

A PROBABILISTIC FRAMEWORK TO IMPROVE MICRORNA TARGET PREDICTION BY INCORPORATING PROTEOMICS DATA

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Due to the difficulties in identifying microRNA (miRNA) targets experimentally in a high-throughput manner, several computational approaches have been proposed. To this date, most leading algorithms are based on sequence information alone. However, there has been limited overlap between these predictions, implying high false-positive rates, which underlines the limitation of sequence-based approaches. Considering the repressive nature of miRNAs at the mRNA translational level, here we describe a probabilistic model to make predictions by combining sequence complementarity, miRNA expression

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level, and protein abundance. Our underlying assumption is that, given sequence complementarity between a miRNA and its putative mRNA targets, the miRNA expression level should be high and the protein abundance of the mRNA should be low. Having identified a set of confident predictions, we then built a second probabilistic model to trace back to the mRNA expression of the confident targets to investigate the mechanisms of the miRNA-mediated post-transcriptional regulation. Our results suggest that translational repression (which has no effect on mRNA level), instead of mRNA degradation, is the dominant mechanism in miRNA regulation. This observation explained the previously observed discordant correlation between mRNA expression and protein abundance.

Keywords: Gene regulation; microRNA target prediction; proteomics data analysis.

1. Introduction

MicroRNAs (miRNAs) are a class of small non-coding RNAs, typically about 22 nucleotides in length, and are known to block protein synthesis of their target genes by binding to the 3' UTR of the mRNA transcripts with perfect (in plants) or imperfect (in animals and *c. elegans*) base pairing.¹ It was estimated that thousands of genes in the mammalian genome are under regulation by miRNA at the post-transcriptional level,² and they have been shown to have many important functional roles.³

Despite microRNAs' importance and prevalence, it has proved difficult to experimentally identify and validate their target genes. To this date, only 40 miRNA targets have been confirmed in mouse and 200 in human.⁴ As an alternative, a number of computational prediction programs had been developed and were widely used to predict miRNA targets *in silico* (see Refs. 2 and 5–7). Most of these computational programs combine two types of data in making predictions: sequence complementarity between the miRNA and the putative target binding sites, and the evolutionary conservation of such sites (for a review, see Ref. 4). Although great progress has been made in improving prediction accuracy, accurate prediction of miRNA targets remains challenging, the major difficulty being the lack of agreement among these algorithms. A recent benchmark study has compared the predicted targets of several leading algorithms and reported significant discrepancy among them.⁴ The disagreement among these algorithms can be largely attributed to the different scoring schemes and weights given to imperfect base pairing between miRNA and binding sites and evolutionary conservation of the binding sites. Moreover, some of these sequence-based algorithms are known to be less robust as slight changes in parameters often result in very different predictions.⁸

Because of the repressive nature of miRNAs' regulatory roles and the availability of the genome-wide mRNA expression data, it was suggested that using gene expressions data could be helpful in predicting true miRNA targets.⁹ The rationale of such approach is the following: if a miRNA is highly expressed and a putative target gene is lowly expressed in a particular tissue type, then it is considered as an additional piece of evidence that the candidate gene is a true target.

Although this algorithm produced encouraging results¹⁰ and such a negative correlation between the expression levels of miRNA and target mRNA has been observed in various experiments (see Refs. 9, 11 and 12), special caution should be taken in this approach for the following reasons: (1) The reported instances of negative correlation between miRNA and mRNA expression levels are limited to a very small fraction of the known miRNAs with anecdotal evidence; it is uncertain whether such negative correlation can be extrapolated to other types of miRNAs and their targets. (2) It is known that, in contrast to the miRNAs in plants, the miRNAs in animals typically have imperfect sequence complementarity to their target sites and function mostly by binding to the target sites to inhibit the translation process, instead of causing degradation of the mRNA transcripts.^{13,14} (3) In some cases, a strong positive correlation has also been observed between the expression levels of miRNA and their target mRNAs,¹⁵ which could be attributed to common regulators shared by the miRNA and their target genes. All the evidence described above suggests that in animals, the repression effect of miRNAs on their target genes is more obviously manifested at the translational level (i.e. protein abundance); thus, identifying miRNA targets solely based on transcriptional data might be less effective. In contrast, regardless of by degradation or by translational repression, the protein abundance of the target genes should be always negatively regulated. Motivated by this observation and by previous work, in this paper we propose a new Bayesian approach to predicting miRNA targets in mouse using proteomic data in a high-throughput manner. In Ref. 16, researchers have used miRNA expression data and proteomics data to identify miRNA targets. However, their predictions are mainly based on biological experiments and no computational algorithm was proposed, and sequence-based methods such as TargetScan, PicTar and miRanda were only used to support their predictions. Besides, their experimental predictions were only performed in the rat kidney. To our knowledge, our approach is the first computational one that incorporates miRNA expression data and proteomics data in multiple tissues and organs to carry out high-throughput miRNA target predictions.

