MicroRNA discovery by similarity search to a database of RNA-seq profiles

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1. INTRODUCTION

MicroRNAs (miRNAs) are small, non-coding RNAs 18–24 nucleotides in length that play important roles in various biological and metabolic processes, including signal transduction, developmental timing, cell maintenance and differentiation (Zhang et al., 2006b). MiRNAs are involved in post-transcriptional regulation of gene expression by directly cleaving targeted mRNAs or repressing translation (Bartel, 2004). Many in-vitro and in-silico based approaches have been developed for the prediction of miRNAs. In-vitro based approaches like genetic screening approach have contributed to many founding members of miRNAs. However, due to low efficiency and high cost, these are limited for wider applications. Many in-silico based approaches have been developed based on major characteristic of miRNAs for example hairpin-shaped stem loop structure integrated with homology search (Wang et al., 2005; Dezulian et al., 2006) or evolutionary conservation (Lai et al., 2003; Lim et al., 2003). Besides, methods based on phylogenetic shadowing (Berezikov et al., 2005), neighbor step loop search (Ohler et al., 2004), minimal folding free energy index (Zhang et al., 2006a) and machine learning approaches have also been developed (Table 1). Various plant and animal miRNAs have been identified using these computational approaches. However, many of these methods have sensitivity problems and give a number of false positive results (Bentwich, 2005). Taken together all search methods aim to reduce the search space in their own respective ways (Lindow and Gorodkin, 2007).

Recent advances in high throughput sequencing have provided a new opportunity for genome annotation including prediction of novel miRNAs. Many tools like miRDeep2 (Friedländer et al., 2012), miRDeep* (An et al., 2013), and miRanalyzer (Hackenberg et al., 2011) exploit the aggregated set of RNA-seq reads along with secondary structure potential to annotate a genomic locus as miRNA. Indeed, these tools have great ability to predict novel miRNA genes (Williamson et al., 2013). However, these tools do not completely exploit the subtle differences in
the arrangement of reads mapped to pre-miRNA. Furthermore, many of these tools start by identifying potential precursor locus for miRNA based on strict rules like fixed precursor size of 110 bp or loop region of size 15 bp. This may result in unconventional miRNA patterns like miRNA-offset RNAs (moRs) that encode for up to four distinct, stable small RNAs (Shi et al., 2009) or plant miRNAs that have different biogenesis mechanism be readily missed by these tools (Lelandais-Briere et al., 2010). Furthermore, many microRNA-sized small RNAs have also been reported to be commonly produced not only from miRNA precursors but also from most other classes of structured RNAs like snoRNA and tRNA (Kawaji et al., 2008; Taft et al., 2010).

Several recent studies have recognized that short RNA-seq data, when mapped back to the host genome form read coverage patterns that are distinct and can be used to distinguish between major non-coding RNAs (ncRNAs) such as miRNA, snoRNA and tRNA (Erhard and Zimmer, 2010; Jung et al., 2010; Langenberger et al., 2010, 2012). These read coverage patterns have been referred to as “read profile” or block group (see also Figure 1) and are composed of distinctive clusters of reads (blocks) with similar start and/or stop position. The read profiles are often influenced by secondary structures of the parent transcript and may convey information about the processing mechanism of the transcript like dicer cleavage in this case (Figure 1).

In this study, we present a novel strategy based on pairwise alignment of two read profiles, deepBlockAlign (Langenberger et al., 2012) to predict putative miRNAs in the human genome. We apply this approach on an extensive dataset of read profiles derived from 18 short RNA-seq experiments from ENCODE (ENCODC Consortium, 2011, 2012), and present some putative miRNAs that showed significant similarity to read profiles of known miRNAs from miRBase. We also show distinct classes of miRNA read profiles identified through alignment and hierarchical clustering of read profiles from human and mouse. Furthermore, we present miRNA read profiles that are specific to animals and plants.

## 2. MATERIALS AND METHODS

### 2.1. DATASETS

We downloaded two RNA-seq datasets for the analysis of read profiles. First dataset is comprised of short-reads mapped to the human (hg19, Feb. 2009) genome assembly and is downloaded in BAM format from the ENCODE database (ENCODC Consortium, 2011, 2012). This dataset is comprised of 18 RNA-seq experiments performed on nine human tissues with each having two biological replicates. It is to be noted that prior to sequencing, these biological replicates have been grown and isolated independently. In the following, we will refer to this dataset as ENCODE dataset (Table 2). Second dataset is comprised of short-reads mapped to 4862 distinct miRNAs from 20 organisms in miRBase (Kozomara and Griffiths-Jones, 2011). The short reads are derived from 244 GEO experiments performed on various tissues. In the following, we will refer to this dataset as miRBase dataset (Table 3).

