

Bioinformatics Toolbox™

Release Notes

How to Contact MathWorks



www.mathworks.com Web
comp.soft-sys.matlab Newsgroup
www.mathworks.com/contact_TS.html Technical Support



suggest@mathworks.com Product enhancement suggestions
bugs@mathworks.com Bug reports
doc@mathworks.com Documentation error reports
service@mathworks.com Order status, license renewals, passcodes
info@mathworks.com Sales, pricing, and general information



508-647-7000 (Phone)



508-647-7001 (Fax)



The MathWorks, Inc.
3 Apple Hill Drive
Natick, MA 01760-2098

For contact information about worldwide offices, see the MathWorks Web site.

Bioinformatics Toolbox™ Release Notes

© COPYRIGHT 2003–2014 by The MathWorks, Inc.

The software described in this document is furnished under a license agreement. The software may be used or copied only under the terms of the license agreement. No part of this manual may be photocopied or reproduced in any form without prior written consent from The MathWorks, Inc.

FEDERAL ACQUISITION: This provision applies to all acquisitions of the Program and Documentation by, for, or through the federal government of the United States. By accepting delivery of the Program or Documentation, the government hereby agrees that this software or documentation qualifies as commercial computer software or commercial computer software documentation as such terms are used or defined in FAR 12.212, DFARS Part 227.72, and DFARS 252.227-7014. Accordingly, the terms and conditions of this Agreement and only those rights specified in this Agreement, shall pertain to and govern the use, modification, reproduction, release, performance, display, and disclosure of the Program and Documentation by the federal government (or other entity acquiring for or through the federal government) and shall supersede any conflicting contractual terms or conditions. If this License fails to meet the government's needs or is inconsistent in any respect with federal procurement law, the government agrees to return the Program and Documentation, unused, to The MathWorks, Inc.

Trademarks

MATLAB and Simulink are registered trademarks of The MathWorks, Inc. See www.mathworks.com/trademarks for a list of additional trademarks. Other product or brand names may be trademarks or registered trademarks of their respective holders.

Patents

MathWorks products are protected by one or more U.S. patents. Please see www.mathworks.com/patents for more information.

R2014a

Functionality being changed or removed	2
--	---

R2013b

Functionality being changed or removed	4
--	---

R2013a

Saving to FASTQ, FASTA, SAM, and BAM files from a BioMap object	8
Sorting unordered BAM files using BioMap objects	8
Quality control plots for unmapped short-read data	8
Select and move behaviors in the Sequence Alignment app	8
Random access of annotation object data, for consistency with BioMap object data access	9
Functionality being changed	9

R2012b

Multiple reference sequences in BioMap objects	12
Mapping single and paired-end short read data to reference genomes	12
Increased performance when reading BAM files	13
Name changes for multialignviewer, phytreetool, and seqtool tools	13
Method removed from neighbor-joining method for phylogenetic tree construction	13
Functionality being changed or removed	14

R2012a

Update to Jmol Functions	18
Enhancements to Objects for NGS Data	18
Enhancements to the NGS Browser	19
Sequence Statistics Functions	19
Demo for DNA Methylation Analysis	20
Functionality Being Changed or Removed	20

R2011b

Visualizing and Investigating Short-Read Alignments and Feature Annotations in the NGS Browser	24
Objects for Genomic Feature Annotations	24
Enhancements to BioRead and BioMap Objects	24
Enhancements to the saminfo and baminfo Functions	26
Conversion of Error and Warning Message Identifiers ...	26
Function Elements Being Removed	27

R2011a

Data Format and Database Functions	30
Sequence Statistics Functions	30
Updates to the BioIndexedFile Class, Properties, and Methods	30
Updates to BioRead and BioMap Classes and Methods ...	31
Demos for High-Throughput Sequence Analysis	32
Support Vector Machine (SVM) Functions	32
Function Elements Being Removed	33

R2010b

Data Format and Database Functions	36
Sequence Conversion Functions	36
Sequence Statistics Functions	36
Pairwise Sequence Alignment Functions	38

Multiple Sequence Alignment Functions	38
Updates to BioMap Class, Methods, and Properties	39
Function Elements Being Removed	40

R2010a

Data Format and Database Functions	44
Pairwise Sequence Alignment Functions	44
Multiple Sequence Alignment Functions	44
Phylogenetic Tree Tools and Methods	45
BioIndexedFile Function, Object, Methods, and Properties	45
BioRead Function, Object, Methods, and Properties	46
BioMap Function, Object, Methods, and Properties	46
Function Elements Being Removed	46

R2009b

Data Format and Database Functions	50
Protein Analysis Functions	50
Data Visualization Functions	50
Sequence Statistics Functions	51
Sequence Utility Functions	51
Sequence Visualization Functions	52
Pairwise Sequence Alignment Functions	52
Multiple Sequence Alignment Functions	53
Phylogenetic Tree Tools and Methods	53
Clustergram Window	54
Clustergram Methods and Properties	54
HeatMap Object, Methods, and Properties	56
DataMatrix Methods	56
Microarray Functions, Objects, Methods, and Properties ..	57
Mass Spectrometry Functions	57
Demos for Sequence Analysis	57
Demos for Microarray Analysis	58

R2009a

Data Visualization Functions	60
Sequence Utility Functions	60
Sequence Conversion Functions	60
Bioanalytic and Mass Spectrometry Functions	60
Microarray Functions	61
Demo for Sequence Analysis	61

R2008b

Data Format and Database Functions	64
Sequence Utility Functions	65
Multiple Sequence Alignment Functions	66
Gene Ontology Functions	66
Protein Analysis Functions	66
Mass Spectrometry Functions	66
Microarray File Format Functions	67
Microarray Functions	68
DataMatrix Object	69
DataMatrix Methods	69
Demo for Visualization Tools	69
Demo for Sequence Analysis	69
Demos for Microarray Data Analysis	69

R2008a

Data Format and Database Functions	72
Sequence Utility Functions	73
Pairwise Sequence Alignment Functions	73
Phylogenetic Tree Tools Function	73
Protein Analysis Functions	74
Microarray File Format Functions	74
Microarray Functions	74
Object	77
Clustergram Methods	77
Demo for Sequence Analysis	77
Demo for Microarray Data Analysis	77
Demo for Visualization Tools	77

Demos for Mass Spectrometry Data Analysis	78
---	----

R2007b

Data Format and Database Functions	80
Microarray File Format Functions	80
Microarray Functions	80
Sequence Conversion, Utility, and Visualization Functions	81
Mass Spectrometry Functions	81
Statistical Learning Functions	81
Gene Ontology Methods	82
Demos for Microarray Data Analysis	82
Demos for Sequence Analysis	83
Demo for Graph Theory Analysis	83

R2007a+

Data Formats and Databases Functions	86
Microarray File Formats Functions	86
Microarray Utility Functions	87
Microarray Normalization and Filtering Functions	87
Mass Spectrometry Functions	87
Demos for Mass Spectrometry Functions	88

R2007a

Data Formats and Database Functions	90
Demo for Data Formats and Database Functions	90
Statistical Learning Functions	91
Protein Analysis and Sequence Utilities Functions	91
Demo for Protein Analysis and Sequence Utilities Functions	92
Sequence Alignment Functions	92
Demo for Sequence Alignment Functions	93
Microarray File Formats Functions	93

Microarray Normalization and Filtering Functions	93
Demo for Microarray File Formats, Normalization, and Filtering Functions	94
Microarray Data Analysis and Visualization Functions . . .	94
Demo for Microarray Data Analysis and Visualization Functions	94
Mass Spectrometry Functions	95
Phylogenetic Tree Tools Functions	95
Demos for Phylogenetic Tree Tools Functions	96
Phylogenetic Tree Methods	96

R2006b

Data Formats and Database Functions	98
Sequence Utilities Functions	98
Sequence Visualization Functions	98
Multiple Sequence Alignment Functions	99
Microarray File Formats	99
Microarray Data Analysis and Visualization Functions . . .	99
Graph Theory Functions	100
Graph Visualization Methods	100
Phylogenetic Tree Methods	101

R2006a+

Data Formats and Databases Functions	104
Sequence Utilities Functions	104
Sequence Visualization Functions	104
Statistical Learning Functions	104
Microarray Functions	105
Demo for Microarray Functions	105

R2006a

No New Features or Changes

R14SP3+

Multiple Sequence Alignment Viewer	110
Microarray Functions for Agilent Software	110
Gene Ontology Database Functions	110
Demo for Gene Ontology Functions	110

R14SP3

No New Features or Changes

R14SP2+

Sequence Alignment Functions	114
Sequence Statistics Functions	114
Sequence Utilities Functions	114
Phylogenetic Tree Functions	114
Phylogenetic Tree Methods	114
Microarray Functions	115
Statistics Functions	115

R14SP2

Updated RBASE Table	118
Expanded Bioperl Demonstration	118

R2014a

Version: 4.4

New Features: No

Bug Fixes: Yes

Functionality being changed or removed

Compatibility Considerations: Yes

Functionality	What Happens When You Use This Functionality?	Use This Instead	Compatibility Considerations
'R2012b' name-value pair input argument for the <code>seqalignviewer</code> function	Warns	The default version of <code>seqalignviewer</code> runs more robustly than the previous version (R2012b), and the default version is recommended to use. This name-value pair is intended only for customers who need the previous version.	See the Compatibility Considerations subheading in “Select and move behaviors in the Sequence Alignment app” on page 8.
<code>bowtierreadd</code> function	Errors	<ul style="list-style-type: none"> • If you have the original FASTQ (or FASTA) file, use the <code>bowtie</code> function (for UNIX® and Mac users only) to remap your files. This will create BAM files that are compatible with the toolbox. • If you have old BOWTIE files without the sequence files, you can read the files using <code>textscan</code>. 	When using other BOWTIE mapper/aligner programs, set appropriate option(s) to create either a SAM or BAM output file. Then use the <code>BioMap</code> object or the <code>samread</code> or <code>bamread</code> function to access the mapped short reads.

R2013b

Version: 4.3.1

New Features: No

Bug Fixes: Yes

Functionality being changed or removed

Compatibility Considerations: Yes

Functionality	What Happens When You Use This Functionality?	Use This Instead	Compatibility Considerations
Index name-value pair argument as input to the <code>bamread</code> function	Errors	—	Remove instances of the Index name-value pair argument. See the Compatibility Considerations subheading in “Increased performance when reading BAM files” on page 13.
'average' as a choice for the Method input argument to the <code>seqneighjoin</code> function	Errors	'equivar'	Replace instances of 'average' as an input to <code>seqneighjoin</code> with 'equivar'.
Changes to tool names: <ul style="list-style-type: none"> • <code>multialignviewer</code> • <code>phytreetool</code> • <code>seqtool</code> 	Errors	<ul style="list-style-type: none"> • <code>seqalignviewer</code> • <code>phytreeviewer</code> • <code>seqviewer</code> 	<ul style="list-style-type: none"> • Replace instances of <code>multialignviewer</code> with <code>seqalignviewer</code>. • Replace instances of <code>phytreetool</code> with <code>phytreeviewer</code>. • Replace instances of <code>seqtool</code> with <code>seqviewer</code>.
'natural' as a choice for the <i>Output</i> name-value pair input argument to these functions:	Errors	'linear'	Replace instances of 'natural' as the value of the <i>Output</i> name-value

Functionality	What Happens When You Use This Functionality?	Use This Instead	Compatibility Considerations
<ul style="list-style-type: none"> • affyrma • affygcma • rmasummary 			<p>pair input argument with 'linear' for these functions:</p> <ul style="list-style-type: none"> • affyrma • affygcma • rmasummary

R2013a

Version: 4.3

New Features: Yes

Bug Fixes: Yes

Saving to FASTQ, FASTA, SAM, and BAM files from a BioMap object

You can write the information of any BioRead/BioMap object to a file using the write function of the object.

Sorting unordered BAM files using BioMap objects

You can pass an unordered BAM file to a BioMap constructor, which then creates a new ordered file.

Quality control plots for unmapped short-read data

You can obtain quality control plots for short-read data using the plotSummary function of the BioRead object. The function creates a figure containing six plots that present summary statistics of the data stored in a FASTQ file.

In addition, you can use the BioReadQualityStatistics object to:

- Parse FASTQ files without creating a BioRead object.
- Interact with the quality data to compare different data sets or filtering options.
- Create customized plots.

Select and move behaviors in the Sequence Alignment app **Compatibility Considerations: Yes**

You can select a block from aligned sequences and move it horizontally if gaps are available.

Compatibility Considerations

To use the previous version of seqalignviewer, set the name-value pair argument 'R2012b' to true.

