A novel scoring schema for peptide identification by searching protein sequence databases using tandem mass spectrometry data

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Abstract

Background: Tandem mass spectrometry (MS/MS) is a powerful tool for protein identification. Although great efforts have been made in scoring the correlation between tandem mass spectra and an amino acid sequence database, improvements could be made in three aspects, including characterization of peaks in spectra, adoption of effective scoring functions and access to the reliability of matching between peptides and spectra.

Results: A novel scoring function is presented, along with criteria to estimate the performance confidence of the function. Through learning the types of product ions and the probability of generating them, a hypothetic spectrum was generated for each candidate peptide. Then relative entropy was introduced to measure the similarity between the hypothetic and the observed spectra. Based on the extreme value distribution (EVD) theory, a threshold was chosen to distinguish a true peptide assignment from a random one. Tests on a public MS/MS dataset demonstrated that this method performs better than the well-known SEQUEST.

Conclusion: A reliable identification of proteins from the spectra promises a more efficient application of tandem mass spectrometry to proteomes with high complexity.

Background

A major goal of proteomics is to study biological processes comprehensively through the identification, characterization, and quantification of expressed protein in a cell or a tissue. Recently, tandem mass spectrometry has been shown to be a powerful tool for sensitive high-throughput identification of proteins [1,2]. Following enzymatic digestion of proteins, the resulting peptides are separated in the mass analyzer according to their mass to charge ratio (m/z-value). Peptides with the same m/z value are mostly broken into two fragments at a single peptide bond, forming N-terminal ions, i.e. a, b or c-ion and C-terminal ions, i.e., x, y or z-ion. All ions are then detected to generate a MS/MS spectrum (See Figure 1). In a mass spectrum, the intensity of a peak is generally proportional to the frequency of ions with the corresponding m/z-value [3].

A number of factors complicate the interpretation of MS/MS data. Neither the peaks corresponding to the a/b/c ver-
on peptide fragmentation [11]. ProbiID [8], a method based on a probabilistic model, adopted a Bayesian approach to interpret mass spectra data.

Random matching between experimental and theoretical masses may bring about false-positive results, giving rise to another key problem of peptide identification—the criteria for evaluation of the reliability of the matching. The difference between the highest and second highest scores [5] and p-value [7] estimate are used to filter false positives. Jan Eriksson [12] and Keller [13] built a model to work out the distribution of scores from random matches, which allowed significance testing under general database searching constraints. Therefore, filtering criteria to distinguish a valid match from all matches should be developed toward being dependent on quantitative estimates rather than on experience.

The present paper aims to tackle the two problems mentioned above.

1. We introduce a new, effective probabilistic scoring function. Adopting a statistical model similar to Dancik et al [8], we have employed relative entropy (i.e., K-L distance) to measure the similarity between hypothetical and experimentally observed spectra. We present a brief proof to show that relative entropy is indeed the simplified form of the conditional probability that the spectrum is generated from the peptide.

2. We present an EVD-based criterion to distinguish valid matches from random ones. Each spectrum will acquire the best score from correlation with all candidate peptides. Such best scores conform to the extreme value distribution, which underlies the quantitative threshold of the significance test.

A system written in C, the Protein Identifier (PI), was implemented adopting the relative entropy scoring function and tested on real data. Tests showed that it performed better than a widely used cross-correlation scoring approach [5]. For the sake of convenience, we have made the program available on our website PI http://www.bioinfo.org.cn/MSMS/papers/supp1/.

Methods

Data set
A publicly available spectrum set from Keller et al [14] was used to test our algorithm. It contains spectra generated by digesting two 18-proteins mixtures with trypsin. To assign a candidate peptide to each spectrum, Keller et al [14] ran SEQUEST against a human peptide database, obtaining 37,044 pairs of peptide and spectrum [14]. Among these assignments, 125 assignments of peptides with a single charge, 1,656 with double charges and 984 with triple...
charges were confirmed manually. For the following two reasons, the original dataset above was subjected to a filtering procedure for refinement. First, because the enzyme had been slightly contaminated (private communication with A. Keller who published the data), peptides with abnormal termini were screened out. Second, the spectra from triply charged ions were filtered out because their fragmentation pattern is not yet fully understood. As for partial proteolysis, peptides with up to 5 trypsin cleavage sites were kept for database searching. The filtering procedure produced a refined query set containing 19,000 spectra, and 1,247 correct peptide assignments. Thereafter the 1,247 correct assignments was divided into a training set derived from 2 proteins and a test set derived from the other proteins for cross validation. The training set was used to study the characteristics of the mass spectrometry while the testing set was to test the performance of our algorithm. (Available at the website PI http://www.bioinfo.org.cn/MSMS/papers/supp1/).

