



Showcase to Illustrate How the Web-Server pLoc_bal-mEuk is Working

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Dear Editor,

Recently, a very powerful web-server predictor has been established for identifying the subcellular localization of a protein based on its sequence information alone for the multi-label systems [1], in which a same protein may occur or move between two or more location sites and hence needs to be marked with the multi-label approach [2].

The web-server predictor is called “pLoc_bal-mEuk”, where “bal” means the web-server has been further improved by the “balance treatment” [3-9], and “m” means the capacity able to deal with the multi-label systems. To find how the web-server is working, please do the following.

Click the link at http://www.jci-bioinfo.cn/pLoc_bal-mEuk/, the top page of the pLoc_bal-mEuk web-server will appear on

pLoc_bal-mEuk: predict subcellular localization of eukaryotic proteins by general PseAAC and quasi-balancing training dataset

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Enter query sequences

Enter the sequences of query proteins in FASTA format ([Example](#)): the number of proteins is limited at 10 or less for each submission.

Or, upload a file for batch prediction

Enter your e-mail address and upload the batch input file ([Batch-example](#)). The predicted result will be sent to you by e-mail once completed; it usually takes 1 minute or so for each protein sequence

Upload file:
Your Email:

Figure 1 A semi screenshot for the top page of pLoc_bal-mEuk (Adapted from [5]).

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Covered by pLoc_bal-mEuk are the following 22 subcellular locations

(1) Acrosome	(2) Cell membran	(3) Cell wall
(4) Centrosome	(5) Chloroplast	(6) Cyanelle
(7) Cytoplasm	(8) Cytoskeleton	(9) Endoplasmic reticulum
(10) Endosom	(11) Extracellular	(12) Golgi apparatus
(13) Hydrogenosome	(14) Lysosome	(15) Melanosome
(16) Microsome	(17) Mitochondrion	(18) Nucleus
(19) Oerxisome	(20) Spindle pole body	(21) Synapse
(22) Vacuole		

Predicted results

Protein ID	Subcellular location or locations
>Q63564	1
>P23276	2, 8
>Q9VVV9	2, 7, 18
>Q673G8	2, 7, 10, 18

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Figure 2 A semi screenshot for the webpage obtained by following Step 3 of Section 3.5 (Adapted from [5]).

your computer screen, as shown in Figure 1. Then by following the Step 2 and Step 3 in [5], you will see the predicted results shown on Figure 2. Nearly all the success rates achieved by the web-server predictor for the eukaryotic proteins in each of the 22 subcellular locations are within the range of 90-100%, which is far beyond the reach of any of its counterparts.

Besides, the web-server predictor has been developed by strictly observing the guidelines of “Chou’s 5-steps rule” and hence have the following notable merits (see, e.g., [4-7,10-33] and three comprehensive review papers [2,34,35]): (1) crystal clear in logic development, (2) completely transparent in operation, (3) easily to repeat the reported results by other investigators, (4) with high potential in stimulating other sequence-analyzing methods, and (5) very convenient to be used by the majority of experimental scientists.

References

1. Chou KC, Shen HB. Recent progresses in protein subcellular location prediction. *Anal Biochem*. 2007; 370: 1-16.
2. Chou KC. Advance in predicting subcellular localization of multi-label proteins and its implication for developing multi-target drugs. *Curr Med Chem*. 2019; 4918-4943.
3. Xiao X, Cheng X, Chen G, Mao Q, Chou KC. pLoc_bal-mVirus: Predict Subcellular Localization of Multi-Label Virus Proteins by Chou’s General PseAAC and IHTS Treatment to Balance Training Dataset. *Med Chem*. 2019; 15: 496-509.
4. Xiao X, Cheng X, Chen G, Mao Q, Chou KC. pLoc_bal-mGpos: predict subcellular localization of Gram-positive bacterial proteins by quasi-



