

iRNA(m⁶A)-PseDNC: Identifying N⁶-methyladenosine sites using pseudo dinucleotide composition

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ABSTRACT

As a prevalent post-transcriptional modification, N⁶-methyladenosine (m⁶A) plays key roles in a series of biological processes. Although experimental technologies have been developed and applied to identify m⁶A sites, they are still cost-ineffective for transcriptome-wide detections of m⁶A. As good complements to the experimental techniques, some computational methods have been proposed to identify m⁶A sites. However, their performance remains unsatisfactory. In this study, we firstly proposed an Euclidean distance based method to construct a high quality benchmark dataset. By encoding the RNA sequences using pseudo nucleotide composition, a new predictor called iRNA(m⁶A)-PseDNC was developed to identify m⁶A sites in the *Saccharomyces cerevisiae* genome. It has been demonstrated by the 10-fold cross validation test that the performance of iRNA(m⁶A)-PseDNC is superior to the existing methods. Meanwhile, for the convenience of most experimental scientists, established at the site [http://lin-group.cn/server/iRNA\(m⁶A\)-PseDNC.php](http://lin-group.cn/server/iRNA(m6A)-PseDNC.php) is its web-server, by which users can easily get their desired results without need to go through the detailed mathematics. It is anticipated that iRNA(m⁶A)-PseDNC will become a useful high throughput tool for identifying m⁶A sites in the *S. cerevisiae* genome.

1. Introduction

Among the ~150 kinds of chemical modifications identified in cellular RNAs, N⁶-methyladenosine (m⁶A) is the most prevalent one in mRNA and noncoding RNA [1]. Since it was first detected in 1970s [2], m⁶A has been observed in a wide range of eukaryotes. As indicated by recent evidences, m⁶A plays fundamental regulatory roles in a series of biological processes, such as affecting mRNA splicing and stability, translation, stem cell pluripotency, as well as immune response [3–7]. Therefore, the transcriptome-wide annotation of m⁶A site will be helpful to understanding its biological functions.

In 2012, the high-throughput sequencing technique termed MeRIP-Seq and m⁶A-seq [8,9], were proposed to detect transcriptome wide

m⁶A sites in *S. cerevisiae*, *Mus musculus*, and *Homo sapiens*. Since this technique relies solely on immunoprecipitation of fragmented RNA, the resolution of MeRIP-Seq and m⁶A-seq is not satisfactory [5]. In 2015, Linder et al. proposed the miCLIP technique [10], which provides the single-nucleotide-resolution profile of m⁶A sites in human transcriptome. Although these experimental techniques promote the research progresses on m⁶A modifications, they are still costly and time consuming in performing transcriptome wide analysis.

During the last several years, many computational methods have been developed for identifying m⁶A sites in the *S. cerevisiae* genome. Based on the m⁶A-seq data, Schwartz et al. proposed the first computational model for identifying m⁶A site in the *S. cerevisiae* genome [11]. Inspired by their work, Chen et al. developed two bioinformatics tools

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termed iRNA-Methyl and m⁶Apred to identify m⁶A sites, respectively [12,13]. Later on, by encoding RNA sequences using both sequence information and the RNA secondary structures, Zhou et al. developed a random forest based method to predict m⁶A sites in *S. cerevisiae* [14]. In order to improve the accuracy of computationally identifying m⁶A site in the *S. cerevisiae* genome, Chen et al. developed RAM-ESVM method that was built by using ensemble classifiers [15]. More recently, by encoding RNA sequences using multi-interval nucleotide pair position specificity, Xing et al. proposed a sequence-based predictor called RAM-NPPS for identifying m⁶A sites [16].

Although these computational methods yielded encouraging results for computationally identifying m⁶A sites in the *S. cerevisiae* [17], further improvement is needed. Particularly, most of those methods were trained by the dataset constructed by Chen et al. [12]. In that dataset, the negative samples were randomly selected from a huge amount of candidates, and hence unavoidably had some arbitrariness. Accordingly, their reliability might be questioned [18]. The present study was initiated in an attempt to develop a new and more powerful method to identify m⁶A sites by refining the benchmark dataset.

