

## **msa**

### **An R Package for Multiple Sequence Alignment**

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## **Scope and Purpose of this Document**

This document provides a gentle introduction into the R package `msa`. Not all features of the R package are described in full detail. Such details can be obtained from the documentation enclosed in the R package. Further note the following: (1) this is not an introduction to multiple sequence alignment or algorithms for multiple sequence alignment; (2) this is not an introduction to R or any of the Bioconductor packages used in this document. If you lack the background for understanding this manual, you first have to read introductory literature on the subjects mentioned above.

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## 1 Introduction

Multiple sequence alignment is one of the most fundamental tasks in bioinformatics. Algorithms like ClustalW [13], ClustalOmega [12], and MUSCLE [3, 4] are well known and widely used. However, all these algorithms are implemented as stand-alone command line programs without any integration into the R/Bioconductor ecosystem. Before the `msa` package, only the `muscle` package has been available in R, but no other multiple sequence alignment algorithm, although the `Biostrings` package has provided data types for representing multiple sequence alignments for quite some time. The `msa` package aims to close that gap by providing a unified R interface to the multiple sequence alignment algorithms ClustalW, ClustalOmega, and MUSCLE. The package requires no additional software packages and runs on all major platforms. Moreover, the `msa` package provides an R interface to the powerful  $\text{\LaTeX}$  package `TeXshade` [1] which allows for a highly customizable plots of multiple sequence alignments. Unless some very special features of `TeXshade` are required, users can pretty-print multiple sequence alignments without the need to know the details of  $\text{\LaTeX}$  or `TeXshade`.

## 2 Installation

The `msa` R package (current version: 1.38.0) is available via Bioconductor. The simplest way to install the package is the following:

```
if (!requireNamespace("BiocManager", quietly=TRUE))
  install.packages("BiocManager")
BiocManager::install("msa")
```

To test the installation of the `msa` package, enter

```
library(msa)
```

in your R session. If this command terminates without any error message or warning, you can be sure that the `msa` package has been installed successfully. If so, the `msa` package is ready for use now and you can start performing multiple sequence alignments.

To make use of all functionalities of `msaPrettyPrint()`, a  $\text{\TeX}/\text{\LaTeX}$  system [5] must be installed. To make use of  $\text{\LaTeX}$  code created by `msaPrettyPrint()` or to use the output of `msaPrettyPrint()` in Sweave [6] or knitr [15] documents, the  $\text{\LaTeX}$  package `TeXshade` (file `texshade.sty`) [1] must be accessible to the  $\text{\LaTeX}$  system too. The file `texshade.sty` is shipped with the `msa` package. To determine where the file is located, enter the following command in your R session:

```
system.file("tex", "texshade.sty", package="msa")
```

Alternatively, `TeXshade` can be installed directly from the Comprehensive  $\text{\TeX}$  Archive Network (CTAN).<sup>1</sup>

---

<sup>1</sup><https://www.ctan.org/pkg/texshade>

### 3 msa for the Impatient

In order to illustrate the basic workflow, this section presents a simple example with default settings and without going into the details of each step. Let us first load amino acid sequences from one of the example files that are supplied with the `msa` package:

```
mySequenceFile <- system.file("examples", "exampleAA.fasta", package="msa")
mySequences <- readAAStringSet(mySequenceFile)
mySequences

## AAStringSet object of length 9:
##      width seq                                names
## [1]   452 MSTAVLENPGLGRKLS...NSEIGILCSALQKIK PH4H_Homo_sapiens
## [2]   453 MAAVLENGVLSRKLS...SEVGILCNALQKIKS PH4H_Rattus_norve...
## [3]   453 MAAVLENGVLSRKLS...SEVGILCHALQKIKS PH4H_Mus_musculus
## [4]   297 MNDRADFVVPDITTRK...LNAGDRQGWADTEDV PH4H_Chromobacter...
## [5]   262 MKTTQYVARQPDDNGF...RLGLHAPLFPPKQAA PH4H_Pseudomonas...
## [6]   451 MSALVLESRALGRKLS...SSEVEILCSALQKLK PH4H_Bos_taurus
## [7]   313 MAIATPTSAAPTPAPA...LNAGTREGWADTADI PH4H_Ralstonia_so...
## [8]   294 MSGDGLSNGPPPGARP...AYATAGGRLAGAAAG PH4H_Caulobacter...
## [9]   275 MSVAEYARDCAAQGLR...VARRKDKALDPATV PH4H_Rhizobium_loti
```

