Abstract. Deep vein thrombosis (DVT) is a disease involving multiple genes and systems. MicroRNAs (miRNAs) represent a class of non-coding small RNAs that post-transcriptionally suppress their target genes. The expression patterns of miRNA and messenger RNA (mRNA) in DVT remain poorly characterized. The aim of the present study was to evaluate miRNA and mRNA expression profiles in a stasis-induced DVT rat model. Male SD rats were randomly divided into three groups as follows: DVT, sham and control. The inferior vena cava (IVC) of rats was ligated to construct stasis-induced DVT models. Rats were sacrificed three days after ligation, and morphological changes in the vein tissues were observed by hematoxylin and eosin and Masson staining. The miRNA and mRNA expression profiles were evaluated by microarrays, followed by bioinformatics analysis. The microarray analysis identified 22 miRNAs and 487 mRNAs that were significantly differentially expressed between the experimental and control groups, and between the experimental and sham groups, but not between the control and sham groups (P≤0.05; ≥2.0 -fold change). By subsequent bioinformatics analysis, a 19 miRNA-98 mRNAs network was constructed in the stasis-induced DVT rat model. Notably, the majority of these miRNAs and mRNAs are reported to be expressed by endothelial cells (ECs) and are associated with the function of ECs. The results provide evidence indicating that the regulatory association of miRNA and mRNA points to key roles played by ECs in thrombosis. These findings advance our understanding of the molecular regulatory mechanisms underlying the pathophysiology of DVT.

Introduction

Deep vein thrombosis (DVT) refers to the formation of a blood clot within a deep vein, predominantly in the legs. DVT and pulmonary embolism (PE) constitute a single disease process, termed venous thromboembolism (VTE), which is the third most common vascular disease in the United States (1). As the disease has an insidious onset and there is no obvious clinical symptom or sign in the early stages, the rate of misdiagnosis is high. When timely diagnosis and effective therapy are missed, DVT may lead to the abnormal swelling and ulceration of lower limbs, post-thrombotic syndrome and PE (1,2). The incidence of VTE is >1,000 per year in the United States (3). Thus, DVT is a major cause of mortality and leads to significant morbidity. In forensic practice, PE is a major cause of sudden death and is attributable primarily to DVT (4). Studies of DVT improve the understanding of the disease, as well as increase the successful rescue rate and detection rate, which is significant in clinical diagnosis treatment and forensic identification.

In recent years, with the deepening of theoretical research into thrombosis, DVT has been recognized as a disease that involves multiple factors and systems (5). Due to the complexity of the disease, traditional Northern blotting and quantitative polymerase chain reaction methods have been unable to fully elucidate its mechanisms. To develop a more comprehensive understanding of DVT, high-throughput microRNA (miRNA) and messenger RNA (mRNA) microarray technology were used in the present study.

miRNAs represent a class of non-coding small RNAs that post-transcriptionally suppress their target genes (6). There is increasing evidence that miRNA expression patterns change in many vascular diseases (7-10). Although DVT has been investigated extensively, the molecular mechanisms underlying the pathophysiological changes remain to be defined. Furthermore, information on changes in miRNA expression within the vessel tissues is limited and, to date, to the best of our knowledge, there are no studies describing miRNA-mRNA interactions in DVT.

In the present study, miRNA and mRNA expression in vessel tissue samples from rat DVT models were assessed by microarray. Furthermore, bioinformatics analyses were used to build and analyze the miRNA-mRNA network. The present findings provide systematic and comprehensive insights
into the molecular mechanisms of DVT. The study design is presented in Fig. 1.

Materials and methods

Animal model of venous thrombosis. All experiments were reviewed and approved by the ethics committee of the Institute of Laboratory Animal Science of Shanxi Medical University (Taiyuan, China).

Adult male Sprague Dawley (SD) rats (n=36; Shanxi Laboratory Animal Center, Taiyuan, China) 8-10 weeks of age and weighing 280-300 g were used in the present study. The rats were divided into three groups as follows: DVT, sham and control (n=12). The rats were anesthetized by 10% chloral hydrate. A midline laparotomy was performed. The inferior vena was explored by moving the small intestine out of the way, and all side branches were ligated. IVC was ligated just below the left renal vein. A microvascular clamp was attached to the confluence of iliac veins for 15 min. The skin was sutured by 3-0 Prolene suture and penicillin powder (Sigma, Washington, DC, USA) covering the incision evenly was used. The sham-surgery rats received anesthesia and all surgical procedures, but without IVC ligation or clamping. The control group received no treatment.

