pLoc_Deep-mAnimal: A Novel Deep CNN-BLSTM Network to Predict Subcellular Localization of Animal Proteins

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ABSTRACT

Current coronavirus pandemic has endangered mankind life. The reported cases are increasing exponentially. Information of animal protein subcellular localization can provide useful clues to develop antiviral drugs. To cope with such a catastrophe, a CNN based animal protein subcellular localization predictor called “pLoc_Deep-mAnimal” was developed. The predictor is particularly useful in dealing with the multi-sites systems in which some proteins may simultaneously occur in two or more different organelles that are the current focus of pharmaceutical industry. The global absolute true rate achieved by the new predictor is over 92% and its local accuracy is over 95%. Both have substantially exceeded the other existing state-of-the-art predictors. To maximize the convenience for most experimental scientists, a user-friendly web-server for the new predictor has been established at http://www.jci-bioinfo.cn/pLoc_Deep-mAnimal/, which will become a very useful tool for fighting pandemic coronavirus and save the mankind of this planet.

1. INTRODUCTION

Knowledge of the subcellular localization of proteins is crucially important for fulfilling the following two important goals: 1) revealing the intricate pathways that regulate biological processes at the cellular level [1, 2]. 2) selecting the right targets [3] for developing new drugs.

With the avalanche of protein sequences in the post-genomic age, we are challenged to develop computational tools for effectively identifying their subcellular localization purely based on the sequence information.

In 2019, a very powerful predictor, called “pLoc_bal-mAnimal [4], was developed for predicting the subcellular localization of animal proteins based on their sequences information alone. It has the following
remarkable advantages. 1) Most existing protein subcellular location prediction methods were developed based on the single-label system in which it was assumed that each constituent protein had one, and only one, subcellular location (see, e.g., [5-7] and a long list of references cited in a review paper [8]). With more experimental data uncovered, however, the localization of proteins in a cell is actually a multi-label system, where some proteins may simultaneously occur in two or more different location sites. This kind of multiplex proteins often bears some exceptional functions worthy of our special notice [2]. And the pLoc_bal-mAnimal predictor [4] can cover this kind of important information missed by most other methods since it was established based on the multi-label benchmark dataset and theory. 2) Although there are a few methods (see, e.g., [9, 10]) that can be used to deal with multi-label subcellular localization for proteins, the prediction quality achieved by pLoc_bal-mAnimal [4] is overwhelmingly higher, particularly in the absolute true rate.

The pLoc_bal-mAnimal predictor [4] has the aforementioned merits; it has not been trained at a deeper level yet [11-14].

The present study was initiated in an attempt to address this problem. As done in pLoc_bal-mAnimal [4] as well as many other recent publications in developing new prediction methods (see, e.g., [15, 16]), the guidelines of the 5-step rule [17] are followed. They are about the detailed procedures for 1) benchmark dataset, 2) sample formulation, 3) operation engine or algorithm, 4) cross-validation, and 5) web-server. But here our attentions are focused on the procedures that significantly differ from those in developing the predictor pLoc_bal-mAnimal [4].

2. MATERIALS AND METHODS

2.1. Benchmark Dataset

The benchmark dataset used in this study is exactly the same as that in pLoc_bal-mAnimal [4]; i.e.,

\[
S = S_1 \cup S_2 \cup \cdots \cup S_9 \cup \cdots \cup S_{20}
\]  

where \( S_1 \) only contains the protein samples from the “Acrosome” location, \( S_2 \) only contains those from the “Cell cortex” location, and so forth; \( \cup \) denotes the symbol for “union” in the set theory. For readers’ convenience, their detailed sequences and accession numbers (or ID codes) are given in Supporting Information S1 that is also available at http://www.jci-bioinfo.cn/pLoc_bal-mAnimal/Supp1.pdf, where none of proteins included has \( \geq 25\% \) sequence identity to any other in the same subset (subcellular location).

2.2. Proteins Sample Formulation

Now let us consider the 2nd step of the 5-step rule [17]; i.e., how to formulate the biological sequence samples with an effective mathematical expression that can truly reflect their essential correlation with the target concerned. Given a protein sequence \( P \), its most straightforward expression is

\[
P = R_1R_2R_3R_4R_5R_6R_7 \cdots R_L
\]  

