Effects of microRNAs polymorphism in cancer progression

Simin SiamiGorji1, Isa Jorjani1, Alireza Tahamtan2, Abdolvahab Moradi*2

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Abstract
MicroRNAs (miRNAs) are known as a new class of small RNAs (18-25 nucleotides) that regulate gene expression at multiple levels from transcription to translation. Considering the important role of miRNAs in cell proliferation, differentiation, and apoptosis, any variations in their expression can contribute to various anomalies, such as tumorigenesis. Single-nucleotide polymorphisms (SNPs) have received much attention as potential genetic markers for diseases due to their advantage of being present at a high frequency in the human genome. SNPs can occur in different parts of the miRNA genes (primary, precursor, and mature) which result in pathological conditions. In this study, recent findings related to the effects of SNPs in miRNAs on their biogenesis and functions and their role in cancer development and progression are discussed. This review was performed using PubMed to search for related reports. The identified effects may be useful for clinical decision-making and providing important new information about the pathophysiology of miRNAs.

Keywords: MicroRNAs, Noncoding RNAs, Single nucleotide polymorphisms, Mutation, Cancer

Introduction
Nearly 1.5% of the human genome is transcribed and translated to protein, and this process leads to creation of a wide variety of noncoding RNAs (ncRNAs), such as microRNAs (miRNAs), long noncoding RNAs (lncRNAs), rRNAs, and tRNAs (1). Also, miRNAs are identified in eukaryotes and attach to the regulatory elements of target mRNA. They tend to negatively regulate the expression of a large number of target genes through the degree of complementarity between the miRNAs and the target genes. Moreover, miRNAs are able to intensify the expression of target genes and regulate transcription and translation. Changes in miRNA s that lead to abnormalities are categorized in 2 ways: The first category is the variation of the cis factors, such as chromosomal changes, epigenetic changes, mutations in the promoter elements, and polymorphism in the miRNA sequence. The second category is changes in trans factors, such as polymorphism in the target genes, mutation in the regulatory and processing elements (2).

Single-nucleotide polymorphisms (SNPs) are the most common and prevalent genetic variation that can affect miRNAs function. It is thought that SNPs in miRNA af-
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miRNAs and tumorigenesis

Cancer is caused by accumulation of genetic changes and subsequently inactivation of tumor suppressor genes (TS-genes) and proto-oncogenes activation. Some special miRNAs serve as a tumor suppressor (TS-miR) and oncogene (onco-miR), thus, their downexpression and overexpression interfere with tumor susceptibility. Deregelation of miRNAs expression through destroying cellular arrangement in normal tissues leads to malignant formation, which results from different mechanisms, e.g., chromosomal abnormalities, epigenetic changes, alterations in the miR-processing machinery, and genomic mutations/polymorphisms.

Chromosomal abnormalities

About half of human miRs genes and clusters exist in (or around) the chromosomal fragile site (FRAs) and cancer-associated genomic regions (CAGRs). FRAs are prone to genomic instability events, such as chromatid exchange, translocation, deletion, amplification, and copy number variation (CNV). These events might change miRs correct sequence and structure or their target dosage, therefore interfere with normal gene regulation and cause abnormalities. Some miRs beside Dicer and Drosha were subjected to CNVs and repeatedly proliferated in NSCLC (nonsmall cell lung cancer). (7).

Epigenetic changes

The regulatory effect of epigenetic on miRs has been achieved through DNA hyper methylation, hypo methylation, and destruction of histone modification patterns in human tumors. In other words, tumor formation might be due to epigenetically silenced TS-miRNAs or activated oncomiRs by aberrant hyper and hypo methylation of CpG islands located around these genes, respectively or by posttranscriptional modifications of nucleosomal histone proteins modifications.

Alterations in the miRNA processing machinery

The biogenesis of miRs in tumor samples is different from normal ones. There is a decreasing expression of Dicer and Drosha in various types of cancer. In fact, the precise location and direction of Drosha cleavage for creating a mature miRs is very important, because the second cleavage occurs at a certain distance from the free end.