Random access of annotation object data, for consistency with BioMap object data access

You can have random access to data in GFFAnnotation and GTFAnnotation objects by using these functions:

- getSubset
- getData
- getIndex

Functionality being changed

Compatibility Considerations: Yes

Functionality	What Happens When You Use This Functionality?	Use This Instead	Compatibility Considerations
bowtierreadd function	Warns	bowtie function for UNIX and Mac users.	When using other BOWTIE mapper/aligner programs, set appropriate option(s) to create either a SAM or BAM output file. Then use the BioMap object or the samread or bamread function to access the mapped short reads.
'natural' as a choice for the <i>OutputValue</i> name-value pair input argument to these functions:	Warns	'linear'	Replace instances of 'natural' as the value of the <i>OutputValue</i> name-value

Functionality	What Happens When You Use This Functionality?	Use This Instead	Compatibility Considerations
<ul style="list-style-type: none"> • affyrma • affygrma • rmasummary 			pair input argument with 'linear' for these functions: <ul style="list-style-type: none"> • affyrma • affygrma • rmasummary
'R2012b' name-value pair input argument for the seqalignviewer function.	Still runs	—	See the Compatibility Considerations subheading in “Select and move behaviors in the Sequence Alignment app” on page 8.

R2012b

Version: 4.2

New Features: Yes

Bug Fixes: Yes

Multiple reference sequences in BioMap objects

Compatibility Considerations: Yes

You can now store information about short reads mapped to multiple references in a BioMap object. The new SequenceDictionary property contains the catalog of references available in a BioMap object.

Compatibility Considerations

For BioMap objects created using R2012b:

- The Reference property is now a cell string of length `obj.NSeqs`, for both BioMap objects with multiple references in the SequenceDictionary and objects with only one reference. For BioMap objects created before R2012b—which can only have a single reference—the Reference property is a string.
- BioMap methods that access data by genomic ranges now accept BioMap objects with multiple references. To use these methods, you must specify the reference or references to operate on. The affected methods are:
 - `getBaseCoverage`
 - `getCounts`
 - `getAlignment`
 - `getCompactAlignment`
 - `getIndex`

Mapping single and paired-end short read data to reference genomes

Two new functions generate an index and map short reads to a reference sequence using the Burrows-Wheeler transform.

- `bowtiebuild` builds index files using an input reference sequence.
- `bowtie` maps single and paired-end short reads to indexed reference files.

Note bowtiebuild and bowtie run on Mac and UNIX platforms only.

Increased performance when reading BAM files

Compatibility Considerations: Yes

The bamread function no longer requires the Index name-value pair argument to provide index information from a structure in the MATLAB® workspace. Indexing happens automatically without a decrease in performance.

Compatibility Considerations

The Index name-value pair argument as input to the bamread function will be removed in a future release. There is no need to replace it, only remove it.

Name changes for multialignviewer, phytreetool, and seqtool tools

Compatibility Considerations: Yes

Three tools in Bioinformatics Toolbox™ are renamed. The old names return a warning and will be removed in a future release.

Compatibility Considerations

- multialignviewer is being replaced with seqalignviewer.
- phytreetool is being replaced with phytreeviewer.
- seqtool is being replaced with seqviewer.

Method removed from neighbor-joining method for phylogenetic tree construction

Compatibility Considerations: Yes**Compatibility Considerations**

The choice of 'average' for the *Method* input argument to the `seqneighjoin` function warns and will be removed in a future release. It is replaced with 'equivar'.

Functionality being changed or removed**Compatibility Considerations: Yes**

Functionality	What Happens When You Use This Functionality?	Use This Instead	Compatibility Considerations
Index name-value pair argument as input to the <code>bamread</code> function	Warns	—	Remove instances of the Index name-value pair argument. See the Compatibility Considerations subheading in “Increased performance when reading BAM files” on page 13.
'average' as a choice for the <i>Method</i> input argument to the <code>seqneighjoin</code> function	Warns	'equivar'	Replace instances of 'average' as an input to <code>seqneighjoin</code> with 'equivar'

Functionality	What Happens When You Use This Functionality?	Use This Instead	Compatibility Considerations
<p>Changes to tool names:</p> <ul style="list-style-type: none"> • multialignviewer • phytreetool • seqtool 	Warns	<ul style="list-style-type: none"> • seqalignviewer • phytreeviewer • seqviewer 	<ul style="list-style-type: none"> • Replace instances of multialignviewer with seqalignviewer • Replace instances of phytreetool with phytreeviewer • Replace instances of seqtool with seqviewer
<p>'natural' as a choice for the <i>Output</i> name-value pair input argument to these functions:</p> <ul style="list-style-type: none"> • affyrma • affygrma • rmasummary 	Still runs	'linear'	<p>Replace instances of 'natural' as the value of the <i>Output</i> name-value pair input argument with 'linear' for these functions:</p> <ul style="list-style-type: none"> • affyrma • affygrma • rmasummary
<p>setName, getName, and getNSeqs methods of BioRead and BioMap objects</p>	Still runs	Dot notation	<p>Replace instances of: setName(<i>BioObj</i>, <i>name</i>) with: <i>BioObj</i>.Name = <i>name</i></p> <p>Replace instances of: getName(<i>BioObj</i>) with: <i>BioObj</i>.Name</p>

Functionality	What Happens When You Use This Functionality?	Use This Instead	Compatibility Considerations
			Replace instances of: <code>getNSeqs(<i>BioObj</i>)</code> with: <code><i>BioObj</i>.NSeqs</code>
isequalwithequalnans for DataMatrix object	Still runs	isequaln	Replace instances of isequalwithequalnans with isequaln
princomp for DataMatrix object	Still runs	pca	Replace instances of princomp with pca

R2012a

Version: 4.1

New Features: Yes

Bug Fixes: Yes

Update to Jmol Functions

The following functions are updated to use Version 12.0.5 of the Jmol molecule viewer:

- `evalrasmolscript` — Send RasMol script commands to Molecule Viewer window.
- `molviewer` — Display and manipulate 3-D molecule structure.

Enhancements to Objects for NGS Data

Compatibility Considerations: Yes

You now can construct and access information in a `BioMap` object (created from a BAM-formatted file) more efficiently. Filtering, binning, counting, and base-coverage calculation operations are now faster because source file scanning is no longer needed.

When using the `BioIndexedFile`, `BioRead`, or `BioMap` constructor to create an object from a FASTA-, FASTQ-, or SAM-formatted file, the source file no longer has a size limit of 4 GB.

Compatibility Considerations

The `BioRead` and `BioMap` constructors are changed as follows:

- When creating a `BioMap` object from a SAM- or BAM-formatted file containing multiple reference sequences, the `BioMap` constructor by default uses the first reference listed in the Dictionary of the source file.
- The following syntaxes, which take a `BioIndexedFile` object as an input, have been removed:

```
BioReadobj = BioRead(BioIFobj)
```

```
BioMapobj = BioMap(BioIFobj)
```

There is no longer a need to use this syntax, as you can create an indexed object directly from the SAM- or BAM-formatted source file. See [Representing Sequence and Quality Data in a BioRead Object](#) or [Representing Sequence, Quality, and Alignment/Mapping Data in a BioMap Object](#).

- The following syntaxes have been removed:

```
BioReadobj = BioRead('SAMFile', File)
```

```
BioReadobj = BioRead('FASTQFile', File)
```

Use this simpler syntax instead:

```
BioReadobj = BioRead(File)
```

- The following syntax has been removed:

```
BioMapobj = BioMap('SAMFile', File)
```

Use this simpler syntax instead:

```
BioMapobj = BioMap(File)
```

- The Indexed name-value pair argument as input to the `getSubset` method of the `BioRead` or `BioMap` class has been removed. Use the `InMemory` name-value pair argument instead.
- The `'SubsetRef'` name-value pair argument of the `BioMap` constructor has been removed. Use the `'SelectRef'` name-value pair argument instead.
- The `getCoverage` method of the `BioMap` class has been removed. Use the `getBaseCoverage`, `getCounts`, or `getIndex` method instead.

Enhancements to the NGS Browser

When you import short-read alignment data from a SAM- or BAM-formatted file into the NGS Browser:

- SAM-formatted files no longer have a size limit of 4 GB. Now, the size of both SAM- and BAM-formatted files is limited only by your operating system and available memory.
- The SAM- or BAM-formatted file can contain alignment data for multiple references. When importing short reads, you can select one reference sequence from those listed in the file header, or scan the file to see a list of the actual reference sequences and the aligned read count for each reference sequence.

Sequence Statistics Functions

Compatibility Considerations: Yes**Compatibility Considerations**

- The `aaccount` and `basecount` functions no longer accept the 'Others' name-value pair. Use the 'Ambiguous' or 'Gaps' name-value pair instead.
- The `aaccount` and `basecount` functions no longer accept the 'Structure' name-value pair. Use the 'Ambiguous' name-value pair with either 'ignore' or 'warn' instead.
- The `aaccount`, `basecount`, `codoncount`, and `dimercount` functions no longer include an `Others` field in the output structure. Use the `Ambiguous` field instead.

Demo for DNA Methylation Analysis

The following demo describes how to identify and compare potential cancer-related methylations at the base-pair level:

Exploring Genome-Wide Differences in DNA Methylation Profiles

Functionality Being Changed or Removed**Compatibility Considerations: Yes**

Functionality	What Happens When You Use This Functionality?	Use This Instead	Compatibility Considerations
SubsetRef name-value pair argument as input to the BioMap constructor function	Errors	'SelectRef' name-value pair argument	Replace instances of SubsetRef with SelectRef.
BioIndexedFile object as input to the BioRead or BioMap constructor function	Errors	A FASTQ-, SAM-, or BAM-formatted file	See the Compatibility Considerations subheading in “Enhancements to Objects for NGS Data” on page 18.

Functionality	What Happens When You Use This Functionality?	Use This Instead	Compatibility Considerations
'FASTQFile', <i>File</i> pair as input to the BioRead constructor	Errors	<i>File</i>	See the Compatibility Considerations subheading in “Enhancements to Objects for NGS Data” on page 18.
'SAMFile', <i>File</i> pair as input to the BioRead or BioMap constructor	Errors	<i>File</i>	See the Compatibility Considerations subheading in “Enhancements to Objects for NGS Data” on page 18.
Indexed name-value pair argument as input to the getSubset method of the BioRead or BioMap class	Errors	InMemory name-value pair argument	Replace instances of 'Indexed', false pair with 'InMemory', true pair.
getCoverage method of the BioMap class	Errors	getBaseCoverage, getCounts, or getIndex method	Replace all instances of getCoverage with getBaseCoverage, getCounts, or getIndex.
'Others' name-value pair as input to aaccount and basecount functions	Errors	'Ambiguous' or 'Gaps' name-value pair as input to aaccount and basecount functions	Replace instances of 'Others' with 'Ambiguous' or 'Gaps'.

Functionality	What Happens When You Use This Functionality?	Use This Instead	Compatibility Considerations
'Structure' name-value pair as input to <code>aaccount</code> and <code>basecount</code> functions	Errors	'Ambiguous' name-value pair with either 'ignore' or 'warn' as input to <code>aaccount</code> and <code>basecount</code> functions	Replace instances of 'Structure' with 'Ambiguous' paired with 'ignore' or 'warn'.
Others field from the output structure returned by <code>aaccount</code> , <code>basecount</code> , <code>codoncount</code> , or <code>dimercount</code> .	Errors	Ambiguous field	Replace instances of Others (as an input) with Ambiguous.

R2011b

Version: 4.0

New Features: Yes

Bug Fixes: Yes

Visualizing and Investigating Short-Read Alignments and Feature Annotations in the NGS Browser

The NGS Browser lets you visually verify and investigate the alignment of short-read sequences to a reference sequence. For more information, see [Visualizing and Investigating Short-Read Alignments and ngsbrowser](#).

Objects for Genomic Feature Annotations

Following are new classes for objects that contain genomic feature annotations for nucleotide sequences:

- `GFFAnnotation` — Contain data from GFF file.
- `GTFAnnotation` — Contain data from GTF file.

These classes have properties and methods that you can use to explore, access, filter, and manipulate all or a subset of the feature annotation data. For more information, see [Storing and Managing Feature Annotations in Objects](#).

Enhancements to BioRead and BioMap Objects

Compatibility Considerations: Yes

You can now construct a `BioMap` object from a BAM-formatted file.

When constructing these objects from source files, by default the data is indexed, which is more efficient for construction and data access. The `BioRead` and `BioMap` constructors now include an `IndexDir` name-value pair argument, which lets you specify the location of the index file.

You can still construct these objects with the data in memory, which lets you modify all the properties of the objects. The `BioRead` and `BioMap` constructors now include an `InMemory` name-value pair argument, which lets you construct the objects with the data in memory.

For details on the previous enhancements, see [Storing and Managing Short-Read Sequence Data in Objects](#).

Compatibility Considerations

The BioRead and BioMap constructors are changed as follows:

- The following syntaxes that take a `BioIndexedFile` object as an input will be removed in a future release:

```
BioReadobj = BioRead(BioIFobj)
```

```
BioMapobj = BioMap(BioIFobj)
```

There is no longer a need to use this syntax, as you can create an indexed object directly from the SAM- or BAM-formatted source file. See [Representing Sequence and Quality Data in a BioRead Object](#) or [Representing Sequence, Quality, and Alignment/Mapping Data in a BioMap Object](#).