- balancing training dataset and PseAAC. *Genomics*. 2019; 111: 886-892.
5. Chou KC, Cheng X, Xiao X. pLoc_bal-mEuk: predict subcellular localization of eukaryotic proteins by general PseAAC and quasi-balancing training dataset. *Med Chem*. 2019; 15: 472-485.
 6. Chou KC, Cheng X, Xiao X. pLoc_bal-mHum: predict subcellular localization of human proteins by PseAAC and quasi-balancing training dataset. *Genomics*. 2019; 111: 1274-1282.
 7. Cheng X, Lin WZ, Xiao X, Chou KC. pLoc_bal-mAnimal: predict subcellular localization of animal proteins by balancing training dataset and PseAAC. *Bioinformatics*. 2019; 35: 398-406.
 8. Cheng X, Xiao X, Chou KC. pLoc_bal-mPlant: predict subcellular localization of plant proteins by general PseAAC and balancing training dataset. *Curr Pharm Des*. 2018; 24: 4013-4022.
 9. Cheng X, Xiao X, Chou KC. pLoc_bal-mGneg: predict subcellular localization of Gram-negative bacterial proteins by quasi-balancing training dataset and general PseAAC. *J Theor Biol*. 2018; 458: 92-102.
 10. Barukab O, Khan YD, Khan SA, Chou KC. iSulfoTyr-PseAAC: Identify tyrosine sulfation sites by incorporating statistical moments via Chou's 5-steps rule and pseudo components. 2019.
 11. Ehsan A, Mahmood MK, Khan YD, Barukab OM, Khan SA, Chou KC. iHyd-PseAAC (EPSV): Identify hydroxylation sites in proteins by extracting enhanced position and sequence variant feature via Chou's 5-step rule and general pseudo amino acid composition. *Curr Genomics*. 2019; 20: 124-133.
 12. Feng P, Yang H, Ding H, Lin H, Chen W, Chou KC. iDNA6mA-PseKNC: Identifying DNA N(6)-methyladenosine sites by incorporating nucleotide physicochemical properties into PseKNC. *Genomics*. 2019; 111: 96-102.
 13. Hussain W, Khan YD, Rasool N, Khan SA, Chou KC. SPalmitoylC-PseAAC: A sequence-based model developed via Chou's 5-steps rule and general PseAAC for identifying S-palmitoylation sites in proteins. *Anal Biochem*. 2019; 568: 14-23.
 14. Hussain W, Khan YD, Rasool N, Khan SA, Chou KC. SPrenylC-PseAAC: A sequence-based model developed via Chou's 5-steps rule and general PseAAC for identifying S-prenylation sites in proteins. *J Theor Biol*. 2019; 468: 1-11.
 15. Ilyas S, Hussain W, Ashraf A, Khan YD, Khan SA, Chou KC. iMethylK-PseAAC: Improving accuracy for lysine methylation sites identification by incorporating statistical moments and position relative features into general PseAAC via Chou's 5-steps rule. *Curr Genomic*. 2019; 20.
 16. Jia J, Li X, Qiu W, Xiao X, Chou KC. iPPI-PseAAC(CGR): Identify protein-protein interactions by incorporating chaos game representation into PseAAC. *J Theor Biol*. 2019; 460: 195-203.
 17. Khan YD, Batool A, Rasool N, Khan A, Chou KC. Prediction of nitrosocysteine sites using position and composition variant features. *Letters in Organic Chemistry*. 2019; 16: 283-293.
 18. Khan YD, Jamil M, Hussain W, Rasool N, Khan SA, Chou KC. pSSbond-PseAAC: Prediction of disulfide bonding sites by integration of PseAAC and statistical moments. *J Theor Biol*. 2019; 463: 47-55.
 19. Lu Y, Wang S, Wang J, Zhou G, Zhang Q, Zhou X, et al. An Epidemic Avian Influenza Prediction Model Based on Google Trends. *Letters in Organic Chemistry*. 2019; 16: 303-310.
 20. Niu B, Liang C, Lu Y, Zhao M, Chen Q, Zhang Y, et al. Glioma stages prediction based on machine learning algorithm combined with protein-protein interaction networks. *Genomics*. 2019.
 21. Pugalenth G, Nithya V, Chou KC, Archunan G. Nglyc: A random forest method for prediction of N-Glycosylation sites in eukaryotic protein sequence. *Protein Pept Lett*. 2019.
 22. Salman K, Mukhtaj K, Nadeem I, Tahir H, Sher AK, KC Chou. A two-level computation model based on deep learning algorithm for identification of piRNA and their functions via Chou's 5-steps rule. *IJPRT*. 2019.
 23. Xiao X, Xu ZC, Qiu WR, Wang P, Ge HT, Chou KC. iPSW(2L)-PseKNC: A two-layer predictor for identifying promoters and their strength by hybrid features via pseudo K-tuple nucleotide composition. *Genomics*. 2019; 111: 1785-1793.
 24. Yang L, Lv Y, Wang S, Zhang Q, Pan Y, Su D, et al. Identifying FL11 subtype by characterizing tumor immune microenvironment in prostate adenocarcinoma via Chou's 5-steps rule. *Genomics*. 2019.
 25. Wiktorowicz A, Wit A, Dziewierz A, Rzeszutko L, Dudek D, Kleczynski P. Calcium Pattern Assessment in Patients with Severe Aortic Stenosis Via the Chou's 5-Steps Rule. *Curr Pharm Des*. 2019.
 26. Vishnoi S, Garg P, Arora P. Physicochemical n-Grams Tool: A tool for protein physicochemical descriptor generation via Chou's 5-steps rule. *Chem Biol Drug Des*. 2019.
 27. Liang Y, Zhang S. Identifying DNase I hypersensitive sites using multi-features fusion and F-score features selection via Chou's 5-steps rule. *Biophys Chem*. 2019; 253: 106227.
 28. Liang R, Xie J, Zhang C, Zhang M, Huang H, Huo H, et al. Identifying Cancer Targets Based on Machine Learning Methods via Chou's 5-steps Rule and General Pseudo Components. *Curr Top Med Chem*. 2019; 19: 2301-2317.
 29. Lan J, Liu Z, Liao C, Merkler DJ, Han Q, Li J. A Study for Therapeutic Treatment against Parkinson's Disease via Chou's 5-steps Rule. *Curr Top Med Chem*. 2019; 19: 2318-2333.
 30. Kabir M, Ahmad S, Iqbal M, Hayat M. iNR-2L: A two-level sequence-based predictor developed via Chou's 5-steps rule and general PseAAC for identifying nuclear receptors and their families. *Genomics*. 2019.
 31. Ju Z, Wang SY. Prediction of lysine formylation sites using the composition of k-spaced amino acid pairs via Chou's 5-steps rule and general pseudo components. *Genomics*. 2019.
 32. Du X, Diao Y, Liu H, Li S, Li M. MsDBP: Exploring DNA-binding Proteins by Integrating Multi-scale Sequence Information via Chou's 5-steps Rule. *Journal of Proteome Research*. 2019; 18: 3119-3132.
 33. Chen Y, Fan X. Use of Chou's 5-Steps Rule to Reveal Active Compound and Mechanism of Shuangshen Pingfei San on Idiopathic Pulmonary Fibrosis. *Current Molecular Medicine*. 2019.
 34. Chou KC. Some remarks on protein attribute prediction and pseudo amino acid composition. *J Theor Biol*. 2011; 273: 236-247.
 35. Chou KC. Impacts of pseudo amino acid components and 5-steps rule to proteomics and proteome analysis. *Current Topics in Medicinal Chemistry*. 2019.