Effect of SNPs on miRNAs

Heterogeneous distribution of SNPs across the genome leads to more aggregation in noncoding regions than coding regions. A large number of SNPs located in the noncoding regions of the genome are functional including the miRNAs and their target sites. Indeed, these SNPs change the level of gene expression instead of altering the protein's performance. mi-SNPs are often found in QTL (quantitative trait loci), chromosomal FRAs, and tumor potential sites, indicating their ability to genetically control complex characteristics. There are several methods to show the effect of human abnormality-related SNPs on miRs expression level: 1. SNPs in miRs biogenesis pathways, which interfere with miRs-processing; II. SNPs in miRs sequence interfere with maturation process; and III. SNPs in miRs' promoter and target genes, which change the expression levels and lose the binding efficiency to miRs.

The consequences of SNPs in miR sequences are aberrant expression of hundreds of genes and pathways. For instance, some studies demonstrated that most SNPs in miRs (pre/mature) changed targets expression level via changing mature-miRs frequency. Mature-miR SNPs are classified into 2 subgroups: 1. SNPs in 5'-seed region affect target recognition and stability of miR-miRNAs interaction; 2. SNPs in 3'-mismatch tolerant region that may withstand mismatch (3). Seed-SNPs interfere with pri to pre-miR conversion and also post-transcriptional regulations via altering miR-target sites (13, 14). Some seed-SNPs that are phenotypically neutral, facilitate miRs' evolution.
toward new miRs with more specialized operation. Principally, miRs variations can eliminate binding to key targets and create a seed sequence which targets new mRNAs, changing the regulatory growth proteins balance (14).

**Effect of SNPs on 3′UTR of miR-target genes in cancers**

The 3′ untranslated regions (3′UTRs) are involved in various levels of mRNA in the regulation of genes expression. Also, 3′UTR-SNPs, as miRs-binding site, by modifying the regulatory loops between them cause genetic changes, and subsequently trigger malignancies. In fact, by changing polyadenylation, mRNA-miRNA and mRNA-protein interaction influence mRNA stability and translation. The effect of 3′UTR-SNPs on changing target sequence and target’s second structure has been surveyed. The degree of alteration in mRNA second structure show the possible availability for miRs-targeting which has been used in predicting the creation or degradation of the miR-binding-site (15). In this regard, 2 categories are provided: 3′UTR-SNPs, which make illegitimate-miR target sites, can cause tumor risk dependent on TS-miRs, or oncomiRs (16-19). 3′UTR-SNPs which destroy miR-binding position are associated with cancer susceptibility (18-21) (Table 1). These achievements indicate the importance of abnormality-associated-SNPs located in UTR within or near the miRs-binding sites in various cancers.

**Effect of SNPs on miRNAs’ sequence in cancer**

Generally, miRs are divided into 4 groups based on genomic location related to intron and exon: (1) intronic miRs in ncRNA transcripts; (2) intronic miRs in protein-coding genes; (3) exonic miRs in ncRNA transcripts, such as miR-22 and miR-198; and (4) exonic miRs in protein-coding genes. For instance, miR-985 is found in the last exon of CACNG8 gene (22). Also, miRs placed in noncoding regions between genes are called intergenic. Noncoding SNPs refer to those which occur in noncoding gene regions, including miRs sequence, regulatory regions, intronic and intergenic regions presence in coding genes. MiRs-related-SNPs that involve in tumorigenesis can also exist in miR’s host genes, which encode tumor-related proteins (23).

**SNPs in miRNAs found in introns of protein-coding genes**

A separate evaluation of miRs based on origin explicitly addresses the SNPs presence in miRs and their possible association with various tumors. Accordingly, rs928508 in intronic-mir-30c1 (24) reduces mature and pre-miR-30c level in NSCLC through heterogeneous genotypes (AG/GG) (25). The homozygous genotype of this polymorphism also results in altered miRNA expression by altering miRNA structure, which has been observed in gastric cancer (26). Table 2 displays other examples of intronic miRs as such. SNP, which interfere with miR maturation (27-30).

