Erratum

A model of random mass-matching and its use for automated significance testing in mass spectrometric proteome analysis

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On Page 262, the corresponding author should read:

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On page 265, equation 1 should read:

\[ p_i = \frac{1}{m_{i+1} - m_i} \sum_{k=0}^{m_i} \frac{\sum_{k=0}^{m_i} p(k)}{H} \]  

(1)

On page 265, equation 4 should read:

\[ f(S) = \left( \sum_{k=0}^{m_i} p(k) \right)^H - \left( \sum_{k=0}^{m_i} p(k) \right)^H \]  

(4)

On page 264, the legend to Fig. 1 should read:

Figure 1. Top left: Mass distribution of the proteins in a genome database (S. cerevisiae) compared with the mass distribution of the proteins identified in simulations using random tryptic peptide maps and ranking by the number of matches. Top right: The theoretical number, \( k_u \), of proteolytic peptides that a protein in a database (S. cerevisiae) can yield when at most \( u \) missed cleavage sites are assumed versus protein mass, \( M_p \). The line represents a least squares fit of a power function to the data. The power equals 1.0. Bottom left: The distribution of \( k_u \) values in the S. cerevisiae database for different values of \( u \). Bottom right: The distribution of \( k_u \) values for proteins identified when using random tryptic peptide maps. The distribution of \( k_u \) values in the whole database is shown for comparison.

On page 264, the legend to Fig. 2 should read:

Figure 2. Tryptic peptide mass distribution peaks in two different mass regions (S. cerevisiae).

On page 265, the legend to Fig. 3 should read:

Figure 3. The frequency of tryptic peptides (within a peptide mass distribution peak) as a function tryptic peptide mass.

On page 267, the legend to Fig. 4 should read:

Figure 4. Comparison of simulated and computed (see Section 3.1) frequency functions \( f(S) \) (left panel) of the score (number of matches) and scores required for significance (right panel) for random protein identification in various genomes. The search constraints are: maximum protein mass, \( M_p < 100 \) kDa, maximum number of missed trypsin cleavages, \( u = 2 \), mass accuracy, \( \Delta m = 0.1 \) Da, unless stated otherwise in the legend.

On page 268, the legend to Fig. 5 should read:

Figure 5. Frequency functions for random protein identification obtained by simulation and model computation for three different cases. Top, (1) all the masses in the maps are between 1396 and 4500 Da; middle, (2) the maps include the entire mass range 800 to 4500 Da; bottom, (3) the maps include masses between 800 and 1396 Da. The model-based computation takes the actual peptide mass distribution as well as all other constraints into account in a direct and rapid way.

On page 268, the legend to Fig. 6 should read:

Figure 6. Simulations demonstrating automated model-based significance testing implemented in a protein identification algorithm that ranks the proteins by their respective number of matches. In each map, a fraction of a total of 35 masses originated from a single randomly chosen protein (correlated masses) and the rest of the masses were each from a different protein (noncorrelated). The correlated masses corresponded to a randomly chosen protein sequence-coverage in the range 15–65%. The significance testing efficiently rejects false results. As a lower frequency of false results is tolerated, more true results become nonsignificant.

On page 269, the legend to Fig. 7 should read:

Figure 7. A comparison between the use of 2, 4 and 8 mass regions in the model computations of frequency functions for random protein identification.

On page 270, the legend to Fig. 8 should read:

Figure 8. The random coverage, \( \delta \), of a peptide mass distribution peak as a function of peptide mass (see Appendix).

On page 270, the legend to Fig. 9 should read:

Figure 9. The mean value of the random coverage, \( \delta \), in four different mass regions (1: 800–1054 Da, 2: 1055–1395 Da, 3: 1396–2055 Da, 4: 2055–4500 Da) as a function of the mass accuracy. The \( \delta(i, \Delta m) \) functions were derived from the S. cerevisiae genome only, but peptide mass distributions are highly conserved between genomes.