia. Sabri et al. found 338 and 41 genes were significantly dysregulated in the growth-restricted and macrosomic placentas through gene expression microarray analysis [11]. Yuan et al. reported that Lep-regulated in the growth-restricted and macrosomic placentas through macrosomia. Sabri et al. found 338 and 41 genes were significantly dysregulated in the growth-restricted and macrosomic placentas through gene expression microarray analysis [11].

2. Materials and methods

2.1. Human material samples and ethics

A total of 91 participants (44 non-diabetes macrosomia pregnant women and 47 normal controls) who delivered in Changzhou Maternal and Child Health Hospital between September 2014 to June 2015 were included. All participants in our study were primigravida. Inclusion criteria of NDFMS: 1. Full-term birth (≥37 weeks and < 42 weeks), birth weight ≥ 4000 g; 2. Normal blood glucose and oral glucose tolerance test (OGTT) result, negative urine sugar; 3. Without pregnant complications (gestational hypertension, placental abruption, placenta previa, intrahepatic cholestasis of pregnancy, infectious diseases, and other pregnancy complications). Inclusion criteria of control: 1. Full-term birth (≥37 weeks and < 42 weeks), birth weight < 4000 g and ≥ 2500 g; 2. Normal blood glucose and oral glucose tolerance test (OGTT) result, negative urine sugar; 3. Without pregnant complications (gestational hypertension, placental abruption, placenta previa, intrahepatic cholestasis of pregnancy, infectious diseases, and other pregnancy complications).

After removal of the fetal membranes, three pieces of the placental tissues with 1 × 1 × 1 cm³ size were randomly taken from the center of the maternal side. Then, these tissues were washed using sterile saline and immediately frozen in liquid nitrogen, and then stored at −80 °C. The study was approved by the Ethics Committee of Nanjing Medical University, and all pregnant women signed the informed consent before participation.

2.2. RNA extraction, miRNA microarray analysis, and quantitative real-time PCR (qRT-PCR)

Total RNA from placental tissues and cultured cells were extracted using TRIzol reagent (Invitrogen, Carlsbad, CA, USA) according to the manufacturer’s protocol. About 50 mg placental tissues were kept in TRIzol for RNA extraction. Ultrasonic cell crusher was used for tissue homogenate (working for 3 s, pause for 3 s with 70% energy about three minutes). Then, total RNA was extracted according to the manufacturer’s instructions. To prevent RNA degradation, all steps were performed on ice. The concentration and purity of RNA were measured using NanoDrop® ND-1000 (samples with A260/A280 ratio > 1.8 and < 2.0 were stored at −80 °C until use), while its integrity was assessed using electrophoresis with 1-5% denaturing agarose gels. RNA was reverse transcribed using the specific Megaplex RT primers (Life Technologies). Thereafter, we screened the expression level of 754 miRNAs using the TaqMan® Array Human MicroRNA A + B Cards Set v3.0 (Life Technologies) on a 7900 Real-Time PCR System (Applied Biosystems, Foster City, CA, USA). Data were analyzed by DataAssist v3.0 (Life Technologies). The amount of target miRNAs was normalized relative to the amount of RNU6B (U6) (ΔCt = Ct_sample – Ct_blank).

For qRT-PCR, PrimeScript RT reagent Kit (Takara, Tokyo, Japan) was used to reverse transcribe RNA sample to cDNA (Takara RR037A), and PCR analysis was performed using SYBR® Premix Ex Taq™ according to the manufacturer’s instructions. The quantity of specific RNA for target genes in each sample was estimated by qRT-PCR using ABI Prism 7900HHT (Applied Biosystems, Foster City, CA, USA). The expression of each miRNA relative to U6 was calculated using the 2^–ΔΔCt method. All the reactions were run in triplicate to control for PCR variation. The primer sequences for qRT-PCR are shown in Table S1.

2.3. Cell culture and transfection

The human placenta trophoblast cell line HTR-8/SVneo was cultured in RPMI 1640 medium containing 10% fetal bovine serum and 10% double antibiotic (100 U/ml penicillin and 100 U/ml streptomycin) at 37 °C in a 5% CO₂ cell incubator. When the cells were in the logarithmic growth phase, they were transfected with miR-141-3p mimic, inhibitor or negative control (GenePharma, Shanghai, China) using transfection reagent Lipofectamine 2000 reagent (Invitrogen, CA, USA) according to the manufacturer’s instructions (5 µl Lipofectamine 2000 reagent per 6-well plate).