**SNPs in miRNAs found in introns of noncoding transcription units**

Sometimes, transcription units, which lack the ability of coding protein, serve as a location for a number of intronic-miRs. ncRNAs transcripts are sometimes known as mRNA-like-ncRNAs (mlncRNAs) because they share common characteristics with mRNAs such as splicing and polyadenylation. A to C SNP in miR-30c-2, as an instance of this group (24), may be effective in target gene-selecting and inefficient miR processing, if located in mature miR and pri-miR, respectively (3). Some examples of miRs in intronic ncRNAs accompanied by related SNPs are presented in Table 2. In fact, many studies have determined the SNPs influence on miR frequency and predisposition to various cancers risk (32-35).

**SNPs in miRNAs located in exons of noncoding transcription units**

Approximately, 10% of miRs are encoded in exonic regions (22). ncRNAs, whose only important activity is acting as miR-host gene, contain many exonic miRs (24). The exonic miRs are transcribed in pri-miRNAs and include 5’-cap and 3’-polyA. A vivid example of this group is miR-155 derived from the third exon of BIC transcript; namely, mlncRNAs. Natural functional SNPs in loop region and miR-155* change the second structure of miR-155 in humans (36). Although the effects of these SNPs have been identified on mature miR expression and immunological response in humans, their relation to cancer has not been studied until now. Also, information on stud-

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**Table 1. Effect of SNP in miR-target genes on cancers**

<table>
<thead>
<tr>
<th>Target gene</th>
<th>miRAs</th>
<th>SNP</th>
<th>Position</th>
<th>Type of SNP</th>
<th>Effect of SNP</th>
<th>Cancer</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>TYRP1</td>
<td>miR-155</td>
<td>rs683/rs910</td>
<td>3′UTR</td>
<td>non-coding</td>
<td>Illegitimate miR-target site</td>
<td>Melanoma</td>
<td>(16)</td>
</tr>
<tr>
<td>KIT</td>
<td>miR-221</td>
<td>rs17084733</td>
<td>3′UTR</td>
<td>non-coding</td>
<td>Illegitimate miR-target site</td>
<td>Thyroid</td>
<td>(17)</td>
</tr>
<tr>
<td>RPA2</td>
<td>miR-3149</td>
<td>rs7356</td>
<td>3′UTR</td>
<td>non-coding</td>
<td>Illegitimate miR-target site</td>
<td>Colorectal</td>
<td>(18)</td>
</tr>
<tr>
<td>ARHGAP26</td>
<td>miR-18a-3p</td>
<td>rs187729</td>
<td>3′UTR</td>
<td>non-coding</td>
<td>Illegitimate miR-target site</td>
<td>ALL, CML</td>
<td>(19)</td>
</tr>
<tr>
<td>TLX1</td>
<td>miR-492</td>
<td>rs2742038</td>
<td>3′UTR</td>
<td>non-coding</td>
<td>Illegitimate miR-target site</td>
<td>ALL</td>
<td>(19)</td>
</tr>
<tr>
<td>IRF8</td>
<td>miR-330</td>
<td>rs10514611</td>
<td>3′UTR</td>
<td>non-coding</td>
<td>Target-site destruction</td>
<td>CML</td>
<td>(20)</td>
</tr>
<tr>
<td>KRAS</td>
<td>miR-155</td>
<td>rs16764370</td>
<td>3′UTR</td>
<td>non-coding</td>
<td>Target-site destruction</td>
<td>Ovarian</td>
<td>(20)</td>
</tr>
<tr>
<td>GTF2H1</td>
<td>miR-518a</td>
<td>rs4596</td>
<td>3′UTR</td>
<td>non-coding</td>
<td>Target-site destruction</td>
<td>Colorectal</td>
<td>(18)</td>
</tr>
<tr>
<td>ET6V</td>
<td>miR-34c</td>
<td>rs1573613</td>
<td>3′UTR</td>
<td>non-coding</td>
<td>Target-site destruction</td>
<td>ALL</td>
<td>(19)</td>
</tr>
<tr>
<td>ESR1</td>
<td>miR-453</td>
<td>rs2747648</td>
<td>3′UTR</td>
<td>non-coding</td>
<td>Target-site destruction</td>
<td>Breast</td>
<td>(21)</td>
</tr>
</tbody>
</table>
ies conducted on exonic miRs and their SNPs related to cancer susceptibility is demonstrated in Table 2 (32, 37, 38).

