

Expression Levels of miR-30c and miR-186 in Adult Patients with Membranous Glomerulonephritis and Focal Segmental Glomerulosclerosis

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Background: Nephrotic syndrome is a common renal problem with different histopathogenesis. MicroRNAs are reported to be involved in the pathophysiology of the syndrome. The aim of this study was to study the levels of miR-30c and miR-186 in NS patients.

Methods: Sixty patients with primary NS (membranous glomerulonephritis (MGN, N=30) and focal segmental glomerulosclerosis (FSGS, N=30)) and 24 healthy volunteers were included. Expression levels of the miR-30c and miR-186 were evaluated in plasma and peripheral blood mononuclear cell (PBMC) samples of adult patients with NS using real-time PCR. Moreover, an in-silico analysis was performed to understand the signaling pathways and biological procedures that may be regulated by these miRNAs.

Results: In the MGN group, significantly elevated levels of miR-30c and miR-186 were observed in PBMC (P= 0.037) and plasma (P= 0.035) samples, respectively. Moreover, there was a significant increase in miR-30c levels in PBMC samples of the FSGS group when compared to healthy controls (P= 0.004). In ROC curve analysis, combined levels of the studied miRNAs could discriminate cases from controls in plasma and blood cells (AUC≥0.72, P<0.05).

Conclusion: A panel of miRNAs may be potential biomarkers in plasma and PBMCs samples of NS patients with different subclasses. More investigations are needed with a large sample size to validate the diagnostic values of the reported miRNAs.

Keywords: nephrotic syndrome, proteinuria, membranous glomerulonephritis, focal segmental glomerulosclerosis, microRNAs, biomarker

Introduction

Nephrotic syndrome (NS) is a common renal problem defined by proteinuria, edema, and hypoalbuminemia.^{1,2} In adults, NS can be caused by different glomerular diseases, including Minimal Change Nephropathy (MCN), Focal Segmental Glomerulosclerosis (FSGS), and Membranous Glomerulonephritis (MGN). In this respect, each type of NS has its specific pathogenesis, for instance, podocytes, specialized kidney glomerular cells, injury is a major cause of FSGS and MCN, while deposited antibodies against phospholipase A2 receptor (PLA2R) on podocytes are the main cause of adult MGN.³⁻⁵ The common clinical method for NS diagnosis is renal biopsy that is an invasive procedure.⁶ Therefore, it is essential to define the underlying histopathological diagnosis with a non-invasive and more reliable method to employ a specific therapy and monitor response to therapy.

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MicroRNAs are small non-coding ribonucleotides that are involved in different cell procedures by binding to 3' UTR (untranslated-region) region of mRNAs and silencing their expression at both transcriptional and translational levels.^{7,8} About 30% of mRNAs are regulated by all 3000 identified human microRNAs.⁹ Based on the stability and function of miRNAs in cells and body fluids and their expression changes during various diseases, they can be considered as possible diagnostic and prognostic biomarkers.¹⁰⁻¹³ In recent studies, altered expression of multiple miRNAs and their target genes are reported in different kidney diseases.¹⁴⁻¹⁶ Likewise, a specific set of miRNAs is described to be deregulated in different biological sources of children and adults with podocyte injury. Although these previous reports offer some insights into the impact of miRNAs in podocytopathy, more efforts need to be done to understand the clinical significance values of the miRNAs involved in the pathophysiology of NS.

Based on the controversial results obtained from different clinical sources of NS patients,¹⁷⁻²¹ there is a lack of definite early non-invasive biomarkers for NS diagnosis. It was the aim of the present study to assess the levels of circulating and cellular levels of miR-30c-5p and miR-186 involved in podocyte homeostasis and podocytopathies in blood samples of adults with NS and to determine their potential diagnostic or prognostic values.

Materials and Methods

Study Subjects

The present cross-sectional study recruited 60 primary NS patients from the Nephrology Ward of Imam Reza Hospital (Tabriz, Iran). Blood samples were collected from November 2017 to March 2019. Nephrotic proteinuria was defined by excretion of protein ≥ 3 g/24 h in urine, as heavy proteinuria. Inclusion criteria for cases were the age range between 20 and 60, primary NS, and normal urinary system without any obstruction. Patients with secondary NS causes such as lupus, diabetes, amyloidosis, viral infection, and drug-related NS were not included. The exclusion criteria for cases were inflammatory diseases, diabetes, a history of cancer, severe bacterial infection, serologic positivity for CMV, HBV, HIV, HCV and BK viruses, obstruction in the renal system, overlap syndrome, any other glomerulopathy, severe organ failure, autoimmune disorder and those with ESRD under dialysis. Histologic detail of patients was collected from the renal

biopsy findings. Further, a group of healthy volunteers (n= 24) with no clinical history of kidney disease was also allocated as controls. Inclusion criteria for normal controls were matched-age and -gender with the cases and having normal kidney function (normal serum creatinine and urine analysis).