### 2.2. PREPROCESSING OF DATASET

Both datasets were subjected to two pre-processing steps. Firstly, the mapped reads were formatted into BED format. The formatting was done for each of the 18 experiments from ENCODE and 20 organisms from the miRBase dataset, separately (Figures 2A,B). Secondly, each BED format file was processed to identify distinct accumulation of reads by assigning two reads to the same locus, when they were separated by less than 40 nt. We chose 40 nt as the threshold to consider two reads from separate loci based on two observations (a) most short reads are less than 40 nt in length and two genomic loci separated by a region of >40 nt with no mapped reads can most likely be considered as distinct. (b) the loop region of most pre-miRNAs...
Table 2 | The ENCODE dataset is comprised of short-reads from 18 RNA-seq experiments performed on nine human tissues with each having two replicates.

| Tiss  | Replicate1 | Replicate2 |
|-------|------------|------------|
|       | # reads   | # BGs    | # reads   | # BGs    |
| Bl    | 49,280,641 | 16,437    | 56,439,584 | 19,609    |
| Br    | 48,773,897 | 15,148    | 48,394,385 | 13,317    |
| Bt    | 26,713,326 | 13,309    | 40,144,816 | 13,555    |
| Cx    | 41,301,918 | 15,890    | 40,798,294 | 15,948    |
| Ep    | 47,775,551 | 13,522    | 44,163,861 | 10,923    |
| Es    | 35,965,377 | 12,692    | 33,651,242 | 14,697    |
| Li    | 31,930,869 | 6158      | 33,939,724 | 10,684    |
| Lu    | 38,877,787 | 14,511    | 43,732,746 | 15,873    |
| Sn    | 37,649,014 | 6370      | 41,022,882 | 11,268    |

The dataset was downloaded from ENCODE (ENCODE Consortium, 2011) and is composed of a uniform read length of 36 nt.

* Tissues with each having two biological replicates (Bl; Blood; Br; Brain; Bt; Breast; Cx; Cervix; Ep; Epithelium; Es; Embryonic stem cell; Li; Liver; Lu; Lung; Sn; Skin).

# Number of mapped reads.

c Total number of block groups retrieved after preprocessing.

Table 3 | miRBase dataset is comprised of short-reads mapped to 4862 distinct microRNAs from 20 organisms.

| Org  | # reads | # miRNAs | # block groups |
|------|---------|----------|---------------|
| Ame  | 475,288 | 159      | 109           |
| Ath  | 472,502 | 159      | 173           |
| Bfl  | 372,171 | 113      | 41            |
| Bmo  | 2,021,309 | 384   | 264           |
| Cbr  | 17,442  | 115      | 25            |
| Cel  | 1,048,509 | 184   | 93            |
| Cqu  | 379,978  | 66       | 29            |
| Cre  | 1082    | 28       | 11            |
| Cmr  | 9988    | 95       | 19            |
| Cte  | 50,659  | 118      | 8             |
| Dme  | 35,664,132 | 237   | 48            |
| Hsa  | 81,138,802 | 1279  | 550           |
| Mmu  | 913,716,590 | 749   | 682           |
| Nve  | 2711    | 34       | 2             |
| Osa  | 1,506,288 | 440   | 288           |
| Ppc  | 11,176  | 113      | 5             |
| Ppt  | 503,573  | 4224     | 148           |
| Rco  | 1472    | 13       | 2             |
| Spu  | 6458    | 38       | 7             |
| Tca  | 4,861,929 | 196   | 189           |
| Total | 1,046,178,299 | 4862 | 2540          |

Dataset was downloaded from miRBase (Kozomara and Griffiths-Jones, 2011).

* Ame, A. mellifera; Ath, A. thaliana; Bfl, B. floridae; Bmo, B. mori; Cbr, C. briggsae; Cel, C. elegans; Cqu, C. quinquefasciatus; Cre, C. reinhardtii; Cmr, C. remanei; Cte, C. teleta; Dme, D. melanogaster; Hsa, H. sapiens; Mmu, M. musculus; Nve, N. vectens; Osa, O. sativa; Ppc, P. pacificus; Ppt, P. patens; Rco, R. communis; Spu, S. purpuratus; Tca, T. castaneum.

b Number of mapped reads. Total number of GEO experiments are given in brackets. Some experiments are comprised of reads from multiple organisms.

c Total number of distinct miRNAs with mapped reads.

d Total number of block groups or miRNAs retrieved after preprocessing.

e Block groups with > 1 block and ≤200 nt in length that are compiled to form a database of miRNA read profiles (miRPPdb).

is <40 nt in length. Consecutive reads within a locus were divided into blocks using blockbuster (with parameters: -distance 40, -minBlockHeight 2, -minClusterHeight 10, -scale 0.5) (Langenberger et al., 2009) (Figure 2C). blockbuster merges mapped reads into blocks based on their location in the reference genome. Thus, stacks of reads are combined to read blocks which is analogous to tags (set of reads) processed from a specific locus. This strategy greatly reduces the size of the data set and allows the application of more costly algorithms while maintaining structural properties such as position, length and approximate read start sites and ends. The obtained set of one or more blocks at a locus are then called block groups (Table 2).

