Abstract. MicroRNAs (miRs) have a major role in the pathogenesis of hepatocellular carcinoma (HCC). As the insulin-like growth factor (IGF) axis is a highly tumorigenic pathway in HCC, the present study attempted to target it with miRs. Potential targeting of crucial members of the IGF axis by miRNAs at the 3'-untranslated region (3'-UTR) was predicted using bioinformatic tools, such as microrna.org, Diana lab and Targetscan, while 5'-UTR targeting was predicted using bibiserv software. Expression profiling of obtained miRNAs was performed using quantitative polymerase chain reaction (qPCR) in 22 non-metastatic HCC biopsy samples and 10 healthy tissues. To investigate the impact of miRNAs on their potential downstream targets, transfection of miRNAs was performed in HuH-7 cells and the targets' expression was quantified using qPCR. Transcripts of insulin-like growth factor-1 receptor (IGF-1R), insulin-like growth factor binding protein-3 (IGFBP-3) and IGF-II were found to be potentially targeted at the 5'-UTR and 3'-UTR regions by the single clustered hepatic metastamiRs miR-96-5p and miR-182-5p. The two miRs showed a similar expression pattern in HCC tissues compared to those in healthy tissues. Forced expression of miR-96-5p and miR-182-5p in the HCC cell line HuH-7 had inducing effects on IGFBP-3 and IGF-II transcripts. Of note, the two miRs had differential effects on IGF-1R, where miR-96-5p induced IGF-1R mRNA expression and miR-182-5p inhibited its expression. The present study revealed the pleiotropic impact of the single clustered hepatic metastamiRs miR-96-5p and miR-182-5p on IGF-1R, and an inducing effect on IGF-II and IGFBP-3 in hepatocellular carcinoma.

Introduction

MicroRNAs (miRs) are a novel class of small, non-coding, single-stranded RNA molecules (~22 nucleotides). They mainly function via base pairing with the 3'-untranslated region (3'-UTR) of their target mRNAs, thus inhibiting their translation or repressing their levels (1-4). The present study attempted to study the regulation of the 5'UTR and 3'-UTR of insulin-like growth factor-1 receptor (IGF-1R), insulin-like growth factor binding protein-3 (IGFBP-3) and IGF-II, which are members of the IGF axis and have a major role in hepatocarcinogenesis, with the metastamiRs miR-96-5p and miR-182-5p. miR-96 and miR-182 were previously shown to have a role in hepatocellular carcinoma (HCC) (5-11). They belong to the same family (miR-96/182/183 family) and are members of a miRNA cluster that resides in the intergenic region between two protein-coding genes in the chromosomal locus (7q31-34) that is frequently amplified in HCC (5,6). Gradual increase in miR-96 expression levels was linked to disease progression from normal liver tissues to HCC (6‑9). miR‑96 was identified as the most strongly associated HCC recurrence‑associated miR in non‑tumor tissues and its expression levels was found to be correlated with the invasive and metastatic potential of HCC cells (7,10). Likewise, miR-182 was significantly upregulated in metastatic HCC tissues compared with that in paired normal tissues. This observed upregulation was significantly associated with intrahepatic metastasis, early recurrence and correlated with the HCC tumor grade (11). However, the expression status of miR‑96 and miR-182 in non-metastatic HCC as well as the underlying mechanism of inducing carcinogenesis has not been investigated previously, to the best of note.
In HCC tissues, the expression of IGF-II ligand protein was shown to be significantly higher than that in normal liver tissues (12). Similarly, the membrane-bound receptor IGF-1R is overexpressed in various HCC cell lines (13). The expression of IGFBP-3, which modulates the bioavailability of IGF-II ligand, was found to be significantly downregulated in HCC tissues (14). This reduced IGFBP-3 expression may increase the availability of biologically active IGF-II potentiating the proliferative effects (15). Therefore, the present study aimed to explore the impact of the single clustered miRNAs on their downstream targets.

To achieve this aim, screening for miR-96 and miR-182 in non-metastatic HCC tissues was performed followed by manipulation of their expression in the HCC cell line HuH-7 in an attempt to reveal their impact on the crucial members of the IGF axis (IGF-1R, IGFBP-3 and IGF-II).

