Overexpression of miR-652-5p in new onset type 1 diabetes

ABSTRACT

Introduction. MicroRNAs (miRNAs) are small non-coding RNA regulating gene expression at the post-transcriptional level. miRNAs have emerged as an important regulators of central and peripheral immune tolerance, therefore study the RNA molecules in the context of type 1 diabetes (T1D) pathogenesis is an important issue. The aim of this study was to investigate miR-652-5p expression level in the new onset T1D and an impact on ADAR and MARCH5, potential target genes.

Material and methods. The miR-652-5p expression was investigated in the peripheral blood mononuclear cell of newly diagnosed T1D pediatric patients (n = 28) and age-matched controls (n = 28) by quantitative reverse-transcription polymerase chain reaction (qRT-PCR). miRNA targets were analyzed by luciferase reporter assays.

Results. Expression analysis revealed upregulation of miR-652-5p in T1D group compared to non-diabetic controls (p < 0.05). Luciferase reporter assay did not indicated ADAR and MARCH5 as miR-652-5p targets.


Key words: T1D, expression, miR-652-5p, ADAR, MARCH5

Introduction

Type 1 diabetes (T1D) is a common autoimmune disorder caused by the T-cell mediated destruction of the insulin-producing pancreatic beta cells. The pathogenesis of T1D is a consequence of complex interplay between genetic, epigenetic and environmental factors [1].

Linkage studies have indicated the HLA class II genes at 6p21 as the major T1D susceptibility locus [2]. The concordance rate for monozygotic twins with high-risk HLA genotypes DR3-DQ2 and DR4-DQ8 and family history of T1D is estimated to be ~50% [3]. To date, over 50 non-HLA susceptibility gene markers have been identified, mostly related to innate and adaptive immune response (http://www.t1dbase.org). These include the cytotoxic T-lymphocyte-associated protein 4 gene (CTLA-4), the protein tyrosine phosphatase, non-receptor type 22 gene (PTPN22), the interleukin 2 receptor subunit alpha gene (IL2RA) and the interferon-induced with helicase C domain 1 gene (IFIH1) [4–7]. Among the environmental risk factors: cow’s milk, vitamin D, glycolotoxin, intestinal gut microbiota and enteroviruses might play a potential role in pathogenesis and progression of T1D [8–11]. Furthermore, epigenetic
agents: DNA methylation, histone modification and microRNA signaling, may contribute to development of autoimmune disorders [12, 13]. MicroRNAs (miRNAs) are short (~22 nucleotides) non-coding RNAs regulating gene expression at the post-transcriptional level. miRNAs binds to 3'UTR of target genes and repress their expression by inhibition of translation or mRNA degradation. Whereas miRNA regulation of particular target results in modest changes in genes expression, the network activity of miRNAs might induce tremendous changes in cell behaviour [14]. Variations in gene expression, influenced by miRNAs are observed in immune system, where miRNAs modulate lymphocytes development and maturation [15–17]. Therefore, deregulation of miRNAs, which are involved in peripheral and central lymphocyte fate decision might promote autoimmunity [18–21]. Last decade demonstrated extensive efforts to examine miRNAs as T1D diagnostic and prognostic markers [22].

Our previous analysis of miRNA expression profile on Affymetrix miRNA 4.1 array in new onset T1D group indicated overexpression of miR-652-5p in patients presenting severe form of autoimmune disease [23]. We observed statistically significant upregulation of miR-652-5p in patients with initial diabetic ketoacidosis (DKA) compared to healthy controls (p = 0.04, fold change 4.72). The patients without DKA did not present overexpression of miR-652-5p. The aim of the study was to validate by qRT-PCR the microarray experiment in enlarged group of newly diagnosed T1D patients and investigate miR-652-5p potential target genes. Expression analysis revealed upregulation of miR-652-5p in T1D group compared to non-diabetic controls (p < 0.05), however we did not confirm overexpression of miR-652-5p in patient with DKA. Luciferase reporter assay did not indicated ADAR and MARCH5 as miR-652-5p targets.