As done in a series of recent reports [12,19–36], here we are to propose the new predictor also according to the Chou's 5-sep rules [37] because doing so will make the entire process more logic and transparent.

2. Materials and methods

2.1. Benchmark dataset

The first step in the 5-step rules [37] is how to construct or select a valid benchmark dataset to train and test the predictor. By following the same procedures as reported in Ref. [12], we first obtained 1307 positive samples and 33,280 negative samples. It was observed via preliminary trials that the optimal sequence length is 51 nt. But instead of using the random selection method to reduce the number of negative samples as reported in Ref. [12], in this study we adopted the subset-balancing treatment according to the Euclidean distance [38], as elaborated below.

First of all, for the reason that will be discussed later, we calculated the average value of each of the 22 features [12] over the 33,280 negative samples; i.e.,

$$\bar{d}_i = \frac{1}{33280} \sum_{j=1}^{33280} d_i^j \quad (i = 1, 2, \dots, 22) \quad (1)$$

where d_i^j is the value of the i -th feature for the j -th negative sample. Thus, the center of the negative samples can be defined as

$$\bar{\mathbf{R}} = [\bar{d}_1 \ \bar{d}_2 \ \dots \ \bar{d}_{16} \ \bar{d}_{17} \ \dots \ \bar{d}_{22}]^T \quad (2)$$

Subsequently, we calculated the Euclidean distance (D^j) between the j -th negative sample and the center $\bar{\mathbf{R}}$; i.e.,

$$D^j = \sqrt{\sum_{i=1}^{22} (d_i^j - \bar{d}_i)^2} \quad (j = 1, 2, \dots, 33280) \quad (3)$$

According to their Euclidean distance values, the 33,280 negative samples were sorted in an ascending order, and the top ranked 1307 negative samples (i.e., they were closest to the center) were picked up to form the refined negative subset.

For reader's convenience, the detailed sequences in the original 1307 positive samples and the refined 1307 negative samples are given in Supporting Information S1 and S2, which can be directly downloaded via the link at [http://lin-group.cn/server/iRNA\(m6A\)-PseDNC.php](http://lin-group.cn/server/iRNA(m6A)-PseDNC.php).

2.2. Sample formulation

The second step in the 5-step rules [37] is how to formulate the biological sequence samples with a discrete model or a vector, yet still keep considerable sequence-order information or key pattern characteristic. This is because all the existing machine-learning algorithms can only handle vector but not sequence samples, as elucidated in a comprehensive review [39]. However, a vector defined in a discrete model may completely lose all the sequence-pattern information. To avoid completely losing the sequence-pattern information for proteins, the pseudo amino acid composition [40] or PseAAC [41] was proposed. Ever since then, it has been widely used in nearly all the areas of computational proteomics (see, e.g. [42–83], as well as a long list of references cited in Ref. [84]). Because it has been widely and increasingly used, recently three powerful open access soft-wares, called 'PseAAC-Builder' [85], 'propy' [86], and 'PseAAC-General' [87], were established: the former two are for generating various modes of special PseAAC [88]; while the 3rd one for those of general PseAAC [37], including not only all the special modes of feature vectors for proteins but also the higher level feature vectors such as "Functional Domain" mode, "Gene Ontology" mode, and "Sequential Evolution" or "PSSM" mode. Encouraged by the successes of using PseAAC to deal with protein/peptide sequences, its idea and approach were extended to PseKNC (Pseudo K-tuple Nucleotide Composition) to generate various feature vectors for DNA/RNA sequences [89] that have proved very successful as well [25,90–95].

As shown in Supporting Information S1 or S2, each of the RNA samples in this study has the form of

$$\mathbf{R} = N_1 N_2 N_3 \dots N_i \dots N_{51} \quad (4)$$

where N_1 represents the 1st nucleotide, N_2 the 2nd nucleotide, and so forth. They can be any of the four nucleotides A, C, G or U. According to PseKNC [89], the RNA sequence can be formulated as a discrete vector

$$\mathbf{R} = [d_1 \ d_2 \ \dots \ d_{16} \ d_{16+\lambda} \ \dots \ d_{16+\lambda}]^T \quad (5)$$

where

$$d_u = \begin{cases} \frac{f_u}{\sum_{i=1}^{16} f_i + w \sum_{j=1}^{\lambda} \theta_j} & (1 \leq u \leq 16) \\ \frac{w \theta_{u-16}}{\sum_{i=1}^{16} f_i + w \sum_{j=1}^{\lambda} \theta_j} & (16 < u \leq 16+\lambda) \end{cases} \quad (6)$$