where \( L \) denotes the protein’s length or the number of its constituent amino acid residues, \( R_1 \) is the 1st residue, \( R_2 \) the 2nd residue, \( R_3 \) the 3rd residue, and so forth. Since all the existing machine-learning algorithms can only handle vectors as elaborated in [3], one has to convert a protein sample from its sequential expression (Equation (2)) to a vector. But a vector defined in a discrete model might completely miss all the sequence-order or pattern information. To deal with this problem, the Pseudo Amino Acid Composition [18] or PseAAC [19] was proposed. Ever since then, the concept of “Pseudo Amino Acid Composition” has been widely used in nearly all the areas of computational proteomics with the aim to grasp various different sequence patterns that are essential to the targets investigated (see, e.g., [20-30] as well as a long list of references cited in [31]). Because it has been widely and increasingly used, recently three powerful open access soft-wares, called “PseAAC-Builder” [32], “propy” [33], and “PseAAC-General” [34], were established: the former two are for generating various modes of special PseAAC [35]; while the
According to the concept of general PseAAC [17], any protein sequence can be formulated as a PseAAC vector given by

$$\mathbf{P} = [\Psi_1, \Psi_2, \ldots, \Psi_u]$$

where \( \mathbf{P} \) is a transpose operator, while the integer \( \Omega \) is a parameter and its value as well as the components \( \Psi_u \) \( (u=1,2,\ldots,\Omega) \) will depend on how to extract the desired information from the amino acid sequence of \( \mathbf{P} \), as elaborated in [4]. Thus, by following exactly the same procedures as described in the Section 2.2 of [4], each of the protein samples in the benchmark dataset can be uniquely defined as a 20-D numerical vector as given in Supporting Information S2, which can also be directly downloaded at http://www.jci-bioinfo.cn/pLoc_bal-mAnimal/Supp2.pdf.

### 2.3. Installing Deep-Learning for Three Deeper Levels

In this study, we use the CNN (Convolutional Neural Network) model to predict the subcellular localization of animal proteins, as illustrated in [Figure 1].

The CNN model consists of input layer, convolutional layer, average-pooling layer and fully connected layer. The input layer represents each animal protein with 6 features. The second layer is convolutional layer which extract dependency relationship between features subsequence of animal proteins. The filter stride is set to one. The activation function is set as "relu". The average-pooling layer down-samples the features and compute the average values of the features. The fully connected layer consists of 2 hidden layers. Finally, the output of connected layer was concatenated into output layer with sigmoid activation function. The label of animal protein was decided by the threshold \( \theta \). If the output is greater than 0.5, the outcome was true; otherwise, false.

The other parameters of CNN model are as follows. 1) The algorithm of Adam was used to train the model and the loss function is set to binary cross-entropy. 2) The activation function of full connected layer and convolutional layer is ReLU [39], and the activation function of output layer is sigmoid. 3) Convolutional Layer used the filter size 2 * 1 to extract features of animal proteins. 4) The batch size is 26. 5)

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[Figure 1. An illustration to show the Architecture of the pLoc-Deep_mAnimal model.]
The model is trained for 120 epochs. The metrics is set as “accuracy”.

The new predictor developed via the above procedures is called “pLoc_Deep-mAnimal”, where “pLoc_Deep” stands for “predict subcellular localization by deep learning”, and “mAnimal” for “multi-label animal proteins”.

3. RESULTS AND DISCUSSION

According to the 5-step rules [17], one of the important procedures in developing a new predictor is how to properly evaluate its anticipated accuracy. To deal with that, two issues need to be considered. 1) What metrics should be used to quantitatively reflect the predictor’s quality? 2) What test method should be applied to score the metrics?

3.1. A Set of Five Metrics for Multi-Label Systems

Different from the metrics used to measure the prediction quality of single-label systems, the metrics for the multi-label systems are much more complicated [40]. To make them more intuitive and easier to understand for most experimental scientists, here we use the following intuitive Chou’s five metrics [41] or the “global metrics” that have recently been widely used for studying various multi-label systems (see, e.g., [42, 43]). For the current study, the set of global metrics can be formulated as:

\[
\begin{align*}
\text{Aiming} & = \frac{1}{N^q} \sum_{k=1}^{N^q} \left( \frac{\|L_k \cap L_k^*\|}{\|L_k\|} \right), \quad [0,1] \\
\text{Coverage} & = \frac{1}{N^q} \sum_{k=1}^{N^q} \left( \frac{\|L_k \cap L_k^*\|}{\|L_k^*\|} \right), \quad [0,1] \\
\text{Accuracy} & = \frac{1}{N^q} \sum_{k=1}^{N^q} \left( \frac{\|L_k \cap L_k^*\|}{\|L_k \cup L_k^*\|} \right), \quad [0,1] \\
\text{Absolute true} & = \frac{1}{N^q} \sum_{k=1}^{N^q} \Delta(L_k, L_k^*), \quad [0,1] \\
\text{Absolute false} & = \frac{1}{N^q} \sum_{k=1}^{N^q} \left( \frac{\|L_k \cup L_k^*\| - \|L_k \cap L_k^*\|}{M} \right), \quad [1,0]
\end{align*}
\]

where \( N^q \) is the total number of query proteins or tested proteins, \( M \) is the total number of different labels for the investigated system (for the current study it is \( L_{\text{total}} = 20 \)), \( \| \| \) means the operator acting on the set therein to count the number of its elements, \( \cup \) means the symbol for the “union” in the set theory, \( \cap \) denotes the symbol for the “intersection”, \( L_k \) denotes the subset that contains all the labels observed by experiments for the \( k \)-th tested sample, \( L_k^* \) represents the subset that contains all the labels predicted for the \( k \)-th sample, and

\[
\Delta(L_k, L_k^*) = \begin{cases} 1, & \text{if all the labels in } L_k \text{ are identical to those in } L_k^* \\ 0, & \text{otherwise} \end{cases}
\]