Tissue harvesting. The rats were sacrificed at day 3 after ligation. The IVC with thrombus was carefully harvested. One part of the tissue was fixed in 10% formalin solution for histological analysis and the rest was stored in RNAsafety (Shanghai Biotechnology Corp., Shanghai, China) for microarray analyses.

Histological analysis. For histological examinations, the IVC tissue was fixed with 4% paraformaldehyde for 48 h and embedded in paraffin wax. The tissues were then cut into 4-μm-thick sections and the sections were dewaxed in xylene twice at 37°C for 15 min each time, rehydrated through decreasing concentrations of ethanol, and washed in distilled water at room temperature for 5 min each time. They were then finally stained with hematoxylin and eosin (H&E) or Masson's trichrome, both following the company's instructions (Nanjing Jiancheng Bioengineering Institute, Nanjing, China). For H&E staining, the slides were dipped into the jar containing hematoxylin for 5 min and with eosin for 30 sec with agitation at room temperature. For Masson staining, slides were stained at room temperature with nuclear staining solution for 1 min, with cytoplasmic stain solution for 45 sec, then washed with phosphomolybic acid for 6 min and last counterstained for 5 min. After staining, specimens were observed under a light microscope (Panoramic SCAN II; 3DHISTECH Kft., Budapest, Hungary) to evaluate the histomorphology of the venous walls.

RNA isolation and quantification. Total RNA was extracted and purified using an Ambion mirVana miRNA isolation kit (cat. no. AM1561; Thermo Fisher Scientific, Inc., Waltham, MA, USA) and RNA integrity was assessed using an Agilent Bioanalyzer 2100 (Agilent Technologies, Inc., Santa Clara, CA, USA). The RNA samples were used for microarrays. To avoid differences between individuals, 200 ng total RNA isolated from three rats per group was pooled into a single sample. Microarray experiments were repeated to produce three independent biological replicates.

miRNA microarray experimental set-up and initial data analysis. miRNA expression analysis was performed using nine Agilent rat miRNA (8x60K) V21.0 microarrays (design ID: 70154; Agilent Technologies, Inc.) at the National Engineering Center for Biochip, Shanghai Biotechnology Corp. (Shanghai, China) miRNA was labeled using the miRNA complete labeling and hybridization kit (cat. no. p/n 5190-0456; Agilent Technologies, Inc.). The kit was used to hybridize each slide with Cy3-labeled RNA in a hybridization oven. Subsequent to hybridization, slides were washed and scanned using a Microarray Scanner (cat. no. G2565CA; Agilent Technologies, Inc.) using Feature Extraction software (version 10.7; Agilent Technologies, Inc.).

Raw data were normalized using the quantile algorithm in the GeneSpring software (version 12.6; Agilent Technologies, Inc.). The results of signal values are presented as means ± standard deviation. Student's t-test was used to identify differences between groups using the R package. Furthermore, the fold change was the ratio of the mean values of two comparative groups. miRNAs with fold differences ≥2.0 and P≤0.05 were considered to indicate a statistically significant difference.

mRNA microarray experimental set-up and initial data analysis. mRNA expression analysis was performed using nine Agilent Whole Rat Genome Microarrays 4x44K (design ID:014879; Agilent Technologies, Inc.) at the National Engineering Center for Biochip, Shanghai Biotechnology Corp. Total RNA was amplified and labeled using the Low Input Quick Amp Labeling kit, One-Color (cat. no. 5190-2305; Agilent Technologies, Inc.). Labeled cRNA was purified using an RNaseasy mini kit (cat. no. 74106; Qiagen GmbH, Hilden, Germany). The left process was similar to miRNA microarray. mRNAs with fold differences ≥2.0 and P≤0.05 were considered to indicate a statistically significant difference.

Integrated analysis of miRNA and mRNA expression profiles. Five prediction tools: TargetMiner (http://www.isical.ac.in/~bioinfo_miu/targetminer20.htm), miRDB (http://mirdb.org/miRDB/index.html), microRNA (http://www.microrna.org/microrna/home.do), TarBase (http://diana.imis.athena-innovation.gr/DianaTools/index.php?r=tarbase/index), and RNA22 (https://cm.jefferson.edu/rna22/) quoted by miRBase database, were used to obtain the target genes of differentially expressed miRNAs. All of the tools were assembled into an online tool (http://www.shbio.com/analysis.html).