Description of algorithm

In order to describe our statistical model, a set of terms similar to Dancik et al [9] and Vineet Bafna et al [10] were defined as follows:

Let \( A \) be the set of amino acids, each amino acid having a molecular mass \( m(a) \), \( a \in A \). A peptide is denoted as a sequence of amino acids \( P = p_1p_2...p_n \) with mass \( m(P) = \sum_i m(p_i) \), \( p_i \in A \). Dissociation at the \( i \)-th peptide bond forms two partial peptides, the N-terminal peptide \( P_i = p_1p_2...p_i \), \( i = 1,2,...,n \), and the C-terminal peptide \( P'_i = p_{i+1}...p_n \). Formally, ion types are represented as a set of numbers indicating the offset of ion to peptide, \( \delta = \{ \delta_1,\delta_2,...,\delta_K \} \), thus, the i-type ion of the partial peptide \( P \) has the mass \( m(P) + \delta_i \). For example, \( \delta = \{1,-27,-17,-16\} \) corresponds to the most frequent N-terminal ions, \( b, a, b' - H_2O, b' - NH_3 \), respectively. The interval from 0 to the precursor ion mass \( M = m(P) \) is discretized into \( N \) bins, hence, a MS/MS spectrum is represented as a vector of intensity, i.e., \( S = \{I_1,I_2,...,I_N\} \), where \( I_i \) is the intensity of the peak with the \( m/z \)-value \( i^*M/N \).

Fragmentation in mass spectrometry is a stochastic process governed by the collision dynamics and the physicochemical properties of a peptide [10]. The fragmentation of a set of the peptide in spectrometry was modeled by a random "ball tossing" trial, each trial tossing an N-terminal ion and/or a C-terminal ion in some bins. Thus, after a series of trials, the number of ions in a bin will be proportional to the intensity of the corresponding peak. The fragmentation of the different peptides, i.e., tossing of different balls, is assumed to be mutually independent. Let random variable \( C \) be the cleavage site, and \( \Pr(C = i,P) \) be the probability of cleavage at the \( i \)-th bond of peptide \( P \). Let \( \Pr(\delta) \) be the probability that a breakage generates an i-type ion in a trial. Since fragmentation is residue dependent [15], we can adopt a more accurate definition, the conditional probability \( \Pr(\delta|p_j) \), where \( p_j \) is the amino acid adjacent to the cleavage site. As mass spectrometry may also generate peaks due to merely "random noise", \( \Pr(R) \) is introduced to denote the probability of such event. For the sake of simplicity, only N-terminal ions are studied here; the considerations would have been similar for C-terminal ions.

Parameter learning and hypothetic spectrum generating

To estimate the above peptide and instrument specific parameters, a learning procedure was modeled according to the offset frequency function [9], with improvements added to take account of intensity. For a set of experimental spectra with known peptide sequences thereof, let \( S = \{I_1,I_2,...,I_N\} \) be a spectrum corresponding to the peptide \( P \), \( \varepsilon \) be the \( m/z \) precision level of the instrument, \( N(S,P,\delta) \) be the sum of the peak intensities within a distance \( \varepsilon \) from \( m(P) \), \( -\delta_i = 1,...,n-1,i = 1,...,|\delta| \), then 

\[
N(S,P_i) = \sum_j N(S,P_i,\delta_j)
\]

is the number of trials during which the \( i \)-th bond was dissociated. Dividing \( N(S,P_i) \) by the total intensity, i.e., \( N(S,P_i)/\sum_j I_j \), gives an estimate of \( \Pr(C = i,P) \) over spectrum \( S \), and \( N(S,P,\delta)/N(S,P) \) is a ditto of \( \Pr(\delta) \). The probability of random noise \( \Pr(R) \) is estimated as the sum of the intensities of the unexplained peaks divided by the number of bins \( N \). Averaging them over all samples will improve the estimation accuracy.

