Deciphering the Role of microRNAs in BRD4-NUT Fusion Gene Induced NUT Midline Carcinoma

Ekta Pathak¹, Bhavya¹, Divya Mishra¹, Neelam Atri¹,², Rajeev Mishra¹*

¹Bioinformatics Department, MMV, Banaras Hindu University, India; ²Botany Section, MMV, Banaras Hindu University, India; Rajeev Mishra - E-mail address: mishrarajeev@gmail.com; rajeev17@bhu.ac.in; Telephone +91 9935338891; *Corresponding author

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Abstract:
NUT midline carcinoma (NMC) is a very aggressive and lethal type of squamous epithelial cell cancer caused due to fusion of BRD4 and NUT genes. The gene fusion results into a new fusion protein that promotes oncogenesis. The detailed molecular mechanisms underlying the NMC are still not clear and new findings are urgently required to complement the current efforts. Abnormal microRNAs (miRNA) expression promotes tumour formation by modulating the functional expression of critical genes other than the parent genes involved in tumour cell proliferation or survival. Here, using Insilco methods, miRNA targeting the transcripts of parent genes (BRD4 and NUT) and the BRD4-NUT fusion gene were predicted. We investigated a situation, wherein abnormal miRNA expression in malignant cells could arise due to deletion and fusion of genomic regions encompassing the target site of miRNA genes. A set of 48 dysregulated miRNAs targeting the critical genes other than the parent genes (BRD4 and NUT) was identified. Functional enrichment analysis of KEGG pathways of target genes of these Ex-miRNAs implicates their role in cancer pathways. Amplification in the expression level of these miRNAs can be used for NMC diagnosis and prognosis.

Keywords: microRNAs, BRD4-NUT, Fusion Gene, NUT midline carcinoma, bioinformatics approach

Background:
NUT midline carcinoma (NMC) is a fatal form of undifferentiated epithelial cancer affecting both children and adults [1]. Commonly, the sites of occurrence of this cancer are head, neck and space in the thoracic cavity between the lungs. In the majority of cases, NMC is due to the fusion of the testis-specific nuclear gene NUT of chromosome 15 and the bromodomain-containing gene BRD4 (bromodomain protein family member 4) on chromosome 19. The gene fusion results into a new fusion protein that markedly disrupts squamous cell differentiation and promotes oncogenesis [2]. The detailed molecular mechanisms underlying the NMC are still not clear and new findings are urgently required to complement the current efforts.

MicroRNAs (miRNAs) play a critical role in regulating target genes and are involved in the initiation and progression of cancer [3]. miRNAs are small, about 21 nucleotides long non-coding RNAs which usually target the miRNAs at their 3'UTR region and inhibit their expression [4]. It has also been shown that miRNAs also target the CDS and 5'UTR region of the miRNAs [5]. miRNAs are not very specific for their targets and a single miRNA can modulate the expression levels of several hundred to thousands of different miRNAs [3]. Only a small portion of the miRNAs, i.e., the about seven nucleotides long seed region, should be complementary to the mRNA sequence to bind with it and inhibit its expression [6]. In recent years, miRNA profiling and deep sequencing provided direct evidences of dysregulated miRNAs that target key genes like oncogenes or tumour repressor genes to induce tumour development. These miRNAs signature could be used to distinguish between tumoral and normal cells, and in some instances prognosis and the progression of cancer [7]. Here, we have identified the miRNA targeting the transcripts of parent genes (BRD4 and NUT) and the BRD4-NUT fusion gene. We hypothesize that some of the miRNAs regulating their respective BRD4 and NUT gene expression may not be required after the BRD4-NUT gene fusion, as the regions targeted by these miRNAs are deleted. This may lead to an apparent amplification in the miRNAs, which in turn may regulate the pathways leading to cancer. A comparative study of miRNAs regulating fused gene and its parent gene was performed and their target genes were discovered. A functional enrichment analysis of KEGG pathway
of target genes of these Ex-miRNAs shows its role in cancer pathways.

