Impact of Acute Bronchiolitis on Cardiac Functions and Serum microRNA-122 and 499

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Abstract: Bronchiolitis is an acute inflammatory injury of the bronchioles that is usually caused by a viral infection, most commonly Respiratory Syncytial Virus (RSV). This study aimed to verify the direct injurious effect of acute bronchiolitis on the myocardium and serum level of microRNA-122 and 499 (miR-122 and miR-499) of infected infants. Serum lactate dehydrogenase (LDH), creatine phosphokinase-isoenzyme MB (CK-MB) and cardiac troponin I (cTnI) were significantly elevated in mild to moderate group as well as severe group except cTnI which showed non-significant elevation in mild to moderate group, compared to control group. Quantitative Reverse Transcriptase Polymerase Chain Reaction (qRT-PCR) was carried out to evaluate the serum level of miR-122 and miR-499. Sharp elevation in serum miR-122 was observed in severe cases, while miR-499 was statistically non-significant as compared to both normal and mild to moderate groups. The flow velocities across the pulmonary, tricuspid and mitral valves were recorded using standard echocardiographic techniques. Echocardiographic data showed significant reduction in pulmonary At/Et ratio in severe bronchiolitis group, compared to both normal and mild to moderate groups, respectively. Parallel reduction was also observed in tricuspid and mitral E/A ratios, compared to both normal and mild to moderate groups, respectively. Severe bronchiolitis causes cardiac dysfunction due to hypoxia, cardiac strain and pulmonary hypertension in addition to up-regulation of serum miRNA-122.

Keywords: Acute Bronchiolitis, Cardiac Troponin I, microRNA-122

Introduction

Bronchiolitis “is a seasonal viral illness, characterized by fever, nasal discharge and dry, wheezy cough. On examination, there are fine inspiratory crackles and/or high-pitched expiratory wheeze” (Lakhanpaul et al., 2009). This is most commonly caused by Respiratory Syncytial Virus (RSV) which is the major respiratory pathogen in young children (<1 year of age) and virtually every individual has been infected with RSV at least once by the age of 3 (Bruce and Alcorn, 2011). The virus is a member of the Paramyxoviridae family, with a single-stranded, negative-sense RNA genome (Brock et al., 2003). It replicates in nasopharyngeal epithelium and then spreads to lower respiratory tract one to three days later (Thorburn and Hart, 2006). Together with influenza virus, RSV is also the most common cause for admissions in adults with chronic cardiac and pulmonary disorders and acute respiratory failure (Carrat et al., 2006). The importance of extra-pulmonary manifestations of RSV infection has become evident (Eisenhut, 2006). Although the most frequent extra-pulmonary manifestations of RSV infection involve the cardiovascular system, the reasons leading to heart involvement during RSV infection are not fully known (Esposito et al., 2010).

MicroRNAs (miRNAs) are endogenous, single-stranded, 22-nucleotide non-coding RNAs. Individual miRNAs modulate the expression of collections of mRNA targets that often have related functions, thereby governing complex biological processes. MiRNAs are generally regarded as negative regulators of gene expression through inhibition of translation and/or promotion of mRNA degradation by base-pairing to complementary sequences within the 3'Untranslated Region (3'UTR) of protein-coding mRNA transcripts (Humphreys et al., 2005; Small and Olson, 2011). Many miRNAs exhibit a tissue-specific
distribution and they appear to play a key role in cell function both under physiological and pathological conditions. The wide ranging functions of microRNAs in the cardiovascular system have provided new perspectives on disease mechanisms and have revealed intriguing therapeutic targets, as well as diagnostics, for a variety of cardiovascular disorders (Small and Olson, 2011). The observation that miRNAs are stable and present in the circulation has led to a rapidly growing number of reports on the use of these molecules as biomarkers for various diseases, for example plasma concentration of miRNA-499 is a useful biomarker of myocardial injury in humans (Esposito et al., 2010; Gidløf et al., 2011).

The objective of this study was to verify the direct injurious effect of acute bronchiolitis on the myocardium of affected infants and on serum level of miRNA-122 and 499 (miR-122 and miR-499).