### Material and methods

#### Study groups

The expression analysis of miR-652-5p was conducted in 28 newly diagnosed T1D subjects (mean age ± SD 11.21 ± 3.3 years) and 28 age-matched healthy donors (mean age ± SD 10.5 ± 4.1 years). Patients were recruited at the Department of Pediatric Diabetes and Obesity, Poznan University of Medical Sciences. The diagnosis of diabetes was based upon the WHO criteria. Autoimmune origin of the disease was confirmed by positive serum autoantibodies to insulin (IAA) and/or glutamic acid decarboxylase (GADA) and/or islet antigen-2 (IA2A). The T1D cohort was further divided into severe and moderate onset, based upon the presence or absence of diabetic ketoacidosis (DKA) at initial presentation. The subgroup with initial DKA presented with more frequent IA2A compared to milder cases of T1D (92% vs. 75%). This finding suggests particularly intense autoimmune reaction and more vigorous beta cell destruction. In line, patients with severe T1D onset presented lower residual insulin synthesis reflected by lower fasting C-peptide, although the difference was statistically non-significant. Clinical characterization of patients is summarized in Table 1. To minimize the interference of the initial metabolic disorder in miRNA expression, PBMCs were collected from fully rehydrated patients with normalized ketonaemia and well-controlled glycaemia, 10 days after introduction of

### Table 1. Clinical features of the studied cohorts of patients with type 1 diabetes

<table>
<thead>
<tr>
<th></th>
<th>T1D</th>
<th>Severe T1D onset</th>
<th>Mild T1D onset</th>
</tr>
</thead>
<tbody>
<tr>
<td>n = 28 (%)</td>
<td></td>
<td>n = 12 (%)</td>
<td>n = 16 (%)</td>
</tr>
<tr>
<td>Gender F/M</td>
<td>4/24</td>
<td>1/11</td>
<td>3/13</td>
</tr>
<tr>
<td>Age, y</td>
<td>11.21 ± 3.33</td>
<td>11.33 ± 3.63</td>
<td>11.63 ± 3.18</td>
</tr>
<tr>
<td>BMI [kg/m²]</td>
<td>17.33 ± 3.05</td>
<td>17.31 ± 3.18</td>
<td>17.62 ± 3.09</td>
</tr>
<tr>
<td>HbA₁₀ (%)</td>
<td>10.66 ± 1.67</td>
<td>10.59 ± 1.15</td>
<td>11.16 ± 1.86</td>
</tr>
<tr>
<td>C-peptide [nmol/l]</td>
<td>0.30 ± 0.08</td>
<td>0.27 ± 0.9</td>
<td>0.32 ± 0.08</td>
</tr>
<tr>
<td>25-OH-D [ng/ml]</td>
<td>18.19 ± 7.68</td>
<td>13.98 ± 3.27*</td>
<td>21.73 ± 8.76</td>
</tr>
<tr>
<td>DKA</td>
<td>12 (43)</td>
<td>12 (100)</td>
<td>16 (0)</td>
</tr>
<tr>
<td>IAA</td>
<td>7 (25)</td>
<td>3 (25)</td>
<td>4 (25)</td>
</tr>
<tr>
<td>GADA</td>
<td>21 (75)</td>
<td>9 (75)</td>
<td>12 (75)</td>
</tr>
<tr>
<td>IA2A</td>
<td>23 (82)</td>
<td>11 (92)</td>
<td>12 (75)</td>
</tr>
</tbody>
</table>

F — female; M — male; BMI — body mass index; HbA₁₀ — glycated haemoglobin A₁₀; 25-OH-D — 25-hydroxyvitamin D; DKA — diabetic ketoacidosis; IAA — antibodies to insulin; GADA — antibodies to glutamic acid decarboxylase; IA2A — antibodies to islet antigen-2; clinical features presented as mean ± standard deviation; number of subjects (%); *p < 0.01 — p values estimated by unpaired t-test, severe T1D onset vs. mild T1D onset subgroup.
insulin therapy. Control samples with negative personal and family history of autoimmunity and no clinical signs of autoimmune disorders were obtained from an outpatient pediatric practice. The study was approved by the local Ethics Committee at the Poznan University of Medical Sciences (decision No 656/15) and all procedures were in accordance with the Declaration of Helsinki. Informed consent was obtained from parents/legal representatives of the minor patients.