Materials and methods

Study patients. The present study included 22 HCC patients who underwent liver transplant surgery at the Kasr El Einy Hospital, Cairo University, Egypt. Ten liver biopsy specimens were obtained from healthy donors during transplantation. Healthy donors were non-diabetic, non-hypertensive and negative for hepatitis B and C viruses. All participants gave their written informed consent.

The study was approved by the ethical review committee of Cairo University (Cairo, Egypt). The institutional ethics committees approving the protocol of the present study comply with the principles set forth in the international reports and guidelines as stated in the Helsinki Declaration and the International Ethical Guidelines For Biomedical Research Involving Human Subjects issued by the Council For International Organizations of Medical Sciences (CIOMS).

All patients were non-metastatic with no extrahepatic manifestations and no vascular invasion. Most of the patients (68.2%) had more than one focal lesion as indicated in the pathology report and were subjected to clinical assessment as shown in Table I.

Cell culture. HuH-7 cells were maintained in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 4.5 g/l glucose, 4 mmol/l L-glutamine, 10% fetal bovine serum and mycozap at 1:500 dilution (all Lonza, Basel, Switzerland) at 37°C in a 5% CO₂ atmosphere.

Transfection of oligonucleotides. For examining the effect of miR-96-5p and miR-182-5p on IGF-1R, IGF-II and IGFBP-3 transcript expression, HuH-7 cells were transfected with mimics and inhibitors of miR-96-5p and miR-182-5p (Qiagen, Hilden, Germany). All transfection experiments were performed in triplicate using HiPerfect Transfection Reagent (Qiagen) according to the manufacturer’s instructions and experiments were repeated three times. Cells that were only exposed to transfection reagent were designated as mock cells, cells transfected with miR-96-5p or miR-182-5p mimics were designated as miR-96-5p cells or miR-182-5p cells, respectively, and cells transfected with miR-96-5p or miR-182-5p inhibitors were designated as anti-miR-96-5p or anti-miR-182-5p cells, respectively.

Table I. Number/size of focal lesions according to Milan criteria.

<table>
<thead>
<tr>
<th>Patient ID</th>
<th>Focal lesions (n)</th>
<th>Size of focal lesions (cm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>3</td>
<td>1.5, 1, 1</td>
</tr>
<tr>
<td>2</td>
<td>1</td>
<td>2.5</td>
</tr>
<tr>
<td>3</td>
<td>3</td>
<td>2, 2.5, 3</td>
</tr>
<tr>
<td>4</td>
<td>3</td>
<td>2, 2, 3.5</td>
</tr>
<tr>
<td>5</td>
<td>1</td>
<td>1.5x2</td>
</tr>
<tr>
<td>6</td>
<td>3</td>
<td>3x4, 1, 1</td>
</tr>
<tr>
<td>7</td>
<td>1</td>
<td>4</td>
</tr>
<tr>
<td>8</td>
<td>3</td>
<td>4, 1, 1</td>
</tr>
<tr>
<td>9</td>
<td>3</td>
<td>1, 1, 1.5</td>
</tr>
<tr>
<td>10</td>
<td>1</td>
<td>2.5</td>
</tr>
<tr>
<td>11</td>
<td>2</td>
<td>1, 1.7</td>
</tr>
<tr>
<td>12</td>
<td>3</td>
<td>1, 1, 1</td>
</tr>
<tr>
<td>13</td>
<td>1</td>
<td>3</td>
</tr>
<tr>
<td>14</td>
<td>3</td>
<td>3, 1.5, 2</td>
</tr>
<tr>
<td>15</td>
<td>3</td>
<td>1, 1, 4</td>
</tr>
<tr>
<td>16</td>
<td>2</td>
<td>3, 1.5</td>
</tr>
<tr>
<td>17</td>
<td>2</td>
<td>1.5, 3</td>
</tr>
<tr>
<td>18</td>
<td>3</td>
<td>2.5, 2.5, 1.5</td>
</tr>
<tr>
<td>19</td>
<td>3</td>
<td>1.5, 1.1</td>
</tr>
<tr>
<td>20</td>
<td>1</td>
<td>2</td>
</tr>
<tr>
<td>21</td>
<td>1</td>
<td>1.5</td>
</tr>
<tr>
<td>22</td>
<td>3</td>
<td>3, 2.5, 1</td>
</tr>
</tbody>
</table>

