MODOMICS: a database of RNA modification pathways. 2017 update

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ABSTRACT

MODOMICS is a database of RNA modifications that provides comprehensive information concerning the chemical structures of modified ribonucleosides, their biosynthetic pathways, the location of modified residues in RNA sequences, and RNA-modifying enzymes. In the current database version, we included the following new features and data: extended mass spectrometry and liquid chromatography data for modified nucleosides; links between human tRNA sequences and MINTBase - a framework for the interactive exploration of mitochondrial and nuclear tRNA fragments; new, machine-friendly system of unified abbreviations for modified nucleoside names; sets of modified tRNA sequences for two bacterial species, updated collection of mammalian tRNA modifications, 19 newly identified modified ribonucleosides and 66 functionally characterized proteins involved in RNA modification. Data from MODOMICS have been linked to the RNAcentral database of RNA sequences. MODOMICS is available at http://modomics.genesilico.pl.

INTRODUCTION

The presence of modified nucleosides in RNA, beyond the basic A, U, C and G, has been recognized for more than half a century. However, their importance to RNA biochemistry and cell biology has been underappreciated, mainly because very few modifications (at defined positions in defined RNA molecules) were found to be truly indispensable for basic biological processes [review: (1)]. Now we know that 163 post-transcriptional modifications of RNA introduce a functional diversity that allows the four basic ribonucleotide residues to gain diverse functions, akin to those of side chains of amino acid residues, which may be e.g., polar, charged, aliphatic or aromatic. Modifications can directly influence RNA structure, by promoting or disrupting certain intramolecular interactions; they can make the RNA molecule more rigid or more flexible. They can also influence RNA interactions with other molecules, in particular proteins. Overall, they contribute strongly to the diversity of functions fulfilled by RNA molecules, especially within complex regulatory networks, where small subtle structural changes can bring about significant changes to cellular metabolism (2).

In the last years, new types of RNA modifications have been found, and biochemical and physiological roles have been elucidated for many known modified ribonucleosides (3–5). Some of these advances were driven by the use of liquid chromatography/mass spectrometry (LC/MS)-based methods, which provide highly precise quantification of changes in the spectrum of modified ribonucleosides in RNA from any organism, facilitating the study of translational control of cellular responses and phenotypes (6). Moreover, a number of previously unknown RNA-modifying enzymes have been identified and characterized.

New important roles for long-known RNA modifications were also discovered. Prominent examples include the involvement of N6-methyladenosine (m6A) in regulating gene expression by influencing transcript stability, splicing, translation efficiency and cap-independent translation, and in promoting circular RNA translation [review: (7)]. Mutations in many human genes encoding RNA modification

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enzymes have been linked to diseases, such as cancer, cardiovascular diseases, metabolic diseases, neurological disorders, and mitochondria-related defects [review: (8)].

To adequately represent the recent accumulation of knowledge, we have added both to the variety and volume of data in the MODOMICS database. The most significant additions are: (i) extensively updated datasets: new modifications, new enzymes, and new RNA sequences with modifications; (ii) a new category of LC/MS data for modifications (Figure 1); (iii) new naming/numbering convention for modified residues in RNA sequences; (iv) replacement of Jmol by JSmol for 3D structure viewing.

**DATABASE CONTENT**

MODOMICS has been developed to house and distribute collections of RNA modification pathways, chemical structures of modified nucleosides, sequences of modified RNAs, enzymes responsible for individual reactions, a catalog of ‘building blocks’ for chemical synthesis of modified RNA, and to be expanded to include new data types. The database was created as a single resource to organize and present all these data in a convenient and straightforward way and is currently the most comprehensive source of information among all existing RNA modification databases. Information about modified residues is also available in the RNAMDB database (9), while information about modifications among all existing RNA modification databases. In- is currently the most comprehensive source of information about 340 functionally characterized proteins involved in RNA modification, both functional enzymes and protein co-factors necessary for multi-protein enzymatic activities.

For each protein a set of detailed information is provided and includes: identifiers and accession numbers from relevant resources and databases such as: NCBI GI, UniProt ID, COG number, PDB ID of structure (if available); amino acid sequence; corresponding ORF; information about catalyzed reaction, the position of modification and modified RNA(s) (if available). For proteins that are parts of enzymatic complexes, the name of the complex is provided.

MODOMICS also contains human and yeast snoRNAs, involved in RNA-guided RNA modification by the C/D box and H/ACA box ribonucleoproteins, linked to the corresponding modification sites in human and yeast RNAs and the catalog of ‘building blocks’ for the chemical synthesis of naturally occurring modified nucleosides.

Several options for database searching and querying are implemented in MODOMICS, including the BLAST (12) search of protein sequences and the PARALIGN (13) search of nucleic acid sequences collected in MODOMICS, as well as a utility that sends a protein sequence from a MODOMICS entry to BLAST on the NCBI web server.

**Updated modifications section**

Since the previous release of MODOMICS (14), 19 new modifications were added to the database. Among those are four types of geranylated nucleosides discovered in bacterial tRNA (3), 5-cyanomethyluridine (cm5U) (15), and 2′-O-methyluridine 5-oxyacetic acid methyl ester (mcm5Um) (5). LC/MS analyses of tRNAs from *Bacillus subtilis*, plants, and *Trypanosoma brucei* revealed the presence of 2-methylthio cyclic N6-threonylcarbamoyladenosine (ms2ct6A), a derivative of N6-threonylcarbamoyladenosine (t6A), at position 37 of tRNAs responsible for recognition of adenosine-starting codons (16).

3D chemical structures of modified nucleosides are now displayed with JSmol (17). It is an open-source JS library and HTML5 viewer for 3D chemical structures which, in contrast to the previously used Jmol tool, does not require the installation of the Java software package. JSmol can also be used on systems that no longer support Java applets due to security concerns or for which Java is not available, like smartphones or tablets, and it does not use hardware graphics acceleration, enabling the software to run in any browser that supports HTML5 standards.