It is to be noted that for the ENCODE dataset, we discarded all blocks that had read count of <10% with respect to the total reads within its block group in order to (a) ensure all blocks are represented by at least one read, since minimum number of reads in a block group is set to 10 (-minClusterHeight 10); and, (b) nullify the effect of sequencing depth during comparison of block groups across 18 RNA-seq experiments from the ENCODE dataset (Figure 2A). Hereafter, we will use the term block group and read profile, synonymously.

Of the 3838 block groups obtained after the preprocessing of miRBase dataset, we filtered 2540 block groups that had more than one block and were ≤200 nt in length. The 2540 block groups were then compiled to from a database of miRNA read profiles (Figure 2B and Table 3) abbreviated hereafter as miRPPdb. Next, for the ENCODE dataset, we derived 58,161 genomic loci where a block group or read profile is observed in at least one tissue (Figure 2A). For each genomic locus, we retrieved one block group corresponding to the tissue in which the block group had the maximum number of blocks leaving us with 58,161 block groups. All the block groups were then compared to known annotation [1049 miRNA from miRBase v16 (Kozomara and Griffiths-Jones, 2011), 513 tRNA loci from gtrNadb (Chan and Lowe, 2009), 402 snoRNA, 1794 scRNA, 2007 snRNA loci and 722 other RNAs from UCSC annotation (Karolchik et al., 2004)]. Block groups were also compared with 8811 ncRNA annotations from RFam (Gardner et al., 2011). All the block groups whose coordinate overlapped at ≥1 nt with that of known annotations were designated as “annotated” block groups (Table 4). Similarly, block groups were compared with coordinates of exon, intron, 5’ UTR and 3’ UTR region downloaded from UCSC (Karolchik et al., 2004) and were annotated accordingly, if overlapping at >50% else designated as from intergenic region. If a block group overlaps to more than two genomic regions, then the region with maximum overlap is assigned to it. Of the 58,161 block groups, we filtered 4795 block groups that had more than one block and were ≤200 nt in length. Out of 4795
In ENCODE dataset, reads mapped to human genome from each of the 18 RNA-seq experiments were subjected to preprocessing to obtain closely spaced set of reads termed here as "block group." Block groups thus obtained were compiled so as to identify a set of distinct genomic loci that have a block group in at least one experiment. Next, for each locus, we retrieved one block group corresponding to the experiment in which the block group has maximum number of blocks leaving us with 58,161 block groups.

In miRBase dataset, reads mapped to microRNAs from each of the 20 organisms were subjected to preprocessing and block groups thus obtained were compiled as miRNA read profile database (miRRPdb).

Given a set of mapped reads in BED format, we derive closely spaced stack of reads termed here as "block group" using blockbuster (Langenberger et al., 2009). Each block group or read profile is composed of one or more blocks of reads.

Table 4 | Annotation status of 58,161 block groups obtained after preprocessing of ENCODE dataset and their alignment to miRRPdb.

| Annotation     | # block groups | # miRRPdb hits(%) |
|----------------|----------------|-------------------|
|                | Alla           | Filterb           |                   |
| miRNA          | 571            | 285               | 223 (78)          |
| snoRNA         | 468            | 255               | 3 (1)             |
| tRNA           | 625            | 496               | 7 (1)             |
| snRNA          | 395            | 143               | 6 (4)             |
| scRNA          | 187            | 46                | 3 (7)             |
| others         | 277            | 136               | 8 (6)             |
| unannotated    | 55,638         | 3434              | 523 (15)          |
| Total          | 58,161         | 4795              | 773 (16)          |

aBlock groups obtained after preprocessing and are overlapping to non-coding RNA annotation.
bBlock groups with > 1 block and ≤ 200 nt in length.
cBlock groups that have significant alignment score (≥ 0.7) to miRNA read profile database (miRRPdb).

2.3. PERFORMANCE EVALUATION

The performance of the proposed method for the prediction of miRNA is evaluated based on sensitivity, specificity and MCC that are computed using a confusion matrix (2 × 2 contingency table). The confusion matrix is essentially composed of four components (a) number of miRNA that are correctly predicted as miRNA (True Positive, TP), (b) number of miRNA that are incorrectly predicted as non-miRNA (False Negative, FN), (c) number of non-miRNA that are incorrectly predicted as miRNA (False Positive, FP); and, (d) number of non-miRNA that are correctly predicted as non-miRNA (True Negatives, TN). The sensitivity measures the proportion of true positives (TP) out of total number of positives (miRNAs) in the benchmark dataset (TP/TP + FN). Like-wise, specificity measures the
proportion of true negatives (TN) out of total number of negatives (non-miRNAs) in the benchmark dataset \([\text{TN}/(\text{TN} + \text{FP})]\). The MCC is a discrete version of Pearson’s correlation coefficient and is widely used in machine learning to measure the quality of (two-class) binary classifications. It is computed as

\[
\text{MCC} = \frac{(\text{TP} \times \text{TN}) - (\text{FP} \times \text{FN})}{\sqrt{(\text{TP} + \text{FP})(\text{TP} + \text{FN})(\text{TN} + \text{FP})(\text{TN} + \text{FN})}} \tag{1}
\]