Fragmentation tends to occur more frequently in the middle than at ends of a peptide. J. Simpson [16] represented the relationship between peak's intensity and cleavage sites as a function of the relative position of a breakpoint, ranging from 0 to 1, whereas David L. Tabb[17] adopted a function based on relative mass of the partial peptide. Here, we used Tabb's approach to work out \( \Pr(C = i,P) \).

Utilizing the above probability, we produced a hypothetic spectrum for a given peptide through simulating its dissociation in the spectrometry. A set of the peptide fragments at the \( i \)-th peptide bond with the probability \( \Pr(C = i,P) \), and generates the \( j \)-type ion with probability \( \Pr(\delta) \). Hence, peaks with mass \( m(P) - \delta \) were assigned with relative intensity \( \Pr(C = i,P)*\Pr(\delta) \), and other peaks were assigned with \( \Pr(R) \) to simulate random noises.
Relative entropy based scoring function

Before comparison to the hypothetic spectrum, an experimental spectrum was pre-processed by dividing the intensity of each peak by the total ion intensity and filling blank bins with the value Pr(R) to simulate random noise. Hence, both the experimental and hypothetic spectra were transformed into distributions of ions over N bins, and relative entropy was employed to measure the similarity between them.

Relative entropy is a pertinent statistical tool to measure distance between two distributions [18]. For two distributions \( U = \langle u_1,u_2,...,u_n\rangle, \sum_i u_i = 1 \) and \( V = \langle v_1,v_2,...,v_n\rangle, \sum_i v_i = 1 \), the relative entropy is defined as \( H(U,V) = \sum u_i \cdot \ln \left( \frac{u_i}{v_i} \right) \). \( H(U,V) \) is always positive, and the more similar the two distributions, the smaller the relative entropy.

The relative entropy measure is essentially a simplified entropy. More similar the two distributions, the smaller the relative entropy was computed as follows:

For each round of test, the frequency of different ions were learnt from a training assignment comprising peptides from two to four proteins (generally less than 300 peptides assignments), including non-peptide bond backbone cleavage, isotope forms, and neutral loss (Figure 2). Here, the precision of an ion-trap mass spectrometry was set to 0.5 Dalton. It could be observed that \( q_i \) is the probability that an ion will be tossed into the \( i \)-th bin, and \( I_i \) the number of ions tossed into this bin after \( I_{total} = \sum_i I_i \) tosses, the count vector \( I = \langle I_1,I_2,...,I_N\rangle \) conforms to a multi-nominal distribution. Hence, \( \Pr(S|P) \) can be computed as follows:

A score matrix was produced after comparing a set of spectra \( S = \{S_1,S_2,...,S_m\} \) with a set of candidate peptides \( C = \{C_1,C_2,...,C_n\} \) (Table 1).

For a specific spectrum \( S_i \), the peptide with the minimal score was thought to be the one most likely to have generated it. The minimal score of a row, recorded in the "min" column, conforms to the extreme value distribution (EVD). In fact, the distribution of items in the min column is the superposition of two EVD distributions because of the following reason. Dividing the spectra in \( S \) into two classes, i.e. \( S = D \cup E \), where \( D \) includes those that were generated from a peptide in \( C \) and \( E \) the rest. Thus, the match of a spectrum with a peptide in \( E \) can occur only by chance, and therefore results in a minimal score higher than the matches with peptides in \( D \). The gap between matches of the two peptide sets, along with the robustness of EVD, forms a foundation to distinguish valid matches from random ones. More important, EVD allows us to quantitatively estimate the confidence for a given criterion. It should be noted that since the score of the peptides depends on the mass of the precursor ions, so does the threshold.