Our method consists of two steps. In the first step, it takes as input a set of putative miRNA target genes derived from sequence information alone; it then applies a probabilistic model using protein abundance data to assign confidence scores to the predicted miRNA-target pairs. In the second step, another probabilistic model is applied to the miRNA and mRNA expression data to predict whether the miRNA-mediated regulation is through translation repression or mRNA degradation. Figure 1 shows a flowchart of our approach.

2. Data Gathering

The mouse protein abundance data were derived from a recently published mass spectrometry study,¹⁷ in which the abundance of 4,768 proteins across six mouse organs (brain, heart, liver, lung, placenta and kidney) was surveyed.

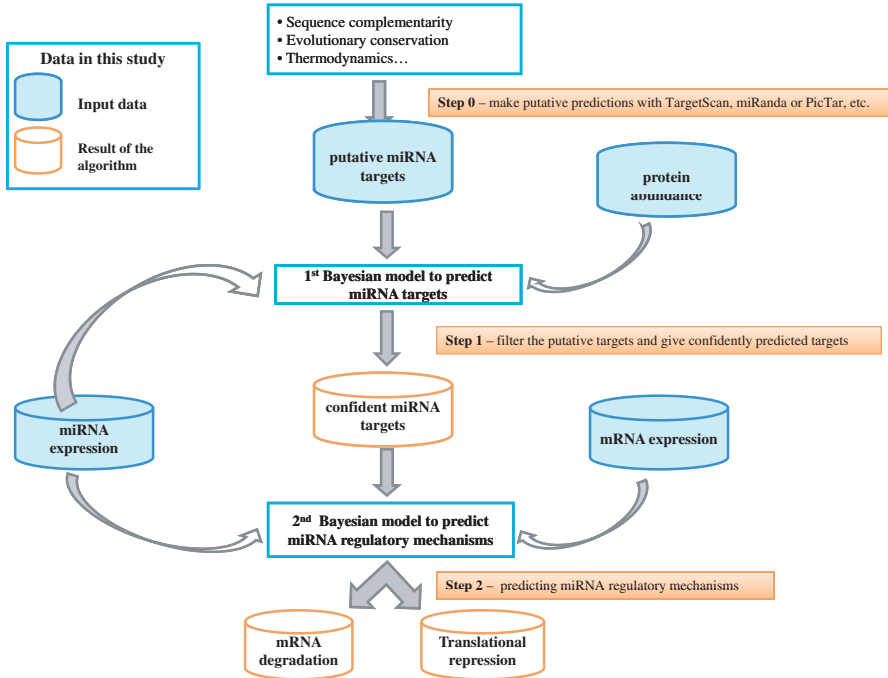


Fig. 1. A flowchart of the algorithms described in this work. The described algorithm takes four types of experimental data: (1) a set of putative miRNA targets, (2) protein abundance, (3) miRNA expression profiles, and (4) mRNA expression profiles. Details can be found in the main text.

After comparing with gene expression data from two microarray studies,^{18,19} 1,758 proteins were confidently cross-mapped to their corresponding mRNAs. The incomplete coverage of the proteomic data was likely due to instrumentation bias, stringent filtering rules of database search or instability of low-level transcripts. The miRNA expression data were extracted from previous published microarray studies.²⁰ These authors also used TargetScan and miRanda separately to derive two lists of putative miRNA targets in mouse. The normalized mRNA expression profiling of 41,699 transcripts in 55 tissues and organs was from Ref. 19. Among these, 1,758 were confidently cross-mapped to the proteins (see above).

We chose to use full Bayesian formalism to make inference so as to take into account all possible uncertainties in our model. Inferences were made based on Gibbs sampling,²¹ which was performed in the WinBugs environment.²²

3. Method and Results

3.1. Deriving a list of putative miRNA targets by sequence data alone

We described the sources of the data in the above section. As described in the Introduction, our current methods take as input a set of putative predictions from

a sequence-based prediction algorithm. We decided to run our procedures twice using two different prediction algorithms: TargetScan²³ and miRanda.⁵ The general results and conclusions are unchanged. Based on the intersection among the four types of datasets (predicted miRNA targets, miRNA expression, protein abundance and mRNA expression), we retained 21,721 putative interactions for TargetScan predictions (75 miRNAs, 1,404 cross-mapped mRNAs) and 17,339 putative interactions for miRanda predictions (70 miRNAs, 1408 cross-mapped mRNAs).

After compiling the datasets, we investigated mRNA or protein expression profiles in six organs (brain, heart, liver, lung, placenta and kidney), among which expression in four organs (brain, heart, liver and lung) were used for model construction and making predictions, while the remaining two organ types (placenta and kidney) were used for blind tests.

3.2. Modeling protein abundance

Instead of looking for negative correlation between miRNAs and mRNA transcripts as previously described in Ref. 10, our method relies on the correlation between miRNA and proteins (see Fig. 1). We start with a set of miRNA and target genes as predicted from a sequence-based approach, before modeling the protein abundance of the putative targets and the miRNA in individual tissues and organs. If we observe a negative correlation between the miRNA and the putative target in one particular tissue or organ, then the algorithm will assign a higher confidence score to this miRNA–protein pair. Conversely, a positive correlation between miRNA expression and protein abundance, especially in the cases where a high miRNA expression coincides with high protein abundance, will result in a low confidence score for the miRNA–protein pair.