**SNPs in intergenic miRNA**

The genomic distribution of 1523 human genes revealed that 57.6% of miRs are intergenic. Intergenic miRs are structurally and functionally similar to intronic miRs and have independent transcriptional regulatory elements, such as promoters and terminators. Also, MIR-125a, as an example of intergenic miR (39), contains rs12975333 in the seed region suppresses pri-miRNA processing to pre-miR-125a (40). The list of other instances of intergenic miRs and their studied SNPs is provided in Table 2 (41-44). Generally, these findings imply the changes in primary transcripts, hairpin-shaped precursors (pre-miRs), or mature miRs (with various origins), which may influence the miRNA properties and increase the risk of carcinogenesis. However, more research in future are needed to approve these findings.

**Effect of SNPs on miRNAs’ regulatory elements in tumors**

In addition to the direct effects of SNPs on various sequences of miRs, the SNPs can indirectly affect the regulatory elements involved in a miRs transcription. More than 90% of disease-related SNPs are located in noncoding regions, such as 5’UTRs, which contain important regulatory factors, e.g., promoters and enhancer or components involved in a translation (12). Noncoding SNPs, situated in the regulatory sites of genome include transcription factors, binding sites, and slice sites, which lead to genes misregulation at transcriptional or posttranscriptional level. There are almost more than 20 000 SNPs in human promoters which can moderate the overall level of pri-miRNAs (6). Taken together, variations in Cis or Trans regulatory elements change the miR host genes expression, and subsequently the miRs regulations are embedded in the host gene (45). Table 3 demonstrates some examples of SNP in regulatory elements related to change in transcription factors, binding tendency, and promoter activity (46-49). All these reports confirmed that SNPs in miRs regulatory regions are engaged in miRs processing and finally predisposition to abnormality.

**SNPs in coding regions of miRNA’s targets and host genes**

The gene coding sequences are between 3’/5’UTRs regions. Coding SNPs only come about in the exonic regions conducted on exonic miRs and their SNPs related to cancer susceptibility is demonstrated in Table 2 (32, 37, 38).