RNA Extraction from Plasma and PBMC

Fresh blood samples (6–8 mL) were collected in EDTA-coated Vacutainer tubes and all processes were performed within 2 hours in the laboratory. For preparing plasma samples, 2 mL of blood sample centrifuged at 1000 \times g for 10 minutes two times at 4°C. Then, supernatant (plasma) was aliquot in microtubes and stored at -70°C for future uses. For peripheral blood mononuclear cells (PBMC) isolation, 4 mL of peripheral blood was diluted with an equal volume of phosphate-buffered saline (PBS) solution and then added into 4 mL Ficoll-Paque. The solution centrifuged at 800 \times g for 20 minutes. PBMCs layer between plasma and Ficoll was taken and transferred into a new RNase-free 15 mL falcon. PBMCs washed with 10 mL PBS and centrifuged at 500 \times g for 10 minutes. The supernatant removed and the pellet was washed with 1mL PBS again and centrifuged at 500 \times g for 10 minutes. Finally, pellet solved in 1mL Trizol and stored at -70°C for later use. Before RNA isolation from plasma samples (250 μ L), 3.5 μ L cel-miR-39 miRNeasy Serum/Plasma Spike-In (1.6 \times 10⁸ copies/ μ L) (Qiagen, Germany) was added to all samples as exogenous control. The total RNA extraction was performed based on the Trizol protocol (RiboExTM, South Korea). Optical Density (OD) was calculated for the detection of RNA integrity, quality, and quantity via A260/A280 and A230/A280 nm by NanoDrop (NanoDropTM One^C Thermo Scientific, USA).

cDNA Synthesis and Quantitative Real-Time PCR (Q-PCR)

Synthesis of the first strand of cDNA in both plasma and PBMCs samples was performed separately in 15 μ L reaction volume; RNA (5 μ L RNA extracted from plasma and 1 μ g RNA extracted from PBMCs), 0.8 μ L RT enzyme, 3 μ L RT buffer, 0.375 μ L Ribolock, 1.5 μ L dNTP, 1.325 μ L DEPC, and 3 μ L primer mix (including equal volumes of customized stem-loop primers of miR-30c, miR-186, U6, spiked in cel-miR-39, Snord-47 and UR (universal reverse)) [Supplementary Table S1](#). Then, cDNA (0.5 μ L) amplified in a reaction of 10 μ L volume using 2 μ L master mix (5 \times HOT FIREPol[®]

EvaGreen[®] qPCR Mix Plus, Biodyne, Austria), 0.5 μ L forward primer, 0.5 μ L reverse primer, and 6.5 μ L DEPC. For each sample, a duplicate run was done. The qPCR was performed at 95°C for 15 min, followed by 40 cycles each at 95°C for 10 s, and annealing at 60°C for 30 s. Melting curves were made at 72–95°C to verify the specificity of reaction after the completion of PCR cycling. Cel-miR-39, SNORD-47, and U6 were used as internal controls for plasma and PBMCs, respectively. The gene expression was calculated with the $2^{-\Delta\Delta Ct}$ method.

Statistical Analysis

To test the normal distribution of variables, the Shapiro–Wilk test was used. Categorical variables presented by numbers and percentages. Mean \pm standard deviation (SD) and median (Max–Min) were used for normally and non-normally distributed variables, respectively. For comparisons of quantitative variables between two groups, Student’s *t*-test or Mann–Whitney *U*-tests were used while the chi-squared test was used for categorical variables. One-way ANOVA test with Tukey’s HSD post hoc test was used for normally and Kruskal–Wallis was used for non-normally distributed variables between more than two groups. Spearman correlation was used for the evaluation of correlations between variables. The receiver operating characteristic (ROC) curve analysis was used for evaluating the potential of each miRNA in discriminating cases from controls or for discriminating cases with FSGS from MGN. The optimal threshold was obtained by Youden’s method. Statistical analyses were performed by the IBM SPSS 17.0 Software (SPSS, Inc.). $P < 0.05$ was considered statistically significant.

