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| Subject Section  maTE: Discovering Expressed MicroRNA - Target Interactions  Malik Yousef1\*, Loai Abddallah2, and Jens Allmer3,4  1Department of Community Information Systems, Zefat Academic College, Zefat, 13206, Israel.  e-mail: [malik.yousef@gmail.com](mailto:malik.yousef@gmail.com)  2Department of Information Systems, The Max Stern Yezreel Valley Academic College, Israel.  e-mail: [Loai1984@gmail.com](mailto:Loai1984@gmail.com)  3Applied Bioinformatics, Bioscience, Wageningen University and Research, Wageningen, the Netherlands. e-mail: [jens@allmer.de](mailto:jens@allmer.de)  4Horticulture, Bioscience, Wageningen University and Research, Wageningen, the Netherlands. e-mail: [jens@allmer.de](mailto:jens@allmer.de)  \*To whom correspondence should be addressed.  Associate Editor: XXXXXXX  Received on XXXXX; revised on XXXXX; accepted on XXXXX  Abstract  **Motivation:** Disease is often manifested via changes in transcript and protein abundance. MicroRNAs (miRNAs) are instrumental in regulating protein abundance and may measurably influence transcript levels, as well. MicroRNAs often target more than one mRNA (for human the average is three) and mRNAs are often targeted by more than one miRNA (for the genes considered in this study, the average is also three). Therefore, it is difficult to deduce the miRNAs that may cause the observed differential gene expression.  We present a novel approach, maTE, based on machine learning, which integrates information about miRNA target genes with gene expression data. maTE depends on the availability of a sufficient amount of patient and control samples. The samples are used to train classifiers to accurately classify the samples on a per miRNA basis. Multiple high scoring miRNAs are used to build a final classifier to improve separation.  **Results:** The aim of the study is to find a set of miRNAs causing regulation of their target genes that best explains the difference between groups (e.g.: cancer vs. control). maTE provides a list of significant groups of genes where each group is targeted by a specific microRNA. For the datasets used in this study, maTE generally achieves an accuracy well above 80%. It is of note, that when the accuracy is much lower (e.g.: ~50%) the set of miRNAs provided is likely not causative for the difference in expression.  This new approach of integrating miRNA regulation with expression data yields powerful results and is independent of external labels and training data. Thereby, it opens up new avenues for exploring miRNA regulation and may pave the way for the development of miRNA-based biomarkers and drugs.  **Availability:** The KNIME workflow, implementing maTE, is available at Bioinformatics online.  **Contact:** Malik Yousef, [malik.yousef@gmail.com](mailto:malik.yousef@gmail.com) and Jens Allmer, [jens@allmer.de](mailto:jens@allmer.de)  **Supplementary information:** Supplementary data are available at *Bioinformatics* online. |

# Introduction

In the past decade it has become clear that microRNAs (miRNAs) are involved in most human diseases (Tüfekci *et al.*, 2014). They are post-transcriptional regulators of protein expression, but recently have been shown to be involved in transcription, as well (Liu *et al.*, 2018). Mature miRNAs are short 18-24 nucleotide long single stranded RNA sequences derived from larger hairpin structures (pre-miRNAs) via a molecular genesis pathway (Erson-Bensan, 2014). These mature miRNAs act as recognition sequences for their target mRNAs within the RISC complex. MicroRNAs can have hundreds of target mRNAs and each of these can be targeted by many miRNAs leading to a many to many regulative relationship. Actual interactions are only possible when both the miRNA and its mRNA target are present in the same space and time (Saçar and Allmer, 2013). Thus miRNA-mRNA interactions are under spatiotemporal control. Transcription of miRNAs seems to be predominantly responsible for controlling possible miRNA-mRNA interactions (Melo and Melo, 2014). A large fraction of human genes are under miRNA control (Jones-Rhoades and Bartel, 2004) and more than 90% of human KEGG pathways contain genes which either harbor miRNAs or are targeted by miRNAs (Hamzeiy *et al.*, 2015, 2017). More than 2000 human miRNAs are available in miRBase (Griffiths-Jones *et al.*, 2006) taken together with their prevalence throughout pathways, makes these post-transcriptional regulators key elements of gene regulation.

High-throughput approaches for identifying and sequencing RNAs and proteins are available via next generation sequencing and mass spectrometry, respectively. For example, the gene expression omnibus (GEO) provides access to microarray measurements (Wheeler *et al.*, 2008), and the sequence read archive (SRA) hosts next generation sequencing data (Leinonen *et al.*, 2011). There are also data repositories for proteomics such as PRIDE (Vizcaíno *et al.*, 2010). Unfortunately, high-throughput measurements encompassing coordinated measurement of protein abundance and miRNA abundance applicable for the present study were not available in PRIDE. Such a dataset would provide a gold standard since one mode of action of miRNAs is translational repression which cannot be queried on the transcriptional level. Another miRNA mode of action, mRNA degradation, however, can be accessed via transcriptomics. For many diseases, measurements of gene expression are available for large patient cohorts. A few examples are collected in Table 1.

MicroRNAs are also transcripts and can be measured via dedicated arrays, short read sequencing, or other more specialized methods such as HITS-CLIP (Saçar Demirci *et al.*, 2019). Some miRNAs are located in transcription units (TUs) and their expression could be inferred from the expression of the enclosing TU. Apart from the miRNA expression levels, their associated targets are important. Experimentally verified miRNA targets are available in databases such as TarBase (Vergoulis *et al.*, 2012) and miRTarBase (Chou *et al.*, 2018). Taken together, transcriptomics data can provide information about miRNAs and their targets’ expression levels.