We chose to use a probabilistic framework to model the relationship between miRNA and the protein abundance. The first challenge in this approach is to find an appropriate background distribution to model the protein abundance data. Unlike mRNA expression profiles, which can be effectively modeled using a Gaussian distribution, the peptide counts are discrete values. A possible choice is to use Poisson to model the count events; however, a simple Poisson model is not suitable for modeling the peptide counts in this study since there are excessive zeros in the dataset and the non-zero counts are also over-dispersed (variance is much greater than the mean). After comparing with other possible models such as zero-inflated Poisson and transformed Gaussian, we chose to use negative binomial model (NB) to characterize the peptide counts, with which the Poisson mean and over-dispersion can be considered simultaneously with lower model complexity. Recent research also suggested that NB is an optimal choice to model the abundance data with excessive zeros.²⁴

The protein abundance data have discrete integer values corresponding to protein counts and a lot of zeros corresponding to no protein abundance. The sample variance is very different from the sample mean, which cannot be effectively modeled

by a Poisson distribution but can be fitted very well by a negative binomial distribution empirically. A negative binomial distribution with a positive real parameter r and a real parameter γ ($0 < \gamma < 1$) is described in the following equation:

$$\begin{aligned}
 p(k|r, \gamma) &= \binom{k+r-1}{r-1} \gamma^r (1-\gamma)^k \\
 &= \frac{\Gamma(k+r)}{k! \Gamma(r)} \gamma^r (1-\gamma)^k.
 \end{aligned}
 \tag{1}$$

In the above equation, k is an integer, and the mean of the distribution is $r \frac{1-\gamma}{\gamma}$. If we re-parameterize the negative binomial distribution using the mean parameter $\lambda = r \frac{1-\gamma}{\gamma}$ and the positive real parameter r , we have the following equation:

$$\begin{aligned}
 p(k|r, \gamma) &= NB(k|\lambda, r) \\
 &= \frac{\lambda^k}{k!} \frac{\Gamma(r+k)}{\Gamma(r) \Gamma(r+\lambda)^k} \frac{1}{1 + \frac{\lambda}{r}}.
 \end{aligned}
 \tag{2}$$

In the above equation, $\Gamma(\cdot)$ is the Gamma function, r controls the over-dispersion of the distribution, and, when the over-dispersion parameter r approaches infinity, $NB(k|\lambda, r)$ approaches a Poisson distribution with mean parameter λ . The above NB model uses r to adjust the variance independently of the mean parameter λ of the distribution, differing from a Poisson distribution which has equal mean and variance. We model protein abundance data by NB using the parameterization in Eq. (2). We assume the abundance of each protein i in tissue type t , W_{it} , follows NB distribution, with two parameters θ_{it} and r_t , $1 \leq i \leq N$ and $1 \leq t \leq T$, where N and T are the total number of genes (proteins) and total number of tissue types, respectively. Thus, the probability of protein i with peptide count k in tissue t can be modeled as the following in Eq. (3).

$$p(W_{it} = k|\theta_{it}, r_t) = NB(k|\theta_{it}, r_t).
 \tag{3}$$

In Eq. (3), θ_{it} represents the Poisson mean for protein i in tissue t and r_t represents the over-dispersion of the data, which was shared by all the proteins in the same tissue t . We then used hierarchical Bayesian negative binomial regression to regress the Poisson mean θ_{it} with miRNA expression in corresponding tissues, M_{jt} , $1 \leq j \leq J$, and $1 \leq t \leq T$, where J is the total number of miRNAs in the dataset. Eq. (4) gives the regression of the Poisson mean in the model. Thus,

$$\ln(\theta_{it}) = \ln(\tau_t) - \rho_t \sum_{j=1}^J \omega_j \delta_{ij} M_{jt}.
 \tag{4}$$

In Eq. (4), τ_t stands for the background protein abundance shared by all the proteins in the same tissue t . As suggested in Ref. 10, we introduced δ_{ij} as a binary latent variable indicating whether or not the miRNA j regulates the gene i ; ω_j is a regression coefficient associated with j th miRNA expression shared by all the

tissue types, and ρ_t is a scaling parameter for tissue t accounting for the measurement difference in different tissues. Since sequence complementarity is a necessary condition for true targets, we use a binary variable S_{ij} as the putative prediction between miRNA j and protein i , which was derived from sequence-based predictions (TargetScanS, miRanda, or PicTar, etc); $S_{ij} = 1$ means that there is a putative prediction between i and j . The probability of a putative prediction being a true positive, p , is formally given in Eq. (5).

$$\begin{aligned} p(\delta_{ij} = 1 | S_{ij} = 1) &= p, \\ p(\delta_{ij} = 1 | S_{ij} = 0) &= 0. \end{aligned} \tag{5}$$