miRNA expression analysis

The miR-652-5p expression was assessed by quantitative reverse-transcription PCR (qRT-PCR). 10 ng of total RNA isolated from PBMCs was used in RT reaction performed with TaqMan Advance miRNA cDNA Synthesis Kit (Applied Biosystem, Thermo Fisher Scientific, US) according to the manufacturer’s protocol. qRT-PCR was performed using 5 µl of diluted 1:10 cDNA samples, TaqMan Fast Advance Master Mix (Applied Biosystem, CA, US) and TaqMan Micro RNA assay (Applied Biosystem, CA, US) in a total volume of 20 µl. PCR reactions were run on BioRad CFX96 Real Time PCR instrument (BioRad Laboratories, CA, US). The thermal cycling conditions were an initial uracil-N-glycosylase activation for 2 minutes at 50°C, polymerase activation for 20 seconds at 95°, followed by 40 cycles of 3 seconds at 95° and 30 seconds at 60°C. All reactions were run in triplicate. RNU6 small nuclear RNA was quantified as control to normalize the differences in RNA levels. The TaqMan Advance assay 479132_mir and NR_002752.2 was used to determine the expression of miR-652-5p and RNU6, respectively. Mean cycle threshold (Ct) values were estimated with BioRad CFX Manager 3.1 software. Relative expression levels were calculated using the 2−ΔΔCt formula.

miRNA target gene prediction

Predicted target genes for miR-652-5p were obtained from miRWalk2.0 (available at http://www.umm.uni-heidelberg.de/apps/zmf/mirwalk). Subsequently the miRNA target genes were verified with regard to their potential role in immune response and T1D pathogenesis. InnateDB (http://www.innatedb.com) and Gene Entrez database with annotation “T1D” were screened. Using the Venn’s diagram (Venny 2.1.0, http://bioinfogp.cnb.csic.es/tools/venny) we identified 22 immune response genes. For further investigation 2 autoimmune related genes: Adenosine Deaminase, RNA Specific (ADAR) [24] and Membrane Associated Ring-CH Finger Protein 5 (MARCH5) [25] were selected. The sequences of miRNA bindings sites in 3’UTR of target genes were verified by PITA-Segal lab software (https://genie.weizmann.ac.il).

Reporter constructs

Binding of ADAR and MARCH5 by miR-652-5p was tested by luciferase reporter assay. The 3’UTRs of selected genes were obtained from the University of California Santa Cruz genome browser on human genome hg38 assembly. 1195 bp and 1031 bp fragments of ADAR and MARCH5 3’UTRs, respectively, were amplified from DNA of Jurkat E6.1 cell line (ATCC®, VA, US) purified with Gentra Puregene Blood kit (Qiagen Sciences, US). 70 ng of DNA was used as a template in 50 µl PCR reaction with 1X Phusion Hot Start II DNA Polymerase (Thermo Scientific, CA, US), 1X Phusion GC buffer, 200 µM dNTP, 500 nM forward and reverse primer. Sgfl restriction site was introduced into forward primer. Purified PCR products were cloned into the pGEM-Teasy vector (Promega, WI, US). Subsequently, the inserts were cleaved out using Sgfl and Noti restriction endonuclease (New England BioLabs, MA, US) and subcloned into psiCHECK2 vector (Promega, WI, US) downstream the Renilla luciferase (RL) reporter gene. The vectors’ sequence was confirmed by sequencing.

Luciferase assay

Luciferase reporter assay was performed using Promega-Dual Luciferase Reporter Assay System (Promega, WI, US). 0.25 × 10⁶ Jurkat cells were co-transfected with 1 µg of psiCHECK2-3’UTR vector and 50 pmol of miR-652-5p mimic or negative control #1 (both Ambion, CA, US) using Neon Transfection System and parameters 1600 V, 10 ms, 3 pulses. Transfected cells were dispensed into 500 µl of RPMI 1640 medium with 10% FBS, 0.25% glucose, 10 mM HEPES, 1 mM sodium pyruvate and incubated in 37°C/5% CO₂. Transfections were performed in three independent experiments in triplicate. Cells lysates were made 24 h post transfection. Renilla and Firefly luciferase activity was measured in duplicate for each transfection on GloMax®96 Microplate Luminometer (Promega, WI, US). The expression of Renilla luciferase were normalized to secondary reporter gene Firefly luciferase. The RL/FL luciferase ratios were compared to negative control (set at 1.00). Significance was estimated with unpaired t-test.