The need for integration of miRNA and target expression data has been identified previously by Gunaratne et al. (Gunaratne *et al.*, 2010). They also identified a need for computational tools facilitating such analyses for novel and publicly available data. Today, although various experimental methods exist for the measurement of miRNA abundance, the need for computational tools still exists because experimental methods are still involved and expensive. A variety of computational tools for the task exist and they use different resources and approaches to accomplish a specific task. One important component for such research is information about miRNA targets. MiRGator (Cho *et al.*, 2013) and mirDIP (Tokar *et al.*, 2018) are two tools which integrate targeting data from multiple resources. Other approaches such as data integration for modeling the miRNA:mRNA regulation have been proposed. NAViGaTing is one such method which explores miRNAs involvement in well-known signaling pathways and their associations with disease (Shirdel *et al.*, 2011). Chen & Yan developed regularized least squares for miRNA-disease association, using semi-supervised learning, to uncover the relationship between diseases and miRNAs (Chen and Yan, 2015). Steinfeld et al. introduced a computational approach (miTEA) that infers miRNA activity from high-throughput data using a novel statistical methodology, called minimum-mHG (Steinfeld *et al.*, 2013). They apply their approach to matched mRNA and miRNA expression profiles for cancer cell lines to achieve mutual enrichment in two ranked lists. MULSEA (Cohn-Alperovich *et al.*, 2016) is a similar approach to miTEA. It features an algorithm collecting factors that can be aggregated into one ranked list which is strongly associated with an input ranked list. Zeng et al. summarized different computational approaches for predicting potential disease-related miRNA based on networks (Zeng *et al.*, 2016). They point out that the main principle of those approaches is the calculation of similarity among disease and miRNA in the expression networks. They further categorized the approaches into two groups: similarity measure-based and machine learning-based. The latter mainly aims to distinguish positive miRNA–disease associations from large-scale negative miRNA–disease associations. The data used for this kind of research are miRNA–disease, disease-phenotype, miRNA association, gene interaction, and protein interaction networks. Such data is transformed into a network and used to compute the similarities among nodes, particularly between a miRNA and a disease to infer associations. For example, mirConnX (Huang *et al.*, 2011) creates a disease specific regulatory network by integrating gene expression data, sequence information, miRNA targeting, and transcription factor binding information. Another tool for the reconstruction of regulatory networks is MAGIA2 (Bisognin *et al.*, 2012). It integrates miRNA target prediction and gene expression data to compile the networks. MiSEA (Çorapçıoğlu and Oğul, 2015) uses gene expression and miRNA-seq data for the enrichment of miRNAs. MiSEA allows further analysis and grouping by, for example, family classification and disease association. Different from the other tools above, CSmiRTar (Wu *et al.*, 2017) allows the mining of gene expression dataset using miRNA and miRNA target filtering.

None of these tools are similar to the system that we present here although the approach employed in this study uses, similarly to the above methods, mRNA expression data. The expression data is integrated via miRNA target association drawn from databases such as miRTarBase. Specifically, the objective is to find a set of miRNAs that best explains differential mRNA expression among samples. To achieve this, we developed a novel machine learning-based approach using two class classification. However, apart from patient and control data no other data annotation is necessary and no additional negative data needs to be created. Instead, we use Monte Carlo cross validation (Xu and Liang, 2001) for repeated random sampling of the dataset and training of predictive models. In each round the miRNAs which generate the most accurate models are combined and an integrated miRNA group model is trained. After at least 25 iterations (here we use 100), an approximation of the set of miRNAs which best explains the difference in mRNA expression is determined.

The results are compared to our previous method SVM-RCE, which is conceptually similar to maTE. The average accuracy for the selecteddatasets is 0.17 points less for maTE. However, it is our contention that maTE, when assigning a lower accuracy indicates that the difference in mRNA expression is not caused by the set of miRNAs and their targets used in the experiment. Furthermore, maTE found, on average, 13 more differentially expressed mRNAs than SVM-RCE and was able to associate them with miRNAs. The much higher variance in average score generated by maTE when compared to SVM-RCE (Table 3) seems to be useful as a quality measure, as well. In the future, we aim to further evaluate the novel algorithm and amend it with an optimization approach to improve on the selection of the best combination of miRNAs explaining the difference in mRNA expression. Taken together with the ability of maTE to assign low scores to cases where miRNA involvement is unlikely, this algorithm will facilitate future research associating miRNAs with disease.

# Materials and Methods

## 2.1 Data

### 2.1.1 Gene Expression Data

10 human gene expression datasets were downloaded from the gene expression omnibus (Clough and Barrett, 2016). For all datasets disease (positive) and control (negative) data was available (Table 1). Additionally, a dataset (GSE19536) containing both mRNA and microRNA measurements (Enerly et al., 2011) was used to validate maTE.

Table 1: Description of the 10 data sets used in our study. The data sets are obtained from GEO. Each entry has the GEO code the name of the data, the number of samples, the number of genes that were measured and the classes of the data.