2.4. Cell proliferation assay

HTR-8/SVneo cells were seeded in 96-well plates at a density of 1 × 10³ cells/well. After transfection for 24 h and 48 h, cell proliferation ability was measured by using the Cell Counting Kit-8 (CCK-8; Dojindo, Kumamoto, Japan). Using culture medium as the blank control, we measured the absorbance at a wavelength of 450 nm by TECAN infinite microplate reader.
2.5. Cell cycle analysis

We collected the treated cells seeded in 6-well plates, washed with cold PBS, fixed in cold 70% ethanol overnight at −20 °C. Propidium iodide (PI) (Sigma, MO, USA) was used to stain the fixed cells for 30 min at room temperature under dark condition. Cell cycle distribution of stained cells was analyzed with FACS Calibur Flow Cytometer afterward, and the fraction of the cell cycle was measured by ModFit LT version 3.0 software (Verity Software House, Toshan, ME, USA).

2.6. Apoptosis assay

We harvested the treated cells seeded in 6-well plates with a density of \(1 \times 10^5\) cells/well, washed twice with PBS, and cells were processed using Annexin V-FITC Kit (BD Pharmingen, NJ, USA) in the dark for 15 min at room temperature. The cells were analyzed using FACS Calibur Flow Cytometer immediately after incubated with Annexin V-FITC.

2.7. Cell invasion analysis

The treated cells were transferred to the transwell chamber with 100 μl empty RPMI 1640 medium, while the 24-well plate contained 600 μl complete medium as the container of the transwell chamber. After 24 h, the transwell chamber was transferred to 95% methanol to fix cells for 30 min, then stained by crystal violet for 30 min, and washed twice with PBS. After drying, we captured the pictures by light microscope. Then, by direct cell counting, we used the cell number in vision as a measure.

2.8. Cell migration analysis

We used 10 μl pipette tip to draw a straight line in each hole of 6-well plate with treated cells and washed twice with PBS. Next, the BioSation IM-Q live cell workstation was used to record the migration situation of cells to the middle real-timely and capture dynamic change graphs. Then we used Image-Pro Plus software to analyze the pictures, which take the horizontal distance of cells migrated as the quantitative indicator.

2.9. C57BL/6j mice feeding and experiment

We used miR-141-3p agomir (microON™ miR-141-3p agomir) (RiboBio, Guangzhou, China) to construct miR-141-3p overexpression model in placental tissues via the tail vein, and used microON™ agomir negative control treated controls. Eight-week-old pregnant C57BL/6j mice were divided into two groups: early pregnant group (treated: control = 5: 5) and late pregnant group (treated: control = 7: 5). Early pregnant group mice were treated twice at pregnant day 5 and day 8, while late pregnant group mice were treated twice at pregnant day 14 and 17, and they were sacrificed afterward at pregnant day 18. Fetal weight, body height, placental weight, and placental diameter were measured. Besides, we treated some non-pregnant mice twice with 3-day interval (number of treated: control = 4: 5), followed by sacrificed at 10 days after the second treatment, and the weight fluctuation was observed (The results of effect on body weight and food intake were shown in Fig. S1). This study was carried out strictly in accordance with the international standards on animal welfare and the guidelines of the Institute for Laboratory Animal Research of Nanjing Medical University.

2.10. Statistical analysis

Data were analyzed by SPSS 18.0 (SPSS Inc., Chicago, IL, USA) and presented with GraphPad Prism 5.01 (GraphPad Software Inc., San Diego, CA, USA). If data were in normal distribution, the student’s t-test was used to compare two groups. Otherwise, the Mann-Whitney U-test was used. All results are expressed as mean ± standard error (SE) without special instructions. \(P < 0.05\) was considered statistically significant. \(P < 0.05: *, P < 0.01: **, P < 0.001: ***\).

3. Results

3.1. Clinical data

Clinical characteristics of the study population are summarized in Table 1. NDFMS and normal controls were matched by maternal age, pre-pregnant body mass index, gestational weeks and infant gender distribution. The birth weight of neonates and weight gain during pregnancy were significantly higher in NDFMS compared with that in normal controls (\(P < 0.05\)).