- The following syntaxes will be removed in a future release:

```
BioReadobj = BioRead('SAMFile', File)
```

```
BioReadobj = BioRead('FASTQFile', File)
```

Use this syntax instead:

```
BioReadobj = BioRead(File)
```

- The following syntax will be removed in a future release:

```
BioMapobj = BioMap('SAMFile', File)
```

Use this syntax instead:

```
BioMapobj = BioMap(File)
```

- The Indexed name-value pair argument as input to the `getSubset` method of the `BioRead` or `BioMap` class will be removed in a future release. Use the `InMemory` name-value pair argument instead.
- The `'SubsetRef'` name-value pair argument of the `BioMap` constructor will be removed in a future release. Use the `'SelectRef'` name-value pair argument instead.
- If you use the `getSubset` method of a `BioRead` or `BioMap` object, and specify the same element more than once, the method errors, even if the object is in memory.

Enhancements to the saminfo and baminfo Functions

Compatibility Considerations: Yes

The saminfo and baminfo functions now include a ScanDictionary name-value pair argument, which controls the return of the reference names and the number of reads aligned to each reference from a SAM- or BAM-formatted file in new fields, ScannedDictionary and ScannedDictionaryCount. This information is needed when constructing a BioMap object from a file with multiple reference sequences. For more information, see Constructing a BioMap Object from a SAM- or BAM-Formatted File.

Compatibility Considerations

The Reference field is no longer returned in the output structure for baminfo. The ScannedDictionary field now includes names of the reference sequences.

Conversion of Error and Warning Message Identifiers

Compatibility Considerations: Yes

For R2011b, some error and warning message identifiers have changed in Bioinformatics Toolbox.

Compatibility Considerations

If you have scripts or functions that use message identifiers that changed, you must update the code to use the new identifiers. Typically, message identifiers are used to turn off specific warning messages, or in code that uses a try/catch statement and performs an action based on a specific error identifier.

For example, the Bioinfo:nwalign:InvalidScoringMatrix identifier has changed to bioinfo:nwalign:InvalidScoringMatrix. If your code checks for Bioinfo:nwalign:InvalidScoringMatrix, you must update it to check for bioinfo:nwalign:InvalidScoringMatrix instead.

To determine the identifier for a warning, run the following command just after you see the warning:

```
[MSG,MSGID] = lastwarn;
```

The preceding command saves the message identifier to the variable MSGID.

To determine the identifier for an error, run the following command just after you see the error:

```
exception = MException.last;
MSGID = exception.identifier;
```

Note Warning messages indicate a potential issue with your code. While you can turn off a warning, a suggested alternative is to change your code so it runs warning-free.

Function Elements Being Removed

Compatibility Considerations: Yes

Function Element Name	What Happens When You Use This Function Element	Use This Instead	Compatibility Considerations
'SubsetRef' name-value pair argument as input to BioMap constructor function	Warns	'SelectRef' name-value pair argument	See the Compatibility Considerations subheading in “Enhancements to BioRead and BioMap Objects” on page 24.
BioIndexedFile object as input to the BioRead or BioMap constructor function	Warns	A FASTQ-, SAM-, or BAM-formatted file	See the Compatibility Considerations subheading in “Enhancements to BioRead and BioMap Objects” on page 24.
'FASTQFile', <i>File</i> pair as input to the BioRead constructor	Warns	<i>File</i>	See the Compatibility Considerations subheading in “Enhancements to BioRead and BioMap Objects” on page 24

Function Element Name	What Happens When You Use This Function Element	Use This Instead	Compatibility Considerations
'SAMFile', <i>File</i> pair as input to the BioRead or BioMap constructor	Warns	<i>File</i>	See the Compatibility Considerations subheading in “Enhancements to BioRead and BioMap Objects” on page 24.
Indexed name-value pair argument as input to getSubset method of the BioRead or BioMap class	Warns	InMemory name-value pair argument	See the Compatibility Considerations subheading in “Enhancements to BioRead and BioMap Objects” on page 24.
Reference field of structure returned by baminfo	Errors	ScannedDictionary field	See the Compatibility Considerations subheading in “Enhancements to the saminfo and baminfo Functions” on page 26.

R2011a

Version: 3.7

New Features: Yes

Bug Fixes: Yes

Data Format and Database Functions

The following functions have a new field, `FilePath`, in their output structure:

- `fastainfo` — Return information about FASTA file.
- `fastqinfo` — Return information about FASTQ file.
- `saminfo` — Return information about Sequence Alignment/Map (SAM) file.

The `fastainfo` function has two additional fields in its output structure: `Header` and `Length`.

Sequence Statistics Functions

Compatibility Considerations: Yes

Compatibility Considerations

In Bioinformatics Toolbox Version 3.6, the `aaccount` and `basecount` functions still allowed 'Others' and 'Structure' name-value pairs, but displayed a warning.

In Bioinformatics Toolbox Version 3.7, the `aaccount` and `basecount` functions do not allow 'Others' and 'Structure' name-value pairs, and return an error if you use them. Now you must use the 'Ambiguous' and 'Gaps' name-value pairs, which specify whether to count or ignore ambiguous characters and gaps, as well as specify how to count ambiguous characters, and whether to display a warning.

Updates to the `BioIndexedFile` Class, Properties, and Methods

The following name-value pairs of the `BioIndexedFile` constructor function are renamed:

- `MapKeys` is now `IndexedByKey`.
- `MemMapIndex` is now `MemoryMappedIndex`.

Note The former name-value pairs are still valid for Bioinformatics Toolbox Version 3.7 (R2011a).

The `MemoryMappedIndex` property of the `BioIndexedFile` class is now editable, which lets you load and unload file indices in memory.

The `BioIndexedFile` class includes the following new methods:

- `getDictionary` — Retrieve reference sequence names from SAM-formatted source file associated with `BioIndexedFile` object.
- `getSubset` — Create object containing subset of elements from `BioIndexedFile` object.

Updates to BioRead and BioMap Classes and Methods

Compatibility Considerations: Yes

The `BioMap` constructor includes a new name-value pair, `SubsetRef`, which lets you specify one reference sequence in the input argument (`BioIndexedFile` object, SAM-formatted file, or structure) when constructing the `BioMap` object.

The following method of the `BioRead` and `BioMap` classes is updated:

`getSubset` — Create object containing subset of elements from object. Updated with addition of the `Indexed` name-value pair, which lets you use the `BioIndexedFile` object when creating a new object, thus saving memory. This name-value pair is ignored if your `BioRead` or `BioMap` object was not created from a `BioIndexedFile` object.

Following are new methods of the `BioMap` class:

- `getBaseCoverage` — Return base-by-base alignment coverage of reference sequence in `BioMap` object.
- `getCounts` — Return count of read sequences aligned to reference sequence in `BioMap` object.

- `getIndex` — Return indices of read sequences aligned to reference sequence in `BioMap` object.

The `getCoverage` method of the `BioMap` class is being removed in a future release. Use the `getBaseCoverage`, `getCounts`, and `getIndex` methods instead.

Compatibility Considerations

In Bioinformatics Toolbox Version 3.6 and earlier, the `BioMap` class included a `getCoverage` method, which computes read coverage in a `BioMap` object.

In Bioinformatics Toolbox Version 3.7, the `getCoverage` method still runs, but displays a warning. Now use the `getBaseCoverage`, `getCounts`, and `getIndex` methods of the `BioMap` class.

Demos for High-Throughput Sequence Analysis

Following are two new high-throughput sequence analysis demos:

- Exploring Protein-DNA Binding Sites from Paired-End ChIP-Seq Data
- Identifying Differentially Expressed Genes from RNA-Seq Data

Support Vector Machine (SVM) Functions

Compatibility Considerations: Yes

The functionality of the `svmsmset` function is incorporated into the `svmtrain` and `statset` functions. Although `svmsmset` is still valid, it is no longer documented.

The `svmtrain` function has been updated:

- The function can now handle NaN values in the training matrix input and performs more checks of parameters you supply.
- The function now includes Sequential Minimal Optimization (SMO) functionality plus four new name-value pairs: `kernelcachelimit`, `kktviolationlevel`, `options`, and `tolkkt`.

- The default training method is SMO, even if you have Optimization Toolbox™ installed.
- The QuadProg_Opts and SMO_Opts name-value pairs have been replaced by the options name-value pair. Although the former name-value pairs are still valid, the recommended ways to perform quadratic programming (QP) training and SMO training are summarized in the following bullets.
- The recommended way to include QP options for svmtrain is to use the QP training method and use the new options name-value pair. For the options value, use a structure you create with optimset.
- The recommended way to include SMO options for svmtrain is to use the default SMO training method and use the new kernelcachelimit, kktviolationlevel, options, and tolkkt name-value pairs. For the options value, use a structure you create with the statset function and its Display and MaxIter name-value pairs.

Compatibility Considerations

In Bioinformatics Toolbox Version 3.6 and earlier, if you had Optimization Toolbox installed, QP was the default training method for the svmtrain function. Now the default training method is SMO.

Function Elements Being Removed

Compatibility Considerations: Yes

Function Element Name	What Happens When You Use This Function Element	Use This Instead	Compatibility Considerations
'Others' name-value pair as input to aaccount and basecount functions	Errors	'Ambiguous' or 'Gaps' name-value pair as input to aaccount and basecount functions	See the Compatibility Considerations subheading in “Sequence Statistics Functions” on page 30.
'Structure' name-value pair as	Errors	'Ambiguous' name-value pair with either 'ignore' or 'warn' as input	See the Compatibility Considerations subheading in

Function Element Name	What Happens When You Use This Function Element	Use This Instead	Compatibility Considerations
input to aaccount and basecount functions		to aaccount and basecount functions	“Sequence Statistics Functions” on page 30.
getCoverage method of BioMap class	Warns	getBaseCoverage, getCounts, and getIndex methods	See the Compatibility Considerations subheading in “Updates to BioRead and BioMap Classes and Methods” on page 31.
svmsmoset function	Still runs	svmtrain and statset functions	svmsmoset is not recommended. Use svmtrain and statset instead.

R2010b

Version: 3.6

New Features: Yes

Bug Fixes: Yes

Data Format and Database Functions

The following new functions allow indexed file access to BAM-formatted files:

- `bamindexread` — Read Binary Sequence Alignment/Map Index (BAI) file.
- `baminfo` — Return information about Binary Sequence Alignment/Map (BAM) file.
- `bamread` — Read data from Binary Sequence Alignment/Map (BAM) file.

The following new functions let you read Bowtie- and SOAP-formatted files:

- `bowtieread` — Read data from Bowtie file.
- `soapread` — Read data from Short Oligonucleotide Analysis Package (SOAP) file.

Sequence Conversion Functions

The following new functions support CIGAR strings for sequence mapping and alignment:

- `align2cigar` — Convert aligned sequences to corresponding Compact Idiosyncratic Gapped Alignment Report (CIGAR) format strings.
- `cigar2align` — Convert unaligned sequences to aligned sequences using Compact Idiosyncratic Gapped Alignment Report (CIGAR) format strings

Sequence Statistics Functions

Compatibility Considerations: Yes

The following functions are updated:

- `aaccount` — Count amino acids in sequence. Updated by adding the `Ambiguous` property, which lets you specify how to count ambiguous amino acid characters. Updated by adding the `Gaps` property, which lets you specify to count or ignore gaps. The `Others` and `Structure` properties still work, but display a warning, indicating that they will be invalid in future versions of Bioinformatics Toolbox. The `Others` field in the output structure is replaced by the `Ambiguous` field.

- **basecount** — Count nucleotides in sequence. Updated by adding the **Ambiguous** property, which lets you specify how to count ambiguous nucleotide characters. Updated by adding the **Gaps** property, which lets you specify to count or ignore gaps. The **Others** and **Structure** properties still work, but display a warning, indicating that they will be invalid in future versions of Bioinformatics Toolbox. The **Others** field in the output structure is replaced by the **Ambiguous** field.
- **codonbias** — Calculate codon frequency for each amino acid coded for in nucleotide sequence. Updated by adding the **Ambiguous** property, which lets you specify how to count codons containing ambiguous nucleotide characters.
- **codoncount** — Count codons in nucleotide sequence. Updated by adding the **Ambiguous** property, which lets you specify how to count codons containing ambiguous nucleotide characters. Updated by adding the **GeneticCode** property, which lets you overlay a grid that groups the synonymous codons on the heat map of the codon counts. The **Others** field in the output structure is replaced by the **Ambiguous** field.
- **dimercount** — Count dimers in nucleotide sequence. Updated by adding the **Ambiguous** property, which lets you specify how to count dimers containing ambiguous nucleotide characters. The **Others** field in the output structure is replaced by the **Ambiguous** field.

Compatibility Considerations

In Bioinformatics Toolbox Version 3.5 and earlier, the **aaccount** and **basecount** functions included 'Others' and 'Structure' property name/property value pairs, which let you specify how to count ambiguous characters and gaps, and whether to display a warning. These functions also returned a structure with an **Others** field.