2.4. PREDICTION OF PUTATIVE miRNA
In an earlier study, we have developed a tool named deepBlockAlign for the alignment of two read profiles (Langenberger et al., 2012). deepBlockAlign normalizes the read counts by the total reads within a block group followed by a two-tier strategy to align read profiles. The alignment score from deepBlockAlign ranges from 0, suggesting perfect dissimilarity, to 1 for perfect similarity between the two read profiles. It is to be noted that in the absence of any statistical power or background distribution to evaluate the significance of deepBlockAlign scores (since “background” transcription still has to be defined) (Langenberger et al., 2012), we have used deepBlockAlign to compare the 1361 annotated read profiles from ENCODE dataset against miRRPdb in order to identify a meaningful discriminative dba score (see section 3.1 below). Based on the derived dba score, we align 3434 unannotated read profiles from ENCODE dataset against miRRPdb and identify novel genomic regions that have read profiles similar to those of known miRNAs (Figure 3A).

2.5. PREDICTION OF miRNA FAMILIES
To predict distinct classes of miRNA based on their read profiles, we performed cluster analysis on 550 human and 624 mouse read profiles from miRRPdb, separately. In cluster analysis, we first perform all vs. all alignment of all block groups using deepBlockAlign to generate a square matrix of alignment scores. Second, the R package pvclust (Suzuki and Shimodaira, 2006) is used for hierarchical clustering of block groups based on their alignment scores (Figure 3B). pvclust computes the \(p\)-value for each cluster in hierarchical clustering using multiscale bootstrap resampling and indicates how strong the cluster is supported by the data. We select all the clusters comprised of at least 15 read profiles at a \(p\)-value of \(< 0.05\), as families of miRNAs that share similar read profiles. To predict any organism specific read profile class, we also performed the cluster analysis on all 2540 read profiles from miRRPdb.

3. RESULTS
3.1. BENCHMARKING
To benchmark the prediction performance of our proposed method for the prediction of putative miRNAs, we aligned 1361 annotated read profiles (285 miRNA and 1076 other ncRNAs; Table 4) from ENCODE dataset against miRRPdb using deepBlockAlign to generate an alignment score matrix. Next, we perform hierarchical clustering of block groups based on their alignment score using pvclust. pvclust computes the \(p\)-value for each cluster in hierarchical clustering using multiscale bootstrap resampling and indicates how strong the cluster is supported by the data. We select all the clusters with \(p \leq 0.05\) that represent families of microRNAs that share similar read profiles.
deepBlockAlign. For each of the 1361 query read profiles, we selected one read profile from miRRPdb which showed highest alignment score as potential hit for the query read profile. We observed two completely distinct distributions of the alignment scores (Figure 4A), one from alignment of both query and subject read profiles as miRNA (miRNA–miRNA) and another from alignment of any other ncRNA except miRNA as query and miRNA as subject read profile (other-miRNA). ROC curve analysis using R package ROCR (Sing et al., 2005) showed a high AUC of 0.93 suggesting that miRNA read profiles have characteristic features that are distinct from read profiles of other ncRNAs and can be employed for confident prediction of miRNA. (Figure 4B).

Indeed, most miRNA–miRNA read profile alignments (223 out of 285) showed an alignment score of ≥0.7 whereas most other-miRNA read profile alignment scores (1049 out of 1076) were <0.7. As the False Positive Rate (FPR) tend to increase above the alignment score of 0.7 (yellow-green intersection) in Figure 4B, we chose 0.7 as the default cut-off alignment score to consider an alignment between unannotated with miRNA read profile as significant and unannotated read profile as putative miRNA candidate.

We further estimated the MCC (Matthews et al., 1975), sensitivity and specificity of the method based on the confusion matrix created for 1361 alignments at a cut-off read profile alignment score of 0.7. The MCC is computed using a confusion matrix that is composed of four components (a) 223 miRNA–miRNA read profile alignments with score ≥0.7 as True Positives (TP), (b) 62 miRNA–miRNA read profile alignments with score <0.7 as False Negatives (FN), (c) 1049 other-miRNA read profile alignments with score <0.7 as True Negatives (TN); and, (d) 27 other-miRNA read profile alignments with score ≥0.7 as False Positives (FP). Based on the confusion matrix, the sensitivity, specificity and MCC of 0.78, 0.97 and 0.80, respectively was observed. We also performed the 5-fold cross validation by splitting the benchmark dataset (1361 annotated read profiles) into five equal and evenly distributed (similar ratio of miRNA and non-miRNA read profiles) datasets. For each of the five rounds of cross-validation, we aligned four of the five datasets against miRRPdb and derived the optimal cut-off of alignment score at which maximum MCC is observed. Using the derived cut-off, the performance of the method is evaluated on the remaining fifth dataset. We observed a mean MCC of 0.80 ± 0.02 and an AUC of 0.93 ± 0.02. A mean cut-off score of 0.7 was observed during the cross-validation and is used hereafter for all the further analysis. In the light of recent reports that many microRNA-sized small RNAs are commonly produced not only from miRNA precursors but also from other classes of structured RNAs like snoRNA and tRNA (Kawaiji et al., 2008; Taft et al., 2009), the above measures can be regarded as a reasonable estimate of the performance of this approach in the prediction of novel miRNA candidates.