### Table 2. Effect of SNP in miRNAs on various cancer

<table>
<thead>
<tr>
<th>miRNA</th>
<th>Location of miRs</th>
<th>SNP</th>
<th>SNP in miR sequence</th>
<th>Cancer type</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>miR-30c-1</td>
<td>Intron of coding region</td>
<td>rs928508</td>
<td>pre-miRNA</td>
<td>Gastric</td>
<td>(26)</td>
</tr>
<tr>
<td>miR-499</td>
<td>Intron of coding region</td>
<td>rs3746444 T&gt;C</td>
<td>Pre-miR-499</td>
<td>Prostate</td>
<td>(27)</td>
</tr>
<tr>
<td>miR-26a</td>
<td>Intron of coding region</td>
<td>rs7372209</td>
<td>pri-miR26a-1</td>
<td>OPL</td>
<td>(28)</td>
</tr>
<tr>
<td>miR-608</td>
<td>Intron of coding region</td>
<td>rs4919510 C&gt;G</td>
<td>mature-miR</td>
<td>ESCC</td>
<td>(30)</td>
</tr>
<tr>
<td>miR-618</td>
<td>Intron of coding region</td>
<td>rs2682818 C&gt;A</td>
<td>pre-miR-618</td>
<td>Lymphoma</td>
<td>(29)</td>
</tr>
<tr>
<td>miR-15a-16-1</td>
<td>Intron of ncRNA transcript (DLEU2)</td>
<td>G&gt;C83</td>
<td>pre-miR-16-1</td>
<td>CLL</td>
<td>(32)</td>
</tr>
<tr>
<td>let-7/miR-98</td>
<td>IntermiR (AK09218)</td>
<td>rs629367</td>
<td>pre-miR-181b-2</td>
<td>Thyroid</td>
<td>(34)</td>
</tr>
<tr>
<td>miR-146a</td>
<td>Exon of ncRNA transcript</td>
<td>rs2910164 G&gt;C</td>
<td>seed stem-loop</td>
<td>HCC</td>
<td>(35)</td>
</tr>
<tr>
<td>miR-101-1</td>
<td>Exon of ncRNA transcript</td>
<td>rs7372209</td>
<td>pri-miR-101-1</td>
<td>Gastric</td>
<td>(37)</td>
</tr>
<tr>
<td>miR-29b-2</td>
<td>Exon of ncRNA transcript</td>
<td>rs7536540 G&gt;C</td>
<td>pri-miR-29b-2</td>
<td>HCC</td>
<td>(41)</td>
</tr>
<tr>
<td>miR-371-372-373</td>
<td>Intergenic</td>
<td>rs359501 C&gt;A</td>
<td>pri-miRNAs</td>
<td>HCC</td>
<td>(41)</td>
</tr>
<tr>
<td>miR-196 cluster</td>
<td>Intergenic</td>
<td>rs1164913 C&gt;T</td>
<td>maturer-miR</td>
<td>Breast</td>
<td>(42)</td>
</tr>
<tr>
<td>miR-125a</td>
<td>Intergenic</td>
<td>rs12976445</td>
<td>pri-miR-125a</td>
<td>Breast</td>
<td>(43)</td>
</tr>
<tr>
<td>miRNA-1268a</td>
<td>Intergenic</td>
<td>rs2859926</td>
<td>pre-miR</td>
<td>Astrocytoma</td>
<td>(44)</td>
</tr>
</tbody>
</table>

### Table 3. Effect of SNP in regulatory element of miRNAs on cancers

<table>
<thead>
<tr>
<th>miRNA</th>
<th>Location of miRs</th>
<th>SNP</th>
<th>Location of SNP</th>
<th>Cancer type</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>miR-148/152 cluster</td>
<td>Intron of coding region</td>
<td>rs4719839</td>
<td>Promoter ~2966bp upper pre-miR</td>
<td>Gastric</td>
<td>(49)</td>
</tr>
<tr>
<td>miR-30a</td>
<td>Intron of coding region</td>
<td>rs763354</td>
<td>Enhancer</td>
<td>NSCLC</td>
<td>(46)</td>
</tr>
<tr>
<td>miR-122</td>
<td>Exon of ncRNA transcript</td>
<td>rs17669 T&gt;C</td>
<td>upstream regulatory region</td>
<td>HCC</td>
<td>(47)</td>
</tr>
<tr>
<td>miR-143/145</td>
<td>Exon of ncRNA transcript</td>
<td>rs41291957 rs353292 rs353293 rs705509</td>
<td>Promoter</td>
<td>Colorectal</td>
<td>(48)</td>
</tr>
<tr>
<td>miR-200b/200a/429 cluster</td>
<td>Intergenic</td>
<td>rs9600710</td>
<td>Promoter Enhancer</td>
<td>NSCLC</td>
<td>(46)</td>
</tr>
</tbody>
</table>
gions. There are 2 types of coding SNPs: synonymous (sSNPs) and nonsynonymous (nsSNPs). Missense and nonsense are 2 subgroups of nsSNPs. Also, sSNPs can directly affect the translation kinetics and alter the folding pattern and the covalent modification of coded proteins. Also, sSNPs could lead to variation in translation regulation by affecting the mRNA-miRs interaction (50). However, nsSNPs as the most recognized genomic variant, alter the structure of the proteins through amino acid changes and contribute to the molecular function diversity in populations (51).