In Equation (4), the first four metrics with an upper arrow \( \uparrow \) are called positive metrics, meaning that the larger the rate is the better the prediction quality will be; the 5\textsuperscript{th} metrics with a down arrow \( \downarrow \) is called negative metrics, implying just the opposite meaning.

From Equation (4) we can see the following: 1) the “Aiming” defined by the 1\textsuperscript{st} sub-equation is for checking the rate or percentage of the correctly predicted labels over the practically predicted labels; 2) the “Coverage” defined in the 2\textsuperscript{nd} sub-equation is for checking the rate of the correctly predicted labels over
the actual labels in the system concerned; 3) the “Accuracy” in the 3rd sub-equation is for checking the average ratio of correctly predicted labels over the total labels including correctly and incorrectly predicted labels as well as those real labels but are missed in the prediction; 4) the “Absolute true” in the 4th sub-equation is for checking the ratio of the perfectly or completely correct prediction events over the total prediction events; 5) the “Absolute false” in the 5th sub-equation is for checking the ratio of the completely wrong prediction over the total prediction events.

3.2. Comparison with the State-of-the-Art Predictor

Listed in Table 1 are the rates achieved by the current pLoc_Deep-mAnimal predictor via the cross validations on the same experiment-confirmed dataset as used in [4]. For facilitating comparison, listed there are also the corresponding results obtained by the pLoc_bal-mAnimal [4], the existing most powerful predictor for identifying the subcellular localization of animal proteins with both single and multiple location sites. As shown in Table 1, the newly proposed predictor pLoc_Deep-mAnimal is remarkably superior to the existing state-of-the-art predictor pLoc_bal-mAnimal in all the five metrics. Particularly, it can be seen from the table that the absolute true rate achieved by the new predictor is over 92%, which is far beyond the reach of any other existing methods [44-49]. This is because it is extremely difficult to enhance the absolute true rate of a prediction method for a multi-label system as clearly elucidated in [4]. Actually, to avoid embarrassment, many investigators even chose not to mention the metrics of absolute true rate in dealing with multi-label systems (see, e.g., [50-56]).

Moreover, to in-depth examine the prediction quality of the new predictor for the proteins in each of the subcellular locations concerned (cf. Table 2), we used the “local metrics” [40] or a set of four intuitive metrics that were derived in [57] based on the Chou’s symbols introduced for studying protein signal peptides [58] and that have ever since been widely concurred or justified (see, e.g., [59-62]). For the current study, the set of local metrics can be formulated as:

\[
\begin{align*}
Sn(i) &= 1 - \frac{N^+_i}{N^+_i} \quad 0 \leq Sn \leq 1 \\
Sp(i) &= 1 - \frac{N^-_i}{N^-_i} \quad 0 \leq Sp \leq 1 \\
Acc(i) &= 1 - \frac{N^+_i + N^-_i}{N^+_i + N^-_i} \quad 0 \leq Acc \leq 1 \\
MCC(i) &= \frac{1 - \left( \frac{N^+_i}{N^+_i} + \frac{N^-_i}{N^-_i} \right)}{\sqrt{\left( \frac{N^+_i}{N^+_i} \right) \left( \frac{N^-_i}{N^-_i} \right) \left( \frac{N^+_i}{N^+_i} \right) \left( \frac{N^-_i}{N^-_i} \right)}} \quad -1 \leq MCC \leq 1
\end{align*}
\]

(6)

where Sn, Sp, Acc, and MCC represent the sensitivity, specificity, accuracy, and Mathew’s correlation coefficient, respectively, and \( i \) denotes the \( i \)-th subcellular location (or subset) in the benchmark dataset. \( N^+_i \) is the total number of the samples investigated in the \( i \)-th subset, whereas \( N^-_i \) is the number of the samples in \( N^+_i \) that are incorrectly predicted to be of other locations; \( N^-_i \) is the total number of samples in any locations but not the \( i \)-th location, whereas \( N^-_i \) is the number of the samples in \( N^-_i \) that are incorrectly predicted to be of the \( i \)-th location.