|  |  |  |
| --- | --- | --- |
| **GEO Accession** | **Title** | **#Samples/Classes/#Genes** |
| GDS1962 | Glioma-derived stem cell factor effect on angiogenesis in the brain | #Samples=180  non-tumor=23 (neg)  astrocytomas=26 (pos)  glioblastomas=131 (pos)  #genes=54613 |
| GDS2519 | Early-stage Parkinson's disease: whole blood | #Samples=105  healthy control=22(neg)  neurodegenerative disease control=33(neg)  Parkinson disease=50 (pos)  #genes=22282 |
| GDS3268 | Colon epithelial biopsies of ulcerative colitis patients | #Samples=202  normal=73 (pos)  ulcerative colitis=129 (neg)  #genes=44289 |
| GDS3900 | Fear conditioning effect on hybrid mouse diversity panel: hippocampus and striatum | #Samples=198  hippocampus=100 (pos)  striatum=98 (neg)  #genes=25696 |
| GDS3929 | Tobacco smoke effect on maternal and fetal cells | #Samples=183  non-smoker=128 (pos)  smoker=55 (neg)  #genes=18253 |
| GDS2547 | Metastatic prostate cancer (HG-U95C) | #Samples=164  normal=75 (pos)  tumor=89 (neg)  #genes=12645 |
| GDS5499 | Pulmonary hypertensions: PBMCs | #Samples=140  control=41 (neg)  idiopathic pulmonary arterial hypertension=30 (pos) scleroderma-associated pulm. arterial hypert.=42(pos) systemic sclerosis (SSc) without pulm. hypert.=19(pos) SSc, interstitial lung disease & pulm. hypert.=8(pos)  #genes=49575 |
| GDS3646 | Celiac disease: primary leukocytes | #Samples=132  healthy control=22(neg)  celiac disease==110 (pos)  #genes=22184 |
| GDS3874 | Diabetic children: peripheral blood mononuclear cells (U133A) | #Samples=117  healthy = 24 (neg)  type 1,2 diabetes =93(pos)  #genes=22282 |
| GDS3837 | Non-small cell lung carcinoma in female nonsmokers | #Samples=120  Lung Cancer = 60 (pos)  Control = 60 (neg)  #genes=30621 |

### 2.1.2 MicorRNA Targets

MicroRNA targeting data was downloaded from miRTarBase release 7.0 (Chou *et al.*, 2016). For compatibility with the gene expression data, only human miRNAs and their targets were considered. All data without experimental evidence from either Reporter assay, Western blot, or both were discarded. In total 740 human miRNAs with 8496 targets remained after filtering (File S1). Table 2 provides a subset of the data for illustration.

Table 2: Part of the miRNA-Target gene table; complete data can be found in the supplement (File S1).

|  |  |
| --- | --- |
| **MicroRNA** | **Target Genes List** |
| HSA-LET-7A-3P | CCND1, CCND2, E2F2 |
| HSA-LET-7D-5P | HMGA2, APP, DICER1, SLC11A2, IL13, MPL, AGO1, TNFRSF10B, COL3A1 |
| HSA-MIR-103A-2-5P | PDCD10 |
| HSA-MIR-129-2-3P | SOX4, UBE2F, CCP110, BCL2L2, MYC, CDK6 |
| HSA-MIR-140-5P | HDAC4, VEGFA, PDGFRA, DNMT1, DNPEP, SOX2, OSTM1, FGF9, TGFBR1, ALDH1A1, SOX9, IGF1R, FZD6, RALA, PAX6, HDAC7, LAMC1, ADA, MMD, PIN1, STAT1, GALC, HMGN5, SOX4, FGFRL1, SMURF1 |
| HSA-MIR-638 | OSCP1, SP2, SOX2, CDK2, STARD10, PLD1, PTEN |
| HSA-MIR-944 | S100PBP, HECW2 |

**Ranking Algorithm -**

**X**s: any subset of the input gene expression data X, the features are gene expression values

**M {** is a list of microRNAs

**Grouping function g(M)**- for each *mi*associate the names of genes (Genes ID) that targeted by microRNA *mi* (See Table 2).

***f*** *is a scalar ()*: split into train and test data

**r:** repeated times (iteration)

**res={}** for aggregation the scores for each *mi*

**Generate Rank for each *mi-Rank(mi):***

For each *mi* in M

*smi*=0;

Perform *r* time (here r=5) steps 1-5:

1. Perform stratified random sampling to split Xs into train Xt and test Xv data sets according to *f* (here 80:20)
2. Remove all genes (features) from Xt and Xv which are not targets of *mi*
3. Train classifier on Xt (here Random Forest)
4. *t* = Test classifier on Xv –calculate performance
5. *smi  = smi + t;*

*Score(mi)*= *smi* /*r* ; Aggregate performance

*res*=

**Output**

*Return res ( res = {Rank(m1),Rank(m2),…,Rank(mp)} )*

Algorithm 1:The Ranking method R(), a main component of the maTE algorithm.

As expected, the number of targets varies among miRNAs and Figure 1 presents the distribution of the number of targets per miRNA. 50% of the miRNAs have 3 or less targets (median: 3) but a few miRNAs have more than 100 assigned targets in miRTarBase: miR-155-5p (223), miR-145-5p (143), miR-21-5p (136), miR-34a-5p (132), miR-125b-5p (119), and miR\_20a-3p (106).

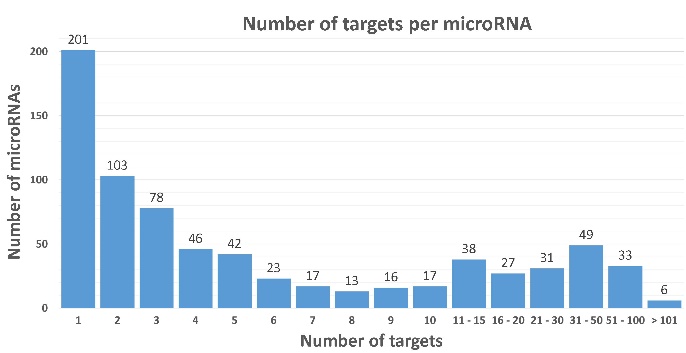


Figure 1: Distribution of the number of target genes per miRNA for human in miRTarBase. Median number of targets is 3, upper quartile is 10 targets and the maximum observed is 223 targets for one miRNA (miR-155-5p).

Unless these miRNAs are extremely abundant, their effect in vivo should be minor (Saçar Demirci *et al.*, 2019).