In-Silico Analysis

An in-silico analysis was performed to understand the underlying signaling pathways and biological procedures regulated by the studied miRNAs. Using miRTarBase 6.0 database (Available at <http://mirtarbase.mbc.nctu.edu.tw/php/index.php>), a list of the confirmed targets of hsa-miR-30c and hsa-miR-186 were obtained and subjected to the ORA (overrepresentation enrichment analysis) based on Kyoto Encyclopedia of Genes and Genomes (KEGG) and gene ontology (GO) pathways using the WebGestalt web server (Available at <http://www.webgestalt.org/option.php>). In these analyses, the reference gene list was set to “genome_protein_coding” and the multiple test adjustment method set to Benjamini–Hochberg (BH). Other factors set as default. The enrichment ratio (R) was considered for each significant category that stands for the ratio of observed versus expected

number of genes. Besides, a literature search was made to find the functions of target genes that may be relevant to NS.

Results

Subjects

Sixty patients with NS, 30 patients with biopsy-proven FSGS, and 30 patients with positive anti-PLA2 receptor antibody and biopsy-proven MGN were included in this study. Table 1 summarizes the clinical and demographic data of patients. Regardless of differences in their histopathology, all of the patients were associated with features of the primary NS and podocyte injury. Most of the patients were refractory to the treatment and they were failed to achieve remission of their proteinuria after the treatment period. There were no significant differences in age and sex between cases and controls ($P = 0.54$). Moreover, no significant differences detected in age, gender, GFR or serum levels of creatinine, uric acid, urea and 24h urine creatinine between the different subtypes of cases ($P > 0.05$) (Table 1).

Among the dysregulated miRNAs in NS, the expression levels of miR-30c and miR-186 were evaluated in both plasma and PBMCs specimens of the cases by q-PCR. Plasma samples were used for the analysis of miRNAs since the true profiling of circulating miRNAs may be altered during the coagulation procedure in serum preparation. Despite significant reported values of exogenous control, neither spiked-in cel-miR-39-5p nor –3p was appropriate for analysis of data; therefore, U6 used as an internal control for normalization of circulating miRNAs in plasma samples. Moreover, PBMCs were used since the evaluation of miRNAs in these cells had not been reported in patients with NS.

Statistically, there was an increase in miR-30c level in the peripheral cells of the NS ($P = 0.005$), MGN ($P = 0.037$), and FSGS ($P = 0.004$) patients when compared to controls. Plasma levels of miR-30c were not significant between the studied groups, Figure 1A and B. Circulating ($P = 0.045$) but not cellular ($P = 0.70$) levels of miR-186 increased in NS groups significantly when compared to controls (Figure 1C and D). Additionally, the increased level of circulating miR-186 was significantly obvious in the MGN group ($P = 0.035$) but not in the FSGS group ($P = 0.183$) as compared to controls. However, increased levels of PBMCs’ miR-186 were not statistically significant in the MGN group in comparison to controls ($P = 0.91$) and FSGS cases ($P = 0.28$), Figure 1C and D.

There were correlations between the studied miRNAs and other clinical variables in NS patients; however, the differences were not statistically significant (data not

shown). There were significant correlations between the studied miRNAs levels in plasma and PBMC (Table 2).

ROC Analysis

To further examine the values of the studied miRNAs to discriminate subtypes of NS, the ROC analysis was done. Fold change values were used to compute the ROC curve. ROC results are presented in Table 3 with more details. PBMC levels of miR-30c showed a sensitivity of 46% and specificity of 96% with AUC=0.65, 95% CI; 0.565 to 0.728 to discriminate NS from controls. Circulating miR-186 with AUC=0.61, 82% sensitivity, and 43% specificity could discriminate FSGS from MGN cases.

The combined effect of two circulating miRNAs with AUC=0.72 [0.64–0.82] ($p<0.001$), 86% sensitivity, and 51% specificity could distinguish NS patients from healthy controls (PPV= 0.806[0.715 to 0.874], NPV= 0.613[0.438 to 0.763], $LR^+ = 1.77$ [1.259 to 2.493], and $LR^- = 0.268$ [0.146 to 0.495]). Cellular levels of these miRNAs with AUC=0.72 [0.62–0.81] ($p<0.001$), 53% sensitivity and 89% specificity could distinguish NS patients from healthy controls with PPV=0.915[0.801 to 0.966], NPV= 0.471[0.359 to 0.587], $LR^+ = 4.98$ [1.928 to 12.824], $LR^- = 0.518$ [0.399 to 0.674]. Moreover, with AUC=0.69 and AUC=0.71, PBMCs levels of the studied miRNAs could discriminate FSGS from MGN, respectively. Although each of the studied miRNAs alone could not discriminate the individuals in the studied groups, their combined levels showed a reliable moderate accuracy. However, both high sensitivity and specificity were not observed (Figure 2A–D).