**Subjects and Methods**

**Study Design**

This study was carried out at the Department of Pediatric, Al-Hussin University hospital, Egypt, during the winter season 2012-2013. Written informed consent to study participation was obtained from the patients’ parents or legal guardians who were well informed in advance about the purpose of the study.

The study involved 26 infants aged 1-9 months who were admitted with acute bronchiolitis. The exclusion criteria were the presence of a chronic disease increasing the risk of complications of respiratory infection, including chronic disorders of the pulmonary or cardiovascular system, chronic metabolic disease, neoplasms, kidney or liver dysfunction, hemoglobinopathies, immunosuppression and genetic or neurological disorders.

The control group (con) consists of 13 apparently healthy children. None of these children were admitted to the pediatric intensive care unit and none were diagnosed with acute bronchiolitis.

Upon admission, the infants’ demographic characteristics and medical history were systematically recorded using standardized written questionnaires and, after a complete physical examination, the infants with a diagnosis of acute bronchiolitis based on well-established criteria (Crowe, 2011) were enrolled. The severity of the disease was defined on the basis of a global evaluation of the signs and symptoms. In particular, on the basis of previously published criteria (Scarfone, 2005), respiratory illness was considered severe in the presence of all of ≤92% pulse oximetry, a respiratory rate of ≥60 breaths/min, marked accessory muscle use, nasal flare or grunting, an inability to feed and a cyanotic appearance.

Consequently, infants with acute bronchiolitis were divided into two groups (13 patients each): Infants with mild to moderate bronchiolitis (mild and moderate) group and infants with severe bronchiolitis (severe) group.

**Hematological and Biochemical Studies**

Venous blood samples were obtained at the time of admission. A part of blood sample was taken on EDTA as whole blood sample for Complete Blood Counting (CBC) and blood gas analysis. Another part was taken in a plain tube without any anticoagulant for separation of serum by centrifugation at 3000 r.p.m for 10 min and then stored at -70°C.

Blood cell counting was carried out by cell counter fully automated (sysmex, Japan). Blood gases were estimated by blood gases and electrolyte analyzer (GEM premier 3000, USA). C-Reactive Protein (CRP) was measured using semi-quantitative CRP latex test kit (Plasmatec, UK) to eliminate bacterial infection. To evaluate myocardial function, serum creatine phosphokinase-isoenzyme MB (CK-MB) and lactate dehydrogenase (LDH) were assessed using colorimetric assay (Abcam, USA) and cardiac troponin I (cTn I) concentration was measured using immunoenzymatic assay (Accubind, USA) according to the manufacturer’s instructions.

**qRT-PCR**

Quantitative Reverse Transcriptase Polymerase Chain Reaction (qRT-PCR) methodologies have been widely applied to miRNA research, especially in assessing the low level of certain serum miRNAs. Therefore, this system was applied to the present study to determine the circulating levels of miRNA-122 and 499. Total RNA was extracted from serum according to the method of Bravo et al. (2007), using high pure miRNA isolation kit (Roche Diagnostics, Switzerland). Reverse transcription was performed using the Transcript or First Strand cDNA Synthesis Kit (Roche Diagnostics). Primer sequences for each miRNA and control gene (Glyceraldehyde-3-phosphate dehydrogenase (GAPDH) are listed in Table 1 (primers were designed and provided by TIB MOLBIOL (Berlin, Germany). cDNA was amplified through PCR using Light Cycler Fast Start DNA Master Plus SYBR Green I (Roche Diagnostics). PCR was performed at 95°C for 10 min followed by 45 cycles of denaturation at 95°C for 10 sec, annealing (primer dependent) for 60 sec and extension at 72°C for 20 sec.