Furthermore, we compared the performance of our method to an already available tool, miRanalyzer that detect miRNAs using short RNA-seq data (Hackenberg et al., 2011). We chose miRanalyzer because it is one of the widely used tool for miRNA prediction using RNA-seq data, has a prediction performance comparable to other miRNA prediction tools such as miRDeep2 (Williamson et al., 2013) and can be readily applied on our benchmark dataset of mapped reads. We evaluated the performance of miRanalyzer on the mapped reads corresponding to 1361 annotated read profiles from our ENCODE dataset using both the default and model mode of miRanalyzer. In default mode, miRanalyzer made predictions by first mapping reads to known miRNAs from miRBase followed by using random forest model for the remaining set of reads. In model mode, all the predictions are exclusively based on random forest model. We
observed an AUC of 0.94 and 0.95 for the former and later mode (Supplementary document available at the web page, http://trh.dk/resources/mirdba). Out of 223 known miRNAs that were correctly predicted by our method, 160 and 208 were also predicted by miRanalyzer using default and model mode. The high performance using the model mode is not surprising since the models have been trained using random forest on the secondary structure features of the same set of known miRNAs within the benchmark dataset. It is to be noted that while computing the AUC, we considered only those ncRNAs, out of 1361 ncRNAs, that have corresponding reads mapped by the miRanalyzer. When computing the AUC for all the 1361 ncRNAs irrespective of their mapping status, we observed a low AUC of 0.64 and 0.68 for the two modes, respectively.

To further evaluate the performance of our method, we analyzed the short RNA-seq data corresponding to Human HeLa cells (GSE10829) that was used to benchmark the performance of miRanalyzer. We observed an AUC of 0.92 for the dataset which is approximately the same AUC of 0.93 observed for 1361 read profiles from the ENCODE dataset. Considering that our method has not implemented any specific miRNA characteristic feature such as hairpin loop size or length of pre-miRNA, it is intriguing that we obtain a high scoring performance of 0.92 in AUC on the original miRanalyzer dataset. In this context, the performance of miRanalyzer on the HeLa dataset has been reported as an AUC of 0.98. Taken together, we observed comparable performance of our method to that of miRanalyzer suggesting that reasonably confident prediction of miRNAs can be performed based on alignment of read profiles. Furthermore, since many of the miRNAs were predicted exclusively by either of the two methods, the diversity in the two prediction approaches can potentially be applied for an enhanced scheme based on the two methods.

3.2. IDENTIFICATION OF NEWLY ADDED miRNA ENTRIES IN miRBase
Since, we used miRBase Release 16 to annotate read profiles from the ENCODE dataset, we checked how many of the newly added miRNAs in miRBase Release 19 are identified by our approach. A total of 558 new human miRNAs has been added in the miRBase since release 16. We observed that out of 558, 49 miRNAs are represented as read profiles in our ENCODE dataset. Of these 49 miRNAs, 35 were observed to have a significant alignment score (≥0.7) with miRRPdb, suggesting that the read profile based search can identify most of the newly added miRNAs given that their expression is sufficiently captured during the RNA-seq experiments.

3.3. PUTATIVE microRNAs
To predict putative miRNA candidates, we aligned 3434 unannotated read profiles from the ENCODE dataset against miRNPdb using deepBlockAlign. We retrieved 523 alignments with an alignment score of ≥0.7 between an unannotated and miRNA read profile from miRNPdb. Figure 5 shows one such alignment between an unannotated read profile from chr1:20841720-20841781(+) and a read profile of hsa-mir-519b. We can observe high similarity between the two read profiles characterized by similar number, size and distance between the read blocks. Furthermore, the relative arrangement of reads is also very similar between the two, as illustrated through the “Block alignment”. We also checked for the consensus RNA secondary structure by extracting the Multiz alignment (Blanchette et al., 2004) corresponding to the unannotated region from the UCSC browser (Figure 5D). The consensus RNA structure is predicted using the PETFold webserver (Seemann et al., 2011), and it showed a highly conserved hairpin loop structure characteristic for miRNA secondary structure. This further suggests that this region potentially can harbor a miRNA. The unannotated status of this region was checked by searching for possible annotations in miRBase, UCSC tracks of ENCODE/GENCODE (Version 7, 10, and 12), RefSeq, tRNA and snoRNAs.