In-Silico Analysis

The ORA identified biological processes that are overrepresented among target genes of miR-30c and miR-186 (Table 4).

GO and KEGG enrichment analysis proposed that the predicted targets may participate in a variety of physiological regulatory functions in NS. This result shows that miR-30c by targeting genes involved in the chromatin modification, gene expression, double-strand RNA (dsRNA) and mRNA metabolic processes, cell cycle and posttranscriptional regulation of gene expression may be involved in NS. Additionally, miR-186 predicted to target chromosome organization, RNA splicing process, gene transcription and translation, peptide biosynthetic, and cellular macromolecule catabolic processes. Moreover, 10 common pathways in the KEGG pathway were found for the studied miRNAs (Supplementary Tables S2 and S3). A literature search indicated that some target genes of miR-30c and miR-186 may be functionally related to NS.

Discussion

To validate the potential value of miRNAs involved in the pathogenesis of NS, the expression levels of miR-30c and miR-186 in plasma and PBMCs samples were compared between NS cases and healthy individuals. The result indicated that PBMCs level of miR-30c was upregulated significantly in NS group compared to healthy groups. Moreover, there was a significant difference in miR-30c levels between different pathology subtypes of NS in comparison to control group. Furthermore, this study demonstrated a significant increase in plasma (not PBMCs) level of miR-186 levels in the MGN group when compared to controls. The combined levels of the studied miRNAs in plasma and PBMCs could discriminate NS patients from controls.

MiR-30 family is a significant epigenetic regulator in podocyte homeostasis and their production can be changed in the course of NS. Evidence shows that the level of miR-30 family is altered with NS progression and therapy.^{22,23} In

Table 1 Demographic and Baseline Clinical Data

Characteristics	Total NS Patients	MGN	FSGS Group	P-value ^a
No. of cases	60	30	30	
Age, mean \pm SD (years)	45.76 \pm 13	46.6 \pm 13	44.5 \pm 12.8	0.404
BMI (kg/m ²)	27.3 \pm 3.4	27.4 \pm 3.6	27.3 \pm 3.4	0.49
GFR(mL/min/1.73m)	80 (10–137)	80 (18–137)	79 (10–130)	0.324 ^b
Proteinuria (mg/24 h)	2497 (85–15,900)	3013 (85–15,000)	1360 (97–15,900)	0.26 ^b
Urea (mg/dL)	40.7 \pm 19.4	38.2 \pm 16.5	44.2 \pm 20	0.18
Uric acid (mg/dL)	6.5 \pm 1.1	6.61 \pm 1.2	6.3 \pm 1.1	0.94
Serum creatinine (mg/dL)	1.2 \pm 0.5	2.21 \pm 0.55	1.2 \pm 0.5	0.17
Urine creatinine (mg/24h)	1124.38 \pm 407	1159.21 \pm 328	1074.63 \pm 500	0.742

Notes: GFR: estimated glomerular filtration rate as measured by GFR [$186 \times (\text{Creatinine}/88.4)^{-1.154} \times (\text{Age})^{-0.203} \times (0.742 \text{ if female})$] formula. The quantity data are expressed as mean \pm SD. ^aMGN versus FSGS. ^bMedian (Min-Max) is presented. P-value is based on Mann–Whitney U-test.