To avoid over-fitting the data and to account for all possible uncertainties, we chose to use full Bayesian approach to infer δ_{ij} so that all the uncertainties and nuisance variables can be integrated out. Thus we assigned the priors to other parameters as follows (most were assigned flat priors):

$$\begin{aligned} p &\sim \text{beta}(1, 1), \\ \tau_t &\sim \text{uniform}(0, 50), \\ \rho_t &\sim \text{gamma}(\alpha, \alpha), \\ \alpha &\sim \text{uniform}(0, +\infty), \\ \omega_j &\sim \text{exponential}(\beta), \\ \beta &\sim \text{uniform}(0, 1000), \\ r_t &\sim \text{exponential}(a), \\ a &\sim \text{uniform}(0, 1000). \end{aligned} \tag{6}$$

With the likelihood and priors defined above, we then implemented Gibbs sampling²¹ to compute marginal distribution of δ_{ij} conditioned on all evidence. All the inferences were made on drawing 5,000 samples after 10,000 iterations.

3.3. Apply Bayesian model to predict miRNA targets

We applied the model described above to the 21,712 putative miRNA–protein interactions derived from TargetScan, and assigned a confidence score to each putative interaction. Then we ranked the 21,712 putative interactions from the highest to the lowest confidence, and grouped them into 44 bins with each bin containing 500 ranked interactions. The results are shown in Figs. 2(a)–2(d) for four different organ types.

As shown in Fig. 2, our model can well capture the miRNA’s repression effects in these four organs. The miRNA–protein pairs that are predicted to have the highest confidence scores have lower protein abundance and higher miRNA expression; conversely, the miRNA–protein pairs with the lowest confidence scores also have higher protein abundance and higher miRNA expression. For the interactions ranked with intermediate confidence scores, the miRNA expression is low, and the

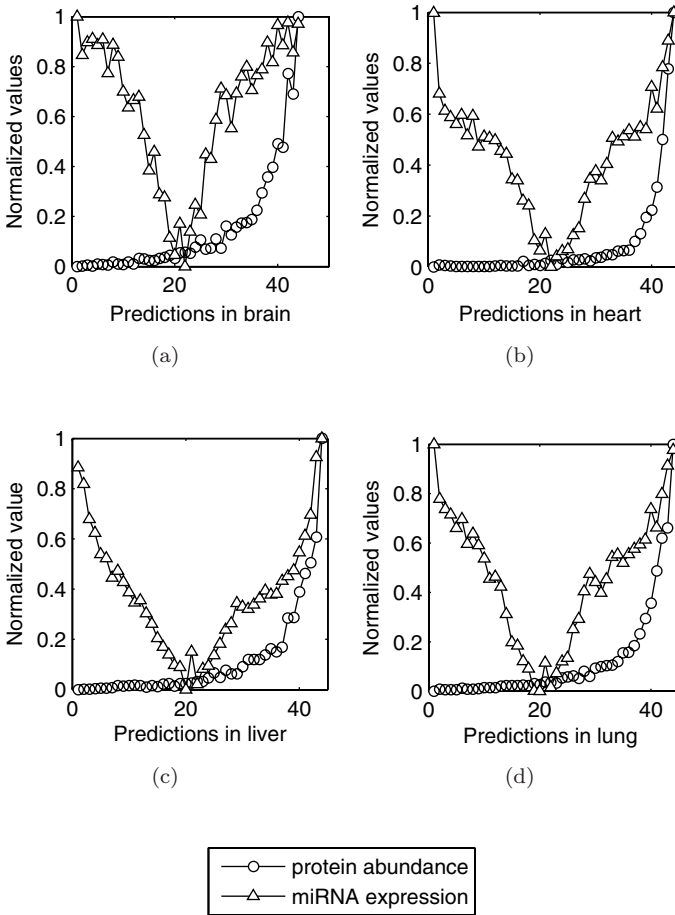


Fig. 2. miRNA targets prediction using miRNA expression and protein abundance. With our model, in the four organs (panel a–d), the most confident predictions (on the left) have the lowest protein abundance and the highest miRNA expression; while the least confident predictions (on the right) are high in both protein abundance and miRNA expression. All the data were scaled between 0 and 1. The putative predictions were from Ref. 20 using TargetScan.

protein abundance can be either low or high. The fact that a large number of TargetScan predictions are located in the right side of the curve, i.e. low confidence score with high miRNA expression and high protein abundance, indicates the extent of possible false-positives in the predictions made from sequence data alone. Because the Bayesian approach is intrinsically evidence-based, a prediction can only be made with high confidence if the miRNA is highly expressed in a certain tissue or organ.

Note that the high-confidence miRNA–protein interaction pairs as shown in Fig. 2 are predictions pooled from all four organs. We do not explicitly model the

tissue specificity of miRNAs in our formulism [see Eq. (3)]; instead, the strengths of the miRNA regulation in specific tissues are inferred from the expression level of miRNAs. For instance, a miRNA can be interpreted as a functional regulator in a given tissue only if it is highly expressed and it has high confidence score with a potential target protein that is lowly expressed in that tissue.