3.4. PUTATIVE miRNAs ARE LOCATED IN REGIONS THAT HAVE SHORT OR POORLY CONSERVED MAF BLOCKS
Secondary structure conservation across evolutionary tree is considered a compelling evidence for the biological function of a RNA. Therefore, many tools like Evofold (Pedersen et al., 2006), RNAAlifold (Bernhart et al., 2008), PETFold (Seemann et al., 2008), and RNAz (Gruber et al., 2010) integrate sequence conservation across multiple organisms (Multiz alignments) with RNA folding algorithms for reliable prediction of RNA secondary structure like hairpin loop structure of miRNA. A Multiz alignment represents multiple sequence alignments across a set of species along with measures of evolutionary conservation (Blanchette et al., 2004). For comparative analysis, we retrieved 46-way and 13-way Multiz alignments corresponding to both 523 miRNA candidates and 23 known miRNAs from UCSC (Karolchik et al., 2004) and our in house compilation of Multiz alignments, respectively. The in house 13-way Multiz alignments have been made on 13 representative organisms in the evolutionary tree (Anthon et al., in preparation). As expected, we observed less MAF block fragmentation over the location of 523 miRNA candidates for the 13-way alignment in comparison to the 46-way alignment, see Figure 6.

For the 13-way alignment, we observed that 272 out of the 523 candidates have too short Multiz alignments (<50 nt) in comparison to 49 out of 223 for known miRNAs (Figure 6). Furthermore, Multiz alignments from putative miRNAs were observed to have a significantly lower average pairwise identity in comparison to that from known miRNAs (Figure 6). A similar pattern of either short or a significantly lower average pairwise identity in Multiz alignments from putative miRNA in comparison to known miRNAs was observed for 46-way Multiz alignments. In this context, absence of well-defined Multiz alignments may well contribute to many of these putative miRNAs not being identified through methods based on a set of aligned RNA sequences for the prediction of non-coding RNAs (Gorodkin et al., 2010).

3.5. SECONDARY STRUCTURE ANALYSIS OF PUTATIVE miRNAs
To independently analyze the 523 miRNA candidate regions for secondary structure, we processed all the corresponding Multiz blocks for quality such as length and average pairwise identity. Interestingly only 278 Multiz alignments have length ≥45 nt and we therefore only carried out prediction on these. Since,
miRNAs usually are \( \sim 22 \) nt in length, a Multiz alignment of length < 45 can seldom harbor a miRNA hairpin loop structure. Therefore, for the remaining 245 Multiz alignments no attempt to predict RNA secondary structures using sequences from multiple organisms was made. To predict the secondary structure using multiple sequences, we employed two widely used tools, CMfinder (Yao et al., 2006) and RNAz (Gruber et al., 2010), respectively. CMfinder is an expectation maximization algorithm that uses covariance models to predict secondary structure motifs for a set of unaligned sequences (Yao et al., 2006). On the other hand, RNAz is a support vector machine (SVM) based method that evaluates evolutionary conserved pre-aligned set of sequences with thermodynamic stability of RNA secondary structure to detect structural ncRNAs (Gruber et al., 2010). Thus by using these two tools, we aimed to predict secondary structure for regions that are relatively conserved (pre-aligned, RNAz) and regions that are not so well conserved in terms of sequence (unaligned, CMfinder). These methods also complement each other (Gorodkin et al., 2010).

We used 13-way Multiz alignments corresponding to 278 miRNA candidates as input and considered a \( P \)-score from CMfinder \( \geq 50 \) (FDR, False Discovery Rate of 0.27) (Seemann et al., in Preparation) and a \( P \)-value from RNAz \( > 0.9 \) (z-score \( < -3 \) and FDR 0.1) (Gruber et al., 2010) as significant. Based on these thresholds, we predicted conserved RNA secondary structure motifs in 42 putative miRNA candidate regions by CMfinder (17 candidates), RNAz (39 candidates), or by both (14 candidates) CMfinder and RNAz. Out of these 42 putative miRNA candidates, 33 were also predicted as miRNA by miR-analyzer (Hackenberg et al., 2011). Interestingly, for 39 putative miRNA regions predicted to have conserved secondary structure by RNAz, we observed a significantly higher pairwise sequence identity in comparison to rest of the 474 putative miRNA regions (Wilcoxon test, \( P \)-value < 0.05). This observation again points to the dependency of tools like RNAz on pre-aligned set of sequences for ncRNA prediction as discussed in the previous section. We also observed 13 candidate regions where despite high average pairwise identify of \( \geq 90\% \) in Multiz alignments, no conserved RNA secondary structure was predicted. In conclusion, we obtain 523 novel miRNA candidates of which 42 are further supported by a predicted conserved hairpin loop RNA secondary structure.
3.6. TWO DISTINCT CLASSES OF READ PROFILES IN HUMAN AND MOUSE

On independent clustering of 550 and 624 human and mouse known read profiles from miRRPdb, we observed two well-separated tree nodes comprised of three constituent clusters of read profiles for both human and mouse, respectively (Figure 7). In mouse, 503 out of 624 (81%) read profiles were represented in the three clusters. In contrast, only 190 out of 550 (38%) read profiles were represented in the three clusters from human. This might well be due to the sequencing depth in RNA-seq experiments from mouse, which is one order of magnitude higher in comparison to human, see Table 3 for details. Higher sequencing depth helps by including only well-defined read blocks while excluding background read noise during preprocessing step (see section 2) thereby leading to well-defined clusters of read profiles.