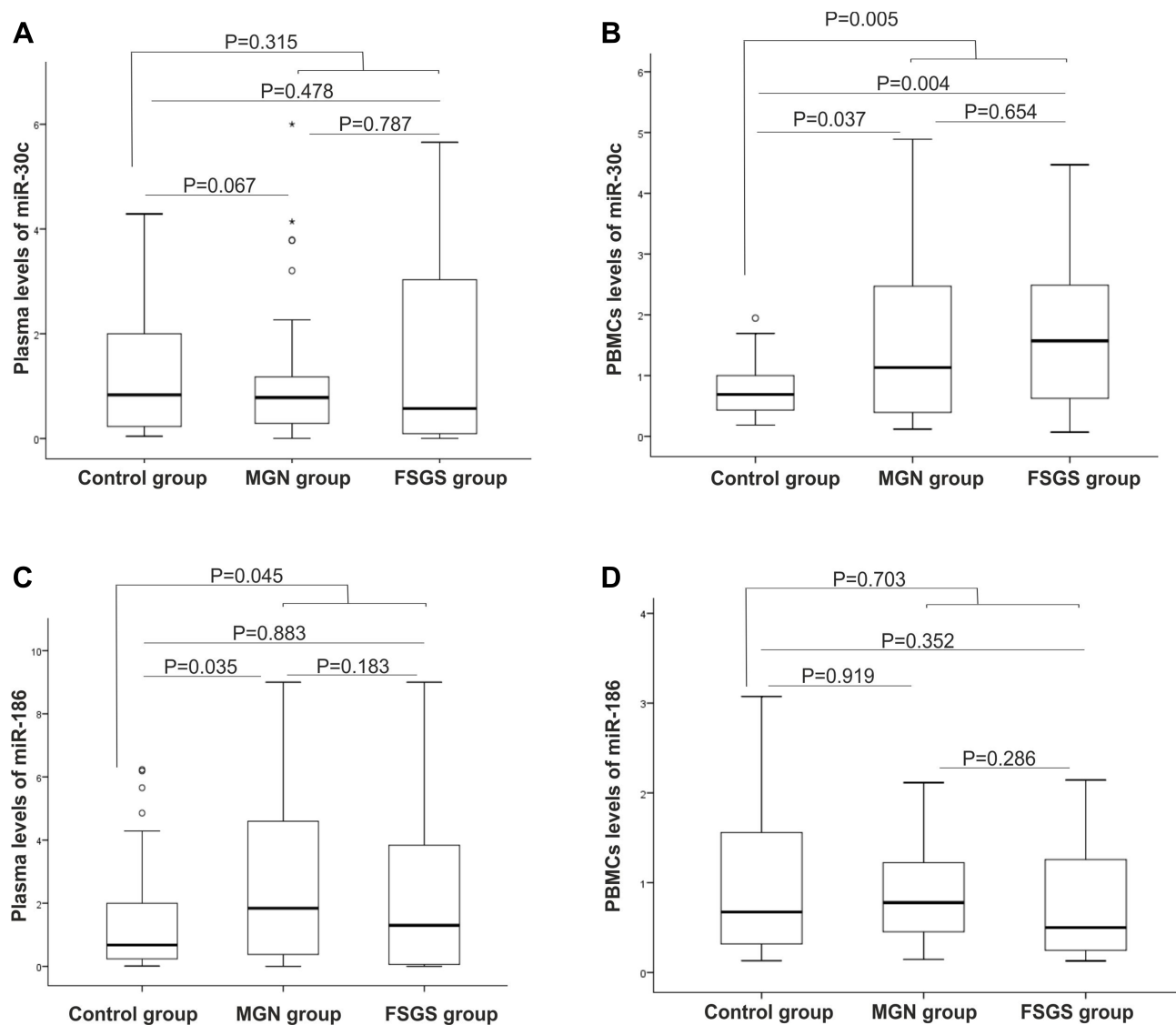


Figure 1 The expression of miRNA-30c-5p and miRNA-186-5p in plasma and PBMCs. The expression of miRNA-30c-5p in (A) Plasma and (B) PBMCs between the studied groups. The expression of miRNA-186-5p in (C) Plasma and (D) PBMCs between the studied groups. The relative expression (fold change) was calculated by the $2^{-\Delta\Delta C_t}$ method. * and \circ are outliers.

Abbreviations: NS, nephrotic syndrome; FSGS, focal segmental glomerulosclerosis; MGN, membranous glomerulonephritis.

serum and urine samples,²³ and urinary exosomes²⁴ of children with active NS, increased concentration of miR-30a-5p was observed, while its level tended to decrease with the remission of patients. In contrast to these results, downregulated levels of miR-30 family were reported in podocytes of FSGS cases¹⁷ and kidney tissue of children with FSGS but not MGN²¹ in comparison to normal controls. Furthermore, the high throughput sequencing of microRNAs showed that miRNA-30 family was downregulated in peripheral blood lymphocyte cells (PBLCs) of MGN patients in comparison to healthy individuals.²² Higher plasma levels of miR-30c were also found in MCN cases compared to FSGS patients

and controls; however, there were no significant differences in miR-30c level between FSGS and healthy controls.¹⁸ In the present study, the PBMCs level of miR-30c significantly elevated in the NS group when compared to healthy controls. Moreover, there was a statistically significant change in the FSGS and MGN groups when compared to controls.