In Eq. (6), f_u ($u = 1, 2, \dots, 16$) is the normalized occurrence frequency of the u -th non-overlapping dinucleotides in the RNA sequence. λ can be viewed as the number of the total pseudo components used to reflect the long-range or global sequence effect, and w is the weight factor. θ_j is the j -th tier correlation factor that reflects the sequence order correlation between all the j -th most contiguous dinucleotide along a RNA sequence as formulated by

$$\theta_j = \frac{1}{L-j-1} \sum_{i=1}^{L-j-1} C_{i,i+j} \quad (j = 1, 2, \dots, \lambda; \lambda < L) \quad (7)$$

where $C_{i,i+j}$ is the correlation function and is defined by

$$C_{i,i+j} = \frac{1}{\mu} \sum_{g=1}^{\mu} [P_g(D_i) - P_g(D_{i+j})]^2 \quad (8)$$

where $\mu = 3$ is the number of RNA physicochemical properties considered, $P_g(D_i)$ is the numerical value of the g -th ($g = 1, 2, 3, \dots, \mu$) RNA local structural property for the dinucleotide $R_i R_{i+1}$ at position i and $P_g(D_{i+j})$ the corresponding value for the dinucleotide $R_{i+j} R_{i+j+1}$ at position $i+j$.

Listed in Table 1 are the three physicochemical properties obtained by standard conversion for the 16 different dinucleotides in RNA. The concrete procedures for how to convert the original 16 physicochemical properties to their standard ones have been elaborated in Ref. [12], and hence there is no need to repeat here.

Table 1

The three physicochemical properties after standard conversion for the 16 different dinucleotides in RNA.

Dinucleotide	Enthalpy (Ka/mol)	Entropy (eU)	Free energy (Ka/mol)
GG	−1.08	−0.88	−1.45
GA	−1.50	−1.82	−0.28
GC	−1.85	−1.72	−1.66
GU	−0.30	−0.32	−0.14
AG	0.71	0.82	0.07
AA	1.10	0.95	1.56
AC	−0.30	−0.32	−0.14
AU	1.44	1.42	1.34
CG	0.55	0.79	−0.29
CA	−0.42	−0.57	0.03
CC	−1.08	−0.88	−1.45
CU	0.71	0.82	0.07
UG	0.71	0.82	0.03
UA	0.51	0.27	1.04
UC	−0.30	−0.32	−0.28
UU	1.10	0.95	1.56

In order to avoid over-fitting or the “high-dimension disaster” problem [96], the search of the two parameters in Eq. (6), namely λ and w , was in the following ranges [3,6] and [0, 1] with the steps of 1 and 0.1, respectively. It was found that the optimal values for λ and w are 6 and 0.9, respectively.

Accordingly, the RNA sequence sample can be formulated by a $(16 + 6)=22$ -D (dimensional) vector as given below

$$\mathbf{R} = [d_1 \ d_2 \ \cdots \ d_{16} \ d_{17} \ \cdots \ d_{22}]^T \quad (9)$$

where the first 16 components are used to incorporate the short-range or local sequence order information of the RNA sample, while the remaining 6 components used to incorporate its long-range or global sequence order information.

2.3. Operation engine: support vector machine

The third step in the 5-step rules [37] is how to introduce or develop a powerful algorithm (or engine) to operate the prediction. Support vector machine (SVM) is a powerful and popular method for pattern recognition and has been widely used in the realm of bioinformatics [31,97–103]. The basic idea of SVM is to transform the input data into a high dimensional feature space and then determine the optimal separating hyperplane. In the current study, the LibSVM package 3.18 (<http://www.csie.ntu.edu.tw/~cjlin/libsvm/>) was used to implement SVM. Owing to its effectiveness and speed in training process, the radial basis kernel function (RBF) was used to obtain the classification hyperplane in the current study. For the SVM operation engine, a grid search approach was employed to optimize the regularization parameter C and kernel parameter γ by using the 5-fold cross validation test in the following ranges $[2^{-5}, 2^{15}]$ and $[2^{-15}, 2^{-5}]$ with the steps of 2 and 2^{-1} , respectively.