Total RNA extraction from liver biopsies and HCC cells. mRNAs and miRs were extracted from liver biopsy specimens and HCC cells. Fresh liver samples (HCC and healthy tissues) were collected during surgery and were snap frozen in liquid nitrogen. The specimens were manually pulverized in liquid nitrogen and ~100 mg tissue powder was used for large and small RNA extraction using the mirVana miRNA Isolation kit (Ambion Life Technologies, Waltham, MA, USA) according to the manufacturer’s instructions. HCC cells were harvested 48 h after transfection according to the HiPerfect Transfection Reagent protocol; 150 ng oligonucleotides were used for HuH-7-cell transfection in a six-well plate.

miR and mRNA quantification. The extracted miRs were reverse transcribed into single stranded complementary DNA (cDNA) using a TaqMan_MicroRNA Reverse Transcription kit (Applied Biosystems Life Technologies) using specific primers for Homo sapiens (hsa)-miR-96-5p, hsa-miR-182-5p as well as RNU6B for normalization. IGF-1R, IGF-II and IGFBP-3 mRNAs were reverse transcribed into cDNA using the High-capacity cDNA Reverse Transcription kit (Applied Biosystems) according to the manufacturer’s instructions. Relative expression of miR-96-5p, miR-182-5p and RNU6B as well as IGF-1R, IGF-II, IGFBP-3 and β-2- microglobulin (β2MG; for normalization) was quantified using TaqMan.
miR-96-5p and miR-182-5p screening in liver tissues and HCC cell lines. The relative expression of miR-96 and miR-182 was assessed in the HCC and healthy liver tissue groups, as well as in a HCC cell line (HuH-7), and its expression was normalized to RNU6B. Expression levels of miR-96 and miR-182 in all HCC biopsy specimens (n=22) were similar to those in healthy tissues (P=0.5555 and P=0.1864, respectively). However, expression of miR-96 in HuH-7 cells was significantly higher than that in healthy tissues (P=0.0225) (Fig. 1A). In a similar manner, the expression of miR-182-5p in HuH-7 cells showed a significantly higher expression compared to that in healthy tissues and HCC tissues (P=0.0002) (Fig. 1A).

Correlation analysis between miR-96-5p and miR-182-5p mRNA expression in HCC tissues and healthy controls. miR-96-5p expression was quantified in all healthy and HCC tissues and correlated to miR-182-5p expression in the same patients and healthy controls. Using Pearson's statistical method of correlation, miR-96-5p expression was found to be directly correlated with miR-182-5p expression in all HCC tissues studied, as indicated by a Pearson correlation coefficient of r=0.5119 (P=0.0149, Fig. 1B). Similarly, in healthy controls, the same direct correlation was observed with a Pearson correlation coefficient of r=0.6708 (P=0.0337) (Fig. 1B).

miR-96-5p and miR-182-5p expression in liver tissues and HCC cell lines. Transfection efficiency of miR-96-5p and miR-182-5p oligonucleotides. HuH-7 cells transfected with miR-96 and miR-182 mimics showed upregulation of miR-96 and miR-182 by up to 47.18- and 98.59-fold, respectively (P=0.0294 and P<0.0001, respectively) compared to mock-transfected cells (Fig. 2). Impact of miR-96-5p and miR-182-5p on IGF-1R. Transfection of HuH-7 cells with miR-96 mimics resulted in a significant
upregulation of IGF-1R mRNA levels (P<0.0001) compared to those in the mock controls. Of note, inhibitors of miR-96 in HuH-7 caused a significant decrease in IGF-1R mRNA levels compared to those in mimic-transfected cells (P<0.0001) (Fig. 3A). Of note, transfection of HuH-7 cells with miR-182 mimics showed a significant downregulation of IGF-1R mRNA levels compared to those in mock controls (P=0.0267). However, inhibitors of miR-182 in HuH-7 did not cause any significant changes in IGF-1R mRNA levels in HuH-7 cells compared with those in mock-transfected controls (P=0.1266) (Fig. 3A).