In Bioinformatics Toolbox Version 3.6, the **aaccount** and **basecount** functions still allow 'Others' and 'Structure' property name/property value pairs, but display a warning. Now the **aaccount** and **basecount** functions include the 'Ambiguous' and 'Gaps' property name/property value pairs, which specify whether to count or ignore ambiguous characters and gaps, as well as specify how to count ambiguous characters, and whether to display a warning. These functions now return a structure with an **Ambiguous** field, which replaces the **Others** field.

In Bioinformatics Toolbox Version 3.6, the `codoncount` and `dimercount` functions return a structure with an optional `Ambiguous` field, which replaces the `Others` field.

Pairwise Sequence Alignment Functions

The following function is updated:

- `nwalign` — Globally align two sequences using Needleman-Wunsch algorithm. Updated to support semiglobal or “glocal” alignments by addition of `Glocal` property.

Multiple Sequence Alignment Functions

Compatibility Considerations: Yes

The following new functions support CIGAR strings for sequence mapping and alignment:

- `align2cigar` — Convert aligned sequences to corresponding Compact Idiosyncratic Gapped Alignment Report (CIGAR) format strings.
- `cigar2align` — Convert unaligned sequences to aligned sequences using Compact Idiosyncratic Gapped Alignment Report (CIGAR) format strings

The following functions are updated:

- `multialign` — Align multiple sequences using progressive method. Updated to include a new property, `'UseParallel'`, which lets you use `parfor`-loops and compute in parallel mode.
- `seqpdist` — Calculate pairwise distance between sequences. Updated to include a new property, `'UseParallel'`, which lets you use `parfor`-loops and compute in parallel mode.

Compatibility Considerations

In Bioinformatics Toolbox Version 3.4 and earlier, the `multialign` and `seqpdist` functions included `'JobManager'` and `'WaitInQueue'` property

name/property value pairs, which let you process in parallel, including support for the MATLAB scheduler for clusters.

In Bioinformatics Toolbox Version 3.5, the `multialign` and `seqpdist` functions allowed the 'JobManager' and 'WaitInQueue' property name/property value pairs, but displayed a warning.

In Bioinformatics Toolbox Version 3.6, the `multialign` and `seqpdist` functions error if you use the 'JobManager' or 'WaitInQueue' property name/property value pair. Instead they include the 'UseParallel' property name/property value pair, which lets you process in parallel, including support for:

- Local workers for multicore machines
- The MATLAB scheduler for clusters
- Third-party schedulers for clusters

Updates to BioMap Class, Methods, and Properties

You can now create a `BioMap` object from a MATLAB structure containing sequence and alignment information, returned by the `bamread` function.

The following method of the `BioMap` class is updated:

`getCoverage` — Compute read coverage in `BioMap` object. Updated to return the coverage of multiple regions of the reference sequence.

The `BioMap` class includes the following new methods:

- `getCompactAlignment` — Construct compact alignment represented in `BioMap` object.
- `getMatePosition` — Retrieve mate positions of read sequences from `BioMap` object.
- `setMatePosition` — Set mate positions of read sequences in `BioMap` object.

The `BioMap` class includes the following new property:

- `MatePosition` — Positions of the mates for all read sequences represented in the `BioMap` object.

Function Elements Being Removed

Compatibility Considerations: Yes

Function Element Name	What Happens When You Use This Function Element	Use This Instead	Compatibility Considerations
'Others' property name/property value pair as input to <code>aaccount</code> and <code>basecount</code> functions	Warns	'Ambiguous' or 'Gaps' property name/property value pair as input to <code>aaccount</code> and <code>basecount</code> functions	See the Compatibility Considerations subheading in “Sequence Statistics Functions” on page 36.
'Structure' property name/property value pair as input to <code>aaccount</code> and <code>basecount</code> functions	Warns	'Ambiguous' property name/property value pair with either 'ignore' or 'warn' as input to <code>aaccount</code> and <code>basecount</code> functions	See the Compatibility Considerations subheading in “Sequence Statistics Functions” on page 36.
'JobManager' property name/property value pair as input to <code>multialign</code> and <code>seqpdist</code> functions	Errors	'UseParallel' property name/property value pair as input to <code>multialign</code> and <code>seqpdist</code> functions	See the Compatibility Considerations subheading in “Multiple Sequence Alignment Functions” on page 38.
'WaitInQueue' property name/property value pair as input to <code>multialign</code> and <code>seqpdist</code> functions	Errors	'UseParallel' property name/property value pair as input to <code>multialign</code> and <code>seqpdist</code> functions	See the Compatibility Considerations subheading in “Multiple Sequence Alignment Functions” on page 38.

Function Element Name	What Happens When You Use This Function Element	Use This Instead	Compatibility Considerations
<p>The following properties of a clustergram object:</p> <ul style="list-style-type: none"> • ColumnMarker • Impute • Ratio • RowMarker • SymmetricRange 	Errors	<p>New properties of a clustergram object:</p> <ul style="list-style-type: none"> • ColumnGroupMarker • ImputeFun • DisplayRatio • RowGroupMarker • Symmetric 	See “Clustergram Methods and Properties” on page 54.
'Dimension' property name/property value pair as input to clustergram function	Errors	'Cluster' property name/property value pair as input to clustergram function	See the Compatibility Considerations subheading in “Microarray Functions” on page 74.
'Pdist' property name/property value pair as input to clustergram function	Errors	Either 'RowPdist' or 'ColumnPdist' property name/property value pair as input to clustergram function	See the Compatibility Considerations subheading in “Microarray Functions” on page 74.
pdbplot function	Errors	molviewer function	See the Compatibility Considerations subheading in “Protein Analysis and Sequence Utilities Functions” on page 91.

Function Element Name	What Happens When You Use This Function Element	Use This Instead	Compatibility Considerations
getpir and pirread functions	Errors	Use getembl, getgenpept, and getpdb to retrieve protein sequences from Web databases. Use emblread, genpeptread, and pdbread to read protein sequence data.	See “Data Formats and Databases Functions” on page 104.
mamadnorm and mameannorm functions	Errors	manorm function	

R2010a

Version: 3.5

New Features: Yes

Bug Fixes: Yes

Data Format and Database Functions

The following functions are new:

- `saminfo` — Return information about Sequence Alignment/Map (SAM) file.
- `samread` — Read data from Sequence Alignment/Map (SAM) file.

The following functions are updated:

- `fastaread` — Read data from FASTA file. Updated to allow trimming of the headers in the output structure by addition of `TrimHeaders` property.
- `fastqread` — Read data from FASTQ file. Updated to allow trimming of the headers in the output structure by addition of `TrimHeaders` property.
- `phytreeread` — Read phylogenetic tree file. Updated to return a second output containing bootstrap values for tree nodes.

Pairwise Sequence Alignment Functions

The following function is updated:

- `fastaread` — Read data from FASTA file. Updated to allow trimming of the headers in the output structure by addition of `TrimHeaders` property.

Multiple Sequence Alignment Functions

Compatibility Considerations: Yes

The following functions are updated:

- `fastaread` — Read data from FASTA file. Updated to allow trimming of the headers in the output structure by addition of `TrimHeaders` property.
- `multialign` — Align multiple sequences using progressive method. Updated to include a new property, `'UseParallel'`, which lets you use `parfor`-loops and compute in parallel mode.
- `seqpdist` — Calculate pairwise distance between sequences. Updated to include a new property, `'UseParallel'`, which lets you use `parfor`-loops and compute in parallel mode.

Compatibility Considerations

In Bioinformatics Toolbox Version 3.4 and earlier, the `multialign` and `seqpdist` functions included 'JobManager' and 'WaitInQueue' property name/property value pairs, which let you process in parallel, including support for the MATLAB scheduler for clusters.

In Bioinformatics Toolbox Version 3.5, the `multialign` and `seqpdist` functions do not include the 'JobManager' and 'WaitInQueue' property name/property value pairs. Instead they include the 'UseParallel' property name/property value pair, which lets you process in parallel, including support for:

- Local workers for multicore machines
- The MATLAB scheduler for clusters
- Third-party schedulers for clusters

Phylogenetic Tree Tools and Methods

The following functions are updated:

- `phytreeread` — Read phylogenetic tree file. Updated to return a second output containing bootstrap values for tree nodes.
- `seqpdist` — Calculate pairwise distance between sequences. Updated to include a new property, 'UseParallel', which lets you use `parfor`-loops and compute in parallel mode.

BioIndexedFile Function, Object, Methods, and Properties

Following is a new class for an object that lets you extract information from large multi-entry text files.

- `BioIndexedFile` — Allow quick and efficient access to large text file with nonuniform-size entries.

This class has properties and methods that are useful for accessing, reading, and parsing data from a large source file.

BioRead Function, Object, Methods, and Properties

Following is a new class for an object that contains data from short-read sequences, including sequence headers, nucleotide sequences, and the quality scores for the sequences.

- BioRead — Contain sequence and quality data.

This class has properties and methods that you can use to explore, access, filter, and manipulate all or a subset of the data, before doing subsequent analyses or sequence alignment and mapping.

BioMap Function, Object, Methods, and Properties

Following is a new class for an object that contains data from short-read sequences, including sequence headers, read sequences, quality scores for the sequences, and data about alignment and mapping to a single reference sequence.

- BioMap — Contain sequence, quality, alignment, and mapping data.

This class has properties and methods that you can use to explore, access, filter, and manipulate all or a subset of the data, before doing subsequent analyses or viewing the data.

Function Elements Being Removed

Compatibility Considerations: Yes

Function Element Name	What Happens When You Use This Function Element	Use This Instead	Compatibility Considerations
'JobManager' property name/property value pair as input to multialign	Warns	'UseParallel' property name/property value pair as input to multialign	See the Compatibility Considerations subheading in “Multiple Sequence

Function Element Name	What Happens When You Use This Function Element	Use This Instead	Compatibility Considerations
and seqpdist functions		and seqpdist functions	Alignment Functions” on page 44
'WaitInQueue' property name/property value pair as input to multialign and seqpdist functions	Warns	'UseParallel' property name/property value pair as input to multialign and seqpdist functions	See the Compatibility Considerations subheading in “Multiple Sequence Alignment Functions” on page 44.

R2009b

Version: 3.4

New Features: Yes

Bug Fixes: Yes

Data Format and Database Functions

Following are new functions:

- `fastainfo` — Return information about FASTA file.
- `fastqinfo` — Return information about FASTQ file.
- `fastqread` — Read data from FASTQ file.
- `fastqwrite` — Write to file using FASTQ format.
- `sffinfo` — Return information about SFF file.
- `sffread` — Read data from SFF file.
- `tgspcinfo` — Return information about SPC file.
- `tgspcread` — Read data from SPC file.

The following functions are updated:

- `affyread` — Read microarray data from Affymetrix® GeneChip® file. Updated to read cell layout files (CLF) and background probe (BGP) files.
- `multialignwrite` — Write multiple alignment to file. Updated to write a file in either ClustalW ALN format (default) or MSF format.

Protein Analysis Functions

Following is a new function:

- `isotopicdist` — Calculate high-resolution isotope mass distribution and density function.

The following function is updated:

- `cleave` — Cleave amino acid sequence with enzyme. Updated to let you specify an exception to the enzyme's cleavage rule and to let you specify a maximum number of missed cleavage sites. Also updated to return the number of missed cleavage sites per peptide fragment.

Data Visualization Functions

Compatibility Considerations: Yes

The following functions are updated:

- `microplateplot` — Display visualization of microtiter plate. Display updated so that first row of input matrix appears at the top and is labeled row A. Updated to return the handle to the axes of the plot, which lets you reverse the order or the rows or columns in the display. Updated to include a new property, 'TextFontSize', which lets you control the font size of text labels.
- `multialignviewer` — Display and interactively adjust multiple sequence alignment. Updated to accept a list of names to label the sequences in the Multiple Sequence Alignment Viewer window.
- `showalignment` — Display color-coded sequence alignment. Updated to control the inclusion or exclusion of terminal gaps from the count of matches and similar residues when displaying a pairwise alignment.

Compatibility Considerations

In Bioinformatics Toolbox Version 3.3, the default layout for the plot returned by `microplateplot` displayed the first row of the input matrix at the bottom.

In Bioinformatics Toolbox Version 3.4, the plot displays the first row of the input matrix at the top.

Sequence Statistics Functions

The following function is updated:

- `seqshowwords` — Graphically display words in sequence. Updated to search for multiple words in a sequence.

Sequence Utility Functions

The following functions are updated:

- `cleave` — Cleave amino acid sequence with enzyme. Updated to let you specify an exception to the enzyme's cleavage rule and to let you specify a maximum number of missed cleavage sites. Also updated to return the number of missed cleavage sites per peptide fragment.
- `rebasecuts` — Find restriction enzymes that cut nucleotide sequence. Updated to use Version 904 of REBASE[®], the Restriction Enzyme Database.
- `restrict` — Split nucleotide sequence at restriction site. Updated to use Version 904 of REBASE, the Restriction Enzyme Database.