Nevertheless, in both human and mouse, we observed two classes of read profiles. The first was comprised of characteristic miRNA-like read profiles which had two read blocks with low entropy (Figure 7A) and the second was comprised of read profiles with a significantly high number of read blocks (p-value < 0.05, Fisher’s exact test) and entropy (p-value < 0.05, Fisher’s exact test). Entropy is a measure of the degree of randomness in the arrangement of reads within a read profile (Langenberger et al., 2012, holds the details). An earlier study showed similar findings in Ciona intestinalis where half of the miRNA loci encoded up to four distinct, stable small RNAs. The additional RNAs were shown to be generated from sequences immediately adjacent to the predicted ~60 nt pre-miRNA (Shi et al., 2009).

Figure 7B illustrates three example of read profiles that have more than two read blocks, first example has expression in loop region that is interestingly even higher in comparison to that of miR* and other two examples with expression from region partially overlapping to miR–miR*, a pattern similar to those of miRNA-offset RNAs (moRs) albeit different in not being completely adjacent to the miR–miR* (Shi et al., 2009).

3.7. READ PROFILES SPECIFIC TO PLANTS AND ANIMALS

We also performed alignment and hierarchical clustering of all 2540 read profiles in miRRPdb leaving us with 11 distinct clusters of miRNA read profiles (Figure 8). We observed an interesting pattern of read profile distribution across different organisms, while four cluster of read profiles (cluster 361, 371, 395, and 396) were mostly observed in animals (hsa, H. sapiens and mmu, M. musculus), three clusters (cluster 389, 386, and 301) were almost exclusively observed in plants (osa, O. sativa and ath, A. thaliana). The rest of the read profiles clusters were observed across both plant and animals. Next, we compared the length and entropy of the read profiles observed specifically in plant and animals. While almost all of the animal specific read profiles were of 60–90 nucleotides in length, plant specific read profiles were either short (<60 nt) or long (≥90 nt) in length. A higher average length of plant pre-miRNA in comparison to human pri-miRNAs has also been observed in an earlier study (Lindow and Gorodkin, 2007) Plant specific read profiles were also observed to have a significantly higher entropy in comparison to those from animals (p-value < 0.05, Kolmogorov–Smirnov test). This observation may be attributed to the different biogenesis mechanism of miRNAs in plants and animals (Lelandais-Briere et al., 2010).

3.8. WEB SERVER FOR miRNA IDENTIFICATION BASED ON SIMILARITY SEARCH TO miRRPdb

Based on our proposed method, we developed a web server to predict putative miRNA candidates. The web server is available at http://rth.dk/resources/mirdba and facilitate users to align a query read profile against the database of miRNA read profiles...
Pundhir and Gorodkin miRNA read profile alignment

(miRRPdb). The standard input is a set of reads mapped to a genomic region of interest in BED format. Users can also adjust several optional parameters that may affect the prediction results, although the default setting is optimal in many cases. The read profile alignment results are presented in a user friendly graphical format and are composed of top two read profiles in miRRPdb that have an alignment score of ≥0.7 (default threshold derived based on benchmarking, see section 3.1) with the query read profile. It should be noted that the alignment score is only an indicator toward similarity of two read profiles. Further checks like similarity in RNA secondary structure should be done to strengthen the prediction results. The alignment of read profiles is visualized using programs developed in PERL, Latex, and R. There is also an option to visualize the read profiles in the UCSC genome browser.

4. DISCUSSION

MiRNAs are important regulators of various biological and metabolic processes and computational prediction of miRNA on a genome wide scale is an active research area. We have presented a novel strategy based on alignment of read profiles generated from short RNA-seq data to predict novel miRNA candidates. The alignment of read profiles was performed using a previously published tool, deepBlockAlign. The applicability of the proposed method has been demonstrated by using two short RNA-seq datasets (ENCODE and miRBase datasets). Totals of 4795 and 2540 read profiles were retrieved after preprocessing of ENCODE and miRBase datasets, respectively. 2540 read profiles from miRBase were then compiled to form a database of miRNA read profiles (miRRPdb). Upon alignment of 1361 annotated (285 miRNA and 1024 other ncRNA) read profiles from

![Figure 7](image-url)