Statistical analysis

Statistical analysis was performed using GraphPad Prism 5 (GraphPad Software Inc., CA, US). D’Agostino and Pearson omnibus normality test was applied to estimate if the values come from a Gaussian distribution. Data normally distributed are presented as mean ± SD. To compare variance within each group the F test was used. Statistical significance of the differences between means was determined with the unpaired t-test with Welch’s correction. Data without Gaussian distribution
are presented as median with interquartile range. Statistical significance of the differences between medians was estimated with the Mann-Whitney test. P values < 0.05 was considered statistically significant.

**Results**

The expression analysis by qRT-PCR revealed statistically significant upregulation of miR-652-5p in new onset type 1 diabetes patients compared to non diabetic controls (fold change 1.45, p = 0.035) (Figure 1). Stratification patients group according to the presence or absence of initial diabetic ketoacidosis did not indicate the significant differences in miR-652-5p expression level compared to controls as well as patients without DKA (p > 0.05). miR-652-5p did not show significant differences between DKA(+)+ and DKA(−)− groups. Horizontal lines indicate median with range; asterisks indicate significance, with P values estimated by Mann-Whitney test. T1D — type 1 diabetes patients; C — controls; DKA(+) — cohort of patients with initial diabetic ketoacidosis, DKA(−) — cohort of patients without initial diabetic ketoacidosis.

**Figure 1.** miR-652-5p expression analysis in T1D patients. A. miR-652-5p is significantly differentially expressed between new onset T1D and controls (*p < 0.05); B. miR-652-5p expression level in T1D group stratified according to the presence of initial diabetic ketoacidosis (DKA). The patients with diabetic ketoacidosis did not present statistically significant differences in miR-652-5p expression level compared to controls as well as patients without DKA (p > 0.05). miR-652-5p did not show significant differences between DKA(+) and DKA(−) groups. Horizontal lines indicate median with range; asterisks indicate significance, with P values estimated by Mann-Whitney test. T1D — type 1 diabetes patients; C — controls; DKA(+) — cohort of patients with initial diabetic ketoacidosis, DKA(−) — cohort of patients without initial diabetic ketoacidosis

**ADAR and MARCH5 are not targeted by miR-652-5p**

*In silico* analysis was performed to indicate target genes of miR-652-5p with potential role in autoimmunity. In the cluster of 22 immune response genes we pinpointed *ADAR* and *MARCH5* for further investigation. 3’UTRs of these genes were cloned into psiCHECK2 luciferase vector.

Co-transfection of psiCHECK2*-ADAR* or psiCHECK2*-MARCH5* and miR-652-5p mimic into Jurkat cells did not indicate significant differences in relative luciferase activity compared to negative control (p >0.05, Figure 2).

**Discussion**

The global incidence of T1D has been increasing by 2–3% per year, with the most rapid increment occurr-
Magdalena Zurawek et al., Overexpression of miR-652-5p in new onset type 1 diabetes.

Figure 2. Luciferase reporter assay did not indicate ADAR and MARCH5 as targets of miR-652-5p. Jurkat cells were co-transfected with luciferase 3’UTR reporter vectors containing 3’UTRs fragment of ADAR and MARCH5 genes in the presence of miR-652-5p mimic or negative control. Average Renilla/Flucyluciferase ratio (RL/FL) from three independent experiments is presented ± SD, values for the negative control were set to 1.00 (t test p > 0.05).

ring among the youngest children (< 5 years of age). A recent decrease in high-risk HLA genetic contribution in new-onset cases versus overall increase in T1D incidence might suggest the changing environmental pressure on disease development and involvement of other genes and epigenetic factors [26].