Sequence Visualization Functions

The following functions are updated:

- `multialignviewer` — Display and interactively adjust multiple sequence alignment. Updated to accept a list of names to label the sequences in the Multiple Sequence Alignment Viewer window.
- `showalignment` — Display color-coded sequence alignment. Updated to control the inclusion or exclusion of terminal gaps from the count of matches and similar residues when displaying a pairwise alignment.

Pairwise Sequence Alignment Functions

Following is a new function:

- `localalign` — Return local optimal and suboptimal alignments between two sequences.

The following functions are updated:

- `multialignviewer` — Display and interactively adjust multiple sequence alignment. Updated to accept a list of names to label the sequences in the Multiple Sequence Alignment Viewer window.
- `showalignment` — Display color-coded sequence alignment. Updated to control the inclusion or exclusion of terminal gaps from the count of matches and similar residues when displaying a pairwise alignment.

Multiple Sequence Alignment Functions

The following functions are updated:

- `multialignviewer` — Display and interactively adjust multiple sequence alignment. Updated to accept a list of names to label the sequences in the Multiple Sequence Alignment Viewer window.
- `multialignwrite` — Write multiple alignment to file. Updated to write a file in either ClustalW ALN format (default) or MSF format.
- `showalignment` — Display color-coded sequence alignment. Updated to control the inclusion or exclusion of terminal gaps from the count of matches and similar residues when displaying a pairwise alignment.

Phylogenetic Tree Tools and Methods

Compatibility Considerations: Yes

The Phylogenetic Tree Tool includes the following updates:

- Includes two new circular print renderings: equal angle and equal daylight
- Updates to **Tools** menu, including commands to select specific branch and leaf nodes based on different criteria, such as distance, common ancestors, leaves only, and descendants.

Following is a new method:

- `cluster` — Validate clusters in phylogenetic tree.

The following method is updated:

- `plot` — Draw phylogenetic tree. Updated to include two new algorithms for circular layouts: equal angle and equal daylight. Updated to let you rotate circular trees from 0 through 360 degrees and to rotate leaf labels of circular trees so that the text is aligned to the root node. Updated the 'LeafLabels' property so that it defaults to `true` for circular layouts and to `false` for square and angular layouts.



Compatibility Considerations

In Bioinformatics Toolbox Version 3.3, the 'LeafLabels' property defaulted to true when the 'Type' property was 'square' or 'angular', and to false when the 'Type' property was 'radial'.

In Bioinformatics Toolbox Version 3.4, the 'LeafLabels' property defaults to false when the 'Type' property is 'square' or 'angular', and to true when the 'Type' property is 'radial'.

Clustergram Window

The Clustergram window has two new toolbar buttons:

- Annotate  button — Shows and hides intensity values for each area of the heat map.
- Show Dendrogram  button — Shows and hides the dendrograms.

Clustergram Methods and Properties

Compatibility Considerations: Yes

The following are new methods of a clustergram object:

- `addTitle` — Add title to clustergram.
- `addXLabel` — Label x -axis of clustergram.
- `addYLabel` — Label y -axis of clustergram.
- `clusterGroup` — Select cluster group.

The following properties of a clustergram object are renamed:

- `ColumnMarker` is now `ColumnGroupMarker`.
- `Impute` is now `ImputeFun`.
- `Ratio` is now `DisplayRatio`.
- `RowMarker` is now `RowGroupMarker`.

- `SymmetricRange` is now `Symmetric`.

Note The former property names are still valid for Bioinformatics Toolbox version 3.4 (R2009b).

Following is a new property related to the display of dendrogram tree diagrams in a clustergram object:

- `ShowDendrogram`

The following are new properties related to the display of row and column labels of a clustergram object:

- `RowLabels`
- `ColumnLabels`
- `RowLabelsLocation`
- `ColumnLabelsLocation`
- `RowLabelsColor`
- `ColumnLabelsColor`
- `LabelsWithMarkers`
- `RowLabelsRotate`
- `ColumnLabelsRotate`

The following are new properties related to annotating data in a clustergram object:

- `Annotate`
- `AnnotColor`
- `AnnotPrecision`

When using clustergram properties with the `get` and `set` methods, the property names are now case sensitive.

Compatibility Considerations

In Bioinformatics Toolbox Version 3.3, the property names of a clustergram object were not case sensitive when used with the `get` and `set` methods.

In Bioinformatics Toolbox Version 3.4, property names of a clustergram object are case sensitive.

HeatMap Object, Methods, and Properties

Following is a new object:

- `HeatMap` object — Object containing matrix and heat map display properties.

The following are methods of a `HeatMap` object:

- `addTitle` — Add title to heat map.
- `addXLabel` — Label x -axis of heat map.
- `addYLabel` — Label y -axis of heat map.
- `plot` — Render heat map for object.
- `view` — Render heat map for object.

A `HeatMap` object includes many properties that control the creation of the heat map, row and column labels, axes labels, title, and data annotation.

DataMatrix Methods

Following is a new method of a `DataMatrix` object:

- `dmwrite` — Write `DataMatrix` object to text file.

Microarray Functions, Objects, Methods, and Properties

Following are new functions to create objects containing data from a microarray gene expression experiment:

- `bioma.ExpressionSet` — Contain data from microarray gene expression experiment.
- `bioma.data.ExptData` — Contain expression data from microarray gene expression experiment.
- `bioma.data.MetaData` — Contain sample or feature metadata from microarray gene expression experiment.
- `bioma.data.MIAME` — Contain experiment information from microarray gene expression experiment.

These objects have properties and methods that are useful for viewing and analyzing the data or a subset of the data.

Mass Spectrometry Functions

Following are new functions:

- `isotopicdist` — Calculate high-resolution isotope mass distribution and density function.
- `tgspcinfo` — Return information about SPC file.
- `tgspread` — Read data from SPC file.

The following function is updated:

- `mspeaks` — Convert raw peak data to peak list (centroided data). Updated to include a new property, 'Style', which lets you specify the style for marking the peaks in the plot.

Demos for Sequence Analysis

Following are two new sequence analysis demos:

- Working with SFF Files from the 454 Genome Sequencer FLX System
- Working with Illumina/Solexa Next-Generation Sequencing Data

Demos for Microarray Analysis

Following are two new microarray analysis demos:

- Working with Objects for Microarray Experiment Data
- Analyzing Illumina Bead Summary Gene Expression Data

R2009a

Version: 3.3

New Features: Yes

Bug Fixes: Yes

Data Visualization Functions

Following is a new function:

- `microplateplot` — Display visualization of microtiter plate.

Sequence Utility Functions

The following functions are updated:

- `rebasecuts` — Find restriction enzymes that cut nucleotide sequence. Updated to use Version 811 of REBASE, the Restriction Enzyme Database.
- `restrict` — Split nucleotide sequence at restriction site. Updated to use Version 811 of REBASE, the Restriction Enzyme Database.

Sequence Conversion Functions

The following function is updated:

- `nt2aa` — Convert nucleotide sequence to amino acid sequence. Updated to include a new property, 'ACGTOnly', to support ambiguous and unknown nucleotide characters.

Bioanalytic and Mass Spectrometry Functions

The following functions are updated to use with data from any separation technique, including mass spectrometry:

- `msalign` — Align peaks in signal to reference peaks.
- `msbackadj` — Correct baseline of signal with peaks.
- `mslowess` — Smooth signal with peaks using nonparametric method.
- `msnorm` — Normalize set of signals with peaks.
- `mspeaks` — Convert raw peak data to peak list (centroided data).
- `msppresample` — Resample signal with peaks while preserving peaks.
- `msresample` — Resample signal with peaks.

- `mssgolay` — Smooth signal with peaks using least-squares polynomial.

Microarray Functions

Compatibility Considerations: Yes

The following functions are updated:

- `cghcbs` — Perform circular binary segmentation (CBS) on array-based comparative genomic hybridization (aCGH) data. Updated to include an optional heuristic stopping rule to improve performance.
- `ilmnbslookup` — Look up Illumina® BeadStudio™ target (probe) sequence and annotation information. Updated to read Illumina microRNA array annotation files.
- `ilmnbsread` — Read gene expression data exported from Illumina BeadStudio software. Updated to read Illumina microRNA array data files.
- `mattest` — Perform two-sample t-test to evaluate differential expression of genes from two experimental conditions or phenotypes. Updated with new property, 'VarType', which lets you specify equal or unequal (default) variance for the test.

Compatibility Considerations

A compatibility consideration related to the `mattest` function was introduced in Bioinformatics Toolbox Version 3.2, but not reported in the Release Notes for Version 3.2 (R2008b). Specifically, in Bioinformatics Toolbox Version 3.1 and earlier, the `mattest` function used equal variance for the test. In Bioinformatics Toolbox Version 3.2, the `mattest` function starting using unequal variance for the test.

Demo for Sequence Analysis

The following is a new sequence analysis demo:

Predicting Protein Secondary Structure Using a Neural Network

R2008b

Version: 3.2

New Features: Yes

Bug Fixes: Yes

Data Format and Database Functions

Compatibility Considerations: Yes

Following are new functions:

- `affygcma` — Perform GC Robust Multi-array Average (GCRMA) procedure on Affymetrix microarray probe-level data.
- `affyrma` — Perform Robust Multi-array Average (RMA) procedure on Affymetrix microarray probe-level data.
- `affysnannotread` — Read Affymetrix Mapping DNA array data from CSV-formatted annotation file.
- `geoseriesread` — Read Gene Expression Omnibus (GEO) Series (GSE) format data.
- `multialignwrite` — Write multiple-alignment to file using ClustalW ALN format.
- `mzcdfread` — Read mass spectrometry data from netCDF file.

The following functions are updated:

- `affyread` — Read microarray data from Affymetrix GeneChip file. Updated so that `Probes` field in the return structure is now a `single`, which reduces memory usage.
- `celintensityread` — Read probe intensities from Affymetrix CEL files. Updated so that `PMIntensities` and `MMIntensities` fields in the return structure are now `singles`, which reduces memory usage.
- `geosoftread` — Read Gene Expression Omnibus (GEO) SOFT format data. Updated to support Platform (GPL) records.
- `getgeodata` — Retrieve Gene Expression Omnibus (GEO) format data. Updated to support Platform (GPL) and Series (GSE) records.
- `goannotread` — Read annotations from Gene Ontology annotated file. Updated to include two new properties, `'Fields'` and `'Aspect'`, which let you read a subset of the data in the annotated file.
- `multialignread` — Read multiple sequence alignment file. Updated to support PHYLIP (Phylogeny Inference Package) multiple-sequence alignment files.

- `mzxmlread` — Read data from mzXML file. Improved to read larger files, faster and without running out of memory. Updated with three new properties, 'Levels', 'TimeRange', and 'ScanIndices', which let you filter and read a subset of the data. Updated with a 'Verbose' property to control the progress display while reading the file.

Compatibility Considerations

In Bioinformatics Toolbox Version 3.1 and earlier, the `Probes` field, in the structure returned by `affyread`, and the `PMIntensities` and `MMIntensities` fields, in the structure returned by `celintensityread`, were doubles. In Bioinformatics Toolbox Version 3.2, these fields are singles.

Sequence Utility Functions

Following is a new function:

- `cleavelookup` — Find cleavage rule for enzyme or compound.

The following functions are updated:

- `blastncbi` — Create remote NCBI BLAST report request ID or link to NCBI BLAST report. Updated to include a 'GapCosts' property, which lets you specify penalties for both opening and extending gaps, and an 'Entrez' property, which lets you limit searches using Entrez query syntax.
- `cleave` — Cleave amino acid sequence with enzyme. Includes a new input argument that specifies the name of an enzyme or compound for which a cleavage rule is specified in the literature.
- `rebasecuts` — Find restriction enzymes that cut nucleotide sequence. Updated to use Version 806 of REBASE, the Restriction Enzyme Database.
- `restrict` — Split nucleotide sequence at restriction site. Updated to use Version 806 of REBASE, the Restriction Enzyme Database.
- `seqlogo` — Display sequence logo for nucleotide or amino acid sequences. Updated to return a figure handle to the sequence logo.

Multiple Sequence Alignment Functions

Following is a new function:

- `multialignwrite` — Write multiple alignment to file using ClustalW ALN format.

The following function is updated:

- `multialignread` — Read multiple sequence alignment file. Updated to support PHYLIP (Phylogeny Inference Package) multiple sequence alignment files.

Gene Ontology Functions

The following function is updated:

- `goannotread` — Read annotations from Gene Ontology annotated file. Updated to include two new properties, 'Fields' and 'Aspect', which let you read a subset of the data in the annotated file.

Protein Analysis Functions

Following are new functions:

- `cleavelookup` — Find cleavage rule for enzyme or compound.
- `pdbsuperpose` — Superpose 3-D structures of two proteins.
- `pdbtransform` — Apply linear transformation to 3-D structure of molecule.