Elevated and diminished levels of miR-186 have been described in previous studies in NS patients.^{20,21} In this research, an overexpression was observed in the circulating miR-186 level in the NS group and MGN. Additionally, no statistically significant changes were observed between the NS group and controls or among

Table 2 Internal Correlations Between miRNA Levels in Plasma and Peripheral Blood Cells

	Plasma miR-30c	Plasma miR-186	PBMCs miR-30c	PBMCs miR-186
Plasma miR-30c	I	r = 0.779* p < 0.001	r = -0.191 p = 0.137	r = -0.099 p = 0.449
Plasma miR-186		I	r = -0.176 p = 0.167	r = -0.141 p = 0.273
PBMCs miR-30c			I	r = 0.60* p < 0.001

Note: *Correlation is significant.

Abbreviation: r, correlation coefficient.

FGSG and MGN groups in PBMC samples. In a literature review, elevated levels of miR-186 are reported in plasma samples of patients with active FSGS relative to healthy controls; however, miR-186 level remained unchanged in steroid-resistant patients before and after steroid treatment.²⁰ It should be also noted that the plasma level of miR-186 was not increased in groups of MGN and diabetic nephropathy patients with nephrotic proteinuria.²⁰ However, down-regulated levels of miR-186 were observed both in human podocytes cell line and renal biopsy of MGN patients.²¹ These results showed that miR-186 might not be involved in the pathogenesis of FSGS.

ROC analysis indicated that each of the studied miRNAs did not have strong power to discriminate NS patients from controls and differentiate histological subtypes of NS in our study. However, their combined levels could distinguish NS patients from healthy control and also discriminate subtypes of NS with either reliable sensitivity or specificity. In our study, there was a positive

strong association between miR-30c and miR-186 in both PBMCs and plasma samples of NS patients, but not between plasma and PBMCs samples.

All these studies along with the present work indicated that levels of microRNA are dysregulated in NS patients with different histopathology; however, the data are not consistent. Several reasons may be accounted for the differences between miRNA levels in different reports, including patients' related factors (age, stage of the disease, genetic background), sample size, different ethnic population, different sample sources (serum, plasma, PBMCs, PBLs, podocyte, and kidney biopsy), and the method of miRNAs evaluation (use of different internal control, control group). Moreover, treatment strategy (the type of drugs and course of treatment at the time of sampling) would be another possible explanation for divergent miRNA expression.

Besides, we performed a bioinformatics analysis to identify the possible targets of the studied genes in the pathogenesis of NS. MiR-30s can guard podocytes against apoptosis by targeting p53 and Notch1 and the loss of miR-30s eases podocyte damage. Moreover, miR-30s stabilize the podocyte cytoskeletons by targeting some other genes.¹⁷ Additionally, miR-30s can regulate calcineurin/calcium signaling in podocytes.²⁵ Overexpression of miR-30c could reduce Notch1 protein levels in the podocytes.¹⁷ Our bioinformatics analysis along with the literature review predicted that miR-30 family may participate in the pathogenesis of NS by targeting different genes involved in TGF- β , Notch1, and p53 signaling pathways. It is also reported that the TGF- β 1 pathway has an important role in the pathogenesis of FSGS.²⁶⁻²⁸ The reduced number of podocytes induced by

Table 3 Results of the ROC Curve Analysis for Discriminating Potential of PBMC and Plasma miR-30c and miR-186 Levels Among Pairs of the Studied Groups

	AUC	95% CI	P-value	Sensitivity	Specificity	Youden Index J	Associated Criterion
NS and controls							
Plasma FC30c	0.50	0.418 to 0.585	0.97	87	30	0.13	>0.02
Plasma FC186	0.53	0.419 to 0.591	0.92	54	29	0.16	≤2.58
PBMC FC30c	0.65	0.565 to 0.728	0.001	46	96	0.4158	>2
PBMC FC186	0.45	0.418 to 0.582	I	0	100	0	>0
FSGS and MGN							
Plasma FC30c	0.52	0.421 to 0.621	0.72	65	54	0.1849	>0.66
Plasma FC186	0.61	0.499 to 0.707	0.07	82	43	0.2472	>0.19
PBMC FC30c	0.54	0.437 to 0.647	0.48	89	30	0.1902	≤3.56
PBMC FC186	0.5	0.400 to 0.600	0.0	0	100	0	>0

Abbreviations: AUC, area under the curve; CI, confidence interval; ROC, receiver operating characteristic; NS, nephrotic syndrome; MGN, membranous glomerulonephritis; FSGS, focal segmental glomerulosclerosis; FC, fold change; PFC, plasma FC.