Result and discussion

We performed the parameter learning, the hypothetic spectrum generation and the search procedure on the chosen dataset as described above (See Methods section)

Parameter learning

For each round of test, the frequency of different ions were learnt from a training assignment comprising peptides from two to four proteins (generally less than 300 peptides assignments), including non-peptide bond backbone cleavage, isotope forms, and neutral loss (Figure 2). Here, the precision of an ion-trap mass spectrometry was set to 0.5 Dalton. It could be observed that \( b, a, b - H_2O, b - NH_3 \), appear more frequently than other product ions, and isotope effect was also dominant. In Figure 3, the relationship between cleavage probability and relative position is shown as a bell-like curve. A conclusion could be made that cleavage tends to occur more frequently at middle than in the ends. The above observations coincide with the characteristic of fragmentation [11]. When the learning procedure was performed on different training sets, no obvious differences were observed for the learnt parameters.
Hypothetic spectrum generation

To visualize the similarity and difference between hypothetic and experimental spectra, the hypothetic spectrum for a peptide VGDANPALQK was generated using the learnt parameters (shown in Figure 4 along with the experimental spectrum). As can be seen from the comparison, the hypothetic spectrum matches the experimental at important ions with respect to relative abundance, including serial ions b, y, and their isotope and group-loss forms (for the clarity of the figure, only b and y ion had been showed).

It should be pointed out that for the positions without hypothetic peaks, random noise learnt from training set was assigned to it. In fact, noise was an intrinsic feature for ion-trap mass spectrometry [19].

However, for some ions, there exists an obvious difference in the intensities between the two spectra. The main reasons for these discrepancies are the residues dependent cleavage and the complicated mobility patterns of the proton. To improve the accuracy of the learning procedure advanced learning techniques, such as EM methods, should be introduced in future work.

Distribution of score and threshold setting

For each spectrum in a training set, similarity with all candidate peptides are measured. The distribution of the best similarity gives a plot conforming to the extreme value distribution (Figure 5). For spectra derived from peptides with no counterpart in the candidate set, the score values are all generated by chance, and one peak with higher score is observed in diagram (Figure 5, dash line). However, for spectra that are derived from peptides with a counterpart in the candidate set, the likely correct matches between spectra and candidate peptides form a peak with lower score. Therefore, two peaks will be observed—one corresponding to the random matches, and the other to the likely correct matches, the latter with lower score values (Figure 5, solid curve). From Figure 5, it is clear that there is a distinct distance between the two peaks, which provides a basis for setting a threshold value to effectively distinguish correct matches from random ones. In our experiment, the threshold was set to 3.8, i.e., the location of the cross-point of the two distributions, which minimizes the sum of false-positives and false-negatives. Therefore, if a match had a score less than 3.8, it was considered highly probable that the spectrum was produced

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**Table 1:** An example of score table. Comparison of spectrum $S_i$ with candidate peptide $C_j$ produces a score in the matrix, and each item of the 'min' column is the minimal score of the same row.

| $S_i$ | $C_1$ | $C_2$ | $C_3$ | $C_n$ | min |
|------|-------|-------|-------|-------|-----|
| $S_1$ | 2.4   | 3.7   | 8.9   | 2.4   |     |
| $S_2$ | 5.3   | 6.8   | 1.9   | 1.9   |     |
| ...  |       |       |       |       |     |
| $S_m$ | 7.7   | 10.0  | 6.1   | 6.1   |     |

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**Figure 2**

Occurrence probability of different N-terminal ions.
In general, serial ions bear a larger probability, however, their variants (like dehydrated form) also often appear. The m/z of the peptide bond is set to 0, and the relative probability is normalized to a total probability of 100.0 (including C-terminal ions, which are not shown).

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**Figure 3**

Breakage preference in the position along peptides.
Breakage tends to occur at middle rather than terminal positions of the peptides. The Abscissa denotes the position of residues relative to the whole sequence. The relative probability is normalized to a total of 100.0.
as result of the matched peptide. Threshold for three different training sets were 3.81, 3.66 and 4.09, respectively, that suggested the relative robustness of the threshold setting.