## 2.2 maTE Algorithm

The maTE algorithm considers miRNAs and their target gene expression for two conditions: control (negative) and disease (positive). Each condition is represented by their gene expression values for the experiment (each sample contributes one feature). The main motivation for the algorithm is that it is not known *a priori* which miRNAs may be involved in causing a disease. Therefore, machine learning is used to learn which miRNAs are associated with gene expression (Figure 2), leveraging knowledge learned in previous studies (Yousef *et al.*, 2007, 2009; AbdAllah *et al.*, 2017). One of the main components of the maTE tool is the ranking stage *R()* (see Algorithm 1).

The expression of each gene (typically thousands) in the gene expression dataset represents a feature. Features are grouped by the miRNAs that can target them according to miRTarBase (median 3 targets per miRNA). For example, one group related to the hsa-let-7a-3p miRNA contains the target genes CCND1, CCND2, E2F2 and another group related to hsa-let7d-5p contains the genes HMGA2, APP, DICER1, SLC11A2, IL13, MPL, AGO1, TNFRSF10B, COL3A1. MicroRNAs may share all or a subset of their targets. The gene DICER1 is targeted by 20 miRNAs, for example, by miR-581 and miR-3928-3p. For the latter it is the only currently known target while miR-581 also targets EDEM1. For another example, about 20 miRNAs target both PTEN and BCL2 among other targets.

In the following, we define the maTE algorithm, which consists of the miRNA-gene ranking (Algorithm 1) and the integration parts (Algorithm 2).

### 2.2.1 Ranking lists of genes associated with miRNAs

Let X denote a two-class gene expression dataset consisting of ℓ covariate samples and *n* genes (Figure 2, input). The classes could be disease and control, or any experimental condition versus a control or another experimental condition.

Let *g()* be a function grouping genes into clusters. Here *g()* can be any algorithm which groups genes. For example, Yousef et al. previously used the k-means algorithm for grouping by gene expression (Yousef *et al.*, 2007). For the maTE algorithm, *g()* is provided by miRNAs and their targets. For example, *g*(hsa-let-7a-3p) groups the genes CCND1, CCND2, and E2F2 (See Table 2). More formally, let *g(mi)* define the grouping based on the *mi* miRNA’s targets (i: index of miRNAs available in miRTarBase). We chose to use miRTarBase, but other databases such as TarBase or computationally predicted targets would provide other/additional valid options. Note, that the number of targets varies with the chosen miRNA (Figure 1).

By iterating over all *g(mi)* the miRNAs are ranked according to their ability to differentiate the two classes based on the test outcomes following training of a random forest classifier (RF) using an 80:20 split into training and testing data (Figure 2: yellow section). First the grouping function *g(mi)* extracts the relevant gene expression data rows from X and then RF is applied. The ranking function is defined as R(Xs, *g(mi), f, r)* where Xs is any subset of X, f defines the data split into training and testing (here 80:20), and *r* the number of repetitions (here 5). *R()* then returns the average accuracy.. The pseudo code is provided as Algorithm 1.

### 2.2.2 Integration

Following the ranking step for each miRNA as grouping factor, the best *j* miRNAs (we set *j* to 2) are selected and their groups (i.e.: their targets) are combined (Figure 2: yellow section). An RF model is trained with *g()* provided by the best *j* miRNAs instead of just *mi*. The model is tested and the performance measures are recorded.

We performed the complete procedure 100 times using Monte Carlo cross validation (Figure 2: N-fold Cross validation loop). For each fold, the input is stratified random sampled and split into training and testing sets. The training set is submitted to t-test analysis. At maximum 2000 differentially expressed genes with a p-value below 0.05 are selected. The selected genes are then used to filter the test dataset so that both datasets contain the same genes. Within each iteration first the miRNAs are ranked and then the best *j* miRNAs are used to train a RF classifier combining the *j* best miRNAs. The pseudo code is available as Algorithm 2.

### 2.2.3 Workflow

**maTE Algorithm**

**Objective**

maTE aims to select j miRNAs with target genes that can best classify samples by expressions.

**Input**

**X**: gene expression data with two-class labels, the features are genes expression.

**M {:** list of microRNAs (here from miRTarBase) where *p* is the number of microRNAs.

**Grouping function *g*(M)**- for each *mi*associate the names of genes (Genes ID) that are targeted by microRNA *mi* (See Table 2).

**Algorithm**

M\*={} empty list

Perform N-fold cross validation (here N = 100):

Randomly split data by samples into train (Xt) and test (Xv) parts,

performs steps 1- 6:

1. Xtf = filter genes (features) from training data by t-test (here p-value ≤ 0.05 and maximum number of filtered genes≤ 2000)
2. Xvf = remove all genes from Xv that are not in Xtf
3. miRp = **R(Xtf, g(M), *f* ,r)** (here *f*=80:20 with stratified random sampling; r=5). R() is the procedure in Algorithm 1, the output will be miRp =*{Rank(m1),Rank(m2),…,Rank(mp)}*
4. M\*= Sort( miRp) according to performance; best first
5. M\* = {m\*1, m\*2,..,m\*j} , Select best j miRNAs (here j=2)
6. Filter Xtf and Xvf by *g*(M\* ),now Xtf and Xvf represented by genes that are targeted by microRNA from M\*.

Train classifier using Xtf and Xvf (here random forest)

Test classifier using Xvf

**Output**

Report performance (e.g.: average accuracy)

Algorithm 2: The overall algorithm of maTE which depends on the R() method (see Algorithm 1).