Now that we have loaded the sequences, we can run the `msa()` function which, by default, runs ClustalW with default parameters:

```
myFirstAlignment <- msa(mySequences)

## use default substitution matrix

myFirstAlignment

## CLUSTAL 2.1
##
## Call:
##   msa(mySequences)
##
## MsaAAMultipleAlignment with 9 rows and 456 columns
##      aln                                names
## [1] MAAVLENGVLSRKLSDF...SINSEVGILCNALQKIKS PH4H_Rattus_norve...
## [2] MAAVLENGVLSRKLSDF...SINSEVGILCHALQKIKS PH4H_Mus_musculus
## [3] MSTAVLENPGLGRKLSDF...SINSEIGILCSALQKIK- PH4H_Homo_sapiens
## [4] MSALVLESRALGRKLSDF...SISSEVEILCSALQKLK- PH4H_Bos_taurus
## [5] -----...GWADTEDV----- PH4H_Chromobacter...
## [6] -----...GWADTADI----- PH4H_Ralstonia_so...
## [7] -----...AYATAGGRLAGAAAG--- PH4H_Caulobacter...
```

```
## [8] -----...----- PH4H_Pseudomonas_...
## [9] -----...----- PH4H_Rhizobium_loti
## Con -----...??????IL??A??-- Consensus
```