The following function is updated:

- `cleave` — Cleave amino acid sequence with enzyme. Includes a new input argument that specifies the name of an enzyme or compound for which a cleavage rule is specified in the literature.

Mass Spectrometry Functions

Following are new functions:

- `mzcdf2peaks` — Convert `mzCDF` structure to peak list.
- `mzcdfinfo` — Return information about `netCDF` file containing mass spectrometry data.
- `mzcdfread` — Read mass spectrometry data from `netCDF` file.
- `mzxmlinfo` — Return information about `mzXML` file.

The following function is updated:

- `mzxmlread` — Read data from `mzXML` file. Improved to read larger files, faster and without running out of memory. Updated with three new properties, `'Levels'`, `'TimeRange'`, and `'ScanIndices'`, which let you filter and read a subset of the data. Updated with a `'Verbose'` property to control the progress display while reading the file.

Microarray File Format Functions

Compatibility Considerations: Yes

Following are new functions:

- `affygcma` — Perform GC Robust Multi-array Average (GCRMA) procedure on Affymetrix microarray probe-level data.
- `affyrma` — Perform Robust Multi-array Average (RMA) procedure on Affymetrix microarray probe-level data.
- `affysnpannotread` — Read Affymetrix Mapping DNA array data from CSV-formatted annotation file.
- `geoseriesread` — Read Gene Expression Omnibus (GEO) Series (GSE) format data.

The following functions are updated:

- `affyread` — Read microarray data from Affymetrix GeneChip file. Updated so that `Probes` field in the return structure is now a `single`, which reduces memory usage.
- `celintensityread` — Read probe intensities from Affymetrix CEL files. Updated so that `PMIntensities` and `MMIntensities` fields in the return structure are now `singles`, which reduces memory usage.

- `geosoftread` — Read Gene Expression Omnibus (GEO) SOFT format data. Updated to support Platform (GPL) records.
- `getgeodata` — Retrieve Gene Expression Omnibus (GEO) format data. Updated to support Platform (GPL) and Series (GSE) records.

Compatibility Considerations

In Bioinformatics Toolbox Version 3.1 and earlier, the `Probes` field, in the structure returned by `affyread`, and the `PMIntensities` and `MMIntensities` fields, in the structure returned by `celintensityread`, were doubles. In Bioinformatics Toolbox Version 3.2, these fields are singles.

Microarray Functions

Compatibility Considerations: Yes

Following are new functions:

- `affysnpintensitiesplit` — Split Affymetrix SNP probe intensity information for alleles A and B.
- `affygcrrma` — Perform GC Robust Multi-array Average (GCRMA) procedure on Affymetrix microarray probe-level data.
- `affyrma` — Perform Robust Multi-array Average (RMA) procedure on Affymetrix microarray probe-level data.
- `DataMatrix` — Create `DataMatrix` object.

The following functions are updated:

- `illumbslookup` — Look up Illumina BeadStudio target (probe) sequence and annotation information. Updated to support BGX and TXT annotation files.
- `mattest` — Perform two-sample t-test to evaluate differential expression of genes from two experimental conditions or phenotypes. Updated to use unequal variance instead of equal variance for the test.
- `probesetlookup` — Look up information for Affymetrix probe set. Updated to accept multiple probe set IDs/names or gene IDs.

Compatibility Considerations

In Bioinformatics Toolbox Version 3.1 and earlier, the `matstest` function used equal variance for the test. In Bioinformatics Toolbox Version 3.2, the `matstest` function uses unequal variance for the test.

DataMatrix Object

Following is a new object:

- DataMatrix object — Data structure encapsulating data and metadata from microarray experiment so that it can be indexed by gene or probe identifiers and by sample identifiers.

DataMatrix Methods

There are many methods that let you create, index into, modify, create subsets, sort, perform operations on, analyze, and plot a DataMatrix object.

Demo for Visualization Tools

The Visualizing the Three-Dimensional Structure of a Molecule demo is updated to use the new `pdbsuperpose` function.

Demo for Sequence Analysis

The following is a new sequence analysis demo:

- Analyzing the Human Distal Gut Microbiome

Demos for Microarray Data Analysis

Following is a new microarray data analysis demo:

- Working with GEO Series Data

The Exploring Gene Expression Data demo is updated to use the new `DataMatrix` object.

The Analyzing Affymetrix SNP Arrays for DNA Copy Number Variants demo is updated to use two new functions: `affysnpannotread` and `affysnpintensitysplit`.

The Preprocessing Affymetrix Microarray Data at the Probe Level demo is updated to use two new functions: `affygcRMA` and `affyRMA`.

R2008a

Version: 3.1

New Features: Yes

Bug Fixes: Yes

Data Format and Database Functions

Compatibility Considerations: Yes

Following is a new function:

- `ilmnbsread` — Read microarray data exported from Illumina BeadStudio software.

The following functions are updated:

- `celintensityread` — Read probe intensities from Affymetrix CEL files. Updated output structure to include a new field, `GroupNumbers`, which contains group numbers of probes.
- `fastawrite` — Write to file using FASTA format. Updated such that if you specify an existing file, new data is appended to the file instead of overwriting it.
- `getgenbank` — Retrieve sequence information from GenBank® database. Updated such that if you use the `'ToFile'` property and specify an existing file, new data is appended to the file instead of overwriting it. Updated to allow you to access a partial sequence by adding new property `'PartialSeq'`.
- `getgenpept` — Retrieve sequence information from GenPept database. Updated such that if you use the `'ToFile'` property and specify an existing file, new data is appended to the file instead of overwriting it. Updated to allow you to access a partial sequence by adding new property `'PartialSeq'`.
- `getgeodata` — Retrieve Gene Expression Omnibus (GEO) SOFT format data. Updated to retrieve both Sample (GSM) and Data Set (GDS) data.

Compatibility Considerations

In Bioinformatics Toolbox Version 3.0 and earlier, when writing to files using the `fastawrite` function or the `getgenbank` or `getgenpept` functions with the `'ToFile'` property, if you specified an existing file, the file was overwritten. In Bioinformatics Toolbox Version 3.1, if you specify an existing file, new data is appended to the file instead of overwriting it.

Sequence Utility Functions

The following functions are updated:

- `evalrasmolscript` — Send RasMol script commands to Molecule Viewer window. Updated to use Version 11.4 of the Jmol molecule viewer.
- `molviewer` — Display and manipulate 3-D molecule structure. Updated to use Version 11.4 of the Jmol molecule viewer.
- `ramachandran` — Draw Ramachandran plot for Protein Data Bank (PDB) data. Updated to handle PDB files with multiple chains and models by adding three properties: 'Chain', 'Plot', and 'Model'. Updated Ramachandran plot to mark glycine residues and display reference regions by adding three properties: 'Glycine', 'Regions', and 'RegionDef'. Updated Ramachandran plot to display amino acid information in ToolTip. Updated to easily determine the names and sequence positions of amino acids corresponding to torsion angles by creating an output structure.
- `rebasecuts` — Find restriction enzymes that cut nucleotide sequence. Updated to use Version 710 of REBASE, the Restriction Enzyme Database.
- `restrict` — Split nucleotide sequence at restriction site. Updated to use Version 710 of REBASE, the Restriction Enzyme Database.

Pairwise Sequence Alignment Functions

The following functions are updated:

- `nwalign` — Globally align two sequences using Needleman-Wunsch algorithm. Updated to improve pairwise sequence performance.
- `swalign` — Locally align two sequences using Smith-Waterman algorithm. Updated to improve pairwise sequence performance.

Phylogenetic Tree Tools Function

The following function is updated:

- `dnds` — Estimate synonymous and nonsynonymous substitution rates. Updated by adding 'AdjustStops' property to control whether stop codons are excluded from calculations.

Protein Analysis Functions

The following functions are updated:

- `evalrasmolscript` — Send RasMol script commands to Molecule Viewer window. Updated to use Version 11.4 of the Jmol molecule viewer.
- `molviewer` — Display and manipulate 3-D molecule structure. Updated to use Version 11.4 of the Jmol molecule viewer.
- `ramachandran` — Draw Ramachandran plot for Protein Data Bank (PDB) data. Updated to handle PDB files with multiple chains and models by adding three properties: 'Chain', 'Plot', and 'Model'. Updated Ramachandran plot to mark glycine residues and display reference regions by adding three properties: 'Glycine', 'Regions', and 'RegionDef'. Updated Ramachandran plot to display amino acid information in ToolTip. Updated to easily determine the names and sequence positions of amino acids by creating an output structure.

Microarray File Format Functions

Following is a new function:

- `ilmnbsread` — Read microarray data exported from Illumina BeadStudio software.

The following functions are updated:

- `celintensityread` — Read probe intensities from Affymetrix CEL files. Updated output structure to include a new field, `GroupNumbers`, which contains group numbers of probes.
- `getgeodata` — Retrieve Gene Expression Omnibus (GEO) SOFT format data. Updated to retrieve both Sample (GSM) and Data Set (GDS) data.

Microarray Functions

Compatibility Considerations: Yes

Following are new functions:

- `affysnpquartets` — Create table of SNP probe quartet results for Affymetrix probe set.
- `cghfreqplot` — Display frequency of DNA copy number alterations across multiple samples.
- `ilmnbslookup` — Look up Illumina BeadStudio target (probe) sequence and annotation information.
- `redbluecmap` — Create red and blue color map.

The following functions are updated:

- `clustergram` — Compute hierarchical clustering, display dendrogram and heat map, and create clustergram object.

Updated properties include:

- `'Linkage'` — Can specify linkage method separately for rows and columns.
- `'Dendrogram'` — Can specify color threshold separately for rows and columns.

Replaced properties include:

- `'Dimension'` — Replaced by the `'Cluster'` property, which lets you cluster along the columns, rows, or both.
- `'Pdist'` — Replaced by `'RowPdist'` and `'ColumnPdist'` properties.

New properties include:

- `'Standardize'` — Specifies the dimension for standardizing the data.
- `'DisplayRange'` — Specifies the display range of standardized values.
- `'LogTrans'` — Controls the \log_2 transform of the data.
- `'Impute'` — Specifies a function and properties to impute missing data.
- `'RowMarker'` — Adds color and text marker to a group of rows.
- `'ColumnMarker'` — Adds color and text marker to a group of columns.

The interactivity of the clustergram figure is enhanced with the following features:

- Select a group of rows or columns and display the group number and genes or samples within.
- Create a new clustergram of only a group of the data.
- Export data as a clustergram object or structure in the MATLAB Workspace.
- `maboxplot` — Create box plot for microarray data. Updated by adding 'BoxPlot' property, which lets you specify arguments to pass to the `boxplot` function, which creates the box plot.
- `mairplot` — Create intensity versus ratio scatter plot of microarray data. Updated by adding 'PlotOnly' property, which lets you display the scatter plot without user interface components.
- `mattest` — Perform two-sample t-test to evaluate differential expression of genes from two experimental conditions or phenotypes. Updated by adding 'Bootstrap' property to run bootstrap tests.
- `mavolcanoplot` — Create significance versus gene expression ratio (fold change) scatter plot of microarray data. Updated by adding 'PlotOnly' property, which lets you display the volcano plot without user interface components.
- `probesetvalues` — Create table of Affymetrix probe set intensity values. Updated by adding 'Background' property to control the background correction.
- `zonebackadj` — Perform background adjustment on Affymetrix microarray probe-level data using zone-based method. Updated to return a third output containing the estimated background values for each probe.

Compatibility Considerations

In Bioinformatics Toolbox Version 3.0 and earlier, the `clustergram` function included 'Dimension' and 'Pdist' properties. In Bioinformatics Toolbox Version 3.1, the 'Dimension' property is replaced by the 'Cluster' property, and the 'Pdist' property is replaced by the 'RowPdist' and 'ColumnPdist' properties.

Object

Following is a new object:

- `clustergram` object — Object containing hierarchical clustering analysis data.

Clustergram Methods

The following are new methods of a `clustergram` object:

- `get` — Retrieve information about `clustergram` object.
- `plot` — Render `clustergram` heat map and dendrograms for `clustergram` object.
- `set` — Set property of `clustergram` object.
- `view` — View `clustergram` heat map and dendrograms for `clustergram` object.

Demo for Sequence Analysis

The following is a new sequence analysis demo:

- Performing a Metagenomic Analysis of a Sargasso Sea Sample

Demo for Microarray Data Analysis

The following is a new microarray data analysis demo:

- Analyzing Affymetrix SNP Arrays for DNA Copy Number Variants

Demo for Visualization Tools

The following is a new visualization tool demo:

- Working with the Clustergram Function

Demos for Mass Spectrometry Data Analysis

- The Batch Processing of Spectra Using Distributed Computing demo is updated to use the latest features of the Parallel Computing Toolbox™ version 3.3, and is now called Batch Processing of Spectra Using Sequential and Parallel Computing
- The Preprocessing Raw Mass Spectrometry Data demo is updated with state-of-the-art examples for peak detection using wavelets denoising, binning by hierarchical clustering, and binning by dynamic programming.