**Echocardiographic Studies**

The echocardiographic studies were made using a real time ultrasound imaging system (GE Vivid 3, China) equipped with 5 and 8 MHz transducers. The
Echocardiographic measurements were made using standard techniques. M-mode measurements were made in accordance with the recommendations of the Committee of M-Mode Standardization of the American Society of Echocardiography. The flow velocities across the pulmonary, tricuspid and mitral valves were recorded from standard pericordial position using pulsed-wave Doppler. Pulmonary arterial blood pressure was assessed by measuring At/Et ratio, where At is the pulmonary acceleration time and Et is the pulmonary ejection time. Right ventricular diastolic function was assessed by measuring tricuspid E/A ratio, where E and A denote early and late tricuspid flow, respectively (Fig. 1). Left ventricular diastolic function was assessed by measuring mitral E/A ratio where E and A denote early and late mitral flow, respectively. Systolic left ventricular function was assessed by determining the Fractional Shortening (FS) and Ejection Fraction (EF) (Gadish et al., 2010). Interpretations of these studies were performed an attending cardiologist blinded as to the cTn I and miRNAs results.

**Statistical Analysis**

All results were expressed as the mean ± Standard Error (SE). Statistical analysis was performed using Statistical Package for the Social Science for Windows (SPSS, version 16.0, Chicago, IL, USA). The data were analyzed by one-way Analysis Of Variance (ANOVA). To compare the difference among the groups, post hoc testing was performed by LSD test. Pearson’s correlation analysis was used to determine the correlation among the parameters assessed. The p-value <0.05 was considered statistically significant.

**Results**

Table 2 illustrates that severe bronchiolitis caused significant increase in respiratory and heart rates (p<0.0001) in addition to the remarkable cyanosis and irritability as compared to normal children. In severe cases, levels of pH, pO$_2$, HCO$_3$ and O$_2$ saturation percent were significantly decreased (p<0.0001), while pCO$_2$ was increased as compared to normal group. Non-significant differences were observed in blood hemoglobin concentration, RBCs count and serum CRP level in both patient groups, compared to normal control.

Table 3 and revealed that cases with severe bronchiolitis showed significant decrease (p<0.0001 and 0.001, respectively) recorded in total WBCs count in both patient groups, lymphocytes showed significant elevations (p<0.0001) in both mild and moderate and severe cases as compared to control group. On the other hand, serum biochemical markers of cardiac function (LDH, CK-MB and cTn I) were significantly elevated in both patient groups except cTn I which showed non-significant elevation in mild and moderate group, compared to control group. Multiple comparison analysis revealed that severe bronchiolitis caused significant elevation in respiratory and heart rates, lymphocyte percent, pCO$_2$, LDH and CK-MB levels as compared to mild and moderate group. Moreover, significant reductions in platelet count, pH, pO$_2$ and O$_2$ saturation percent were observed in severe cases, compared to control group.

Figure 1 shows sharp elevation (p<0.0001) in the logarithm of serum mir-122 level in severe cases as compared to both normal and mild and moderate groups. In spite of the observed down regulation of mir-499 in serum of severe group, it was statistically non-significant, compared to the other two groups.

**Echocardiographic Data**

Table 3 and revealed that cases with severe bronchiolitis showed significant decrease (p<0.0001 and 0.001) in pulmonary At/Et ratio, compared to both normal and mild and moderate groups, respectively.
Fig. 2. Two-dimensional transthoracic echocardiography. (A) Pulsed Doppler echocardiographic through the tricuspid valve of normal case. (B) Pulsed Doppler echocardiographic through the tricuspid valve of severe case showing right ventricular diastolic dysfunction as a result of A wave more than E wave. (C) Pulsed Doppler echocardiographic through the pulmonary valve showing the pulmonary indices (At and Et)
Fig. 3. Pearson correlations of serum miRNA-122 with (A) cTp I \( (r = 0.73, p \leq 0.01) \), (B) pulmonary At/Et \( (r = -0.37, p \leq 0.02) \) (C) tricuspid E/A \( (r = -0.61, p \leq 0.01) \)
Table 2. Characteristics of normal children and patients with mild, moderate and severe bronchiolitis (Mean ± SE)