The predictor thus obtained is called “iRNA(m6A)-PseDNC”, where “i” stands for “identify”, “RNA(m6A)” for “RNA N6-methyladenosine site”, and “PseDNC” for “via Pseudo Dinucleotide Composition”.

2.4. Cross-validation

The fourth step in the 5-step rules [37] is how to properly perform cross-validation tests to objectively evaluate the anticipated accuracy of the predictor. To address this problem, we need to consider the two issues. (1) What metrics should be used to quantitatively measure the predictor's quality? (2) What test method should be utilized to score the metrics?

2.4.1. A set of four intuitive metrics

To measure the quality of a predictor, four metrics [104] are often

used in literature; they are (1) overall accuracy or Acc, (2) Mathew's correlation coefficient or MCC, (3) sensitivity or Sn, and (4) specificity or Sp. But their conventional formulations directly copied from math books are difficult to understand for most experimental scientists, particularly the one for MCC. Fortunately, by using the symbols introduced by Chou [105,106] in studying the signal peptide cleavage sites, a set of intuitive metrics were derived [107–109], as given below

$$\begin{cases} \text{Sn} = 1 - \frac{N^+}{N^+} & 0 \leq \text{Sn} \leq 1 \\ \text{Sp} = 1 - \frac{N^+}{N^-} & 0 \leq \text{Sp} \leq 1 \\ \text{Acc} = \Lambda = 1 - \frac{N^+ + N^+}{N^+ + N^-} & 0 \leq \text{Acc} \leq 1 \\ \text{MCC} = \frac{1 - \left(\frac{N^+}{N^+} + \frac{N^+}{N^-} \right)}{\sqrt{\left(1 + \frac{N^+ - N^+}{N^+} \right) \left(1 + \frac{N^+ - N^+}{N^-} \right)}} & -1 \leq \text{MCC} \leq 1 \end{cases} \quad (10)$$

where represents the total number of positive samples investigated, while N^+ is the number of positive samples incorrectly predicted to be of negative one; N^- the total number of negative samples investigated, while N^+ the number of the negative samples incorrectly predicted to be of positive one. Because of its merit in intuitiveness, the set of metrics (Eq. (10)) has been increasingly and widely used in computational biology (see, e.g., [12,20–33] [56,59] [91–95] [110–144]). It is instructive to point out, however, that both the original four metrics [104] in math books and the intuitive ones in Eq. (7) are valid only for the single-label systems (where each sample belongs to one and only one class). For the multi-label systems (where a sample may simultaneously belong to several classes), whose existence has become more frequent in system biology [145–151], system medicine [152,153] and biomedicine [154], a completely different set of metrics as defined in Ref. [155] is absolutely needed.”

2.4.2. Cross-validation

In statistical prediction, the following three cross-validation methods are often used to check the performance of a predictor: independent dataset test, subsampling (or K-fold cross-validation) test, and jackknife test [38]. Of the three test methods, the jackknife test is deemed the least arbitrary that can always yield a unique result for a given benchmark dataset as elaborated in Ref. [37] and demonstrated by Eqs.28–30 therein. However, to reduce the computational time, in this study we adopted the 10-fold cross validation test, as done by many investigators with SVM as the operation engine.

2.5. Web-server for iRNA(m6A)-PseDNC

The last, but not the least important, step of the 5-step rules is how to establish a user-friendly web-server for the predictor that is accessible to the public. As pointed out in Ref. [156] and demonstrated in a series of recent publications (see, e.g. [20,24,25,92,94,95,136] [138–142] [144–146] [148–152] [157–162]), user-friendly and publicly accessible web-servers represent the future direction for developing practically more useful prediction methods and computational tools. Actually, many practically useful web-servers have significantly increased the impacts of bioinformatics on medical science [39], driving medicinal chemistry into an unprecedented revolution [84]. For the convenience of the majority of the experimental scientists, the web-server for the iRNA(m6A)-PseDNC predictor has also been established at [http://lin-group.cn/server/iRNA\(m6A\)-PseDNC.php](http://lin-group.cn/server/iRNA(m6A)-PseDNC.php).