3.4. *Blind tests for the Bayesian predictions method*

As described above, we only used the protein abundance and miRNA expression in brain, lung, heart and liver to train our model and make predictions; the data in the remaining two organs (placenta and kidney) were left out during the model construction stage. To further validate our method, we subsequently conducted a blind test on the placenta and kidney datasets.

Figure 3(a) shows the results of the blind test in placenta. On the x -axis, we sorted the miRNA–protein pairs according to the confidence scores predicted by using the four training organs; on the y -axis, we plotted the protein abundance and miRNA expression level that are observed in placenta. The results indicate that, as a general trend, the predicted interactions can also reflect the desired tendency in placenta. The predicted interactions with high confidence usually have low protein abundance and high miRNA expression. The least confident predictions also have highly expressed proteins and highly expressed miRNAs, indicating those proteins are unlikely to be repressed by the miRNAs in placenta.

In kidney as the second blind test, shown in Fig. 3(b), although the miRNA expression data were not available for this organ type, clearly our predictions were also effective and the most confident predictions have the lowest protein abundance and *vice versa*. All the above analyses were based on the sequence-based predictions from TargetScan. The same results also hold true after we repeated the analysis using predictions from another program miRanda.⁵ Next, to test the robustness of our method, we shuffled the gene labels to randomize the data. The results from the shuffled data appear clearly random [Figs. 2(c)–2(d)], which strongly suggest that our prediction did not occur by chance.

3.5. *Comparison with TarBase and other methods*

We further searched for published experimental evidence for our predicted interactions in TarBase,⁴ which is a comprehensive database containing experimentally verified miRNA targets in a number of organisms. However, to this date, there are only 41 experimentally verified miRNA targets for mouse in the database. Since in the database all targets used gene symbol, we then converted the Swiss-Prot protein names in our study to corresponding gene symbols via <http://idconverter.bioinfo.cnio.es/>.²⁵ However, except for the gene Arid3a (ARI3A_MOUSE), all other genes were not included in our dataset as they do not have protein abundance data compiled in this study. For Arida3a, in TarBase, it was annotated to be regulated by miR-125b. From our predictions based on miRanda