Obviously, the default printing function shortens the alignment for the sake of compact output. The `print()` function provided by the `msa` package provides some ways for customizing the output, such as, showing the entire alignment split over multiple blocks of sub-sequences:

```
print(myFirstAlignment, show="complete")

##
## MsaAAMultipleAlignment with 9 rows and 456 columns
##   aln (1..39)                               names
## [1] MAAVLENGVLSRKLSDFGQETSYIEDNSNQNGAISLIF PH4H_Rattus_norve...
## [2] MAAVLENGVLSRKLSDFGQETSYIEDNSNQNGAVSLIF PH4H_Mus_musculus
## [3] MSTAVLENPGLGRKLSDFGQETSYIEDNCNQNNGAISLIF PH4H_Homo_sapiens
## [4] MSALVLESRALGRKLSDFGQETSYIEGNSDQN-AVSLIF PH4H_Bos_taurus
## [5] ----- PH4H_Chromobacter...
## [6] ----- PH4H_Ralstonia_so...
## [7] ----- PH4H_Caulobacter_...
## [8] ----- PH4H_Pseudomonas_...
## [9] ----- PH4H_Rhizobium_loti
## Con ----- Consensus
##
##   aln (40..78)                               names
## [1] SLKEEVGALAKVLRRLFEEENDINLTHIESRPSRLNKDEYE PH4H_Rattus_norve...
## [2] SLKEEVGALAKVLRRLFEEENEINLTHIESRPSRLNKDEYE PH4H_Mus_musculus
## [3] SLKEEVGALAKVLRRLFEEENDVNLTHIESRPSRLKKDEYE PH4H_Homo_sapiens
## [4] SLKEEVGALARVLRRLFEEENDINLTHIESRPSRLRKDEYE PH4H_Bos_taurus
## [5] ----- PH4H_Chromobacter...
## [6] ----- PH4H_Ralstonia_so...
## [7] ----- PH4H_Caulobacter_...
## [8] ----- PH4H_Pseudomonas_...
## [9] ----- PH4H_Rhizobium_loti
## Con ----- Consensus
##
##   aln (79..117)                              names
## [1] FFTYLDKRTKPVLSGSIKSLRNDIGATVHELSDRDKKNT PH4H_Rattus_norve...
## [2] FFTYLDKRSKPVLSGSIKSLRNDIGATVHELSDRDKKNT PH4H_Mus_musculus
## [3] FFTHLDKRSLPALTNIKILRHDIGATVHELSDRDKKDT PH4H_Homo_sapiens
## [4] FFTNLDQRSVPALANIKILRHDIGATVHELSDRDKKDT PH4H_Bos_taurus
## [5] ----- PH4H_Chromobacter...
## [6] ----- PH4H_Ralstonia_so...
## [7] ----- PH4H_Caulobacter_...
## [8] ----- PH4H_Pseudomonas_...
## [9] ----- PH4H_Rhizobium_loti
```

```

## Con ----- Consensus
##
##     aln (118..156)                               names
## [1] VPWFPRTIQELDRFANQILSYGAELDADHPGFKDPVYRA PH4H_Rattus_norve...
## [2] VPWFPRTIQELDRFANQILSYGAELDADHPGFKDPVYRA PH4H_Mus_musculus
## [3] VPWFPRTIQELDRFANQILSYGAELDADHPGFKDPVYRA PH4H_Homo_sapiens
## [4] VPWFPRTIQELDNFANQVLSYGAELDADHPGFKDPVYRA PH4H_Bos_taurus
## [5] -----MNDRADFVVPD-----ITTRKNVG PH4H_Chromobacter...
## [6] -----MAIATPTSAAPTAPAGFTGTLTDKLEQ PH4H_Ralstonia_so...
## [7] -----MSG-----DGLSNG PH4H_Caulobacter_...
## [8] -----MKTQY PH4H_Pseudomonas_...
## [9] -----MSVAEYAR-----DCAAQG PH4H_Rhizobium_loti
## Con -----?????????Y????D?????D????? Consensus
##
##     aln (157..195)                               names
## [1] RRKQFADIAYNYRHGQPIPRVEYTEEEKQTWGTVFRTLK PH4H_Rattus_norve...
## [2] RRKQFADIAYNYRHGQPIPRVEYTEEEKKTWGTVFRTLK PH4H_Mus_musculus
## [3] RRKQFADIAYNYRHGQPIPRVEYMEEEKKTWGTVFRTLK PH4H_Homo_sapiens
## [4] RRKQFADIAYNYRHGQPIPRVEYTEEEKKTWGTVFRTLK PH4H_Bos_taurus
## [5] LSHDAN-----DFTLPQPLDRYSAEDHATWATLYQRQC PH4H_Chromobacter...
## [6] FAEGLDGQTLRDPDFMEQPVHRYTAADHATWRTLYDRQE PH4H_Ralstonia_so...
## [7] PPPGAR-----PDWTIDQGWEYTYTQAEHDVWITLYERQT PH4H_Caulobacter_...
## [8] VARQPD-----DNGFIHYPETEHQVWNTLITRQL PH4H_Pseudomonas_...
## [9] LRGDYS--VCRADFTVAQDYD--YSDEEQAVWRTLCDRQT PH4H_Rhizobium_loti
## Con ?R?Q????????????P?P???YTEEE??TW?TL??RQ? Consensus
##
##     aln (196..234)                               names
## [1] ALYKTHACYEHNHIFPLLEKYCGFREDNIPQLEDVSQFL PH4H_Rattus_norve...
## [2] ALYKTHACYEHNHIFPLLEKYCGFREDNIPQLEDVSQFL PH4H_Mus_musculus
## [3] SLYKTHACYEHNHIFPLLEKYCGFHEDNIPQLEDVSQFL PH4H_Homo_sapiens
## [4] SLYKTHACYEHNHIFPLLEKYCGFREDNIPQLEEVSQFL PH4H_Bos_taurus
## [5] KLLPGRACDEFMEGL----ERLEVDADRVPDFNKLNQKL PH4H_Chromobacter...
## [6] ALLPGRACDEFQGL----STLGMSREGVPSFDRLNETL PH4H_Ralstonia_so...
## [7] DMLHGRACDEFMRGL----DALDLHRSGIPDFARINEEL PH4H_Caulobacter_...
## [8] KVIEGRACQEYLDGI----EQLGLPHERIPQLDEINRVL PH4H_Pseudomonas_...
## [9] KLTRKLAHHSYLDGV----EKLGL-LDRIPDFEDVSTKL PH4H_Rhizobium_loti
## Con ?L????AC?E???G?----??LG???D?IPQLE?VSQ?L Consensus
##
##     aln (235..273)                               names
## [1] QTCTGFRLRPVAGLLSSRDFLGGLAFRVFHCTQYIRHGS PH4H_Rattus_norve...
## [2] QTCTGFRLRPVAGLLSSRDFLGGLAFRVFHCTQYIRHGS PH4H_Mus_musculus
## [3] QTCTGFRLRPVAGLLSSRDFLGGLAFRVFHCTQYIRHGS PH4H_Homo_sapiens
## [4] QSCTGFRLRPVAGLLSSRDFLGGLAFRVFHCTQYIRHGS PH4H_Bos_taurus
## [5] MAATGWKIVAVPGLIPDDVFFEHLANRRFPVTWVWLRPH PH4H_Chromobacter...
## [6] MRATGWQIVAVPGLVPDEVFFEHLANRRFPASWWMRRPD PH4H_Ralstonia_so...
## [7] KRLTGWTVAVPGLVPDDVFFDHLANRRFPAGQFIRKPH PH4H_Caulobacter_...