R2007b

Version: 3.0

New Features: Yes

Bug Fixes: Yes

Data Format and Database Functions

Following are new functions:

- `blastformat` — Create local BLAST database.
- `blastreadlocal` — Read data from local BLAST report.
- `cytobandread` — Read cytogenetic banding information.

The following function was updated:

- `affyread` — Read microarray data from Affymetrix GeneChip file. Updated the structure returned when reading a CDF library file. The structure contains three new subfields: `GroupNumber`, `Direction`, and `GroupName`.

Microarray File Format Functions

Following is a new function:

- `cytobandread` — Read cytogenetic banding information.

The following function was updated:

- `affyread` — Read microarray data from Affymetrix GeneChip file. Updated the structure returned when reading a CDF library file. The structure contains three new subfields: `GroupNumber`, `Direction`, and `GroupName`.

Microarray Functions

Following are new functions:

- `chromosomeplot` — Plot chromosome ideogram with G-banding pattern.
- `cghcbs` — Perform circular binary segmentation (CBS) on array-based comparative genomic hybridization (aCGH) data.

The following function is updated:

- `probesetvalues` — Create table of Affymetrix probe set intensity values. Updated return matrix, which contains intensity values for probe-level

data, to include two new fields: `GroupNumber` and `Direction`. Updated to return a second output containing the column names for the return matrix, which contains intensity values for probe-level data.

Sequence Conversion, Utility, and Visualization Functions

Following are new functions:

- `blastlocal` — Perform search on local BLAST database to create BLAST report.
- `rnaconvert` — Convert secondary structure of RNA sequence between bracket and matrix notations.
- `rnafold` — Predict minimum free-energy secondary structure of RNA sequence.
- `rnaplot` — Draw secondary structure of RNA sequence.

Mass Spectrometry Functions

The following function is updated:

- `mssalign` — Align mass spectra from multiple peak lists from LC/MS or GC/MS data set. Updated to include a new property, `ShowEstimation`, which controls the display of an assessment plot relative to the estimation method and the vector of common mass/charge (m/z) values.

Statistical Learning Functions

Compatibility Considerations: Yes

The following function is updated:

- `svmsmoset` — Create or edit Sequential Minimal Optimization (SMO) options structure. Updated default values for the `MaxIter` and `KernelCacheLimit` properties. Changed the `Display` property so that when set to `iter`, a report displays every 500 iterations instead of 10.

Compatibility Considerations

In Bioinformatics Toolbox Version 2.6 and earlier, the `svmsmoset` function used a `'MaxIter'` property with a default of 1500 and a `'KernelCacheLimit'` property with a default of 7500. In Bioinformatics Toolbox Version 3.0, the defaults are 15000 and 5000, respectively. Also, when you set the `'Display'` property to `'iter'`, a report displays every 500 iterations instead of 10.

Gene Ontology Methods

The following methods of a gene ontology object are updated:

- `geneont.getancestors` — Find terms that are ancestors of specified Gene Ontology term. Updated to also return the number of times each ancestor is found. Updated to include two new properties, `'Relationtype'`, which specifies a relationship type to search for in the gene ontology, and `'Exclude'`, which controls excluding the original queried term(s) from the output, unless the term was reached while searching the gene ontology.
- `geneont.getdescendants` — Find terms that are descendants of specified Gene Ontology term. Updated to also return the number of times each descendant is found. Updated to include two new properties, `'Relationtype'`, which specifies a relationship type to search for in the gene ontology, and `'Exclude'`, which controls excluding the original queried term(s) from the output, unless the term was reached while searching the gene ontology.
- `geneont.getrelatives` — Find terms that are relatives of specified Gene Ontology term. Updated to also return the number of times each relative is found. Updated to include three new properties, `'Levels'`, which specifies the number of levels up and down to search in the gene ontology, `'Relationtype'`, which specifies a relationship type to search for in the gene ontology, and `'Exclude'`, which controls excluding the original queried term(s) from the output, unless the term was reached while searching the gene ontology.

Demos for Microarray Data Analysis

The following are two new microarray data analysis demos:

- Detecting DNA Copy Number Alteration in Array-Based CGH Data
- Analyzing Array-Based CGH Data Using Bayesian Hidden Markov Modeling

Demos for Sequence Analysis

The following are two new sequence analysis demos:

- Predicting and Visualizing the Secondary Structure of RNA Sequences
- Identifying Over-Represented Regulatory Motifs

The Investigating the Bird Flu Virus demo was updated to demonstrate how to write KML-formatted files, which can be used by Google Earth™ to display geospatial data.

Demo for Graph Theory Analysis

The following is a new graph theory demo:

- Working with Graph Theory Functions

R2007a+

Version: 2.6

New Features: Yes

Bug Fixes: Yes

Data Formats and Databases Functions

Compatibility Considerations: Yes

The following functions are updated:

- `affyread` — Read microarray data from Affymetrix GeneChip file. Updated to read Affymetrix files from expression, genotyping, or resequencing assays on all platforms, except Solaris™.
- `celintensityread` — Read probe intensities from Affymetrix CEL files. Updated to read Affymetrix CEL and CDF files from expression or genotyping assays on all platforms, except Solaris.
- `mzxmlread` — Read mzXML file into MATLAB as structure. Updated to read mzXML files that conform to the mzXML 2.1 specification or earlier specifications.

Compatibility Considerations

In Bioinformatics Toolbox Version 2.6, the structure returned by `affyread` when reading a CHP file from an expression assay no longer contains a `ProbePairs` field. The `ProbePairs` field still exists in the structure returned by `affyread` when reading a CDF file.

Microarray File Formats Functions

Compatibility Considerations: Yes

The following functions are updated:

- `affyread` — Read microarray data from Affymetrix GeneChip file. Updated to read Affymetrix files from expression, genotyping, or resequencing assays on all platforms, except Solaris.
- `celintensityread` — Read probe intensities from Affymetrix CEL files. Updated to read Affymetrix CEL and CDF files from expression or genotyping assays on all platforms, except Solaris.

Compatibility Considerations

In Bioinformatics Toolbox Version 2.6, the structure returned by `affyread` when reading a CHP file from an expression assay no longer contains a `ProbePairs` field. The `ProbePairs` field still exists in the structure returned by `affyread` when reading a CDF file.

Microarray Utility Functions

Compatibility Considerations: Yes

The following function is updated:

- `probesetplot` — Plot Affymetrix probe set intensity values. Updated to accept structures created from CEL and CDF files, instead of a structure created from a CHP file.

Compatibility Considerations

In Bioinformatics Toolbox Version 2.5 and earlier, the `probesetplot` function accepted a structure created from a CHP file as input. Currently it requires two structures: one created from a CEL file and one created from a CDF library file. If you have any scripts that call the `probesetplot` function, you need to update them to provide the correct input arguments.

Microarray Normalization and Filtering Functions

Following is a new function:

- `zonebackadj` — Perform background adjustment on Affymetrix microarray probe-level data using zone-based method.

Mass Spectrometry Functions

The following function is updated:

- `mzxmlread` — Read mzXML file into MATLAB as structure. Updated to read mzXML files that conform to the mzXML 2.1 specification or earlier specifications.

Following is a new function you can use to calibrate and/or synchronize multidimensional mass spectrometry data:

- `samplealign` — Align two data sets containing sequential observations by introducing gaps.

Demos for Mass Spectrometry Functions

The following are two new mass spectrometry demos:

- Visualizing and Preprocessing Hyphenated Mass-Spectrometry Data Sets for Metabolite and Protein/Peptide Profiling
- Differential Analysis of Complex Protein and Metabolic Mixtures Using Liquid Chromatography/Mass Spectrometry (LC/MS)

R2007a

Version: 2.5

New Features: Yes

Bug Fixes: Yes

Data Formats and Database Functions

Compatibility Considerations: Yes

Following are new functions for reading and creating files:

- `affyprobeseqread` — Read data file containing probe sequence information for Affymetrix GeneChip array.
- `pdbwrite` — Write to file using Protein Data Bank (PDB) format.

The following functions were updated:

- `celintensityread` — Read probe intensities from Affymetrix CEL files (Windows® 32). Updated so that the order of columns (CEL files) in return matrices `PMIntensities` and `MMIntensities` matches the order of CEL files in the `CELFiles` input argument.
- `pdbread` — Read data from Protein Data Bank (PDB) file. Updated so that the six fields containing coordinate information (`Atom`, `AtomSD`, `AnisotropicTemp`, `AnisotropicTempSD`, `Terminal`, and `HeterogenAtom`) are now subfields within the `Model` field of the MATLAB structure. Updated to include a new property, `ModelNum`, which reads only the specified model from a PDB-formatted text file.

Compatibility Considerations

In Bioinformatics Toolbox Version 2.4 and earlier, the `celintensityread` function ordered the columns (CEL files) of return matrices `PMIntensities` and `MMIntensities` alphabetically.

In Bioinformatics Toolbox Version 2.4 and earlier, the `pdbread` function stored coordinate information in six fields (`Atom`, `AtomSD`, `AnisotropicTemp`, `AnisotropicTempSD`, `Terminal`, and `HeterogenAtom`) within the MATLAB structure. These six fields are now subfields within the `Model` field of the MATLAB structure.

Demo for Data Formats and Database Functions

The Accessing NCBI Entrez Databases with E-Utilities demo illustrates how to programatically search and retrieve data.

Statistical Learning Functions

Compatibility Considerations: Yes

Following are new functions:

- `optimalleaforder` — Determine optimal leaf ordering for hierarchical binary cluster tree.
- `svmsmset` — Create or edit Sequential Minimal Optimization (SMO) options structure.

The following function was updated:

- `svmtrain` — Train support vector machine classifier. Updated to include a new SMO method and a new property, `SMO_Opts`, which provides options for the SMO method. The `BoxConstraint` property has changed, including a new default value.

Compatibility Considerations

In Bioinformatics Toolbox Version 2.4 and earlier, the `svmtrain` function used a `BoxConstraint` property with a default of $\frac{1}{\sqrt{\epsilon}}$. In Bioinformatics Toolbox Version 2.5, the default is 1, which can lead to slightly different results.

Protein Analysis and Sequence Utilities Functions

Compatibility Considerations: Yes

Following are new functions:

- `evalrasmolscript` — Send RasMol script commands to molecule viewer.
- `molviewer` — Display and manipulate 3-D molecule structure.
- `proteinpropplot` — Plot properties of amino acid sequence.
- `seqinsertgaps` — Insert gaps into nucleotide or amino acid sequence.

The following functions were updated:

- `featuresparse` — Parse features from GenBank, GenPept, or EMBL data. Updated to include a new property, `Sequence`, which controls the extraction, when possible, of the sequences.
- `oligoprop` — Calculate sequence properties of DNA oligonucleotide. Updated to handle ambiguous N characters in a sequence.

The following function is removed:

- `pdbplot` — Plot 3-D protein structure. This function was replaced by the `molviewer` function.

Compatibility Considerations

In Bioinformatics Toolbox Version 2.5, the `pdbplot` function was replaced by the `molviewer` function. If you have any scripts that call the `pdbplot` function, you need to update them to call the `molviewer` function.

Demo for Protein Analysis and Sequence Utilities Functions

The Visualizing the Three-dimensional Structure of a Molecule demo illustrates the `molviewer` function.

Sequence Alignment Functions

Compatibility Considerations: Yes

The following function was updated:

- `seqpdist` — Calculate pairwise distance between sequences. Updated to assume that all input sequences are aligned if they have the same length, regardless of the presence of gaps. If you know your input sequences are not aligned, you can align them before passing them to `seqpdist` (for example, using `multialign`), or set `PairwiseAlignment` to `true` when using `seqpdist`.

Compatibility Considerations

In Bioinformatics Toolbox Version 2.4 and earlier, the `seqpdist` function assumed all input sequences were aligned if they had the same length and at least one gap.

Demo for Sequence Alignment Functions

The Comparing Whole Genomes demo illustrates how to compare features of organisms on a genomic evolution scale.

Microarray File Formats Functions

Compatibility Considerations: Yes

Following is a new function:

- `affyprobeseqread` — Read data file containing probe sequence information for Affymetrix GeneChip array.

The following function was updated:

- `celintensityread` — Read probe intensities from Affymetrix CEL files (Windows 32). Updated so that the order of columns (CEL files) in return matrices `PMIntensities` and `MMIntensities` matches the order of CEL files in the `CELFiles` input argument.

Compatibility Considerations

In Bioinformatics Toolbox Version 2.4 and earlier, the `celintensityread` function ordered the columns (CEL files) of return matrices `PMIntensities` and `MMIntensities` alphabetically.

Microarray Normalization and Filtering Functions

Following are new functions:

- `affyprobeaffinities` — Compute Affymetrix probe affinities from their sequences and MM probe intensities.
- `gcrmabackadj` — Perform GC Robust Multi-array Average (GCRMA) background adjustment on Affymetrix microarray probe-level data using sequence information.
- `gcrma` — Perform GC Robust Multi-array Average (GCRMA) background adjustment, quantile normalization, and median-polish summarization on Affymetrix microarray probe-level data.