**Effect of coding SNPs on miRNAs’ target genes in malignancies**

There are about 40% to 60% of miRs binding sites in the coding region of genes (50). miRs binding site on the CDS is related to targets’ posttranscriptional regulation and are more effective in translation inhibition. In addition, the synergistic action of miRs-target site in CDS are confirmed in contributing to the recognition of seeds in 3’UTR or in elevating mRNA-mRNA interactions (52). Both SNPs (sSNP and nsSNP) in the coding regions occur frequently and alter the miR regulation (50). A number of SNPs on CDS in target genes were found to create new miR-matching sites (15, 53) or conversely disrupt the binding site (17, 54) (Table 4). As an illustration, an nsSNP rs41427445 in CDS of MMP9 create the desired matching in miR-671-3p and miR-657 binding sites and reduced the protein secretion. Also, the missense SNP changes the N38S amino acid and removes the N-glycosylation site (55).

**The effect of coding-SNPs in miRNAs’ host genes on cancers**

SNPs in the coding regions of miRs may alter the expression and maturation of miRs and lead to abnormalities (50). Some examples of SNPs on the CDS in the host genes are presented in Table 4 (49, 56). As a result, coding SNPs in miR’s host genes and target genes may alter the processing and binding site on miR-target genes, respectively, and have vital effects on miRs function.

**SNPs in miRNAs processing genes**

As mentioned earlier, if variants exist in miR-biogenesis genes and genes related to silencing-machinery (AGO family and GEMIN proteins), they can alter the overall silencing efficiency (3). Some of SNPs in miRNAs processing genes are listed in Table 5 (28, 30, 57-60). According to the recent reports, the miR-biogenesis-degradation may interfere with the miR-maturation or even mRNA stability and protein expression due to the SNPs and may have a profound impact on cancer risk.

**Conclusion**

According to numerous studies, noncoding RNAs act as the main regulators of switching on and silencing the genes in cells, and the gene misexpression is one of the main pillars of the disease-making mechanisms. The emergence of miRs as one of the various noncoding RNAs and their widespread and critical interference in various cell and molecular parts have been considered as an amazing selection of nature in multicellular life. Here, 2 distinct classes of miRs have been reviewed: miRs resulting from intron overlapping with other transcripts and miRs, which are coded from exon transcripts. The new genomic sources of miRs, including transposable elements, pseudogenes, genome duplication, Denovo, and the antisense miRs loci, exhibit the miRs biosynthesis and maturation process and have many unknown dimensions. Despite all the advances made in identifying the biogenesis pathway and activity of miRs, more research is needed. Determining the exact origin of miRs (exon, intron, and

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Table 4. Effect of coding SNP on cancers

<table>
<thead>
<tr>
<th>miRNA</th>
<th>SNP</th>
<th>Location of SNP</th>
<th>Effect of SNP</th>
<th>Cancer type</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>miR-196</td>
<td>sSNP (c.313C&gt;T)</td>
<td>Coding region of miR-target gene</td>
<td>Destroying the miR-196 binding</td>
<td>Crohns</td>
<td>(54)</td>
</tr>
<tr>
<td>miR-146a/miR-146b</td>
<td>sSNP (2607G&gt;C) rs7843542</td>
<td>Exon of KIT miR-target gene</td>
<td>Changing miR-146a/b binding site</td>
<td>Thyroid</td>
<td>(17)</td>
</tr>
<tr>
<td>miR-638</td>
<td>sSNP</td>
<td>rs799917 C/T</td>
<td>Exon of BRCA1 miR-target gene</td>
<td>More interaction between</td>
<td>Breast</td>
</tr>
<tr>
<td>miR-671-3p</td>
<td>rs41427445.N38S</td>
<td>Exon of MMP-9 miR-target gene</td>
<td>Changing miR-target site</td>
<td>Gastrointestinal</td>
<td>(53)</td>
</tr>
<tr>
<td>miR-34a</td>
<td>rs72631823 G&gt;A</td>
<td>Coding region of pre-miR-34a</td>
<td>miR-34a down-regulation</td>
<td>osteosarcoma</td>
<td>(56)</td>
</tr>
<tr>
<td>miR-148b</td>
<td>rs11170877 A&gt;G</td>
<td>Exon of COP2 miR-host gene</td>
<td>Changing methionine to Val</td>
<td>Gastric</td>
<td>(49)</td>
</tr>
</tbody>
</table>