```

```

## [8] QATTGWRVARVPALIPFQTFPELLASQQFPVATFIRTPE PH4H_Pseudomonas_...
## [9] RKLTGWEIIAVPGLIPAAPFFDHLANRRFPVTNWLRTQ PH4H_Rhizobium_loti
## Con Q??TGWR??VPGL?P??FF??LA?R?FP?TQ?IR??? Consensus
##
##     aln (274..312)                                names
## [1] KPMYTPEPDICHELLGHVPLFSDRSFAQFSQEIG-LASL PH4H_Rattus_norve...
## [2] KPMYTPEPDICHELLGHVPLFSDRSFAQFSQEIG-LASL PH4H_Mus_musculus
## [3] KPMYTPEPDICHELLGHVPLFSDRSFAQFSQEIG-LASL PH4H_Homo_sapiens
## [4] KPMYTPEPDICHELLGHVPLFSDRSFAQFSQEIG-LASL PH4H_Bos_taurus
## [5] QLDYLQEPDVFHDLFGHVPLLINPVFADYLEAYGKGGVK PH4H_Chromobacter...
## [6] QLDYLQEPDGFHDIFGHVPLLINPVFADYMQAYQGGLK PH4H_Ralstonia_so...
## [7] ELDYLQEPDIFHDVFGHVPLMLTDPVFADYMQAYGEGRR PH4H_Caulobacter_...
## [8] ELDYLQEPDIFHEIFGHCPLLTNPWFAEFTHTYKGLGLK PH4H_Pseudomonas_...
## [9] ELDYIVEPDMFHDFFGHVPVLSQPVFADFMQMYGKKAGD PH4H_Rhizobium_loti
## Con ?LDY??EPDIFHELFGHVPLSDP?FA?F?Q?YG?LA?? Consensus
##
##     aln (313..351)                                names
## [1] GAPDEYIEKLATIIYWFTVEFGLCKEG-DSIKAYGAGLLS PH4H_Rattus_norve...
## [2] GAPDEYIEKLATIIYWFTVEFGLCKEG-DSIKAYGAGLLS PH4H_Mus_musculus
## [3] GAPDEYIEKLATIIYWFTVEFGLCKQG-DSIKAYGAGLLS PH4H_Homo_sapiens
## [4] GAPDEYIEKLATIIYWFTVEFGLCKQG-DSIKAYGAGLLS PH4H_Bos_taurus
## [5] AKALGALPMLARLYWYTVEFGLINTP-AGMRIYGAGILS PH4H_Chromobacter...
## [6] AARLGALDMLARLYWYTVEFGLIRTP-AGLRIYGAGIVS PH4H_Ralstonia_so...
## [7] ALGLGRLANLARLYWYTVEFGLMNTP-AGLRIYGAGIVS PH4H_Caulobacter_...
## [8] ASKE-ERVFLARLYWMTIEFGLVETD-QGKRIYGGGILS PH4H_Pseudomonas_...
## [9] IIALGDEMITRLYWYTAEYGLVQEAGQPLKAFGAGLMS PH4H_Rhizobium_loti
## Con ?A?????E?LARLYW?TVEFGL????-???KAYGAGLLS Consensus
##
##     aln (352..390)                                names
## [1] SFGELQYCLSD-KPKLLPLELEKTACQEYSVTEFQPLY Y PH4H_Rattus_norve...
## [2] SFGELQYCLSD-KPKLLPLELEKTACQEYTVTEFQPLY Y PH4H_Mus_musculus
## [3] SFGELQYCLSE-KPKLLPLELEKTAIQNYTVTEFQPLY Y PH4H_Homo_sapiens
## [4] SFGELQYCLSD-KPKLLPLELEKTAVQEYTITEFQPLY Y PH4H_Bos_taurus
## [5] SKSESIYCLDSASPNRVGFDLMRIMNTRYRIDTFQKTYF PH4H_Chromobacter...
## [6] SKSESVYALDSASPNRIGFDVHRIMRTRYRIDTFQKTYF PH4H_Ralstonia_so...
## [7] SRTESIFALDDPSNRI GFDLERVMRTLYRIDDFQQVYF PH4H_Caulobacter_...
## [8] SPKETVYSLSD-EPLHQAFNPLEAMRTPYRIDILQPLYF PH4H_Pseudomonas_...
## [9] SFTELQFAVEGKDAHHPFDLETVMRTGYEIDKFQRAYF PH4H_Rhizobium_loti
## Con SF?ELQYCLSD-?P??PF?LE??M?T?Y?ID?FQPLYF Consensus
##
##     aln (391..429)                                names
## [1] VAESFSDAKEKVRTFAATIPRPFSVRYDPYTQRVEVL DN PH4H_Rattus_norve...
## [2] VAESFNDAKEKVRTFAATIPRPFSVRYDPYTQRVEVL DN PH4H_Mus_musculus
## [3] VAESFNDAKEKVRNFAATIPRPFSVRYDPYTQRIEVL DN PH4H_Homo_sapiens
## [4] VAESFNDAKEKVRNFAATIPRPFSVHYDPYTQRIEVL DN PH4H_Bos_taurus
## [5] VIDSFKQLFDATA-PDFAPLYLQLADAQPWGAGDVAPDD PH4H_Chromobacter...