**Methodology:**
BRD4 gene (Refseq: NM_058243.2), NUT gene (GenBank: AF482429.1) and BRD4-NUT fusion gene (GenBank: AY166680.1) transcript sequences were retrieved from the NCBI Nucleotide database. In order to locate the deleted regions in the fusion gene transcript, the mRNA sequences of parent (BRD4 and NUT) were compared against the fusion gene (BRD4-NUT) using align two sequences option of NCBI- BlastN tool. Prediction of miRNAs targeting the BRD4, NUT and BRD4-NUT fusion gene transcripts was performed using MIRDB (http://mirdb.org/miRDB) [8]. For each miRNAs, mirWalk (http://www.umm.uni-heidelberg.de/apps/zmf/mirwalk/) was used to predict miRNA targets and analyse plausible KEGG pathway enrichment of dysregulated miRNA targets. mirWalk dataset hosts experimentally verified miRNA-mRNA interactions as well as the information on the published pathway targets from KEGG (http://www.genome.jp/kegg/) [9]. Since miRNAs can target 3’UTR, CDS and 5’UTR region of the mRNAs [5], therefore parameter for target search was set to include 3’UTR, 5’UTR and CDS regions of transcript. Minimum seed length was chosen as seven with p-value equal to 0.05.

**Results and Discussion:**
It is generally accepted that fusion genes are translated into fusion proteins, which are drivers of cancer pathway activation. On the other hand, it is well established that alterations in miRNA expression promote tumour formation. The dysregulated miRNAs modulate the functional expression of critical genes other than the parent genes involved in tumour cell proliferation or survival [3]. Recent findings suggested that abnormal miRNA expression in malignant cells could arise due to amplification or deletion of specific genomic regions encompassing miRNA genes [10]. Here, we investigated miRNAs targeting the parent BRD4 and NUT genes and BRD4-NUT fusion transcript. We assume that in case of BRD4-NUT gene fusion, transcripts of parent gene will no more available to be targeted by their corresponding miRNAs termed as “Ex-miRNA”. This will lead to a situation where an apparent amplification and dysregulation by these miRNAs are possible.

| Gene transcript (NCBI Accession No.) | Length of mRNA (bases) | Aligned region onto BRD4-NUT fusion gene (bases) | Sequence region deleted due to fusion (bases) |
|-------------------------------------|------------------------|------------------------------------------------|---------------------------------------------|
| BRD4 (Refseq NM_058243.2)           | 5198                   | 1-2380                                         | 2818 (3’ end)                               |
| NUT GenBank: AF482429.1             | 3795                   | 170-3795                                       | 169 (5’end)                                 |

**Figure 1.** Schematic representation showing the alignment of BRD4 (yellow) and NUT (green) transcripts onto BRD4-NUT fused transcript. Deleted regions during the gene fusion event are marked with dashed line.
We predicted and compared the miRNA targeting the BRD4 and NUT transcripts region before and after the gene fusion events. Sequence alignment of parent BRD4, NUT and fusion BRD4-NUT mRNA sequences, using NCBI Blast tool, revealed that 2818 bases (54%) from 3' region of BRD4 and 170 bases from 5' region of NUT were deleted in the BRD4-NUT gene fusion transcript (Figure 1). miRNAs binding sites were identified in parent BRD4 and NUT transcripts using miRDB tool (see methods). 34, 33 and 19 miRNAs were predicted to target BRD4 mRNA, NUT mRNA and fused BRD4-NUT mRNA, respectively. Except, Has-miR-5193 targeting the fusion gene transcript, 18 miRNAs were common to target the parent BRD4 or NUT and fused BRD4-NUT transcript (Table 1). As evident in Figure 1, due to fusion transcript formation, the deleted regions will no more be available as targets for their respective miRNAs. Therefore, we compared the predicted miRNAs before and after the gene fusion and listed 48 such Ex-miRNAs (Table2). We speculate that many of the these Ex-miRNA will be apparently amplified in cancer cells, as they are not required by their respective BRD4 and NUT targets in fusion gene condition, and hence show dysregulation in cancer.