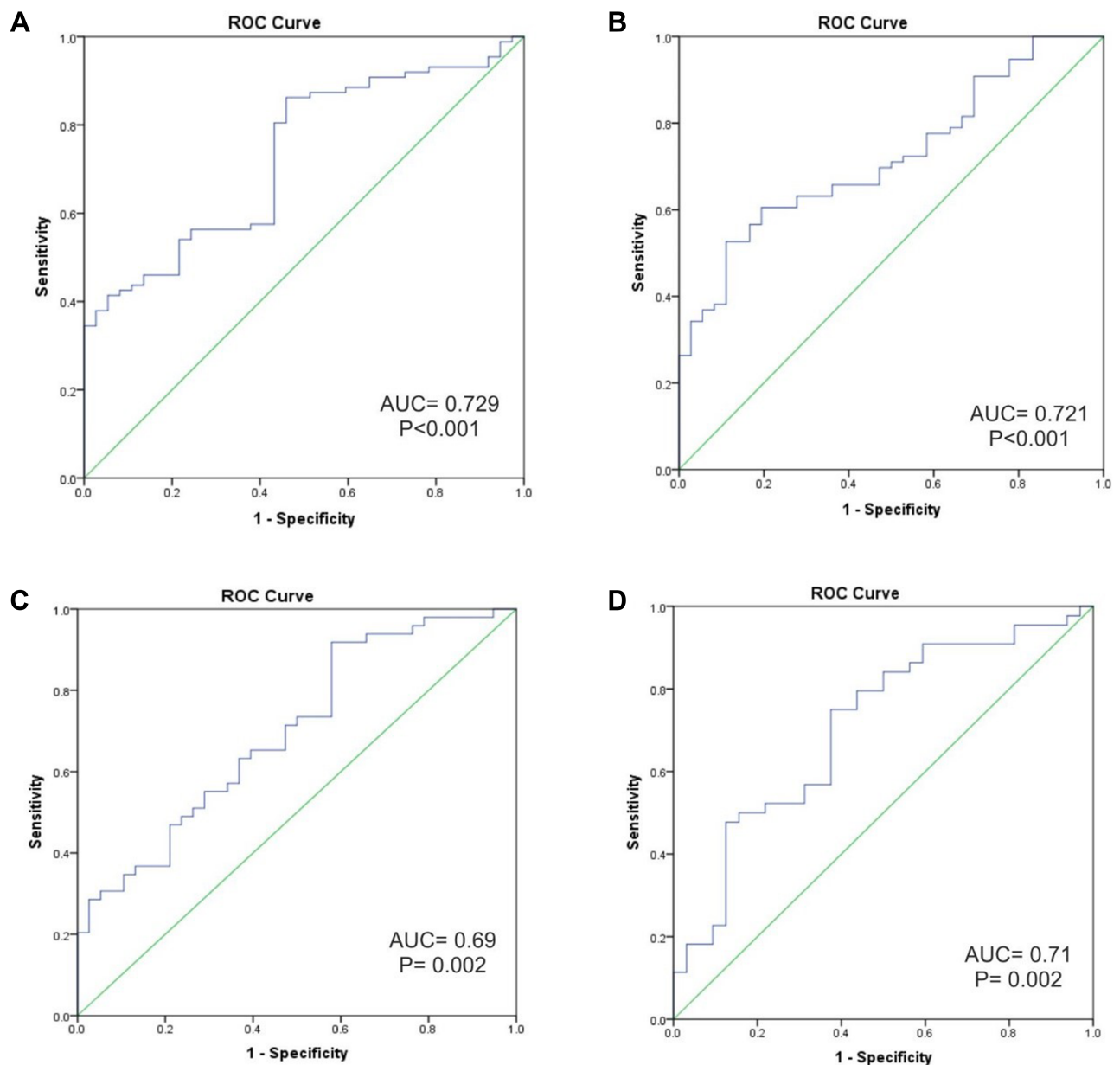


Figure 2 The combined ROC curves analysis of miR-30c and miR-186 in plasma and PBMCs samples. **(A)** Plasma and **(B)** PBMCs levels of the studied miRNAs could discriminate NS patients from controls. **(C, D)** ROC curve for the potential of the studied miRNAs in discriminating FSGS from MGN patients in **(C)** Plasma and **(D)** PBMCs. AUCs and the corresponding statistics of each miRNA between the groups are shown in [Table 3](#). **Abbreviations:** ROC, receiver operating characteristic; AUC, area under the curve.

apoptosis has a serious role in the development of MGN and decreased levels of miR-186 leads to an increase in its target genes Toll-like receptor 4 (TLR4), caspase-3, and P2X7 that are involved in apoptosis²¹ and may contribute to the development of FSGS.