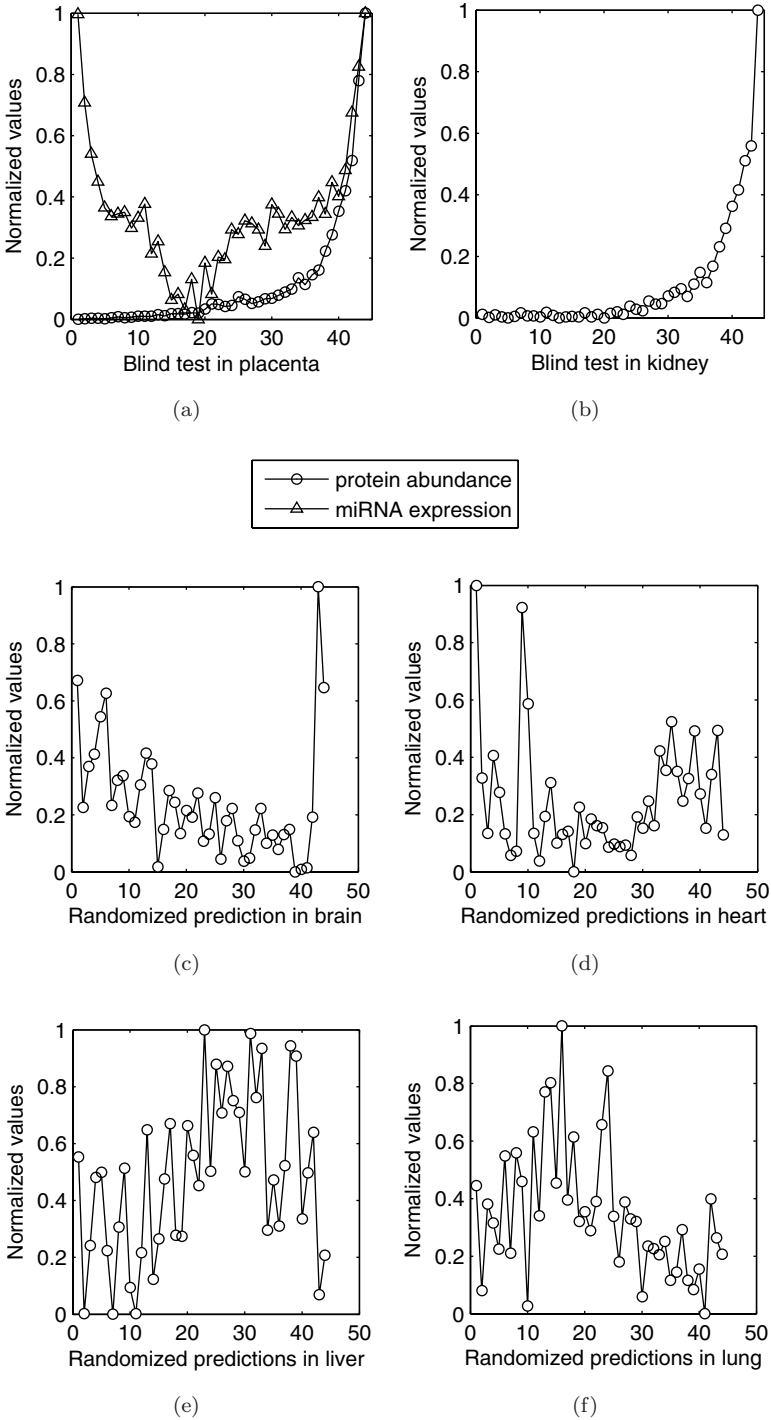


Fig. 3. Blind test of our predictions on (a) placenta and (b) kidney. (c)–(f) Result from randomization while protein labels are shuffled.

predictions, the interactions between miR-125b and ARI3A_MOUSE were ranked among top 5% in all the 17,339 putative interactions, suggesting ARI3A_MOUSE is likely to be a true target. However, this interaction was not detected by TargetScan as compiled in Ref. 20.

We also compared our prediction results to the results obtained using the method in Ref. 10. Although most of the predictions by both models are consistent, we found that proteomics data can further remove a lot of false-negative predictions,^a and we believe that modeling proteomics data is the most reliable way of filtering miRNA target predictions when large-scale proteomics data become available. In details, the miRNA/target interactions such as mmu-mir-214/Q8R399_MOUSE, mmu-mir-211/Q8BYX4_MOUSE, mmu-miR-292-5p/KCNN3_MOUSE, and mmu-miR-298/RRAS2_MOUSE all ranked among top 10% in all the putative interactions in both models. However, the miRNA/target interactions such as mmu-miR-298/PLF3_MOUSE, mmu-miR-210/Q8BSZ8_MOUSE, and mmu-miR-92/8BZZ4_MOUSE all ranked among top 1% in all the putative interactions in our model, but they all ranked among bottom 15% in all the putative interactions in the model by Ref. 10. We found that these miRNA/target pairs all have very good relatively high miRNA expression versus relatively low protein abundance patterns, but they do not have very clear relatively high miRNA expression versus relatively low mRNA expression patterns. Since miRNAs can either degrade mRNAs or repress mRNA translation, which will be discussed later in this paper, we believe that these interactions are very likely to be false-negatives predicted by the model in Ref. 10.

3.6. *Discordant correlation between mRNA expression and protein abundance*

We next investigated whether these target genes are regulated by translational repression or by mRNA degradation. This can be achieved by comparing the protein abundance and mRNA expression data of predicted target genes. If miRNAs predominantly regulate their targets by degradation, then for target genes we would expect to see a good correlation between mRNA expression and the protein abundance (low mRNA concentration leads to low protein abundance). For all the putative targets, Fig. 4 plots their average mRNA expression level and their total protein abundance across four organs (brain, heart, liver and lung); the x -axis is the predicted targets sorted with the highest confidence on the left and lowest confidence on the right.

In Fig. 4, interestingly, among the top ranked predicted targets, their mRNA expression fluctuates greatly, while among the bottom ranked predicted targets their mRNA expression is well correlated with their protein abundance. Such a lack of

^aOur Bayesian model filters sequence-based predictions and removes a lot of false positives predicted by sequence models.

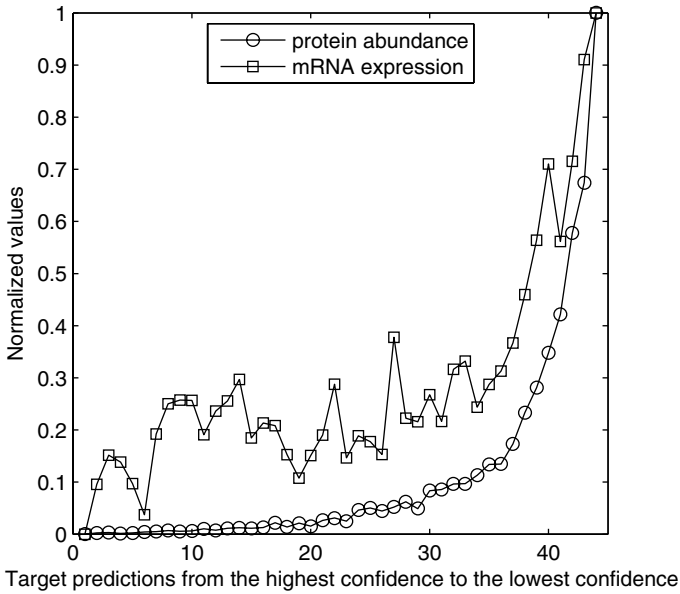


Fig. 4. mRNA expression and protein abundance for the gene targets of predicted interactions from the highest confidence (left) to the lowest confidence (right). The mRNA expression was averaged over the five organ types (brain, heart, kidney, liver and lung) and protein abundance is the total peptide counts across five organs (brain, heart, kidney, liver and lung). All the experimental data were scaled between 0 and 1.

correlation between protein abundance and gene expressions suggested that degradation is unlikely to be the dominant mechanism for miRNA-mediated regulation, which is consistent with previous observations.^{20,26} Such pronounced discordance between mRNA expression and protein abundance was also reported in a previous study.¹⁷ In the next step, we describe another Bayesian model that allows us to trace back to the mRNA expression data, and calculate the probability of miRNA regulation by mRNA degradation or by transcriptional inhibition.