```



```

## [6] VIDSFELFDATR-PDFTPLYEALGTLPTFGAGDVVDGD PH4H_Ralstonia_so...
## [7] VIDSIQTLQEVTL-RDFGAIYERLASVSDIGVAEIVPGD PH4H_Caulobacter_...
## [8] VLPDLKRLFQLAQ-EDIMALVHEAMRLG-LHAPLFPPKQ PH4H_Pseudomonas_...
## [9] VLPSFDALRDAFQTADFEAIVARRKDQKALDPATV---- PH4H_Rhizobium_loti
## Con V??SF??L?E??R??D?T????????????P??????V?D? Consensus
##
##     aln (430..456)                names
## [1] TQQLKILADSINSEVGILCNALQKIKS PH4H_Rattus_norve...
## [2] TQQLKILADSINSEVGILCHALQKIKS PH4H_Mus_musculus
## [3] TQQLKILADSINSEIGILCSALQKIK- PH4H_Homo_sapiens
## [4] TQQLKILADSISSVEILCSALQKIK- PH4H_Bos_taurus
## [5] LVLNAGDRQGWADTEDV----- PH4H_Chromobacter...
## [6] AVLNAGTREGWADTADI----- PH4H_Ralstonia_so...
## [7] AVLTRGT-QAYATAGGRLAGAAAAG--- PH4H_Caulobacter_...
## [8] AA----- PH4H_Pseudomonas_...
## [9] ----- PH4H_Rhizobium_loti
## Con ?????????????????IL??A???--- Consensus