Demo for Microarray File Formats, Normalization, and Filtering Functions

The Preprocessing Affymetrix Microarray Data at the Probe Level demo illustrates the `affyprobeseqread`, `affyprobeaffinities`, `gcrmabackadj`, and `gcrma` functions.

Microarray Data Analysis and Visualization Functions

Following is a new function:

- `mafdr` — Estimate false discovery rate (FDR) of differentially expressed genes from two experimental conditions or phenotypes.

The following function was updated:

- `mattest` — Perform two-tailed t-test to evaluate differential expression of genes from two experimental conditions or phenotypes. Updated to include a new property, `Permute`, which controls whether permutation tests are run.

Demo for Microarray Data Analysis and Visualization Functions

The Exploring Gene Expression Data demo illustrates the `mattest` and `mafdr` functions.

Mass Spectrometry Functions

Following are new functions:

- `msdotplot` — Plot set of peak lists from LC/MS or GC/MS data set.
- `mssalign` — Align mass spectra from multiple peak lists from LC/MS or GC/MS data set.
- `mspeaks` — Convert raw mass spectrometry data to peak list (centroided data).
- `mssppresample` — Resample mass spectrometry signal while preserving peaks.
- `mzxml2peaks` — Convert mzXML structure to peak list.

The following function was updated:

- `msheatmap` — Create pseudocolor image of set of mass spectra. Updated to handle LC/MS and GC/MS data.

Phylogenetic Tree Tools Functions

Compatibility Considerations: Yes

Following is a new function:

- `seqinsertgaps` — Insert gaps into nucleotide or amino acid sequence.

The following functions were updated:

- `dnds` — Estimate synonymous and nonsynonymous substitution rates. Updated to include two new properties, `Verbose`, which controls the display of the codons considered in the computations and their amino acid translations, and `Window`, which performs the calculations over a sliding window.
- `dndsm1` — Estimate synonymous and nonsynonymous substitution rates using maximum likelihood method. Updated to include a new property, `Verbose`, which controls the display of the codons considered in the computations and their amino acid translations.

- `seqpdist` — Calculate pairwise distance between sequences. Updated to assume that all input sequences are aligned if they have the same length, regardless of the presence of gaps. If you know your input sequences are not aligned, you can align them before passing them to `seqpdist` (for example, using `multialign`), or set `PairwiseAlignment` to `true` when using `seqpdist`.

Compatibility Considerations

In Bioinformatics Toolbox Version 2.4 and earlier, the `seqpdist` function assumed all input sequences were aligned if they had the same length and at least one gap.

Demos for Phylogenetic Tree Tools Functions

The following demos illustrate the `nwalign`, `seqinsertgaps`, `dnds`, and `multialign` functions:

- Analyzing Synonymous and Nonsynonymous Substitution Rates
- Investigating the Bird Flu Virus

The Reconstructing the Origin and the Diffusion of the SARS Epidemic demo presents an analysis of the SARS epidemic.

Phylogenetic Tree Methods

Following is a new method of a `phytree` object:

- `reorder` — Reorder leaves of phylogenetic tree.

R2006b

Version: 2.4

New Features: Yes

Bug Fixes: Yes

Data Formats and Database Functions

Following is a new function for getting data into the MATLAB environment:

- `mzxmlread` — Read mzXML file into the MATLAB software as structure.

The following functions were updated:

- `celintensityread` — Read probe intensities from Affymetrix CEL files (Windows 32). Updated to include a new property, `Verbose`, which controls the display of a progress report showing the name of each CEL file as it is read.
- `fastaread` — Read data from FASTA file. Updated to include a new property, `Blockread`, which controls reading a single entry or block of entries from a file.
- `geosoftread` — Read Gene Expression Omnibus (GEO) SOFT format data. Updated to read Data Set (GDS) files as well as Sample (GSM) files.
- `getblast` — BLAST report from NCBI Web site. Updated to include a new property, `WaitTilReady`, which pauses the MATLAB software and waits a specified time (minutes) for a report from the NCBI Web site.
- `scfread` — Read trace data from SCF file. Updated to include more output options.

Sequence Utilities Functions

Following is a new function for parsing sequence data:

- `featuresparse` — Parse features from GenBank, GenPept, or EMBL data.

Sequence Visualization Functions

The following function was updated:

- `seqtool` — Open tool to interactively explore biological sequences. Updated to download sequences from the EMBL database, interactively move the viewing frame in the Sequence Viewer by pressing and holding **Ctrl** while

click-dragging, and export an amino acid translation as a FASTA file or to the MATLAB Workspace.

Multiple Sequence Alignment Functions

The following function was updated:

- `multialignviewer` — Open viewer for multiple sequence alignments. Updated to export consensus sequences.

Microarray File Formats

The following function was updated:

- `celintensityread` — Read probe intensities from Affymetrix CEL files (Windows 32). Updated to include a new property, `Verbose`, which controls the display of a progress report showing the name of each CEL file as it is read.

Microarray Data Analysis and Visualization Functions

The following functions were updated:

- `clustergram` — Create dendrogram and heat map. Updated to include a new property, `OptimalLeafOrder`, which enables or disables the optimal leaf ordering calculation, which determines the leaf order that maximizes the similarity between neighboring leaves.
- `mairplot` — Create intensity versus ratio scatter plot for microarray signals. Updated to include a new property, `Type`, which creates either an IR plot or MA plot, changing the plot axes to log scale, and adding plot interactive features such as displaying gene labels, changing factor lines, normalizing data, and exporting data.
- `mapcaplot` — Create Principal Component plot of expression profile data. Updated by adding an export feature.
- `redgreenmap` — Create red and green colormap. Updated to include a new property, `Interpolation`, which sets the method for color interpolation.

Graph Theory Functions

Following are new functions for applying basic graph theory algorithms to sparse matrices:

- `graphallshortestpaths` — Find all shortest paths in graph.
- `graphconncomp` — Find strongly or weakly connected components in graph.
- `graphisdag` — Test for cycles in directed graph.
- `graphisomorphism` — Find isomorphism between two graphs.
- `graphisspantree` — Determine if tree is spanning tree.
- `graphmaxflow` — Calculate maximum flow and minimum cut in directed graph.
- `graphminspantree` — Find minimal spanning tree in graph.
- `graphpred2path` — Convert predecessor indices to paths.
- `graphshortestpath` — Solve shortest path problem in graph.
- `graphtopoorder` — Perform topological sort of directed acyclic graph.
- `graphtraverse` — Traverse graph by following adjacent nodes.

Graph Visualization Methods

Following are new methods for applying basic graph theory algorithms to a `biograph` object:

- `allshortestpaths` — Find all shortest paths in `biograph` object.
- `conncomp` — Find strongly or weakly connected components in `biograph` object.
- `getmatrix` — Get connection matrix from `biograph` object.
- `isdag` — Test for cycles in `biograph` object.
- `isomorphism` — Find isomorphism between two `biograph` objects.
- `isspantree` — Determine if tree created from `biograph` object is spanning tree.
- `maxflow` — Calculate maximum flow and minimum cut in `biograph` object.

- `minspantree` — Find minimal spanning tree in biograph object.
- `shortestpath` — Solve shortest path problem in biograph object.
- `topoorder` — Perform topological sort of directed acyclic graph extracted from biograph object.
- `traverse` — Traverse biograph object by following adjacent nodes.

Phylogenetic Tree Methods

Following is a new method for the `phytree` object:

- `getmatrix` — Convert `phytree` object into a relationship matrix.

R2006a+

Version: 2.3

New Features: Yes

Bug Fixes: No

Data Formats and Databases Functions

The following functions are removed:

- `getpir` — Sequence data from PIR-PSD database. This function retrieved data from the PIR-PSD database. This database has been discontinued and this function no longer retrieves data. See <http://pir.georgetown.edu/pirwww/dbinfo/nref.shtml> for more details.
- `pirread` — Read data from Protein Information Resource (PIR) file. This function supported the data format of the PIR-PSD database. This database has been discontinued. See <http://pir.georgetown.edu/pirwww/dbinfo/nref.shtml> for more details.

Sequence Utilities Functions

The following function was updated to include five new databases, including `refseq_rna`, `refseq_genomic`, `env_nt`, `refseq_protein`, and `env_nr`:

- `blastncbi` — Generate remote BLAST request.

Sequence Visualization Functions

Following is a new function for visualizing sequence data:

- `featuresmap` — Draw linear or circular map of features from GenBank structure.

Statistical Learning Functions

The following function was updated to include three new properties, including `RBF_Sigma`, `BoxConstraint`, and `Autoscale`:

- `svmtrain` — Train support vector machine classifier.

Microarray Functions

The following function is supported on the Windows 32 platform only:

- `affyread` — Read microarray data from Affymetrix GeneChip file (Windows 32).

Following are new functions for preprocessing Affymetrix probe-level microarray data:

- `celintensityread` — Read probe intensities from Affymetrix CEL files (Windows 32).
- `rmabackadj` — Perform background adjustment on Affymetrix microarray probe-level data using Robust Multi-array Average (RMA) procedure.
- `rmasummary` — Calculate gene (probe set) expression values from Affymetrix microarray probe-level data using Robust Multi-array Average (RMA) procedure.
- `affyinvarsetnorm` — Perform rank invariant set normalization on probe intensities from multiple Affymetrix CEL or DAT files.

Following is a new function for two-color microarray normalization:

- `mainvarsetnorm` — Perform rank invariant set normalization on gene expression values from two experimental conditions or phenotypes.

Following are new functions for microarray differential expression analysis:

- `mattest` — Perform two-sample, two-tailed t-test to evaluate differential expression of genes from two experimental conditions or phenotypes.
- `mavolcanoplot` — Create significance versus gene expression ratio (fold change) scatter plot of microarray data.

Demo for Microarray Functions

New demo of the new microarray functions (Analyzing Affymetrix Microarray Gene Expression Data).

R2006a

Version: 2.2.1

New Features: No

Bug Fixes: No

No New Features or Changes

R14SP3+

Version: 2.2

New Features: Yes

Bug Fixes: No

Multiple Sequence Alignment Viewer

- `multialignviewer` — Interactively view, explore alignments, and make manual modifications.

Microarray Functions for Agilent Software

- `agferead` — Read an Agilent® Feature Extraction Software file.
- `magetfield` — Utility function to extract data from a microarray.

Gene Ontology Database Functions

- `geneont` — Import the Gene Ontology database from the Web.
- `geneont.getancestors`, `geneont.getdescendants`, `geneont.getrelatives` — Get a subset of the ontology.
- `goannotread` — Parse Gene Ontology Annotated files.
- `num2goid` — Convert numbers to Gene Ontology IDs.

Demo for Gene Ontology Functions

New demo for the new Gene Ontology functions (`geneontologydemo`) and working with whole genomes (`biomemorymapdemo`).

R14SP3

Version: 2.1.1

New Features: No

Bug Fixes: No

No New Features or Changes

R14SP2+

Version: 2.1

New Features: Yes

Bug Fixes: No

Sequence Alignment Functions

- `multialign` — Align multiple sequences using a progressive method with Distributed Computing Toolbox™ support.
- `multialignread` — Read multiple sequence alignment file.
- `profalign` — Align two profiles using Needleman-Wunsch global alignment.
- `showalignment` — Updated to show multiply aligned sequences.
- `seqpdist` — Updated to calculate pairwise distances between observations with Distributed Computing Toolbox support.

Sequence Statistics Functions

- `codonbias` — Calculate codon frequency for each amino acid in a DNA sequence.
- `cpgisland` — Locate CpG islands in a DNA sequence.

Sequence Utilities Functions

- `rebasecuts` — Find restriction enzymes that cut a protein sequence.
- `seqtool` — Graphical User Interface (GUI) for single sequence analysis.

Phylogenetic Tree Functions

- `dnds`, `dndsm1` — Estimate synonymous and nonsynonymous substitutions rates.
- `seqneighjoin` — Reconstruct a phylogenetic tree with a Neighbor-joining method.

Phylogenetic Tree Methods

- `getcanonical` — Calculate the canonical form of a phylogenetic tree.
- `getnewickstr` — Create a Newick formatted string.
- `reroot` — Change the root of a phylogenetic tree.

- `subtree` — Extract a subtree.
- `weights` — Calculate weights for a phylogenetic tree.

Microarray Functions

`probesetplot` — Plot values for an Affymetrix CHP file probe set.

Statistics Functions

`rankfeatures` — Renamed function. The previous name was `sqlfeatures`.

R14SP2

Version: 2.0.1

New Features: Yes

Bug Fixes: No

Updated RBASE Table

RBASE is the enzyme table that the function `restrict` uses to locate sequence patterns.

Expanded Bioperl Demonstration

Example of calling the MATLAB software from Perl scripts now includes several examples of passing various types of data (both directly and by variant variable) back and forth between Perl and a MATLAB Automation Server. To view the demo, type `bioperldemo`.