3.2. Expression pattern of miRNAs in placentas of NDFMS

We randomly selected 18 cases of NDFMS and 18 normal controls, and pooled RNA samples of the 18 cases and 18 controls, respectively and subjected them to miRNA microarray screening. A total of 754 miRNAs were detected by miRNA microarray, and 264 and 318 miRNAs could be detected in placental tissues of normal controls and NDFMS, respectively (\(Ct < 30\)). One hundred and thirty-three miRNAs showed dysregulated in NDFMS placentas (fold change ≥2), including 63 upregulated and 70 down-regulated (Table S2). There were 21 miRNAs that expressed four times higher in NDFMS group (miR-523-3p, miR-519a-3p, miR-518f-3p, miR-331-3p, miR-204-5p, miR-106b-5p, miR-574-3p, miR-518a-3p, miR-370-3p, miR-518c-3p, miR-205-5p, miR-141-3p, miR-126-3p, miR-424-5p, miR-194-5p, miR-411-5p, miR-376c-3p, miR-200b-3p, miR-374-5p, miR-200c-3p, and miR-199a-3p), and 11 miRNAs expressed eight times lower compared with normal controls (miR-1290, miR-1227-3p, miR-660-5p, miR-1275, miR-320b, miR-584-5p, miR-34b-3p, miR-656-3p, miR-15a-5p, miR-1291, and miR-1283).

3.3. qRT-PCR validation of miRNAs in placentas

Considering the bioinformatic analysis results, species conservativity and feasibility, we selected twelve miRNAs (miR-331-3p, miR-204-5p, miR-106b-5p, miR-370-3p, miR-205-5p, miR-141-3p, miR-126-3p, miR-424-5p, miR-194-5p, miR-411-5p, miR-376c-3p, miR-200b-3p, miR-374-5p, miR-200c-3p, and miR-199a-3p) listed in Table S3 for further validation using qRT-PCR in the remaining placental samples (26 of NDFMS, and 29 of normal controls). The twelve miRNAs listed in Table S3 are highly conservative and homologous between human and mouse, thus which have a strong practical significance in the following in vitro and in vivo experiments. The results showed that the expression levels of miR-331-3p, miR-106b-5p, miR-370-3p, miR-205-5p, miR-141-3p, miR-126-3p, miR-424-5p, miR-194-5p, miR-411-5p, and miR-1283 (Table S3) for further validation using qRT-PCR in the remaining placental samples (26 of NDFMS, and 29 of normal controls). The twelve miRNAs listed in Table S3 are highly conservative and homologous between human and mouse, thus which have a strong practical significance in the following in vitro and in vivo experiments. The results showed that the expression levels of miR-331-3p, miR-106b-5p, miR-370-3p, miR-205-5p, miR-141-3p, miR-126-3p, miR-424-5p, miR-194-5p, miR-411-5p, and miR-1283 were statistically different between NDFMS and normal controls. However, only miR-141-3p was significant with false discovery rate (FDR) adjustment (Table S4, Fig. 1). Additionally, the infant birth weight gradually increased along with the elevated expression of miR-141-3p (\(P = 0.019, r^2 = 0.101\)) (Fig. S2B). Therefore, miR-141-3p was determined as the key miRNA for further functional analyses (Fig. 2A).

3.4. Effects of miR-141-3p on proliferation, apoptosis, invasion, and migration in HTR-8/SVneo

Cell proliferation, cell cycle, apoptosis, invasion, and migration assays were performed with HTR-8/SVneo cells transfected with miR-
**Table 1**
Clinical characteristics of the study population.

| Characteristic                        | Total (n = 91) | Control (n = 47) | NDFMS (n = 44) | P-Value |
|---------------------------------------|----------------|-----------------|----------------|---------|
| Maternal age (years)                  | 26·42 ± 2·87   | 26·15 ± 0·36    | 26·70 ± 0·48   | NS      |
| Gestational age (weeks)               | 39·71 ± 1·01   | 39·70 ± 0·15    | 39·72 ± 0·15   | NS      |
| BMI before pregnancy (kg/m²)          | 20·71 ± 3·32   | 20·05 ± 0·33    | 21·43 ± 0·62   | NS      |
| Placental weight (g)                  | 672·36 ± 15·03 | 587·77 ± 14·66  | 762·73 ± 19·08 | < 0·001 |
| Weight gain during pregnancy (kg)     | 18·21 ± 4·77   | 16·60 ± 0·59    | 19·97 ± 0·75   | < 0·001 |
| Birth weight (g)                      | 3764·68 ± 58·63| 3305·57 ± 47·01| 4250·82 ± 41·41| < 0·001 |
| Infant gender, n (%)                  |                |                 |                | NS      |
| Male                                  | 43 (47)        | 23 (49)         | 20 (45)        |         |
| Female                                | 48 (53)        | 24 (51)         | 24 (55)        |         |

Values are mean ± SD. BMI, body mass index. NS, not significant.

**Fig. 1.** Heat map of miRNA expression levels in placental tissue screened out by miRNA microarray in NDFMS compared with controls.