Performance of the score function
Searching the query spectra against the same human peptide database as Keller et al [14] had been performed three times by selecting different proteins as testing sets. Take the first round as an example (The first two proteins in the mixture were composed of the training set and the rest test set). Our program PI reported 1,053 potential assignments, having 801 assignments overlap with the test set. For the 252 assignments inconsistent with the test set, manual inspection showed that 171 of them were correct. The difference in performance arose from SEQUEST neglecting some correct assignment, as reported by Andrew. Finally, PI achieved an average sensitivity of 0.87 and an average error rate of 0.07. Table 2 shows the comparison between the performances of SEQUEST and PI. As for the stability of our program, when PI was tested on a different dataset from OPD [20] it still displayed a sound performance (sensitivity 0.89 and error rate 0.06). It is interesting that performance of PI on OPD dataset was better than that on Keller’s mixture dataset although the parameter training was done on the latter. This may be attributed to the different quality of these datasets. All the assignments are presented on our website at PI http://www.bioinfo.org.cn/MSMS/papers/supp1/.

Comparison of score functions
Roughly speaking, factors influencing the dissociation of peptides come in two categories. One is the chemical property of a peptide, and the other is the collision energy. In recent years, much effort has been made to unravel the fragmenting mechanism [15,21-26]. Based on theoretical computation, experimental studies, and statistical analyses, models for prediction of product ions have become increasingly sophisticated and accurate, taking into consideration the chemistry of the ion rearrangements, gas-phase basicity of the individual residues, and the location effect. However, there is apparently still a way to go before the fragmentation mechanism is completely understood. Therefore, due to the incomplete understanding of the fragmentation chemistry, analyzers still have to rely on statistical approaches. A probabilistic formulation was originally brought forth by Dancik [9] and provoked several subsequent studies [10,27]. However, the principle of "a premium for present ions" and "a penalty for non-present ions" [9] aims only at the hypothetic but not at the experimental spectrum. In other words, that formulation concentrates on what should appear in a hypothetic spectrum and neglects what emerges in the observed spectrum. Hence the formulation has the substantial defect that there is no penalty for emergence of unexplainable ions in an experimental spectrum. Our sys-
Table 2: Performance of SEQUEST and our program PI.
SEQUEST achieved different sensitivities with different filtering criteria. Here the best one is listed.

|       | Sensitivity | Error rate |
|-------|-------------|------------|
| SEQUEST | 0.78        | 0.09       |
| PI    | 0.87        | 0.07       |

tem describes the correlation between theoretical and experimental spectra in a bi-directional manner, taking account of non-coherence in a homogeneous way.

Conclusion
In this paper we present a novel method for correlating peptides with tandem mass spectra. The collision-induced dissociation is analyzed as a random event and occurrence probabilities for characteristic ions are calculated. By using these parameters, a statistic model to predict the theoretical spectra for peptides is built. Based on the hypothetical spectra, the relative entropy is presented to correlate the experimental spectra with the hypothetical ones. Then it is pointed out that the score of spectra obey-the superposition of two extreme value distributions, which allows the quantitative estimate for the confidence of peptide assignments. Computational experiment was done on two public databases and it showed that the performance of the present method was superior in comparison with SEQUEST.

Authors’ contributions
R. S. Chen and D.B. Bu conceived and organized the research; D. B. Bu proposed the relative entropy scoring function and EVD-based threshold setting, and wrote the main part of program PI; Z. Zhang characterized the model of peptide fragmentation, carried out the computational study, analyzed the result and wrote the paper; S. W. Sun proposed the statistical model for theoretical spectrum prediction and learning procedure; X. P Zhu analyzed the characteristic of ions; S. H Chang helped Zhang test the program; X. F. Liu and C. G. Yu wrote the website and interface the of software.