The algorithms we developed can be implemented in many systems. In order to test our approach and to provide a proof of principle as well as an interface for users, we developed the approach using the Konstanz Information Miner (KNIME). The resulting KNIME workflow (Figure 3) is available for download from Bioinformatics online and <https://malikyousef.com/>. The workflow in Figure 3 consists of processing nodes and data connections (lines/edges). Data travels along the edges through the workflow. For better readability, and to increase modularity, meta-nodes (grey nodes; e.g.: Preprocess incl. t-test) encapsulate sub workflows. Workflow control includes programming constructs such as loops (blue nodes) and branching. The maTE workflow in Figure 3 contains user input in the orange boxes and presents the results in the green box. Processing is performed in the yellow and blue box. The green dots under nodes indicate that the process has successfully succeeded.

**2.3 Classification Approach**

We used the random forest (RF) classifier implemented by the platform KNIME (Berthold *et al.*, 2008). The classifier was trained and tested with a split into 80% training and 20% testing data. We have considered the under-sampling balancing approach to deal with imbalanced data. The under-sampling balancing approach is reducing the size of the abundant class by keeping all samples in the rare class and randomly selecting an equal number of samples in the abundant class. This approach is repeated during each round of cross-validation. We implement 100-fold Monte Carlo cross-validation (MCCV) (Xu and Liang, 2001) for model training. We used the default RF parameters where the split criterion is information gain ratio. We didn’t limit the number of levels (tree depth) and the number of models was set to 100. Slight changes to these values did not change overall performance.

### 2.3.1 Model Performance Evaluation

For each established model, we calculated a number of statistical measures such as sensitivity, specificity, and accuracy to evaluate model performance. The following formulations were used to calculate the statistics (with TP: true positive, FP: false positive, TN: true negative, and FN referring to false negative classifications):

Sensitivity (SE, Recall) = TP / (TP + FN)

Specificity (SP) = TN / (TN + FP)

Accuracy (ACC) = (TP + TN) / (TP + TN + FP + FN)

All reported performance measures refer to the average of 100-fold MCCV. The positive class and negative class for each data is described in Table 1.

## 2.4 Recursive Cluster Elimination

We have previously developed a different method with a similar aim, SVM-RCE (Yousef *et al.*, 2007), and later compared the methodology with other approaches (Yousef *et al.*, 2009)(AbdAllah *et al.*, 2017). While there is similarity in the general idea of using classification, the methods differ in the way genes are grouped. The SVM-RCE algorithm groups gene expression based on the k-means clustering algorithm’s grouping of the gene expression data (intrinsic information). Our novel approach (maTE) groups gene expression data based on information about miRNAs and their target genes (extrinsic information). The SVM-RCE algorithm performs three steps: 1) the clustering step which groups the genes into clusters based on k-means, 2) the scoring step that evaluates the importance of each cluster of genes by internal cross validation, and 3) the RCE step that removes the clusters with lower scores which is repeated until reaching a desired number of clusters. In order to benchmark our novel approach, we performed SVM-RCE analysis on all datasets used in this study using the default settings as in (Yousef *et al.*, 2007).