Table 5. Effect of SNP in miR-processing genes on cancers

<table>
<thead>
<tr>
<th>Gene</th>
<th>SNP</th>
<th>Location of SNP</th>
<th>Type of SNP</th>
<th>Cancer type</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>GEMIN4</td>
<td>rs78123</td>
<td>Cys1033Arg</td>
<td>nsSNP</td>
<td>ESCC</td>
<td>(30)</td>
</tr>
<tr>
<td></td>
<td>rs3744741</td>
<td>Gin684Arg</td>
<td>nsSNP</td>
<td>Oral</td>
<td>(28)</td>
</tr>
<tr>
<td>TRBP</td>
<td>rs784567</td>
<td>Promoter</td>
<td>non-coding</td>
<td>Larynx</td>
<td>(59)</td>
</tr>
<tr>
<td>XPO5</td>
<td>rs11077</td>
<td>3’UTR</td>
<td>non-coding</td>
<td>Thyroid</td>
<td>(60)</td>
</tr>
<tr>
<td>Dicer</td>
<td>rs3742330</td>
<td>3’UTR</td>
<td>non-coding</td>
<td>OPL</td>
<td>(28)</td>
</tr>
<tr>
<td></td>
<td>rs13078, rs3742330</td>
<td>3’UTR</td>
<td>non-coding</td>
<td>Larynx</td>
<td>(59)</td>
</tr>
<tr>
<td>Drosha</td>
<td>rs640831</td>
<td>promoter</td>
<td>non-coding</td>
<td>Lung</td>
<td>(58)</td>
</tr>
<tr>
<td>RAN</td>
<td>rs14035</td>
<td>3’UTR</td>
<td>non-coding</td>
<td>Oral</td>
<td>(57)</td>
</tr>
</tbody>
</table>

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http://mjiri.iums.ac.ir

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Effects of microRNAs polymorphism in cancer progression

intergenic), with respect to specific cell and tissue function of miRs, contributes to recognizing the precise cellular pathways and effective therapeutic interventions with tumor tissues and cells.

SNPs are present in a variety of genomic sequences that are related to miRs directly or indirectly and have a high capability of interfering with the miRs biological activity and controlling various cellular pathways. The phenotype of the cancers occurs during the deregulation of several genes and unusual cell divisions. Since miRs have remarkable effects on cellular processes, their misregulation due to mutation, polymorphism, etc., similar to coding protein genes, will influence the cells pathogenesis pathway. A large number of case-control studies were investigated in this review, most of which indicated the involvement of genetic SNPs in altering the miR expression levels and the miR biogenesis process genes, miR activity, and cancer susceptibility. The fundamental step in the study of association of miRs and tumorigenesis (early stages to metastasis) is identifying the SNPs related to miRs and its involvement in abnormalities. Here, it was aimed to obtain in-depth understanding of how the SNPs work in forming the cellular pathology and recognizing the miRs-controlled pathways. Any change in cancers requires designing a more complex experiments and more precise methods for early diagnosis and even therapeutic purposes. In addition to participating in tumorigenesis through sequential and structural changes, miR polymorphisms act as a double sword in pathogenesis and cancers. Subsequently, miR level deregulation will interfere with the target gene-binding. Based on studies conducted in the field of miR pharmacogenomics, the genetic heredity of miR-SNPs in each person impresses the body’s therapeutic response to drugs.

Conflict of Interests

The authors declare that they have no competing interests.

References