3.7. Two possible mechanisms: mRNA degradation versus translational repression

For the top miRNA–protein pairs that are predicted to be true regulators and targets, we can distinguish between these two possible regulatory mechanisms by analyzing the correlation between the miRNA expression and the mRNA expression. For example, if a top-ranked miRNA Cprotein pair has high miRNA expression and high mRNA expression, then it is a strong indicator that the protein target is regulated by translational expression. In contrast, if a predicted miRNA target has low mRNA expression, then it is likely to be regulated by mRNA degradation.

There are two common concerns in modeling the mRNA expression data: (1) the intrinsic low signal-to-noise ratio of microarray data, and (2) the potential problem

of missing values since a large number of the genes have expression levels measured as 0.¹⁹ To overcome these difficulties, we discretize the mRNA expression data by using a cutoff of 0.1 to binarize the expression level to either low or high. For a given mRNA i in tissue t , $1 \leq i \leq L$ and $1 \leq t \leq T$, where L is the total number of mRNAs in the confident predictions derived from the first model, its mRNA expression R_{it} can be either low ($R_{it} = 0$) or high ($R_{it} = 1$). Let the probability of degradation for mRNA i in tissue t be q_{it} , we assume,

$$P(R_{it} = k) = q_{it}^{(1-k)}(1 - q_{it})^k, \quad k = 0 \text{ or } 1. \quad (7)$$

We next used logistic regression to regress q_{it} with the expression of miRNAs that regulates gene i , in tissue t . Then we have the following equation,

$$\log it(q_{it}) = \log \left(\frac{q_{it}}{1 - q_{it}} \right) = \sum_{j=1}^H \Phi_j b_{ij} M_{jt}, \quad (8)$$

in which H is the total number of miRNAs in the miRNA–mRNA interactions, M_{jt} is the expression of the j th miRNA in tissue t , b_{ij} is a binary latent variable indicating whether or not the gene i is degraded by miRNA j , and Φ_j is a scaling parameter associated with the j th miRNA, shared by all tissue types. The rationale behind Eq. (8) is that for a given gene i , if its expression is low in tissue t , i.e. $R_{it} = 0$, then from the perspective of maximum likelihood, we need to maximize q_{it} so that the interactions between gene i and its regulating miRNAs that are highly expressed in tissue t should be assigned a higher degradation score. In this sense, the observed low expression of mRNA and high expression of miRNA together lead to the assignment of a high degradation probability. Similarly, if in tissue t , $R_{it} = 1$, then q_{it} needs to be minimized, implying those highly expressed miRNAs should be associated with a low degradation score, so the highly expressed miRNAs and mRNAs indicate that such regulation is more likely to be through translational repression than through degradation.

Regarding the latent variable b_{ij} , we further required that:

$$\begin{aligned} p(b_{ij} = 1 | \delta_{ij} = 1) &= h, \\ p(b_{ij} = 1 | \delta_{ij} = 0) &= 0, \end{aligned} \quad (9)$$

in which δ_{ij} indicates whether or not mRNA i is targeted by miRNA j . If $\delta_{ij} = 1$, then the miRNA j has a probability h to cause degradation to its target mRNA i . We then used a full Bayesian approach to estimate the parameters in the model to avoid overfitting the data and to account for all potential uncertainties. In the Bayesian framework, we then assigned priors to other parameters in the model as follows:

$$\begin{aligned} \Phi_j &\sim \text{exponential}(\Psi), \\ \Psi &\sim \text{uniform}(0, +\infty), \\ h &\sim \text{beta}(1, 1). \end{aligned} \quad (10)$$

Having defined the likelihood and the priors, we then inferred the posterior marginal distribution of $p(b_{ij} = 1 | \mathbf{S}, W, M)$, conditioned on all the evidence. By implementing Gibbs sampling in the environment of WinBugs,²² all the inferences were based on drawing 5,000 samples after 10,000 iterations.

3.8. Apply the Bayesian model to mRNA data

By implementing the model described above, we calculated the confidence scores for mRNA degradation for each miRNA–mRNA interaction pair, which indicated the likelihood that miRNA causes degradation to their mRNA targets. The lower degradation score implies higher probability of being translationally repressed. Then, we ranked the scores from the highest to the lowest, and grouped them into 50 bins, each containing 100 ranked interactions. Figure 5 shows the mRNA expression level of the ranked miRNA targets across four organ types. The miRNA targets near the top of Fig. 5(a) have the highest probability of being regulated by mRNA degradation, as demonstrated by their low mRNA expression level [details shown in Fig. 5(b)]. Conversely, the targets near the bottom have the highest probability of being regulated by translational repression [details shown in Fig. 5(c)].