Impact of miR-96-5p and miR-182-5p on IGFBP-3. Transfection of HuH-7 cells with miR-96 mimics resulted in a significant upregulation of IGFBP-3 mRNA levels (P<0.0001) compared to those in mock-transfected controls. While inhibitors of miR-96 caused a significant downregulation of IGFBP-3 mRNA levels in HuH-7 cells compared to those in mock-transfected controls (P<0.0001) and mimic-transfected cells (P<0.0001) (Fig. 3B). Transfection of HuH-7 cells with miR-182 caused a significant upregulation of IGFBP-3 mRNA levels compared to those in mock-transfected controls (P=0.0357). However, inhibitors of miR-182 did not cause any significant change in IGFBP-3 mRNA levels in HuH-7 cells compared to those in mock cells (P=0.7937) (Fig. 3B).

Discussion

miR-96-5p and miR-182-5p, which belong to the same family of single clustered miRs, were recently shown to be oncomiRs that promote oncogenesis with an essential role in HCC progression and metastasis (5,10,11,16). To the best of our knowledge, the present study was the first to perform expression profiling of miR-96 and miR-182 in non-metastatic HCC tissues. The two miRNAs were found to have similar expression levels in non-metastatic HCC tissues as compared to that in healthy liver tissues. However, when screened in the HuH-7 cell line, expression levels of the two miRNAs were significantly upregulated. Alongside these results, miR-96 and miR-182 were previously reported to be upregulated in metastatic tissues (10,11). Furthermore, there was a significant direct correlation between the expression of miR-96 and miR-182 in 22 HCC biopsy samples and in 10 healthy controls. This similar expression pattern may be explained by the fact that they belong to
The aberrant expression pattern of IGF axis members in human HCC tissues with overexpression of IGF-1R and IGF-II and downregulation of IGFBP-3 was reported to promote HCC progression and metastasis (14,18,19).

Bioinformatics tools showed that miR-96-5p and miR-182-5p have the same potential binding sites in two regions of the 3'-UTR of IGF-1R mRNA. The two miRNAs were also predicted to target two different regions in the 5'-UTR of IGF-1R mRNA. In spite of the fact that miR-96 and miR-182 share an identical seed region (UUGGCA, nucleotides 2-7), the IGF-1R expression levels obtained upon transfection of HuH-7 cells with miR-96 and miR-182 mimics were not the same: In HuH-7 cells, IGF-1R mRNA was upregulated by miR-96 mimics and downregulated by miR-182 mimics compared to that in their respective mock-transfected cells. These findings are in agreement with those of another study, in which the glypican-3 (GPC3) 3'-UTR was predicted as a common target for the two miRs due to exhibiting one miR-96/182 binding site. miR-96-5p downregulated GPC3 expression by targeting its mRNA 3'-UTR; however, miR-182 failed to regulate it despite the seed similarity between the two miRNAs (20).

The second potential target, IGFBP-3 mRNA, was predicted to be targeted by the two miRs at the 5'- and 3'-UTRs. IGFBP-3 mRNA was found to be significantly upregulated upon ectopic expression of miR-96 in HuH-7 cells compared to that in mock-transfected cells, and the same results were observed upon transfection of HuH-7 cells with miR-182 mimics. This overexpression of the downstream target was restored to normal levels upon antagonizing miR-96 and miR-182 using specific antagomirs in HuH-7 cells. The finding that the two miRNAs have the same effect on their target IGFBP-3 mRNA consorts with another study where the 3'-UTR of chloride intracellular channel 5, which contains a single miR-96/-182 binding site, was found to be directly targeted; however, in that case it was downregulated by the two miRs at the mRNA and protein level (21). The observed induction of IGFBP-3 mRNA by miR-96 and miR-182 may be due to 5'-UTR binding leading to stabilization of the mRNA, thus activating gene expression, which is considered a novel mechanism of action for miRs (22). This inducing impact was previously reported when miR-122 directly targeted two sites in the 5'-UTR of Hepatitis C virus RNA stimulating its translation and increasing its abundance (23-25).

Following the exploration of the effect of miR-96 and miR-182 on IGF-1R and IGFBP-3 mRNA, the present study examined the effect of the two miR mimics on IGF-II mRNA, which was predicted to be targeted by both miRs at the 5'-UTR but not the 3'-UTR. Unexpectedly, it was found that IGF-II mRNA levels were significantly upregulated following transfection of HuH-7 cells with each of the miR mimics