LC/MS data for modified nucleosides

This new MODOMICS release features a new section on modification detail page created to host the LC/MS data of the modified nucleoside. The new fields include information concerning the product ions, the protonated mass [M+H]+, the LC elution order and its characteristics, the normalized LC elution time and their literature references. The LC elution time is normalized to guanosine (G), measured with an RP C-18 column with acetonitrile/ammonium acetate as mobile phase and the elution order is based on the retention times of C, U, G, A and m6A to cover all areas of the chromatogram. The LC data is intended to provide the
Figure 1. Example of MS/LC data for N2,N2-dimethylguanosine (m2,2G). The display of a new data type: LC/MS information on the modification detail page.

novice LC-MS user with guidance on the relative hydrophobicity of modified nucleosides and an estimated elution region using the denoted stationary and mobile phases. Currently, only 48 modifications, for instance, m1A, m3C, and m2,2G are associated with LC information, while 138 modifications have been associated with MS information. The new system is in place to be extended to all the modifications present in the database as soon as the new data become available. LC/MS based methods allow RNA modification profiling of different organisms in a semi-quantitative manner for the newly-detected modifications along with known modifications, and one can expect that approach will be further extended. LC/MS data for MODOMICS collection of modified nucleosides provides a comprehensive source of information for mapping of the identity and position of modified residues in RNA sequences.

New nomenclature for modified nucleosides

The old systems used to encode modified residues in RNA sequences have been very cumbersome for automated data processing, especially in the case of special characters (often interpreted as special actions) or names that contained letters such as ‘c’ or ‘i’ that could be confused with different bases. Thus, we developed a new naming convention that uses only digits in addition to standard letters (A, G, C, U), and which makes names distinct from one another. In the new proposed system, a number is introduced before the modified residue, so the software for sequence processing can recognize the original residue type before modification, as well as identify the specific modification(s) introduced. For the most common modification type, i.e., simple methylations, we use single digit numbers that indicate, whenever possible, methylated positions, with 0 representing the 2’-OH group. Consequently, modifications Am, m1A, m6A, m5C, are indicated as 0A, 1A, 6A, 5C, respectively. Residues with several methylations list all methylated positions, sorted in ascending order, e.g., m1Am, m2,2,7G, m4,4Cm are indicated as 01A, 227G, 044C, respectively. Some other modifications also use single digits for convenience, e.g., I and /Psi1 are indicated as 9A and 9U. Other modifications are indicated with additional numbers, usually following the position of the modification. For example, i6A, io6A, k2C, are indicated as 61A, 60A, 21C, respectively. Some naming decisions, especially in case of very complex modifications, were arbitrary, keeping in mind that ambiguity in the numbering must always be avoided, i.e., that a given sequence of digits preceding a letter corresponds to a unique modification. For each modified nucleoside the code in new nomenclature is available on the nucleoside site in the Modifications section of the database. The nomenclature was also implemented in the Sequences section by providing the option to display modifications in sequences using this nomenclature instead of one-letter symbols. As the next step we intend to develop and provide format conversion tools to allow for exporting and importing RNA sequences with modifications, and e.g., to run sequence searches that take modifications into account.
tRNA sequences section update and development

For this database release, 102 new tRNA sequences were added, and a major update of mammalian tRNA modifications was performed based on (18). Among new sequences are sets of tRNAs from two bacterial species: *Streptomyces griseus* and *Lactococcus lactis* (19,20), 60 and 26 sequences, respectively. As technology and methods improve, well-studied sequences continue to undergo revision (21). We have also introduced links of human tRNA sequences to MINTbase (22), which is a framework for the interactive exploration of mitochondrial and nuclear tRNA fragments. The MINTbase link in MODOMICS opens a page with a list of the latest profile of expressed tRNA fragments, aligned against the sequence from MODOMICS.

Updated collection of proteins, enzymatic activities, and pathways

The MODOMICS collection of functionally characterized proteins involved in RNA modification is under constant development. Since the previous release, 66 new proteins have been added. The collections of protein sequences and enzymatic activities are updated in parallel, which resulted in 105 new enzymatic activities. Among new proteins that were added in this release there is a collection of human RNA modification enzymes, including: dimethyladenosine transferase TFB1M (23), tRNA pseudouridylate synthase PUS3 (24), tRNA m^5C methyltransferases NSUN3 (25) and NSUN6 (26), and tRNA m^5C methyltransferase NSUN4 (27). Apart from the addition of newly characterized enzymes, data entries for many enzymes and associated pathways were updated.

Future prospects

The number of experimentally identified modifications and RNA modifying enzymes keeps growing. New modified nucleosides are being discovered in particular in RNAs from recently adopted model systems, such as extremophilic prokaryotes. Though there is a considerable amount of information available about the enzymes responsible for introducing specific modifications, there are still many modified positions in well-characterized RNA molecules, for which the responsible enzymes are not known, e.g., m^5Am at the 5’ end of human mRNAs or m^3U, m^4C in 12S mitochondrial rRNA. To help us keep up with new discoveries, we encourage the users of MODOMICS to submit suggestions for additions to be included in the database. We also encourage developers of other computational resources to contact us to have our databases mutually linked to each other.

For the next release of MODOMICS, we plan to update the visualization options and to refurbish the website, to keep up with the changing trends in web design. We also intend to renew data structures, to make MODOMICS more compatible with other databases and web servers, to facilitate automated data exchange, and to introduce the ability to search sequences by taking modifications into account.

AVAILABILITY

The data are accessible freely for research purposes at http://modomics.genesilico.pl.

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