**Fig. 2.** The expression level of miR-141-3p in placental tissues and the correlation with infant birth weight. (A) The expression level of miR-141-3p in NDFMS placental tissues compared with normal controls. (B) The correlation between the expression level of miR-141-3p and infant birth weight.
141-3p mimic or inhibitor. Cell proliferation was significantly increased in HTR-8/SVneo cells transfected with miR-141-3p mimic and significantly decreased in HTR-8/SVneo cells transfected with miR-141-3p inhibitor at 48 h (Fig. 3A), whereas no significant difference was found at 24 h (Fig. 3A). Additionally, no significant differences of cell apoptosis and cell cycle were found in miR-141-3p mimic-treated or inhibitor-treated group compared with negative controls (Fig. 3B, C).

Overexpressed miR-141-3p in HTR-8/SVneo cells resulted in increased cell invasion and migration (Fig. 4A, B), miR-141-3p inhibition in HTR-8/SVneo cells resulted in decreased cell invasion (Fig. 4A). However, no significant difference of cell migration was found in HTR-8/SVneo cells treated with miR-141-3p inhibitor (Fig. 4B).

3.5. Roles of miR-141-3p in birth weight of C57BL/6J mice

To confirm the role of miR-141-3p in the development of NDFMS, we constructed miR-141-3p overexpression mouse pregnancy models at early pregnancy and at late pregnancy (Fig. S2). There was no significant difference of birth weight between the treated group and control group in early pregnancy (Fig. 5A, B, C, D), while the birth weight of mice in the treated group was significantly higher than that of controls in late pregnancy (P = 0.006) (Fig. 5E, F, G, H). In addition, no statistically significant difference of placental weight, placental diameter, and body length of offspring was observed between the treated group and control group either in early pregnancy or in late pregnancy (Fig. 5).

4. Discussion

From the miRNA microarray data, we found many miRNAs were dysregulated in placenta of NDFMS (Table S2). miR-17, miR-19a, miR-20a were up-regulated, which is consistent with the finding of Li et al. in NDFMS [18]. In contrast to previous studies in macrosomia [15,17], we found miR-483-3p and miR-21 were down-regulated in microarray screening results. The inconsistent results may be attributed to the differences of sample size and detection methods. Besides, these miRNAs were only examined in the microarray with no further verification in large sample size. Among all the miRNAs expression, we found only the miR-141-3p expression level was significantly increased in NDFMS placental tissues, suggesting aberrant expression of miR-141-3p may be related to the occurrence of NDFMS. miR-141-3p is a member of the miR-200 family, which is highly conservative in vertebrates [19]. The study of miR-200 family in rodent animal model is common while its regulating in human fetal growth and development of population-based is rare [20–22]. Yao et al. [22] found miR-200 family expression in the uterus and offspring (F2 generation) brain tissues was altered when the parental generations (F0) exposed to stress during pregnancy. Chim et al. [23] detected known 157 miRNAs by miRNA microarray and found 17 miRNAs were 10 times higher in placental tissues than in maternal blood, which could not be detected in maternal blood after delivery. miR-141, miR-149, miR-299-5p, and miR-135b were the four most abundant miRNAs in the placenta, of which the expression level of miR-141 in maternal blood significantly increased in the third trimester [23]. Prior investigations have documented miR-141-3p participates in the process of cell proliferation, especially dysregulated in a variety of cancers, including exerting a growth-promoted activity in ovarian cancer [24], cervical cancer [25], and a growth-suppressive activity in hepatocellular carcinoma [26]. Similarly, it also showed different effects on fetal growth. Tang et al. found miR-141 up-regulated in fetal growth restriction (FGR) by down-regulating its target genes [27]. Baker et al. reported low maternal folate status is associated with higher incidence of small for gestation age (SGA) infants, with elevated level of miR-141-3p in placental tissues [28]. Interestingly, they also found that trophoblast proliferation was increased in low maternal folate status, to some extent that is consistent with our results i.e. miR-141-3p elevates trophoblast proliferation. It is possible that the elevated miR-141-3p expression is a compensatory increase in the placenta of SGA and FGR. But the specific in-depth mechanism demands further research to clarify in the future.

The cell experiments suggest miR-141-3p may participate in the development of NDFMS by affecting the proliferation and invasion of trophoblast cells. Ospina-Prieto et al. [29] suggested the expression level of miR-141 in HTR-8/SVneo cells was significantly lower than that in normal and preeclampsia placenta. The expression level of miR-141 in normal placental tissues was 42,394 times higher than that in HTR-8/SVneo.
cells. Morales-Prieto et al. [30] also found the expression levels of miR-141 in all placental lineage cells were low, suggesting that the basal level of miR-141 in human chorionic trophoblast cells was very low, which may explain why the proliferation of HTR-8/SVneo cells treated by miR-141-3p inhibitor did not decrease as expected. Moreover, further studies are still needed to confirm this conclusion.