The predicted mRNA targets were then compared with experimentally determined mRNAs by microarray. Subsequently, according to Pearson's correlation coefficients and the associations of mRNAs in STRING (http://string-db.org/), a regulatory network was determined, which was comprised of 19 miRNAs and 98 mRNAs.

These 98 mRNAs were uploaded to Database for Annotation, Visualization and Integrated Discovery (version 6.7; https://david.ncifcrf.gov/) for Gene Ontology (GO) functional annotation. The Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway database (http://www.genome.jp/kegg/kegg1.html) was used to identify the enriched pathways of targets. Additionally,
19 miRNAs were subjected to a two-way unsupervised hierarchical clustering analysis using MultiExperiment Viewer software 4.7.4, a part of the TM4 Microarray software suite (Dana-Farber Cancer Institute, Boston, MA, USA) and ranked according to fold change and degrees in the network.

**Statistical analysis.** Data are presented as the means ± SD. Statistical difference (P≤0.05) was calculated by Student’s t-test and correlations were evaluated by Pearson’s correlation coefficients using the R package. The P-value <0.05 was considered statistically significant.

**Results**

*Histological changes in veins following ligation.* Sections of the IVC were stained with H&E and Masson’s trichrome and observed under a light microscope. Clear differences were observed among the three groups (Fig. 2). With discontinuous vascular endothelial cells (ECs), numerous inflammatory cells infiltrated and entered the blood vessels, demonstrating a stable mixed thrombus following treatment in the DVT group. The staining results indicated the success of the SD DVT models.
miRNA expression profiling. To evaluate the role of miRNAs in thrombosis, the miRNA profiling in the DVT animal models were investigated, along with the sham-surgery and control rats (n=9) by microarray. Among 758 detectable miRNAs, 65 miRNAs (29 upregulated and 36 downregulated) significantly difference between the DVT and control groups, 34 miRNAs (14 upregulated and 20 downregulated) between the DVT and sham groups, and 48 miRNAs (26 upregulated and 22 downregulated) between the sham and control groups using a cut-off with an adjusted P≤0.05 and a fold-change of ≥2.0 (Fig. 3A). To exclude any effects of the surgical procedure, only 22 differentially expressed miRNAs in the overlapping areas (Fig. 3C) were focused upon, indicating differences between the DVT and the two other groups, but no difference between the control and sham groups. These 22 miRNAs were subjected to further analyses.

mRNA expression profiling. More than 41,000 rat genes and transcripts were investigated in the present study. Of them, 3,695 genes (2,028 upregulated and 1,667 downregulated) differed significantly between the DVT and control groups, 2,627 genes (1,568 upregulated and 1,059 downregulated) between the sham and control groups, and 1,071 genes (242 upregulated and 829 downregulated) between the DVT and sham groups using a cut-off with an adjusted P≤0.05 and a fold-change of ≥2.0 (Fig. 3A). To exclude any effects of the surgical procedure, only 22 differentially expressed miRNAs in the overlapping areas (Fig. 3C) were focused upon, indicating differences between the DVT and the two other groups, but no difference between the control and sham groups. These 22 miRNAs were selected for further analyses.

miRNA-mRNA network. The 22 miRNAs were integrated to the experimentally determined 487 DEGs to obtain the miRNA-mRNA network (Fig. 4), which was composed of 19 miRNAs (6 upregulated and 13 downregulated) and 98 differentially expressed mRNAs (13 upregulated and 85 downregulated).

Functional classification of 98 genes. GO assigns biological processes (BP), cellular components (CC) and molecular function (MF). The BP analysis involved 83 GO terms; five of the top 10 were associated with adenylate cyclase activity, and two were associated with cell growth (Fig. 5A). Seven were found in CC, indicating that the DEGs were predominantly extracellular genes (45.66%) and plasma membrane genes (34.78%) (Fig. 5B). In seven MF annotations, the majority of genes belonged to binding activity genes (18.4%), while active protein genes represented 8.7% (Fig. 5C).