Table 1. List of miRNA targeting BRD4, NUT and BRD4-NUT fusion gene transcripts

| miRNA targeting BRD4 | miRNA targeting NUT | miRNA targeting BRD4-NUT |
|----------------------|---------------------|--------------------------|
| hsa-miR-4447         | hsa-miR-6861-3p     | hsa-miR-4279             |
| hsa-miR-7159-5p      | hsa-miR-1291        | hsa-miR-4441             |
| hsa-miR-3065-5p      | hsa-miR-4306        | hsa-miR-663b             |
| hsa-miR-765          | hsa-miR-4713-3p     | hsa-miR-4303             |
| hsa-miR-608          | hsa-miR-4535        | hsa-miR-4683             |
| hsa-miR-4492         | hsa-miR-6728-5p     | hsa-miR-6861-3p           |
| hsa-miR-4516         | hsa-miR-3202        | hsa-miR-1291             |
| hsa-miR-4731-5p      | hsa-miR-1285-3p     | hsa-miR-4627             |
| hsa-miR-4651         | hsa-miR-4483        | hsa-miR-4535             |
| hsa-miR-4843         | hsa-miR-6722-3p     | hsa-miR-6728-5p           |
| hsa-miR-6870-5p      | hsa-miR-1909-3p     | hsa-miR-4731-5p           |
| hsa-miR-4227         | hsa-miR-4441        | hsa-miR-5193             |
| hsa-miR-663b         | hsa-miR-654-5p      | hsa-miR-1285-3p           |
| hsa-miR-4279         | hsa-miR-541-3p      | hsa-miR-654-5p            |
| hsa-miR-5787         | hsa-miR-4443        | hsa-miR-541-3p            |
| hsa-miR-185-3p       | hsa-miR-330-5p      | hsa-miR-6751-3p            |
| hsa-miR-4472         | hsa-miR-326         | hsa-miR-1199-5p               |
| hsa-miR-637          | hsa-miR-6751-3p     | hsa-miR-6764-5p           |
| hsa-miR-1277-5p      | hsa-miR-1199-5p     | hsa-miR-1915-3p           |
| hsa-miR-4784         | hsa-miR-660-3p      | hsa-miR-4279             |
| hsa-miR-3150b-3p     | hsa-miR-7106-3p     | hsa-miR-4441             |
| hsa-miR-92a-2-5p     | hsa-miR-6745        | hsa-miR-663b             |
| hsa-miR-6812-5p      | hsa-miR-363-5p      | hsa-miR-4303             |
| hsa-miR-149-3p       | hsa-miR-4446        | hsa-miR-5787             |
| hsa-miR-5703         | hsa-miR-1231        | hsa-miR-4434             |
| hsa-miR-4454         | hsa-miR-6884-5p     | hsa-miR-4303             |
| hsa-miR-4267         | hsa-miR-5196-5p     | hsa-miR-4267             |
| hsa-miR-2861         | hsa-miR-4747-5p     | hsa-miR-4266             |
| hsa-miR-4266         | hsa-miR-328-3p      | hsa-miR-6819-5p           |
| hsa-miR-4266         | hsa-miR-6764-5p     | hsa-miR-6737-5p           |
| hsa-miR-1275         | hsa-miR-1915-3p     | hsa-miR-4279             |
| hsa-miR-4505         | hsa-miR-3123        | hsa-miR-4441             |

In order to predict the Ex-miRNA, BRD4 (1-2830 region) and NUT (170-3795 region) of fused gene were subjected to miRNA prediction using miRDB. Out of 34 miRNAs of BRD4, 28 miRNA were no more required to target the same gene, when compared to its fused form. This situation arises because of the deletion of 2818 bases from 3’end of BRD4 (Table 2). Similarly, out of 33 predicted miRNAs, 20 miRNA were predicted not to target NUT transcript compared to its fused form. The deletion of 170 bases from 5’end of NUT justifies such situation. Overall, we have identified a list of 48 dysregulated “Ex-miRNAs” that target critical genes other than the parent genes. Functional enrichment analysis of KEGG pathway using miRWalk 2.0 tool revealed that Ex-miRNAs targeting 3’UTR region of the target genes belongs to Pathways in cancer (Table 2). Also, miRNAs targeting the genes on their promoter, 5’UTR, CDS and 3’UTR regions were shown to be enriched for “Pathways in cancer” (Table 2). We suggest that an apparent amplification in Ex- microRNA can be used as molecular markers to diagnose NMC.
Table 2. miRNA targeting the critical genes other than the parent genes show functional enrichment in cancer pathways.