Because miR-141-3p affected proliferation and invasion ability of HTR-8/SVneo cells, we constructed the miR-141-3p overexpression model of the placenta in early pregnancy and late pregnancy of C57BL/6J mice to verify the role of miR-141-3p in the development of NDFMS. The miR-141-3p agomir reagent from Guangzhou RiboBio can be maintained for about three days, so as to ensure the treatment at early pregnancy will not be maintained until late pregnancy, which ensured the accuracy of modeling. Interestingly, we found that the fetal birth weight of the treated group was significantly higher only in the late pregnant group. Therefore, we conclude that miR-141-3p may affect the proliferation at late pregnancy by participating in the

Fig. 4. Effects of miR-141-3p on invasion, and migration of HTR-8/SVneo cells. (A) The invasion ability of HTR-8/SVneo transfected with miR-141-3p mimic or inhibitor detected by crystal violet staining in transwell. (B) The migration ability of HTR-8/SVneo transfected with miR-141-3p mimic or inhibitor detected by BioSation IM-Q live cell workstation.

Fig. 5. The effects of miR-141-3p treated at early pregnancy and late pregnancy on offspring and placenta of pregnant C57BL/6j mice. (A) Birth weight of offspring at early pregnancy. (B) Body length of offspring at early pregnancy. (C) Placental weight at early pregnancy. (D) Placental diameter at early pregnancy. (E) Birth weight of offspring at late pregnancy. (F) Body length of offspring at late pregnancy. (G) Placental weight at late pregnancy. (H) Placental diameter at late pregnancy.
occurrence of NDFMS, rather than by influencing the invasion at early pregnancy. Previous researches have demonstrated the role of miR-141 in fetal growth [29,31], Ospina-Prieto et al. [29] found miR-141 was highly expressed in placentas of preeclampsia, and overexpression of miR-141 enhanced the invasion ability of HTR-8/SVneo cells, which is consistent with their findings. They did not find any significant changes of cell proliferation ability in HTR-8/SVneo cells after 24 h of transfection with either miR-141-mimic or miR-141-inhibitor. This result is also consistent with our findings that no significant difference of cell proliferation was found in HTR-8/SVneo cells after 24 h of transfection. However, we found cell proliferation was significantly higher in the HTR-8/SVneo cells transfected with miR-141-3p mimic at 48 h. Morales-Prieto et al. [32] demonstrated that the reduction of miR-141 caused by leukemia inhibitory factors inhibited the proliferation of choriocarcinoma cell line JEG-3. In addition, the study team also found miR-141 expression level in placental tissues at the late pregnancy was higher than the early pregnancy [32]. This may be due to placenta grows fastest along with the growth of fetus in late pregnancy, which also suggested miR-141 in placenta is related to proliferation function and confirmed the rationality of our results.

A limitation of this study is the lack of follow-up mechanism of how miR-141-3p promotes the proliferation of trophoblasts. It is confirmed that miR-141-3p has many target genes [24,33] and miRNA can regulate target genes via partial complementary binding to the 3′-UTR of mRNAs [34]. For example, miR-141-3p targets FOXA2 to foster the growth, invasion, and tumorigenesis of cervical cancer cells [25]. Another study showed miR-141 targeting PLAG1 to participate FGR [27]. We speculated that there are one or more specific target genes of miR-141-3p to regulate the proliferation of trophoblasts in NDFMS and follow-up experiments are required to confirm. Besides, it is well known that there was excess nutrient transfer to the baby through placenta in macrosomia, therefore effect of miR-141-3p on nutrient transporters can’t be ruled out. Future studies are required to reveal the specific mechanism of miR-141-3p participating in the occurrence of NDFMS.

In conclusion, aberrant high expression level of miR-141-3p may contribute to the pathogenesis of NDFMS through regulating trophoblast proliferation in late pregnancy as shown in Fig. 6. Our study provided a foundation for the explanation of miRNA changes related to NDFMS and expanded the current understanding of its pathogenesis. Future studies are needed to illustrate the exact molecular mechanism of miR-141-3p in NDFMS.

Fig. 6. A model was presented for miR-141-3p mediated NDFMS. Up-regulation of miR-141-3p in placenta contributes to the pathogenesis of NDFMS through regulating trophoblast proliferation in late pregnancy.
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