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References
1. Zhu H, Bilgin M, Snyder M: Proteomics. Annu Rev Biochem 2003, 72:783-812.
2. Yates JR 3rd: Mass spectrometry and the age of the proteome. J Mass Spectrom 1998, 33(1):1-19.
3. Aebersold R, Goodlett DR: Mass spectrometry in proteomics. Chem Rev 2001, 101(2):269-295.
4. Rappsilber J, Mann M: What does it mean to identify a protein inproteomics? Trends Biochem Sci 2002, 27(2):74-78.
5. Yates JR 3rd, Eng JK, McCormack AL: An approach to correlate tandem mass spectral data of peptides with amino acid sequences in the protein database. J Am Soc Mass Spectrom 1994, 5:976-989.
6. Yates JR 3rd, Eng JK, McCormack AL, Schietz D: Method to correlate tandem mass spectra of modified peptides to amino acid sequences in the protein database. Anal Chem 1995, 67(8):1426-1436.
7. Perkins DN, Pappin DJ, Creasy DM, Cottrell JS: Probability-based protein identification by searching sequence databases using mass spectrometry data. Electrophoresis 1999, 20(18):3551-3567.
8. Zhang N, Aebersold R, Schwikowski B: ProbiD: a probabilistic algorithm to identify peptides through sequence database searching using tandem mass spectral data. Proteomics 2002, 2(10):1406-1412.
9. Dancik V, Addona TA, Clauser KR, Vath JE, Pavznzer PA: De novo peptide sequencing via tandem mass spectrometry. J Comput Biol 1999, 6(3–4):327-342.
10. Bafna V, Edwards N: SCOPE: a probabilistic model for scoring tandem mass spectra against a peptide database. Bioinformatics 2001, 17(Suppl 1):S13-21.
11. Havilio M, Haddad Y, Smilansky Z: Intensity-based statistical scorer for tandem mass spectrometry. Anal Chem 2003, 75(3):435-444.
12. Eriksson J, Chait BT, Fenyo D: A statistical basis for testing the significance of mass spectrometric protein identification results. Anal Chem 2000, 72(5):999-1005.
13. Keller A, Nesvizhskii AI, Kolker E, Aebersold R: Empirical statistical model to estimate the accuracy of peptide identifications made by MS/MS and database search. Anal Chem 2002, 74(20):3383-3392.
14. Keller A, Purvins S, Nesvizhskii AI, Stolyar S, Goodlett DR, Kolker E: Experimental protein mixture for validating tandem mass spectral analysis. Omics 2002, 6(2):207-212.
15. Dongre AR, Jones JL, Somogyi A, Wysocki VH: Influence of peptide composition, gas-phase basicity, and chemical modification on fragmentation efficiency: evidence for the mobile proton bond in protonated peptides. J Mass Spectrom 2001, 36(12):1382-1390.
16. Schutz F, Kapp EA, Simpson RJ, Speed TP: Deriving statistical models for predicting peptide tandem MS product ion intensities. Biochem Soc Trans 2003, 31(6):1479-1483.
17. Tabb DL, Smith LL, Breci LA, Wysocki VH, Lin D, Yates JR 3rd: Statistical characterization of ion trap tandem mass spectra from doubly charged tryptic peptides. Anal Chem 2003, 75(5):155-163.
18. Baldi P, Brunak S: Bioinformatics-The machine learning approach, The MIT Press: Massachusetts. In Bioinformatics-The machine learning Approach The MIT Press: Massachusetts; 2001.
19. Knutichsky AN, Chait BT: On the nature of the chemical noise in MALDI mass spectra. J Am Soc Mass Spectrom 2002, 13(2):129-134.
20. Prince JT, Carlson MW, Wang R, Lu P, Marcotte EM: The need for a public proteomics repository. Nat Biotechnol 2004, 22(4):471-472.
21. Wysocki VH, Tsapralis G, Smith LL, Breci LA: Mobile and localized protons: a framework for understanding peptide dissociation. J Mass Spectrom 2000, 35(12):1399-1406.
22. Shukla AK, Futrell JH: Tandem mass spectrometry: dissociation of ions by collisional activation. J Mass Spectrom 2000, 35(9):1069-1090.
23. Schlosser A, Lehmann WD: Five-membered ring formation in unimolecular reactions of peptides: a key structural element controlling low-energy collision-induced dissociation of peptides. J Mass Spectrom 2000, 35(12):1382-1390.
24. Police MJ, Ren D, Wiesemesiotis C: Dissociation of the peptide bond in protonated peptides. J Mass Spectrom 2000, 35(12):1391-1398.
25. O’Hair RA: The role of nucleophile – electrophile interactions in the unimolecular and bimolecular gas-phase ion chemistry.
of peptides and related systems. J Mass Spectrom 2000, 35(12):1377-1381.
26. Paizs B, Suhai S: Fragmentation pathways of protonated peptides. Mass Spectrom Rev 2004.
27. Chen T, Kao MY, Tepel M, Rush J, Church GM: A dynamic programming approach to de novo peptide sequencing via tandem mass spectrometry. J Comput Biol 2001, 8(3):325-337.