Listed in Table 2 are the results achieved by pLoc_Deep-mAnimal for the animal proteins in each of 20 subcellular locations. As we can see from the table, nearly all the success rates achieved by the new predictor for the animal proteins in each of the 20 subcellular locations are within the range of 90% - 100%.
Table 1. Comparison with the state-of-the-art method in predicting animal protein subcellular localization.

| Predictor                | Aiming (↑)  | Coverage (↑)  | Accuracy (↑)   | Absolute true (↑) | Absolute false (↓) |
|--------------------------|-------------|---------------|----------------|-------------------|--------------------|
| pLoc_bal-mAnimal         | 88.31%      | 85.06%        | 84.34%         | 78.78%            | 0.07%              |
| pLoc_Deep-mAnimal        | 96.21%      | 97.77%        | 95.46%         | 92.26%            | 0.00%              |

*See Equation (4) for the definition of the metrics. *See [4], where the reported metrics rates were obtained by the jackknife test on the benchmark dataset of Supporting Information S1 that contains experiment-confirmed proteins only. *The proposed predictor; to assure that the test was performed on exactly the same experimental data as reported in [4] for pLoc_bal-mAnimal.

Table 2. Performance of pLoc_Deep-mAnimal for each of the 20 subcellular locations.

| i  | Location               | Sn(\(i\)) | Sp(\(i\)) | Acc(\(i\)) | MCC(\(i\)) |
|----|------------------------|------------|------------|------------|------------|
| 1  | Acrosome               | 1.0000     | 1.0000     | 1.0000     | 1.0000     |
| 2  | Cell cortex            | 0.9821     | 0.9957     | 0.9920     | 0.9798     |
| 3  | Cell membrane          | 0.9799     | 1.0000     | 0.9997     | 0.9897     |
| 4  | Centriole              | 0.9745     | 0.9997     | 0.9985     | 0.9833     |
| 5  | Centrosome             | 0.9155     | 0.9996     | 0.9984     | 0.9428     |
| 6  | Cytoplasm              | 0.9534     | 0.9779     | 0.9693     | 0.9317     |
| 7  | Cytoskeleton           | 0.9517     | 0.9986     | 0.9918     | 0.9666     |
| 8  | Endoplasmic reticulum  | 0.9662     | 0.9984     | 0.9929     | 0.9747     |
| 9  | Endosome               | 0.9744     | 0.9988     | 0.9974     | 0.9771     |
| 10 | Extracellular space    | 0.9170     | 0.9994     | 0.9967     | 0.9464     |
| 11 | Golgi apparatus        | 0.9716     | 0.9991     | 0.9957     | 0.9800     |
| 12 | Lysosome               | 0.9818     | 0.9997     | 0.9913     | 0.9876     |
| 13 | Melanosome             | 0.9585     | 0.9973     | 0.9995     | 0.9663     |
| 14 | Microsome              | 1.0000     | 0.9999     | 0.9999     | 0.9825     |
| 15 | Mitochondrion          | 0.9802     | 1.0000     | 0.9995     | 0.9897     |
| 16 | Nucleus                | 0.9408     | 0.9889     | 0.9782     | 0.9371     |
| 17 | Peroxisome             | 1.0000     | 1.0000     | 1.0000     | 1.0000     |
| 18 | Plasma membrane        | 0.9822     | 0.9957     | 0.9920     | 0.9798     |
| 19 | Spindle                | 0.9544     | 0.9994     | 0.9974     | 0.9705     |
| 20 | Synapse                | 0.9685     | 1.0000     | 0.9988     | 0.9834     |

*See Table 1 and the relevant context for further explanation. *See Equation (6) for the metrics definition.
which is once again far beyond the reach of any of its counterparts. Meanwhile, as a byproduct, the present paper has also stimulated some kinds of provocative or challenging but very intriguing papers (see, e.g., [63-68]).

3.3. Web Server and User Guide

As pointed out in [69], user-friendly and publicly accessible web-servers represent the future direction for developing practically more useful predictors. Actually, user-friendly web-servers will significantly enhance the impacts of theoretical work because they can attract the broad experimental scientists [31]. In view of this, the web-server of the current pLoc_Deep-mAnimal predictor has also been established at http://www.jci-bioinfo.cn/pLoc_Deep-mAnimal/, by which users can easily get their desired data without the need to go thru the mathematical details.

4. CONCLUSION

It is anticipated that the pLoc_Deep-mAnimal predictor holds very high potential to become a useful high throughput tool in identifying the subcellular localization of animal proteins, particularly for finding multi-target drugs that is currently a very hot trend in drug development. Most important is that the predictor will become a very useful tool for fighting against the pandemic coronavirus to save the mankind on this planet.

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CONFLICTS OF INTEREST

The authors declare no conflicts of interest regarding the publication of this paper.

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