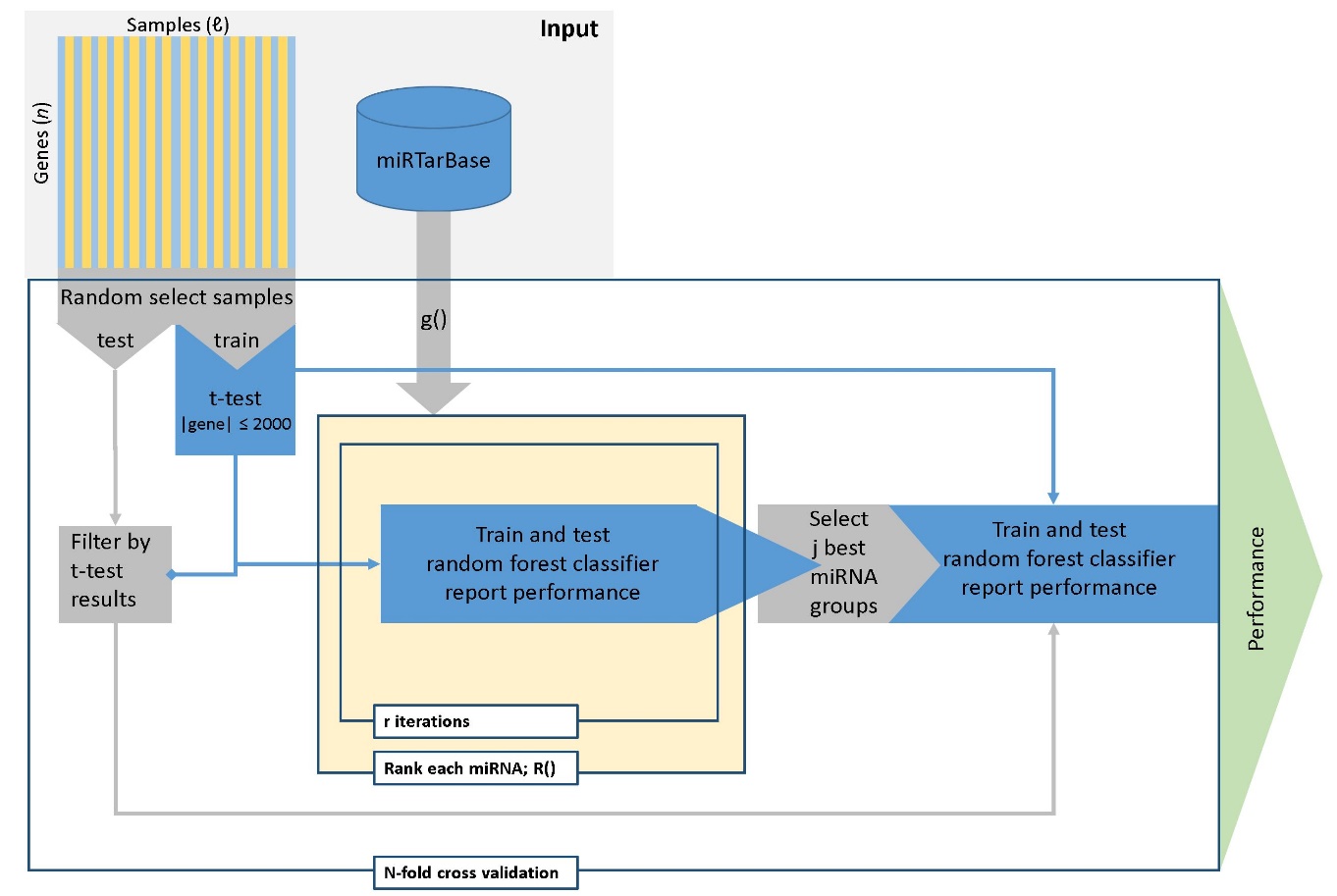


Figure 2: maTE work flow. The two main steps of the workflow are creating models for each miRNA (yellow) and then combining multiple miRNAs into one model and training a classifier using these miRNAs. Input: samples are horizontal with the two classes represented in yellow and blue. Genes are represented by the vertical bars. miRTarBase depicts the miRNA target data from miRTarBase. Loops are represented by rectangles with a tag (e.g.: N-fold cross validation). T-test calculations are based on the training data but filtering is applied to the genes in both training and testing data.

# Results and Discussion

We previously showed that for categorizing miRNAs into species, using machine learning, a minimum of 100 examples was needed (Yousef, Khalifa, *et al.*, 2017)(Yousef, Nigatu, *et al.*, 2017). Therefore, we selected datasets with large number of samples (Table 1). In these datasets, patient and control samples are indicated, but miRNAs that lead to changes in mRNA expression and miRNAs that do not are unknown and unlabeled *a priori*. However, classification, in general, depends on annotated positive and negative data. Here we use classification such that the annotation of which miRNA is significantly different between the two classes can be learned without the need for annotated examples. The trick allowing this feat is creating an abundance of machine learning models from the data. The model performance on the withheld test data indicates whether it effectively separates between classes. The better the performance the more likely it points to a biological explanation. Here our interest was to determine miRNAs that best describe the differential mRNA expression between patient and control samples. However, the same approach could be used to tackle many other biological questions such as pathway enrichment.

We applied both our novel methodology maTE and our previous, related, algorithm: SVM-RCE to the selected data in Table 1 (Table 3). Our novel approach, maTE, is searching for significant miRNAs and their targets, thereby limiting the search space to significantly differentially expressed ones, while SVM-RCE is searching for significant genes in the complete space, not taking into account extrinsic grouping factors like miRNAs. Table 3 summarizes the result of hundreds of thousands of trained models. It is obvious that SVM-RCE (avg. acc.: 0.94) outperforms maTE (avg. acc.: 0.78) for all datasets. However, SVM-RCE seems relatively indiscriminate and leads to similar results for all datasets which is due to the missing extrinsic grouping factor. Therefore, SVM-RCE or other approaches focusing on DE analysis might compile effects of different regulatory mechanisms; whereas maTE focuses on effects caused by miRNAs. maTE seems to discriminate between datasets where known miRNAs may not be the main cause of the observed difference in gene expression. Specifically, the differential gene expression for datasets GDS2519, GDS3929, and GDS3646 does not seem to be caused by miRNAs and the contribution of miRNAs seems to be low for the dataset GDS3268. Except for datasets GDS3646, GDS3929, and GDS2547 maTE generally collects more genes explaining the overall difference among states. The first two datasets are unlikely to be caused by miRNAs and the latter one’s difference in expression may only have a limited contribution via miRNAs.

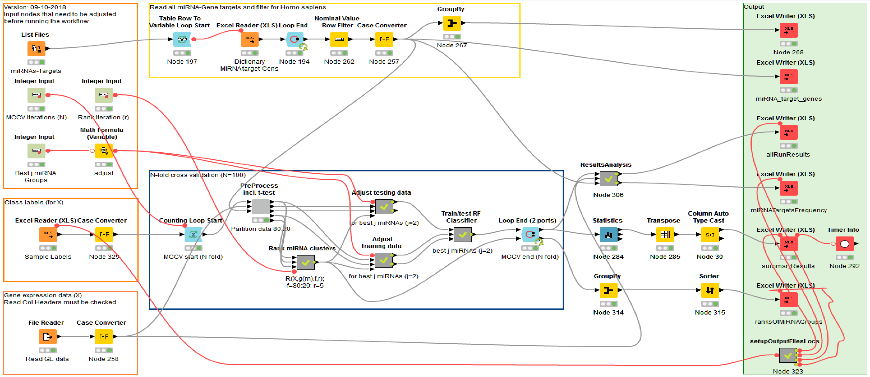


Figure 3: maTE work flow. Overview of the KNIME workflow available at Bioinformatics online. Input that needs to be adjusted is in the orange boxes to the left. The blue central box contains the MCCV and further logic is encapsulated in meta-nodes such as PreProcess and R(). Results are stored within the green box based on the location of the files with the class labels which can be adjusted in Node 323 (bottom of green box) if desired.