Figure 3. Impact of miR-96-5p and miR-182-5p on IGF-1R, IGFBP-3 and IGF-II transcript expression. (A) Transfection with mimics of miR-96-5p in HuH-7 cells resulted in a significant upregulation of IGF-1R mRNA levels compared to those in mock-transfected controls, while inhibitors of miR-96-5p caused a significant downregulation in the expression of IGF-1R transcripts compared to that in mimic-transfected cells. Mimicking of miR-182-5p in HuH-7 cells resulted in a significant downregulation of IGFBP-3 mRNA levels compared to those in mock-transfected controls, while inhibitors of miR-182-5p did not significantly change the expression of IGF-1R transcript levels compared to those in mock-transfected controls. However, inhibitors of miR-182-5p caused a significant downregulation in the expression of IGFBP-3 transcript levels compared with those in mock-transfected controls. (B) Transfection of HuH-7 cells with miR-96-5p mimics resulted in a significant upregulation of IGFBP-3 mRNA levels compared to those in mock-transfected controls. However, inhibitors of miR-182-5p did not cause any significant changes in the expression of IGFBP-3 transcript levels compared to those in mock-transfected controls. (C) Transfection of HuH-7 cells with miR-96-5p mimics resulted in a significant upregulation of IGFBP-3 mRNA levels compared to those in mock-transfected controls. Of note, inhibitors of miR-96-5p caused a significant downregulation in the expression of IGF-II transcript levels compared to those in mock-transfected cells. miR-182-5p mimics caused a significant upregulation of IGF-II mRNA levels in HuH-7 cells compared to those in mock-transfected controls. However, inhibitors of miR-182-5p did not cause any significant change in the expression of IGF-II transcript levels compared to those in mock-transfected cells. The Mann-Whitney U test was performed. **P<0.001, *P<0.05. RQ, relative quantitation, IGFBP, insulin-like growth factor binding protein; IGF-1R, insulin-like growth factor-1 receptor; miR, microRNA.
compared with those in their respective mock-transfected cells. The same enhancing effect on target mRNA by a miR was reported by a previous study on the receptor-interacting protein 140 (RIP140) mRNA, where miR-346 targeted the 5'-UTR of the RIP 140 transcript, leading to an upregulation of its protein levels (26).

This expression variability of IGF axis components upon miR-96 mimic-transfection raises a question about the simultaneous increase of IGF-1R and IGF-II, which are mitogenic, as well as IGFBP-3, which is anti-tumorogenic. This may be explained as a compensatory mechanism where the upregulation of IGF-1R and IGF-II may be neutralized by the upregulation of IGFBP-3, counteracting the tumorogenic effect of the mitogens and mediating their effect. This mediation by upregulated IGFBP-3 is realized through its binding to the overexpressed IGF-II, which inhibits the binding and consequent activation of IGF-1R and subsequent downstream signaling. By contrast, miR-182 has a dual role where it upregulates the mitogenic IGF-1R and subsequent downstream signaling. By contrast, miR-182 has a dual role where it upregulates the mitogenic IGF-1R and IGF-II may be neutralized by the upregulation of IGFBP-3, counteracting the tumorogenic effect of the mitogens and mediating their effect. This mediation by upregulated IGFBP-3 is realized through its binding to the overexpressed IGF-II, which inhibits the binding and consequent activation of IGF-1R and subsequent downstream signaling. By contrast, miR-182 has a dual role where it upregulates the mitogenic IGF-1R and IGF-II may be neutralized by the upregulation of IGFBP-3, counteracting the tumorogenic effect of the mitogens and mediating their effect. This mediation by upregulated IGFBP-3 is realized through its binding to the overexpressed IGF-II, which inhibits the binding and consequent activation of IGF-1R and subsequent downstream signaling.

In conclusion, the present study showed for the first time, to the best of our knowledge, that miR-96 and miR-182 have similar expression levels in non-metastatic HCC tissues and in healthy controls, contradicting previous studies reporting their upregulation in metastatic tissues (10,11). This highly implies that miR-96-5p and miR-182-5p are metastatic miRs in HCC. The present study also shows for the first time, to the best of our knowledge, the pleiotropic effect of miR-96-5p and miR-182-5p in targeting various components of the oncogenic IGF signaling pathway in HCC. Thus, these data provide the rationale for potential targeting of miR-96-5p and miR-182-5p in HCC therapy.

Acknowledgements

The authors acknowledge the patients and healthy volunteers for their valuable participation in this study. Furthermore, the authors would like to thank the Science and Technology Development Fund (STDF; grant no. 4242) for funding this work.

References