<table>
<thead>
<tr>
<th>Groups</th>
<th>Con</th>
<th>Mild and Moderate</th>
<th>Severe</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (mon)</td>
<td>4.31±0.59</td>
<td>4.15±0.44</td>
<td>3.35±0.41</td>
</tr>
<tr>
<td>Weight (Kg)</td>
<td>6.54±0.43</td>
<td>5.97±0.21</td>
<td>5.90±0.25</td>
</tr>
<tr>
<td>Respiratory rate (c/m)</td>
<td>34.38±0.81</td>
<td>59.31±1.95&lt;sup&gt;a&lt;/sup&gt;</td>
<td>79.38±1.38&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
<tr>
<td>Heart rate (beat/min)</td>
<td>116.23±1.49</td>
<td>120.46±2.74</td>
<td>135.38±3.24&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
<tr>
<td>Cyanosis</td>
<td>NO</td>
<td>NO</td>
<td>YES</td>
</tr>
<tr>
<td>Irritability</td>
<td>NO</td>
<td>YES</td>
<td>YES</td>
</tr>
<tr>
<td>pH</td>
<td>7.40±0.01</td>
<td>7.37±0.02</td>
<td>7.28±0.003</td>
</tr>
<tr>
<td>pO&lt;sub&gt;2&lt;/sub&gt; (mmHg)</td>
<td>75.23±1.03</td>
<td>79.99±5.64</td>
<td>55.92±1.01&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
<tr>
<td>pCO&lt;sub&gt;2&lt;/sub&gt; (mmHg)</td>
<td>39.92±1.10</td>
<td>33.51±2.53&lt;sup&gt;a&lt;/sup&gt;</td>
<td>51.46±1.00&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
<tr>
<td>HCO&lt;sub&gt;3&lt;/sub&gt; (mmol/L)</td>
<td>26.00±0.97</td>
<td>19.63±0.57&lt;sup&gt;a&lt;/sup&gt;</td>
<td>19.42±0.24&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>O&lt;sub&gt;2&lt;/sub&gt; sat. (%)</td>
<td>93.23±0.56</td>
<td>94.11±0.99</td>
<td>85.23±1.83&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
<tr>
<td>Hb (g/dL)</td>
<td>12.48±0.33</td>
<td>11.67±0.54</td>
<td>12.28±0.54</td>
</tr>
<tr>
<td>RBCs (10&lt;sup&gt;6&lt;/sup&gt; cell/mL)</td>
<td>4.82±0.19</td>
<td>4.48±0.26</td>
<td>5.02±0.10</td>
</tr>
<tr>
<td>WBCs (10&lt;sup&gt;3&lt;/sup&gt; cell/mL)</td>
<td>9.86±1.28</td>
<td>6.02±0.21&lt;sup&gt;a&lt;/sup&gt;</td>
<td>5.48±0.10&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Eosinophyl (%)</td>
<td>2.08±0.24</td>
<td>1.23±0.26&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.08±0.24&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Lymphocyte (%)</td>
<td>27.31±0.87</td>
<td>56.11±2.33&lt;sup&gt;a&lt;/sup&gt;</td>
<td>62.00±1.83&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
<tr>
<td>Monocyte (%)</td>
<td>3.92±0.42</td>
<td>0.54±0.14&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.00±0.28&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Platelet (10&lt;sup&gt;3&lt;/sup&gt; cell/mL)</td>
<td>304.08±18.78</td>
<td>321.38±24.17</td>
<td>257.38±14.35&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
<tr>
<td>CRP (mg/L)</td>
<td>-ve</td>
<td>-ve</td>
<td>-ve</td>
</tr>
<tr>
<td>LDH (mU/mL)</td>
<td>1.95±0.07</td>
<td>2.36±0.09&lt;sup&gt;a&lt;/sup&gt;</td>
<td>2.94±0.02&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
<tr>
<td>CK-MB (ng/mL)</td>
<td>0.99±0.07</td>
<td>1.52±0.07&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.73±0.04&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>cTp I (ng/mL)</td>
<td>1.91±0.14</td>
<td>1.93±0.09</td>
<td>3.06±0.05&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