To evaluate the roles of the target genes in DVT, six specific terms (angiogenesis, cell proliferation, adhesion, inflammatory response, apoptosis and hypoxia) mediating vascular function were extracted from multiple levels within the GO hierarchy (Table I). Notably, the majority of these genes were closely associated with endothelial function (marked in bold in Table I) and the majority demonstrated a decreased expression level in DVT. For example, the expression levels of EC proliferative and anti-apoptosis genes, growth arrest specific 6 (GAS6), vascular endothelial growth factor B (VEGFB), the most well-known inducer of angiogenesis, and endothelial inflammation factor, Kruppel like factor 4 (KLF4) were reported to be reduced in atherosclerosis (11). In addition, adhesion-associated genes, including junctional adhesion molecule 2 (JAM2), which affects EC junctions (12), C-X-C motif chemokine ligand 12 (CXCL12), which mediates...
recruitment of inflammatory and thrombotic cells (13), and apolipoprotein L domain containing 1 (APOLD1) (14), a novel EC early response protein associated with hypoxia, were equally reduced in DVT.

KEGG pathway analysis demonstrated that four of the top 15 signal pathways were associated with cardiovascular disease, and three to metabolism. Among them, the following three had enriched gene numbers >3: dilated cardiomyopathy, vascular smooth muscle contraction and focal adhesion pathways (Fig. 5D).

Clustering of 19 miRNA expression. The clustering analysis of the nine samples and 19 differentially expressed miRNAs revealed a distinct miRNA signature during DVT (Fig. 6A). The six upregulated miRNAs grouped together and the 13 downregulated miRNAs were divided into two categories, in which miR-133b-3p, miR-218a-5p and miR-204-5p were grouped together, while the remaining 10 miRNAs all gather in a second cluster.

Ranking. To demonstrate the value of the 19 miRNAs in DVT, they were positioned according to the degree of co-expression from network (Table II) and fold change form profiling (Tables III and IV). The co-expression was ranked according to the degree of co-expression, the highest degree scored 19 points, and the lowest 1 point. The fold change score was ranked in the same way. The most differentially expressed upregulated miRNA received 6 points (for a total of 6 upregulated miRNAs) and downregulated miRNAs received 13 points (for a total of 13 downregulated miRNAs), and the lowest 1 point (data not shown). Bar charts of the rankings are presented in Fig. 6B and C. The top upregulated miRNA was miR-92a-3p, while miR-218a-5p was ranked first among the downregulated miRNAs.
Table I. Genes associated with specific GO terms (angiogenesis, cell proliferation, adhesion, inflammatory response, apoptosis and hypoxia) in DVT. Genes related to endothelial cells are marked in bold.

| Specific Go terms                  | Upregulated genes | Downregulated genes |
|-----------------------------------|-------------------|---------------------|
| Angiogenesis                      |                   | APOLD1, CALCRL, KLF4, DLL1, CXCL12, EMP2, GATA2, VEGFB |
| Cell proliferation                | PTH, KCNH1, ITGA2 | ARC, CALCRL, CAV2, CHP2, CXCL12, DLL1, EFEMP1, EMP2, GAS6, GATA2, KLF4, NACC2, NDRG4, PBX1, PPARG, PTHLH, RERG, SPRY2, TNFRSF11B, VEGFB, WDFC1, WISP2, ZFP36L1 |
| Adhesion                          | ITGA2             | PPAP2B, HSPA2, GAS6, ZFP36L1, CXCL12, JAM2, KLF4, EMP2, FLRT3, WISP2, DLL1, CAV2, HAS1, ITGA, RND3, CNTNAP1 |
| Inflammatory response             | ITGA2             | PPARG, CALCRL, WDFC1, STK39, TNFRSF11B, IL17RE |
| Apoptosis                         | PTH               | ARC, HSPA2, GAS6, TNFRSF11B, NACC2, PPARG, SPRY2, CXCL12, VEGFB, GULP1, KLF4, ZFP36L1, DLL1, STK39 |
| Hypoxia                           | ITGA2             | APOLD1, CYGB, ARC, CXCL12 |

GO, Gene Ontology.