3. Results and discussion

The success rates achieved by the proposed predictor by 10-fold cross validation test on the benchmark datasets as described in Section 2.1 are given in Table 2, where for facilitating comparison, the corresponding results by the existing state-of-the-art method RAM-NPPS

Table 2

Comparison with the existing state-of-the-art method in identifying m6A sites for *S. cerevisiae* genome.

Methods	Sn(%) ^a	Sp(%) ^a	Acc(%) ^a	MCC ^a
RAM-NPPS ^b	74.29	69.93	72.11	0.44
iRNA(m6A)-PseDNC ^c	86.84	95.64	91.24	0.83

^a See Eq. (10) for the metrics definition.

^b See [16] for RAM-NPPS, which is the existing state-of-the-art method in identifying m6A sites for *S. cerevisiae* genome.

^c The predictor proposed in this paper with $C = 2^{15}$ and $\gamma = 2^{-7}$.

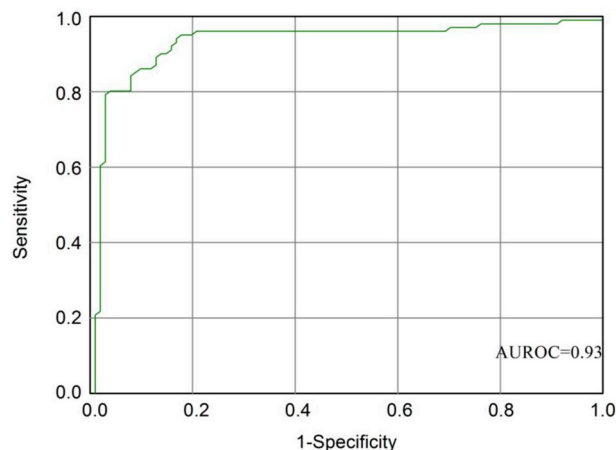


Fig. 1. A graphical illustration to show the performances of iRNA(m6A)-PseDNC in identifying m6A sites. See the text in Section 3 for further explanation.

[16] are also listed. As we can see from the table, for the existing best method in identifying m6A sites for the *S. cerevisiae* genome: its Sn (sensitivity) is over 12% lower than that of iRNA(m6A)-PseDNA, its Sp (specificity) is over 25% lower, its Acc (accuracy) is about 20% lower, and its MCC (stability) is 39% lower. These compelling results indicate that the iRNA(m6A)-PseDNA predictor is indeed very powerful.

Since graphic analysis is a very useful vehicle for studying complicated biological systems, as demonstrated by a series of previous studies (see, e.g., [163–171]). Shown in Fig. 1 is the graph of Receiver Operating Characteristic (ROC) widely used to reflect the quality of a predictor [172]. The area under the ROC curve, also called AUROC, is used to check the performance quality of the classifier: the value 0.5 of AUROC is equivalent to the outcome obtained by the random guess while 1 of AUROC represents the perfect prediction with 100% correctness. For the case of the current predictor, it is 0.93 clearly indicating its performance is really very good.

4. Conclusion

iRNA(m6A)-PseKNC is a powerful predictor for identifying the m⁶A sites in *S. cerevisiae* genome. It was developed by using the Euclidean distance to balance out the size of the negative training subset with that of the positive one. In order to consider both the local and global effects, each of the statistical samples in this predictor is formulated with a 22-D PseKNC vector. The learning machine implemented in the new predictor is SVM. It has been observed by the 10-fold cross-validation test that iRNA(m6A)-PseKNC is superior to RAM-NPPS, the state-of-the-art method in this regard. A public-accessible web-server for the new predictor has been established. We anticipate that it will become a very useful high throughput tool for genome analysis.

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Appendix A. Supplementary data

Supplementary data related to this article can be found at <https://doi.org/10.1016/j.ab.2018.09.002>.

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