a: Significance Vs control group, b: Significance Vs mild and moderate group

Table 3. Echocardiographic measurements of children in different studied groups

<table>
<thead>
<tr>
<th>Groups parameters</th>
<th>Con</th>
<th>Mild and Moderate</th>
<th>Severe</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pulmonary At/Et</td>
<td>0.39±0.02</td>
<td>0.36±0.02</td>
<td>0.31±0.02&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
<tr>
<td>Tricuspid E/A</td>
<td>1.56±0.06</td>
<td>1.30±0.11&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.85±0.09&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
<tr>
<td>Mitral E/A</td>
<td>2.15±0.07</td>
<td>1.59±0.04&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.89±0.11&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
<tr>
<td>EF</td>
<td>61.62±1.22</td>
<td>59.92±0.88</td>
<td>61.62±1.04</td>
</tr>
<tr>
<td>FS</td>
<td>31.38±0.40</td>
<td>32.69±0.68</td>
<td>32.77±0.58</td>
</tr>
</tbody>
</table>

a: Significance Vs control group, b: Significance Vs mild and moderate group. EF: Ejection fraction, FS: Fraction shortening

Parallel reduction was also observed in tricuspid and mitral E/A ratios with p values <0.0001 and 0.001 for tricuspid and p<0.001 for mitral blood flow compared to both normal and mild and moderate groups, respectively. Mild and moderate cases showed remarkable reduction in tricuspid and mitral blood flow with p values <0.049 and 0.0001, respectively as compared to normal group (Fig. 2A-C).

Echocardiographic data showed non-significant differences in ejection fraction and fraction shortening in all the studied groups.

To evaluate the significance of increased levels of circulating mir-122, we analyzed whether the level of this miRNA correlated with a set of biochemical and clinical parameters. Figure 3 revealed that the level of circulating mir-122 correlated significantly (p<0.01) with serum cTp I levels (r = 0.73). While, negative correlations were observed between serum mir-122 and both the pulmonary and tricuspid ratios (r = -0.37, p<0.02 and -0.61, p<0.01, respectively).

Discussion

Acute bronchiolitis produces significant morbidity and mortality worldwide every year. RSV bronchiolitis is the most important cause for admission to the pediatric intensive care unit in infants with lower respiratory tract infection (Eisenhut, 2006). In recent years the importance of extra-pulmonary manifestations of RSV infection has become evident.

Cardiac troponin I (cTp I) is an isofrom of a thin filament contractile protein present in high concentrations in the myocardium. It is the only known molecular marker of myocardial injury and is detectable within 6 h of the damage (Thiru et al., 2000). A number of studies have suggested a possible association between cTp I and myocardial injury in patients with non-cardiac diseases (Khan et al., 1999; ver Elst et al., 2000; Checchia et al., 2000; Ammann et al., 2001). Gurkan et al. (2004) reported that the relationship between cTp I as a marker of cell death and myocardial function suggests that “cell death” may have a role in the
pathogenesis of myocardial dysfunction in sepsicaemia. Furthermore, it has been demonstrated in other lung diseases, such as bacterial pneumonia, that severe lung involvement can be accompanied by a significant increase in cTnI and T concentrations (Weinberg et al., 2002; Labugger et al., 2004).

Increased serum level of cTnI in severe group of the present study is in line with previous studies that demonstrated elevation in cTnI in RSV bronchiolitis (Checchia et al., 2000; Moynihan et al., 2003). Initially, increased levels of cTnI were presumed to be due to myocardial necrosis, as happens in myocardial infarction (Martins et al., 2009). Pulmonary embolism is also a relatively frequent cause of increased plasma troponin (Meyer et al., 2000). In the case of pulmonary embolism, myocardial necrosis is not a prominent phenomenon and right ventricular strain could be the cause of troponin release (Martins et al., 2009). Previous work by Feng et al. (2001) illustrated that preload induces troponin I degradation independently of myocardial ischemia. In addition, Chronic Obstructive Pulmonary Disease (COPD) patients frequently have hypoxia and thus the hypothesis that hypoxia also plays a role in troponin release in COPD patients cannot be ruled out (Martins et al., 2009). Moreover, the RSV may play a direct role in causing heart disease since the virus itself was detected in the myocardial tissue and the occurrence of significant pericardial effusion in children with severe RSV bronchiolitis (Bowles et al., 2003; Eisenhut, 2006; Esposito et al., 2010).