Table II. Degrees of differential expression and targets of the 19 differentially expressed miRNAs.

| miRNA   | Degree | Targets                                                                 |
|---------|--------|-------------------------------------------------------------------------|
| miR-125a-5p | 22     | ENTPD2, CXCL12, GAS6, YPEL4, ARC, CHP2, GRB10, ITGA7, CYGB, SLC4A10, HAS1, WISP2, OLFM1L2A, LZTS1, TNFRSF11B, ADAMTS1, EMP2, APOLD1, RHOV, VEGFB, CLIC5, ITPKC |
| miR-195-5p | 20     | NKD1, DLL1, PLCXD2, ADRA1D, GK, SLC16A6, WIFI, ITGA2, SNX33, PLA2G15, FLRT3, TPPP3, MMP17, ADCY5, RUNX1T1, ISLR, DYRK1B, PTHLH, PTPN4, PTH |
| miR-497-5p | 19     | DLL1, ITGA2, NKD1, GK, PTH, ADRA1D, ISLR, PLA2G15, PTHLH, RUNX1T1, SNX33, TPPP3, MMP17, SLC16A6, PLCXD2, ADCY5, DYRK1B, WIFI1, PTPN4 |
| miR-27b-3p | 18     | PPARG, RND3, ZFP36L1, PPAP2B, FAM171A1, EPB41L1, HEG1, FLRT3, GATA2, NOVA1, STK39, EEVPD1, WIPF3, SPRY2, PLCXD2, STNM2, DCX, PKIA |
| miR-15a-5p | 12     | ISLR, RUNX1T1, DLL1, PTH, PTHLH, TPIPP3, GK, NKD1, ITGA2, ADRA1D, PLA2G15, MMP17 |
| miR-10a-5p | 9      | ADRA1D, RBMS3, LZTS1, KAZALD1, SH3D19, KLF4, EEPD1, LTBP1, PP1R14C |
| miR-218a-5p | 9      | FLRT3, SH3D19, PLA2G15, CALCRL, KCNH1, PLCXD2, NDRG4, FAM63B, LHF |
| miR-92a-3p | 9      | FAM110B, GATA2, NOVA1, SLC32A1, STK39, KLF4, PRKAR1B, KLF6, GPM6A |
| miR-128-3p | 8      | EFEMP1, DCX, SCLM4, RND3, IL17RE, ITPKC, KLF4, ABCB9 |
| miR-19b-3p | 8      | GULP1, SH3D19, HSPA2, RBMS3, HEG1, GRB10, ENC1, ARC |
| miR-23b-3p | 7      | KCNH1, DLG2, CXCL12, LOC499602, CAV2, NACC2, NAP1L5 |
| miR-204-5p | 6      | NAT8L, LRRN4CL, DHH, NOVA1, HEG1, RGD1562629 |
| miR-143-3p | 5      | DYRK1B, CXCL12, CLIC5, WDFC1, PTPN4 |
| miR-133b-3p | 4      | EMP2, ENC1, CNTNAP1, GPM6A |
| miR-142-5p | 4      | RERG, TAM2, CNTLN, GCH1 |
| miR-196a-5p | 4      | PLCXD2, RBMS3, PBX1, ABCB9 |
| miR-196b-5p | 4      | RBMS3, PLCXD2, PBX1, ABCB9 |
| miR-10b-5p | 4      | GMP6A |
| miR-486 | 4      | BTBD3 |

Genes mentioned in the main text: miR, microRNA; GAS6, growth arrest specific 6; VEGFB, vascular endothelial growth factor B; JAM2, junctional adhesion molecule 2; CXCL12, C-X-C motif chemokine ligand 12; APOLD1, apolipoprotein L domain containing 1; HSPA2, heat shock protein family A (Hsp70) member 2; DLL1, delta like canonical Notch ligand 1; GATA2, GATA binding protein 2; KLF4, Kruppel like factor 4; ITGA2, integrin subunit α2.
Figure 5. Bioinformatics analysis of 98 mRNAs in DVT. GO analysis assigns high-level terms from each of the following three major GO terms: (A) Biological processes, (B) cellular components, and (C) molecular functions. (D) The top 15 canonical pathways [with score >1.7, P<0.02]. mRNA, messenger RNA; DVT, deep vein thrombosis; GO, gene ontology; ACA, adenylate cyclase activity; ARVC, arrhythmogenic right ventricular cardiomyopathy.

