rTANDEM, an R/Bioconductor package for MS/MS protein identification

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ABSTRACT
Summary: rTANDEM is an R/Bioconductor package that interfaces the X!Tandem protein identification algorithm. The package can run the multi-threaded algorithm on proteomic data files directly from R. It also provides functions to convert search parameters and results to/from R as well as functions to manipulate parameters and automate searches. An associated R package, shinyTANDEM, provides a web-based graphical interface to visualize and interpret the results. Together, those two packages form an entry point for a general MS/MS-based proteomic pipeline in R/Bioconductor.

Availability and implementation: rTANDEM and shinyTANDEM are distributed in R/Bioconductor, http://bioconductor.org/packages/release/bioc/. The packages are under open licenses (GPL-3 and Artistic-1.0).

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1 INTRODUCTION
Protein identification and analysis by tandem mass spectrometry relies mostly on matching spectra to a database of protein sequences and scoring those matches. The resources of the R statistical language and its platform Bioconductor (Gentleman et al., 2004) have proven to be well suited for this kind of data manipulation and statistical scoring with regard to genomic data. However, those resources remain largely untapped with regard to proteomic data. This is partly explained by the lack of tools to perform protein identification directly from R. rTANDEM fills this gap by implementing in R the tandem algorithm (Craig and Beavis, 2004) and various associated scoring functions (Keller et al., 2005; Kertesz-Farkas et al., 2014; MacLean et al., 2006). rTANDEM also provides functions for conversion to/from R. This brings to proteomics the many advantages of building an analysis pipeline in the R/Bioconductor statistical platform: easy deployment on high-performance computing and cloud computing through Bioconductor Cloud Amazon Machine Image (AMI), fully open-source workflows, interconnectivity of annotation and analytic packages, full reproducibility of analysis, etc.

Fig. 1. Building processing pipelines around rTANDEM

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resources of R and Bioconductor (Fig. 1). For example, protein accession can be retrieved and passed to biomaRt (Durinck et al., 2005, 2009) to retrieve cross-references or annotation, which in turn can be passed to packages like topGO (Alexa and Rahnenfuhrer, 2010) to calculate annotation enrichment and RamiGO (Schröder, 2013) to display the Gene Ontology (GO) tree. The online Supplementary Material provides a short tutorial that demonstrates this kind of workflow.

3 shinyTANDEM

The shinyTANDEM package is a web-based graphical interface that provides an easy way to visualize results from rTANDEM. It is based on the newly released ‘shiny’ package from the RStudio team. This package adds web-server functionalities to R, making it possible to visualize the results of computations in a web browser. shinyTANDEM extends the basic use of ‘shiny’ to provide a full Graphical User Interface (GUI) from a single R command. The interface will display in the user’s default web browser.

The interface comprises a series of tabs representing various aspects of the search results: overview, statistics, protein view, peptide view. The ‘Overview’ tab presents the search parameters as well as raw lists of protein and peptide identifications. The ‘Statistics’ tab plots the protein and peptide matches according to their score and expectation values (Fig. 2). The ‘Protein’ tab presents filters to select specific proteins and present their coverage and associated peptides. The ‘Peptide’ tab presents filters to select specific peptide sequences and see their associated MS2 spectra and protein sequence.

shinyTANDEM also features tabs that act as entry points for R. A loading tab allows the user to change the R object that is represented. Tabs allowing the user to create parameters objects, launch analysis, start conversion process or link to other R packages directly from the interface are currently being implemented.

4 RESULTS

rTANDEM was tested on a recently published dataset of breast cancer tissues (Liu et al., 2013) and compared with the reported results obtained with MaxQuant (v.1.1.1.36). We used raw data obtained from four whole tissue lysate of breast cancer samples. rTANDEM obtained an average of 8044 unique peptide-spectrum matches at an expect value $<0.01$ for those samples (Supplementary Table S1). This represents a noticeable increase compared with the original results from MaxQuant, which reported an average of 6254 peptides per sample at $P<0.01$.

5 CONCLUSION

The Bioconductor package rTANDEM and its associated graphical interface, shinyTANDEM, form an entry point to build complete proteomics workflows in the R statistical language. They provide ways to perform protein identification directly from R and to convert search results to/from R object. The S4 result data structure makes it easy to use R/Bioconductor to build complex processing pipelines around proteomic datasets. Further statistical tests for quantification in rTANDEM are currently under development.

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REFERENCES