4. Discussion

4.1. miRNA regulation by translational repression

In this paper we described two novel formulisms in the computational analysis of miRNA regulation. We first introduced a Bayesian approach to identify miRNA targets based on protein abundance data. After having selected high confidence predictions, we then introduced a second Bayesian model to further distinguish the two possible regulatory mechanisms, i.e. mRNA degradation versus translational repression. We showed that our model is very effective in describing the three intertwining genomics datasets, i.e. miRNA expression, mRNA expression, and protein abundance. Our results demonstrated that protein abundance is a very useful resource in predicting miRNA targets, especially with the emerging evidence that translational repression is more prevalent than mRNA degradation as a regulatory mechanism in mammals. Our work also suggested that such repression mechanism probably contributed to the previously observed discordance between mRNA expression level and protein abundance. We would like to point out that although in this paper our model takes as input the predictions from TargetScan and miRanda, essentially results from any other sequence-based predictions can be used in our formulism.

4.2. Potential limitations and future directions

Even though our framework has obtained encouraging results, it certainly has limitations. We envision that it can be improved in the following areas: (1) As we

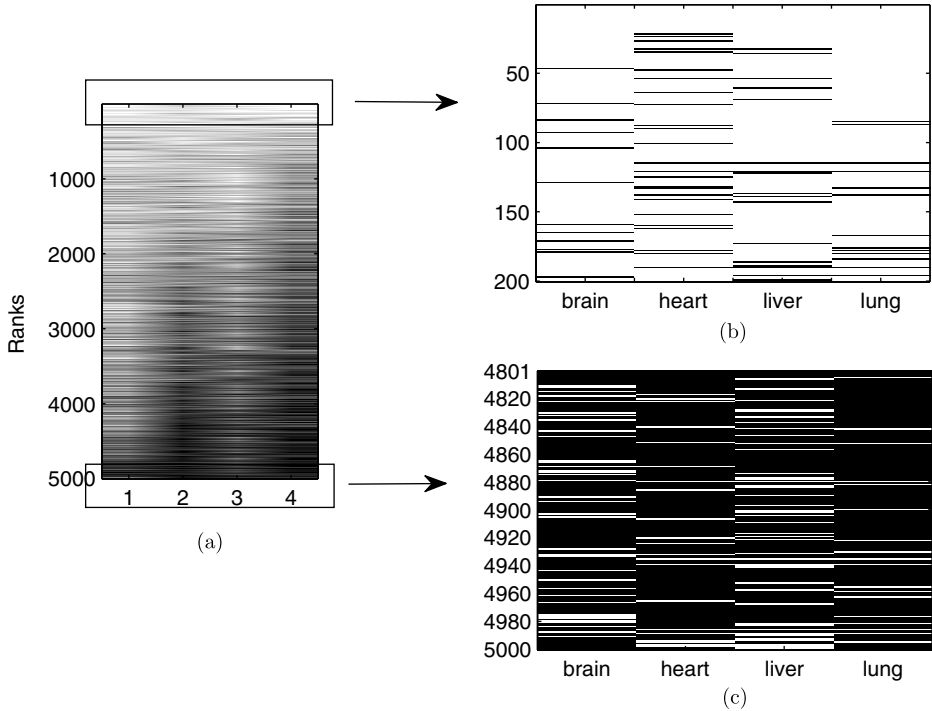


Fig. 5. (a) Ranking of miRNA targets according to the probability of being regulated by the mechanism of mRNA degradation; the targets were ranked from the highest degradation probability (top) to the lowest degradation probability (bottom). It also showed mRNA expression of the targets across four organ types. Black color denotes high expression and white color denotes low expression. (b) The top ranked interactions have the highest degradation probability, and are associated with the low mRNA expression. (c) The bottom ranked interactions have the lowest degradation probability, and thus are associated with high mRNA expression.

noted, miRNA is not the only mechanism of gene regulation. Some of the observed variations in protein abundance across organs are likely the result of regulation at the transcription level by transcription factors, or at the post-transcriptional level by mRNA degradation pathways. (2) Although it has been reported that the mechanism of translational repression by miRNAs has little impact on mRNA level, the mRNA expression might still be helpful in predicting miRNA targets, and recent research suggested that targeted mRNA showed strong correlation (positive and negative) with miRNAs.^{13,15} In the future, we could incorporate the mRNA expression data with the proteomic data to build an integrated predictive model. (3) At this stage, our model takes as input the sequence-based predictions from other prediction programs such as TargetScan or miRANDA, therefore our algorithm does not explicitly consider the sequence complementarity and evolutionary conservation. As a future work, it would be interesting to extend our model to incorporate these properties into a unified probabilistic framework. (4) In our model, similar to

Ref. 10, we assumed a single baseline distribution of protein abundance for all the genes in each tissue type. However, this is a significant simplification since different genes could have distinct baseline expression levels. The next step in this work is to take this into account and develop a more realistic expression baseline model. For example, it would be possible to consider the codon usage of the genes to infer the possible baseline expression of a given gene.²⁷

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