Consequently, elevated serum cTnI in the studied severe cases could be a reflection of hypoxaemia that result from alveolar hypoventilation, indicated by elevated pCO2 and decreased arterial blood oxygenation. In addition, tachycardia and subsequent myocardial strain on the right ventricle associated with pulmonary hypertension, that indicated by decreased At/Et ratio, could be another cause for troponin degradation. Regarding mild to moderate cases, serum cTnI, arterial blood oxygenation and pulmonary At/Et ratio were normal confirming our previous assumption. However, both diseased groups suffered from right and left ventricular diastolic dysfunctions as indicated by decreased E/A ratios for tricuspid and mitral valves, respectively.

To our knowledge, no previous studies have been conducted on circulating miRNA-122 and 499 in infants with acute bronchiolitis. miR-122 is abundantly expressed in the liver and plays a role in Hepatitis C virus (HCV) replication by binding to sites within the HCV 5'-UTR and stimulating HCV accumulation in vivo (Jopling et al., 2005). Conversely, overexpression of miR-122 blocked replication of Borna disease virus in oligodendroglial cells (Qian et al., 2010) and inhibited HBV expression in hepatoma cells (Qiu et al., 2010).

These data suggest that miR-122 can have contrasting effects on expression of different viruses. In addition, Bruce and Alcorn (2011) studied the differential miRNA expression profile of fetal human type II epithelial cells to RSV infection. Cells were treated in the absence and presence of RSV for two hours. RSV infection resulted in a significant decrease in miRNA-122. The present study revealed that serum level of miRNA-122 was highly expressed in severe bronchiolitis group as compared to both control and mild and moderate cases. While miRNA-499 was not significantly altered in all groups. These results disagree with Bruce and Alcorn (2011). D'Alessandra et al. (2010) reported that acute myocardial infarction up-regulated miRNA-499 plasma levels, both in humans and mice; whereas miRNA-122 was lower than control patients and these miRNAs represent novel biomarkers of cardiac damage.

It has been suggested that miR-122 down-regulates the expression of Cationic Amino Acid Transporter-1 (CAT-1) mRNA (Chang et al., 2004) through binding its 3'-UTR resulting in decreased protein abundance of CAT-1 (Jopling et al., 2006). CAT-1 is a membrane transporter for the essential amino acid arginine, thus supporting important functions, such as synthesis of Nitric Oxide (NO) (Hatzoglou et al., 2004), which acts as an antiviral agent. The study of Tsutsumi et al. (1999) revealed that in spite the up-regulation of the Inducible Nitric Oxide Synthase (iNOS) in RSV-infected respiratory epithelial cells, NO production specific for RSV infection itself could not be detected by nitrite assay. Moreover, Gadish et al. (2010) investigated the Fractional exhaled Nitric Oxide (FeNO) levels in infants during acute RSV bronchiolitis and during convalescence. They found that the FeNO levels were temporarily reduced during acute RSV bronchiolitis and increased during convalescence to normal levels and higher. Thus the up-regulation of miR-122 in the present study may down-regulate the expression of CAT-1 or/and prevent the translation of iNOS mRNA.

According to Pearson’s correlation, miRNA-499 did not correlate with cTnI and EF, indicating absence of cardiac damage. Whereas, the positive correlations of miRNA-122 with cTnI, pulmonary hypertension and diastolic dysfunction of the right ventricle indicating presence of cardiac dysfunction.

**Conclusion**

Infants with severe acute bronchiolitis suffer from cardiac dysfunction due to hypoxia, cardiac strain and pulmonary hypertension. In spite the up-regulation of serum miRNA-122 in bronchiolitis patients, it cannot be considered as a biomarker. Further research is needed to improve current knowledge of the role of miRNAs particularly 122 in RSV.
Acknowledgement

Authors thank Prof. Dr. Hassan S. Abu Saif, Head of Pediatric Cardiology Unit, Pediatric Department, Al-Azhar University, who performed the echocardiography and explained the data.

Funding Information

The authors have no support or funding to report.

Author’s Contributions

Mona A. Mohamed: Conceived and designed the study. Performed the experiments and analyzed the data.

Khalid M.S. Zayed: Conceived and designed the study. Selected and followed up the included cases. Authors wrote, read and approved the manuscript.

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