| Sl. No. | miRNAs                        | No. of gene targets implicated in cancer pathways |
|---------|-------------------------------|-------------------------------------------------|
| 1.      | hsa-miR-1231                  | 235                                             |
| 2.      | hsa-miR-1275                  | 255                                             |
| 3.      | hsa-miR-1277-5p               | 183                                             |
| 4.      | hsa-miR-149-3p                | 270                                             |
| 5.      | hsa-miR-185-3p                | 265                                             |
| 6.      | hsa-miR-1909-3p               | 245                                             |
| 7.      | hsa-miR-2861                  | 237                                             |
| 8.      | hsa-miR-3065-5p               | 192                                             |
| 9.      | hsa-miR-3123                  | 182                                             |
| 10.     | hsa-miR-3144-5p               | 181                                             |
| 11.     | hsa-miR-3150b-3p              | 238                                             |
| 12.     | hsa-miR-3202                  | 241                                             |
| 13.     | hsa-miR-326                   | 196                                             |
| 14.     | hsa-miR-328-3p                | 187                                             |
| 15.     | hsa-miR-330-5p                | 238                                             |
| 16.     | hsa-miR-363-5p                | 213                                             |
| 17.     | hsa-miR-4266                  | 172                                             |
| 18.     | hsa-miR-4272                  | 166                                             |
| 19.     | hsa-miR-4306                  | 216                                             |
| 20.     | hsa-miR-4434                  | 194                                             |
| 21.     | hsa-miR-4443                  | 70                                              |
| 22.     | hsa-miR-4447                  | 83                                              |
| 23.     | hsa-miR-4456                  | 148                                             |
| 24.     | hsa-miR-4472                  | 148                                             |
| 25.     | hsa-miR-4492                  | 159                                             |
| 26.     | hsa-miR-4505                  | 201                                             |
| 27.     | hsa-miR-4516                  | 159                                             |
| 28.     | hsa-miR-4651                  | 100                                             |
| 29.     | hsa-miR-4713-3p               | 188                                             |
| 30.     | hsa-miR-4747-5p               | 194                                             |
| 31.     | hsa-miR-4784                  | 188                                             |
| 32.     | hsa-miR-5196-5p               | 188                                             |
| 33.     | hsa-miR-5703                  | 179                                             |
| 34.     | hsa-miR-5787                  | 188                                             |
| 35.     | hsa-miR-606                   | 188                                             |
| 36.     | hsa-miR-637                   | 117                                             |
| 37.     | hsa-miR-660-3p                | 105                                             |
| 38.     | hsa-miR-6722-3p               | 85                                              |
| 39.     | hsa-miR-6737-5p               | 188                                             |
| 40.     | hsa-miR-6745                  | 93                                              |
| 41.     | hsa-miR-6812-5p               | 188                                             |
| 42.     | hsa-miR-6819-5p               | 150                                             |
| 43.     | hsa-miR-6870-5p               | 176                                             |
| 44.     | hsa-miR-6884-5p               | 188                                             |
| 45.     | hsa-miR-7106-3p               | 188                                             |
| 46.     | hsa-miR-7159-5p               | 137                                             |
| 47.     | hsa-miR-765                   | 188                                             |
| 48.     | hsa-miR-92a-2-5p              | 116                                             |

Conclusion:
A set of 18, 33, and 34 miRNAs were predicted to target the BRD4-NUT fusion gene transcript, BRD4 and NUT mRNA respectively. Comparative analysis of these miRNAs and functional enrichment of pathway revealed that a set of 48 miRNA might be dysregulated to target the critical genes other than the parent genes (BRD4 and NUT), causing the cancer. Amplification in the expression level of these miRNAs can be used for NMC diagnosis and prognosis.
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