A small sample size was the possible limitation of the present study. Different results from biological resources were considered as another limitation. We used strict criteria for the collection of the patients to validate the reported

candidate miRNAs and to lessen potential unclear factors on the miRNAs expression. Since NS (especially FSGS subtype) has heterogeneous pathogenesis,²⁸ it is difficult to identify an ideal biomarker for specific pathology of NS in all cases. This heterogeneity needs to study a large cohort to validate the reported microRNAs as diagnostic and prognostic biomarkers in patients with different subclasses of NS. Additionally, a potential diagnostic/prognostic value of miRNAs in MGN and FSGS requires further studies.

Table 4 The Ten Top Enriched Biological Processes Among Target Genes of miR-30c and miR-186

GO ID	Biological Process	C	O	E	R	p-value ^a
miR-30c						
GO:0070671	Response to interleukin-12	49	9	1.518943743	5.925170068	1.67E ⁻⁰⁵
GO:0040029	Regulation of gene expression, epigenetic	258	22	7.997703789	2.750789549	1.82E ⁻⁰⁵
GO:0001763	Morphogenesis of a branching structure	196	18	6.075774971	2.962585034	3.96E ⁻⁰⁵
GO:0002009	Morphogenesis of an epithelium	480	32	14.87944891	2.150617284	4.08E ⁻⁰⁵
GO:0045023	G0 to G1 transition	45	8	1.394948335	5.734979424	6.30E ⁻⁰⁵
GO:0031050	dsRNA processing	48	8	1.487,944,891	5.37654321	1.02E ⁻⁰⁴
GO:0016569	Covalent chromatin modification	468	30	14.50746269	2.067901235	1.43E ⁻⁰⁴
GO:0104004	Cellular response to environmental stimulus	320	23	9.919632606	2.318634259	1.67E ⁻⁰⁴
GO:1,990,823	Response to leukemia inhibitory factor	95	11	2.94489093	3.735282651	1.69E ⁻⁰⁴
GO:0010608	Posttranscriptional regulation of gene expression	486	30	15.06544202	1.9913123	2.74E ⁻⁰⁴
miR-186						
GO:0016071	mRNA metabolic process	765	79	30.80383	2.564616	8.88E ⁻¹⁵
GO:0010629	Negative regulation of gene expression	1733	127	69.78175	1.81996	1.01E ⁻¹¹
GO:0006412	Translation	613	62	24.68333	2.511817	2.06E ⁻¹¹
GO:0006396	RNA processing	887	78	35.71633	2.183875	5.11E ⁻¹¹
GO:0043043	Peptide biosynthetic process	636	62	25.60946	2.420981	9.58E ⁻¹¹
GO:0051276	Chromosome organization	1143	92	46.02454	1.998933	1.00E ⁻¹⁰
GO:0044265	Cellular macromolecule catabolic process	1108	88	44.61522	1.972421	5.32E ⁻¹⁰
GO:0008380	RNA splicing	417	46	16.79111	2.739545	5.62E ⁻¹⁰
GO:0043604	Amide biosynthetic process	766	68	30.8441	2.204636	6.64E ⁻¹⁰
GO:0006397	mRNA processing	487	50	19.60976	2.549751	1.20E ⁻⁰⁹

Note: ^aThe Benjamini–Hochberg adjusted *p*-value.

Abbreviations: C, total number of genes in the category; O, number of observed target genes in the category; E, number of expected genes in the category; R, fold enrichment.

Conclusion

We investigated the expression levels of miR-30c and miR-186 in PBMC and plasma samples of patients with NS. There were significant differences between the NS group and its subtypes when compared to controls. The combined levels of the studied miRNAs could distinguish NS patients from healthy control with a moderate accuracy. Further investigations are essential to validate a panel of miRNAs with diagnostic and prognostic values in NS patients with different histopathology.

Ethical Statement

This study was conducted in accordance with the Declaration of Helsinki. This study was certified by the Ethics Committee of Tabriz University of Medical Sciences, Tabriz, Iran (IR.TBZMED.REC.1396.912). By providing written informed consent, subjects agreed to participate in this study.

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Author Contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

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Disclosure

The authors declared no conflicts of interest.

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