Figure 6. Bioinformatics analysis of 19 miRNAs in DVT. (A) Hierarchical cluster analysis of the 19 differentially expressed miRNAs. Hierarchical clustering analysis separated the nine samples into three groups, which were consistent with the DVT, control and sham groups. Red bars indicate high expression levels, green bars indicate low expression levels, and black bars indicate similar expression levels. (B) Scores of 6 upregulated and (C) 13 downregulated miRNAs. The scores are based on the degree of co-expression and the fold change. miRNA, micro RNA; DVT, deep vein thrombosis; FD-DVS, fold change in the DVT vs. sham groups; FD-DVC, fold change in the DVT vs. control groups.
Discussion

DVT and PE are significant public health concerns, representing major sources of mortality and morbidity. Animal models are important for understanding the pathophysiology of these thrombogenesis-associated diseases (15). To minimize the error caused by surgery, a rat IVC ligation model was used in the present study, which provides a total stasis environment and a consistent thrombus size after 3 days of ligation. This model has been widely used in previous studies (1,2,16-19).

Although miRNA screening methodologies have become widely available, and large studies of the role of miRNAs were performed in the pathogenesis of various cardiovascular diseases (20-24), to the best of our knowledge, only three studies reported miRNA profiling in vein thrombosis. Xiao et al (25) identified markedly higher plasma levels of various miRNAs, (including miR-134, miR-410 and miR-520 amongst others) in patients suffering from acute PE (25). The miR-320a and miR-320b were upregulated in plasma samples from VTE patients compared with healthy controls (26). Qin et al (27) demonstrated an increased serum level of three miRNAs (miR-582, miR-195 and miR-532) in patients with postoperative DVT versus control subjects (27).

All previous studies identified differentially expressed miRNAs in the thrombosis group, consistent with the present study, but the specific miRNA profiles differed, which may be associated with differences in the subsets of selected diseases, species and types of samples that were evaluated. Beyond this, all of the known miRNA profiles associated with thrombosis were detected in plasma or serum as potential markers, while, to the best of our knowledge, there are no studies evaluating the miRNA profiles in venous tissue. The miRNA profiles from biological fluid are used primarily for diagnoses, while miRNA profiles in venous tissues may be more conducive to investigating the underlying mechanism of DVT, which has rarely been the focus of study. Therefore, the present study was designed to investigate this further.

Table III. Differential expression of 6 upregulated miRNAs in the DVT, the control and the sham group.

| miRNA    | DVT group | Control group | sham group | Fold change (P-value) |
|----------|-----------|---------------|------------|-----------------------|
| miR-19b-3p | 573.02±42.12 | 229.30±24.47 | 273.82±34.41 | 2.50 (1.14E-03) 2.09 (5.09E-03) |
| miR-142-5p | 104.09±22.38 | 45.03±11.02 | 42.85±2.04 | 2.31 (2.5E-02) 2.43 (3.16E-02) |
| miR-15a-5p | 185.14±32.59 | 89.34±11.84 | 72.66±56.24 | 2.07 (1.40E-02) 2.55 (1.44E-02) |
| miR-92a-3p | 769.03±111.47 | 190.20±9.65 | 258.05±14.72 | 4.04 (3.17E-03) 2.98 (5.12E-03) |
| miR-486   | 826.37±182.19 | 303.27±55.64 | 263.43±112.57 | 2.72 (1.21E-02) 3.14 (2.93E-02) |
| miR-128-3p | 396.74±52.95 | 52.95±25.23 | 88.78±4.53 | 2.91 (3.89E-03) 4.47 (1.61E-03) |

miRNA, micro RNA; DVT, deep vein thrombosis.

Table IV. Differential expression of 13 downregulated miRNAs in the DVT, the control and the sham group.