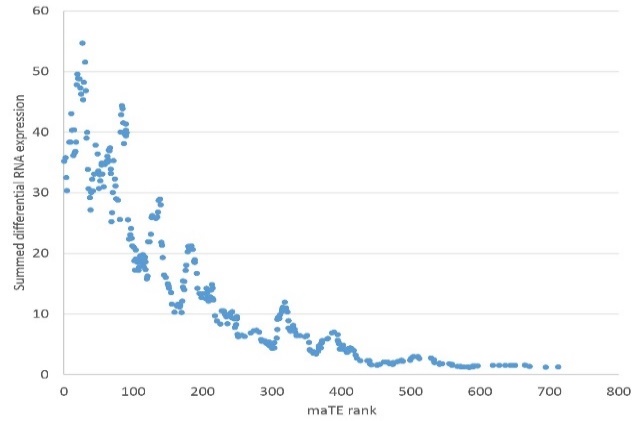


Figure 4 The maTE rank for miRNAs versus the sum of their absolute target differential expressions

For the data analyzed above, no ground truth is known and, therefore, it is difficult to assign a confidence measure to our new approach. Some studies have measured mRNA and miRNA differential expression (Enerly *et al.*, 2011). It was our aim to use such data to benchmark our new method. Naively, the miRNAs selected by our algorithm should have high differential expression between conditions. Unfortunately, this is not the case since many mRNAs are targeted by multiple miRNAs (Figure 1) so that additionally a combined effect should be taken into account. Our current approach is, however, miRNA centric and selects miRNAs that maximally explain the differential mRNA expression. Combined effects of miRNAs using our method are found by selecting the top *j* miRNAs (see maTE algorithm step 7). Here we report results with j =2. However, we have also tested different values of j such as 3, 4, and 5 leading to little improvement. In the future we aim to employ optimization to select the set of miRNAs best able to separate between classes. The motivation for this are the spikes in the trend in Figure 4. MicroRNAs with lower rank but high impact on the DE can cause such spikes. An optimization approach would be able to combine these miRNAs into a minimal set explaining a large part of the differential RNA expression.

We applied maTE and SVM-RCE to a miRNA-mRNA Breast Tumors dataset (Enerly *et al.*, 2011) considering the mRNA expression of 15 basal-like and 41 luminal-A samples. These are the subtypes with strongest reciprocal mRNA expression profiles (GO identifier GSE19536). We will refer to this experiment as LumA\_vs\_Basal.

SVM-RCE ranks the importance of each gene by the number of times it appears on each level of the RCE levels. For example, if we start the process with the top 1000 genes selected by t-test from the training data and start with 100 clusters, then we will have 27 levels of RCE (each time we reduce the number of clusters by 10%). We track the frequency of each gene in each level over the 100 iteration. The score is the total number of frequencies divided by 2700 (100 iteration \* 27 levels). The complete results for the top 1000 genes are available in supplementary file S2. The top gene is GATA3 that is required for the development of the mammary gland and has been implicated in breast cancer. MLPH (rank 2) is also a marker for breast cancer survival. Other genes in the top 10 such as BCMP11 are also implicated in breast cancer. Thus, SVM-RCE collected relevant genes from the dataset. Since it does not assign any relevance to miRNAs, it cannot be compared with maTE on this dataset that consists of coordinated miRNA and mRNA measurements. It is of note, that GATA3 was also part of three miRNA target sets in the top 10 miRNAs deduced by maTE. MLPH and BCMP11 were not found as targets since they were not available as targets in miRTarBase.

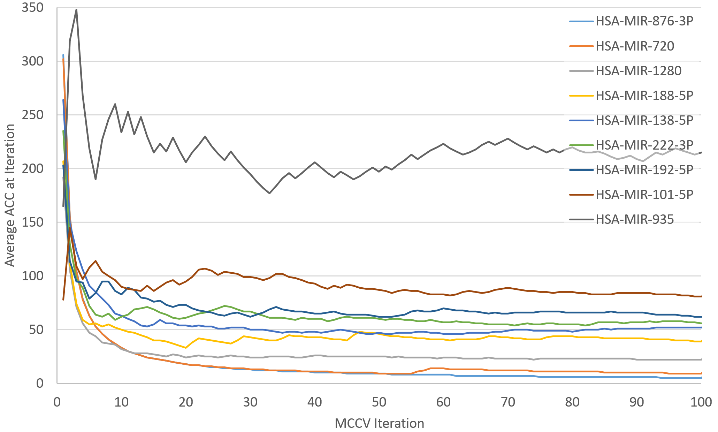


Figure 5: The development of the average rank for selected miRNAs for 100 MCCV iterations for the experiment GDS3837.

Table 3: Accuracy results for both methods, SVM-RCE and maTE. SE: sensitivity, SP: specificity, ACC: accuracy, stdev: standard deviation and #G is the number of genes. We consider the top 2 clusters for SVM-RCE and top two miRNAs for the maTE.

|  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | **SVM-RCE** | | | | | **maTE** | | | | |
| **Dataset** | **SE** | **SP** | **ACC** | **stdev** | **#G** | **SE** | **SP** | **ACC** | **stdev** | **#G** |
| GDS1962 | 0.97 | 1.00 | 0.98 | 0.06 | 44 | 0.96 | 1.00 | 0.98 | 0.05 | 66 |
| GDS2519 | 0.87 | 0.90 | 0.88 | 0.14 | 24 | 0.64 | 0.57 | 0.61 | 0.10 | 62 |
| GDS3268 | 0.89 | 0.88 | 0.88 | 0.08 | 42 | 0.78 | 0.71 | 0.74 | 0.07 | 84 |
| GDS3900 | 1.00 | 1.00 | 1.00 | 0.00 | 64 | 1.00 | 0.95 | 0.98 | 0.01 | 86 |
| GDS3929 | 0.98 | 0.96 | 0.97 | 0.05 | 81 | 0.50 | 0.57 | 0.54 | 0.10 | 26 |
| GDS2547 | 0.89 | 0.81 | 0.85 | 0.08 | 54 | 0.87 | 1.00 | 0.83 | 0.07 | 34 |
| GDS5499 | 0.96 | 0.95 | 0.95 | 0.07 | 59 | 0.79 | 0.97 | 0.88 | 0.09 | 90 |
| GDS3646 | 0.96 | 0.93 | 0.95 | 0.10 | 29 | 0.42 | 0.63 | 0.53 | 0.16 | 29 |
| GDS3874 | 0.97 | 0.97 | 0.97 | 0.00 | 17 | 0.77 | 0.90 | 0.84 | 0.15 | 52 |
| GDS3837 | 0.97 | 0.96 | 0.96 | 0.05 | 63 | 0.76 | 0.99 | 0.88 | 0.04 | 79 |