**FIGURE 7** | Clusters of read profiles obtained after hierarchical clustering of 550 and 624 read profiles from Human and Mouse, respectively. (A) Two distinct and well-separated tree nodes were observed for both plant and animals (red and blue). For better representation, the count of read profiles within each cluster are shown on top of the corresponding branch. In comparison to read profiles in first node, read profiles in second node were characterized by a significantly high number of read blocks (p-value = 4.3e-115, Fisher’s exact test) and entropy (p-value = 0.005, Fisher’s exact test, blue shade bars). An earlier study has shown similar finding in *Ciona intestinalis* where half of the miRNA loci encode up to four distinct, stable small RNAs (Shi et al., 2009). (B) Three example read profiles with more than two read blocks, (i) beside expression in miR–miR*, loop region also showed expression even higher in comparison to miR*. (ii) and (iii) many reads are observed from region partially overlapping to miR–miR*, a pattern similar to those of miRNA-offset RNAs (moRs) albeit different in not being completely adjacent to the miR–miR* (Shi et al., 2009).
we observed comparable performance between the two methods on both our dataset as well as the dataset used to benchmark miRanalyzer. Since our method is only based on alignment of read profiles, it has some notable advantages: (a) it can predict putative miRNAs in genomic regions that are devoid of RNA secondary structure information either due to low sequence conservation across multiple organisms that many tools like RNAz (Gruber et al., 2010) require or due to inherent limitation of tools based on single sequence to predict a RNA secondary structure. In this context, many mRNA regions that were predicted to form large, single stranded loops by RNAfold (Gruber et al., 2008) have been shown to form highly base-paired regions using experimental methods (Zheng et al., 2010; Li et al., 2012). (b) it also provides the most similar read profile to a query read profile with respect to arrangement and expression of reads. This can be a useful information, especially when the most similar read profile from miRRPdb is comprised of reads derived from only one experiment, and; (c) similarity search of read profiles can be applied to identify lineage specific miRNA read profiles based on arrangement of reads (see section 3.6).

In total, our method based on alignment of read profiles can be a suitable complement to the other tools such as miRanalyzer and miRDeep that also use short RNA-seq data for miRNA prediction.

Using the proposed method on 3434 unannotated read profiles from ENCODE data set, we predicted 523 unannotated read profiles as putative miRNAs. On RNA secondary structure analysis, 42 of these putative miRNAs were observed to have conserved RNA secondary structure. Furthermore, many of the 523 putative miRNAs were characterized by either short or remarkably low average pairwise identity in corresponding Multiz alignments. Since many RNA secondary structure prediction methods use a set of pre-aligned sequences for ncRNA prediction, absence of well-defined Multiz alignments can contribute to the lack of secondary structure predictions overlapping with many of our miRNA candidates. Global screen of ncRNA candidates using multiple sequence alignment has also been suggested to fail in regions of low sequence similarity (Torarinsson et al., 2006, 2008).

We have shown an unannotated genomic region that share read profile similar to a miRNA along with a characteristic hairpin loop RNA secondary structure with a 3′ overhang. On cluster analysis of read profiles from human and mouse separately, we observed two distinct clusters of read profiles, one with read profiles that have two read blocks and low entropy corresponding to miR–miR* and the other with read profiles that have more than two blocks and a significantly higher entropy in comparison to the former. Many of these additional read blocks were observed in regions adjacent or partially overlapping to miR–miR* and can be possible candidates for miRNA-offset RNAs (moRs).

On cluster analysis of all 2540 read profiles from 18 different organisms in miRRPdb, we observed 11 distinct clusters of read profiles. Interestingly, four of these read profile clusters were mostly observed in animals and another three read profile clusters were mostly observed in plants. The remaining four read profile clusters were observed in both plant and animals. Furthermore, most of the plant specific read profiles were either too
short (<60 nt) or long (≥90 nt) in comparison to animal specific read profiles. Besides, plant read profiles were also observed to have a significantly higher entropy in comparison to the animal read profiles.

Further work includes analysis of miRNA read profiles that have more than two read blocks for potential miRNA-offset miRNAs. In this context, these miRNAs can also be wrongly annotated as miRNA in miRBase. A meta tool integrating the predictions of methods such as miRanalyzer, miRDeep2 and our method would also be useful for the identification of miRNA candidates that are supported by diverse set of prediction approaches. It would also be of wide interest to classify miRNAs based on read profile patterns that are specific to a given tissue, pathological condition or organism. In this context, closer inspection of plant and animal specific read profiles may also reveal novel and organism specific features in their miRNA read profiles.

FUNDING
This work is funded in part by the Danish Strategic Research Council (Strategic Growth technologies), the Danish Independent Research Council (Technology and Production), and the Danish Center for Scientific Computation.

ACKNOWLEDGMENTS
We would like to thank the reviewers for their helpful comments and suggestions, Christian Anthon for the help in development of the web server and Betina Wingreen Jensen for proof-reading the manuscript.

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Received: 01 February 2013; paper pending publication: 26 March 2013; accepted: 21 June 2013; published online: 11 July 2013.
Citation: Pundhir S and Gorodkin J (2013) MicroRNA discovery by similarity search to a database of RNA-seq profiles. Front. Genet. 4:133. doi: 10.3389/fgene.2013.00133
This article was submitted to Frontiers in Bioinformatics and Computational Biology, a specialty of Frontiers in Genetics.
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