Last decade demonstrate extensive efforts in understanding the role of specific miRNAs in pathogenesis of T1D, which pinpoint miRNAs as diagnostic and prognostic biomarkers of autoimmune disorders.

In current study we observed overexpression of miR-652-5p in newly diagnosed T1D patients compared to healthy controls. T1D patients presented ongoing autoimmune process indicated by positive serum autoantibodies to insulin (IAA) and/or glutamic acid decarboxylase (GADA) and/or islet antigen-2 (IA-2A). Expression analysis did not display significantly higher miR-652-5p expression level in patients with severe form of T1D compared to control group. The observation stays in opposite to our previous microarray experiment [23]. The discrepancy in expression level between microarray and validation analysis might be partially explained by the relative small sample size of microarray cohort (patients with DKA, n = 7). Patients with initial diabetic ketoacidosis presented trend of miR-652-5p upregulation compared to patients without DKA, although the differences did not reach statistical significance. miR-652-5p has not been previously reported in the context of autoimmune disease and it is likely to be a potential biomarker of new onset T1D. Limitation of current investigation is small sample size, therefore further extension of study group is required to affirm miR-652-5p as epigenetic T1D risk factor.

Furthermore, studies concerning miRNAs expression analysis in PBMCs and serum of adult type 1 diabetes patients, revealed other miRNAs that might be associated with autoimmunity [27–29]. miR-326 was found to be upregulated in PBMCs from autoantibody-positive T1D patients [29]. In mice, overexpression of miR-326 during experimental autoimmune encephalomyelitis leads to increased Th17 cell differentiation [30]. In contrast, Salas-Perez et al. observed downregulation of miR-21a and miR-93 in PBMC from T1D patients [31]. Interestingly, the in silico analysis of their target genes revealed pro-apoptotic and pro-inflammatory Caspase 8, Caspase 7 and IL-8 for miR-93 as well as Fas ligand, MTPN, APAF1, IL-12A, IL-22 and IL-1B genes for miR-21.

Recent studies demonstrated miRNAs as potential tool for disease staging. It was indicated that decreased miR-146 expression in PBMCs correlate with ongoing islet autoimmunity and high serum GADA titers [28]. In addition, dysregulation of miR-125b-5p, miR-365a-3p, miR-5190 and miR-770-5p was associated with hyperglycemia in patients with T1D and might contribute to the development of diabetes complications [32]. Pathway analysis revealed that 50 Kyoto Encyclopedia of Genes and Genomes pathways were significantly enriched by genes targeted by these four miRNAs [32].

An important strategy for understanding the role of miRNAs in pathogenesis of autoimmune disease is to identify their target genes. We are particularly interested in the potential mechanism linking innate immunity and development of type 1 diabetes. Therefore, we validated ADAR and MARCH5 as potential miR-652-5p targets by luciferase reporter assay. The mutations in ADAR were indicated to cause the autoimmune disorder Aicardi-Goutières syndrome [24, 33, 34]. As in Adar1-null mice, the human disease state is associated with upregulation of interferon-stimulated genes [33]. The study of Domínguez-Gutierrez et al. showed overexpression of ADAR in Systemic Lupus Erythematosus (SLE) patients [35]. In addition, decreased MARCH5 protein might account for the constant Mitochondrial Antiviral-Signaling Protein aggregations and abnormalities of type I interferon levels in SLE patients [25]. Current study did not indicate ADAR and MARCH5 as miR-652-5p targets in luciferase reporter assay. Therefore, we hypothesize that miR-652-5p is unlikely to regulate ADAR and MARCH5 expression.

In conclusion, current study revealed miR-652-5p as potential marker of new onset type 1 diabetes. Investigation miRNAs in context of autoimmunity is important issue as improved biomarkers are required for the detection of developing type 1 diabetes, prior to critical loss of pancreatic beta cells.
Conflicts of interests
All authors declare that there are no conflicts of interest associate with this publication.

Acknowledgments
The study was supported by Polish Diabetes Association (Polskie Towarzystwo Diabetologiczne), Artur Czyzyk Research Grant 2015.

Disclosure statement: The authors have nothing to disclose

REFERENCE