Some outcomes of the breast cancer study (Enerly *et al.*, 2011) used here have also been confirmed in (Sandhu *et al.*, 2014). Especially, miR-146a is overexpressed in basal-like breast cancer cells. However, some p53-dependent changes, including expression of miR-134, miR-146a and miR-181b, were found to be subtype specific. maTE also assigns some importance to miR-146a (rank 30) but it is not within the top 10 of miRNAs explaining the difference between luminal and basal.

Since miRNAs can have several targets, we wanted to see whether the summed absolute differential expression of these targets versus the rank assigned by maTE has any correlation. The expectation that with better ranks more differential expression is explained in general holds true (Figure 4). This result shows that maTE fulfills the expectation and at least for mir-146a agrees with previous results. Other miRNAs found to be important in separating luminal from basal type are mir-128, miR-17 (part of miR-17~92 family) and the miR~30 family (Iorio *et al.*, 2005); see (Enerly *et al.*, 2011) and (Sandhu *et al.*, 2014). These miRNAs, or representatives of their families are found in the top 20 of maTE assignments. It can be expected, that some results differ among algorithms and the top assignments by maTE do not overlap with findings by (Enerly *et al.*, 2011). For the top assignments we submitted the deregulated targets to Reactome analysis and found that many of the miR-93 (rank 1) targets are (also) under p53 control and involved in the PTEN pathway. Furthermore, a role of miR-93 (rank 2) in breast cancer has previously been confirmed (Liang *et al.*, 2017; Hao *et al.*, 2018). The same is also true for miR-24 (Khodadadi-Jamayran *et al.*, 2018; Yu *et al.*, 2018). The targets of miR-24 (rank 3) are involved in senescence control and their downregulation will likely lead to avoidance of cell death. Among the top 10 there is only miR-510 with a single target (SPDEF). Interestingly, SPDEF has been implicated with various cancer types among them breast cancer (Sood *et al.*, 2017) as well as miR-510 (Guo *et al.*, 2013). These findings confirm all top assignments of maTE to be implicated in breast cancer and thereby qualitatively validate the strategy employed.

In our experiments, we used 100 MCCV iterations, which can take a few hours on a regular personal computer. Therefore, we were interested whether 100 iterations are necessary. To test that, we recorded all miRNA ranks for the 100 iterations for the GDS3837 experiment. We then calculated the average rank for development per iteration i.e.: the average of all ranks for each miRNA until the iteration. The development of the average rank is plotted for nine miRNAs (Figure 5) including the highest ranked one (hsa-miR-876-3p) and the lowest ranked one (hsa-miR-935).

From Figure 5 it can be deduced, that 100 fold MCCV is not necessary. Additionally, we calculated how many iterations were needed for each miRNA to reach its average rank. Averaged over all miRNAs the result is about 24 iterations. Therefore, using much less than 100-fold MCCV seems adequate for future calculations.

# Conclusion

The analysis of differential gene expression is employed in various biological scenarios. For instance, to differentiate between control and disease state. It has become clear that regulation occurs on many levels and that some regulatory switches cause a large downstream response while others lead to more subtle changes. MicroRNAs are both, master switches and fine tuners of protein expression. One mode of action of miRNAs leading to transcript degradation is accessible on the transcriptomic level.

One of the novelties of our approach, called maTE, is that we provide not just a significant list of deregulated genes, but group them by their targeting miRNAs. To the best of our knowledge, this is the first account of such an approach. The generated information is very valuable to the biology community and it will allow addressing novel biological questions.

We applied our approach to breast cancer data (Enerly *et al.*, 2011) and were able to confirm some of the previous findings. However, the top assignments made via maTE point to different miRNAs and their targets than in the original assessment. Interestingly, the maTE assignments have clear associations with breast cancer, something missing in the original study.

In the future maTE can be extended with existing tools. For example, MetaMirClust (Chan *et al.*, 2012) deduces miRNA Clusters and a grouping function build on such clusters instead of single miRNAs would be worth considering for maTE, assuming coordinated transcription of miRNA clusters. MAGIA2 (Bisognin *et al.*, 2012) and CSMirTar (Wu *et al.*, 2017) can be employed as an alternative to miRTarBase to have a more comprehensive list of target genes per miRNA. MAGIA2 and miRConnX (Huang *et al.*, 2011) can further be utilized to construct regulatory circuits and perform pathway enrichment following relevant detection of miRNAs by maTE. In order to filter maTE input, miSEA (Çorapçıoğlu and Oğul, 2015) can be employed to reduce the number of miRNAs for datasets where both miRNA and mRNA expression is available.

While maTE selects the top j (here 2) miRNAs; in the future, it would be beneficial to determine a minimal network of miRNAs and their targets which maximizes the amount of differential expression among states. For this we aim to add an optimization step embedding the yellow part of the algorithm (Figure 2) using for example a genetic algorithm.

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