| miRNA    | DVT group | Control group | Sham group | Fold change (P-value) |
|----------|-----------|---------------|------------|-----------------------|
| miR-218a-5p | 1.03±0.73 | 58.88±9.69 | 40.44±6.74 | 56.94 (2.92E-02) 39.11 (3.41E-02) |
| miR-204-5p | 0.73±0.87 | 29.89±18.58 | 28.45±2.91 | 40.75 (3.01E-02) 38.79 (3.93E-02) |
| miR-133b-3p | 1.79±1.71 | 1631.01±1898.14 | 47.74±22.25 | 912.16 (1.00E-02) 26.70 (4.85E-02) |
| miR-10b-5p | 75.88±9.55 | 488.90±0.00 | 333.18±53.56 | 6.44 (2.48E-03) 4.39 (9.00E-04) |
| miR-10a-5p | 113.96±6.70 | 569.35±34.06 | 365.22±26.42 | 5.00 (1.11E-05) 3.20 (9.26E-05) |
| miR-143-3p | 807.53±91.71 | 5490.83±1852.49 | 2341.94±438.29 | 6.80 (1.54E-02) 2.90 (6.51E-03) |
| miR-195-5p | 169.80±23.48 | 793.40±215.98 | 456.61±0.00 | 4.67 (5.20E-03) 2.69 (1.06E-02) |
| miR-497-5p | 121.73±10.93 | 331.66±56.00 | 308.19±39.63 | 2.72 (4.81E-03) 2.53 (2.33E-03) |
| miR-27b-3p | 245.33±14.03 | 768.64±108.02 | 605.37±8.52 | 3.13 (2.67E-03) 2.47 (1.26E-03) |
| miR-196a-5p | 32.02±5.81 | 157.60±58.56 | 74.12±5.96 | 4.92 (1.21E-02) 2.31 (9.65E-03) |
| miR-23b-3p | 421.90±49.45 | 1712.50±358.92 | 972.26±53.57 | 4.06 (2.97E-03) 2.30 (3.16E-03) |
| miR-196b-5p | 31.71±4.44 | 120.13±38.66 | 70.53±3.39 | 3.79 (1.75E-02) 2.22 (8.43E-03) |
| miR-125a-5p | 70.06±2.92 | 142.10±10.40 | 150.30±9.76 | 2.03 (1.14E-03) 2.15 (4.62E-04) |

miRNA, micro RNA; DVT, deep vein thrombosis.
By integrative methodology, miRNA/mRNA pairs in DVT were used to construct the regulatory network, indicating that these miRNAs may be important in DVT by regulating their target genes. Further GO and KEGG analyses of 98 genes indicated that DVT was associated with specific biological processes, such as angiogenesis or inflammation.

It is worth noting that the changes in these genes were reported to affect endothelial function (28,29). For example, heat shock protein family A (Hsp70) member 2 (HSPA2) in the adult corneal endothelium were demonstrated to be sensitized to mediators of cell death (30). In addition, delta like canonical Notch ligand 1 (DLL1) is an essential Notch ligand in the vascular endothelium, and activates Notch1 to maintain arterial integrity (31). Another example is the angiogenic factor, GATA binding protein 2 (GATA2). Previous studies (32,33) have indicated that GATA2 is important for vascular integrity. In addition, KLF4 functions as an important regulator of EC inflammation (34). Integrin subunit α2 (ITGA2), the major collagen-binding α-integrin subunit in ECs, is involved in inflammatory processes and cell invasion (35,36). Increased expression levels of ITGA2 increases joint inflammation (36).

Clustering analysis of 19 miRNAs revealed a distinct miRNA signature during DVT. Notably, the majority of the 19 miRNAs have also been reported in the endothelium. For example, miR-10b-5p and miR-195 have been identified as diagnostic markers of VTE (26,27). In addition, miR-92a, miR-15a, miR-196a/196b and miR-19b have been reported to be involved in the processes of EC dysfunction, proliferation, apoptosis, migration and angiogenesis, which ultimately influence diseases, such as thrombosis and atherosclerosis, and tumors (37-41).

Maintenance of the functional integrity of the endothelium is important to preserve blood flow and prevent thrombosis (42). The endothelium secretes factors that control vascular relaxation and contraction, thermogenesis, fibrinolysis, and platelet activation and inhibition (43). While EC injury and dysfunction are considered to be the initial events in the development of thromboembolism, atherosclerosis, postangioplasty restenosis and plaque erosion contribute to macrovascular complications (44). The present findings indicate that endothelial dysfunction is particularly significant in ligation-induced thrombosis. A flowchart of the role of endothelial miRNAs in DVT was generated by integrating the expression and regulation of endothelial-associated miRNAs and target genes (Fig. 7).

In conclusion, profiling of miRNA and mRNA expression was performed in parallel to integrate the two biological levels to better understand DVT. The use of this integrated approach elucidates the associations between the expression levels of miRNAs and mRNAs. This novel systematic study provides information on miRNA target regulation and provides a foundation for future studies.

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