```

The `msa` package additionally offers the function `msaPrettyPrint()` which allows for pretty-printing multiple alignments using the  $\text{\LaTeX}$  package `TeXshade`. As an example, the following R code creates a PDF file `myFirstAlignment.pdf` which is shown in Figure 1:

```

msaPrettyPrint(myFirstAlignment, output="pdf", showNames="none",
               showLogo="none", askForOverwrite=FALSE, verbose=FALSE)

```

In the above call to `msaPrettyPrint()`, the printing of sequence names has been suppressed by `showNames="none"`. The settings `askForOverwrite=FALSE` and `verbose=FALSE` are necessary for building this vignette, but, in an interactive R session, they are not necessary.

Almost needless to say, the file names created by `msaPrettyPrint()` are customizable. By default, the name of the argument is taken as file name. More importantly, the actual output of `msaPrettyPrint()` is highly customizable, too. For more details, see the Section 7 and the help page of the function (`?msaPrettyPrint`).

The `msaPrettyPrint()` function is particularly useful for pretty-printing multiple sequence alignments in Sweave [6] or knitr [15] documents. More details are provided in Section 7. Here, we restrict to a teasing example:

```

msaPrettyPrint(myFirstAlignment, y=c(164, 213), output="asis",
               showNames="none", showLogo="none", askForOverwrite=FALSE)

```

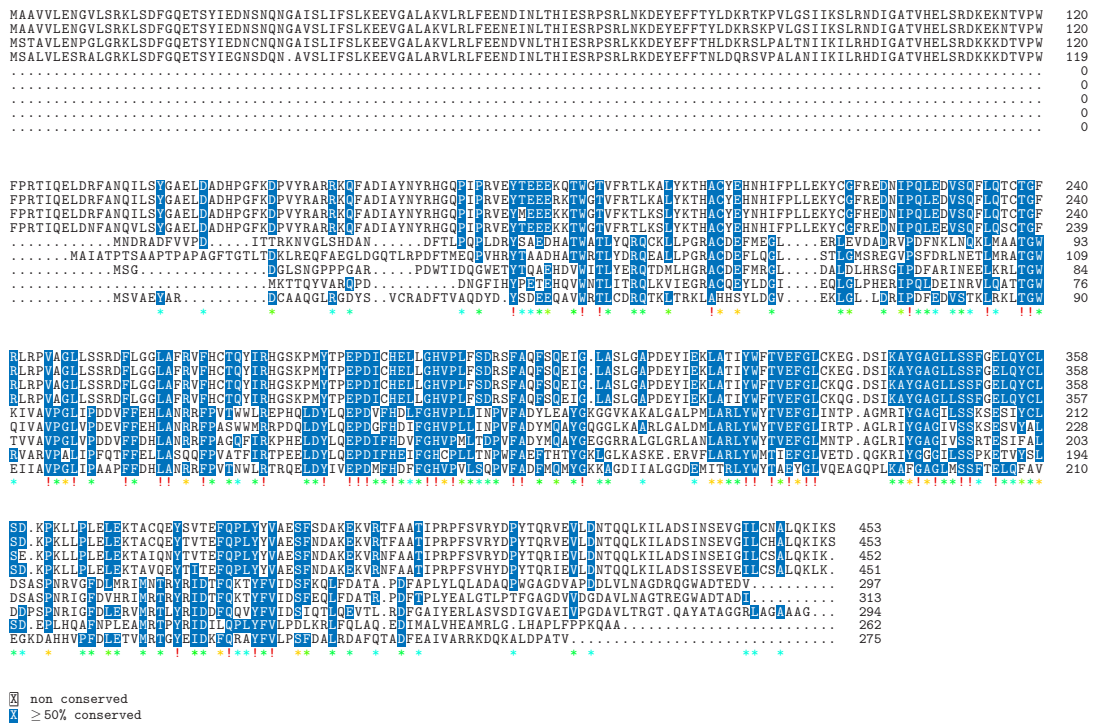


Figure 1: The PDF file myfirstAlignment.pdf created with msaPrettyPrint().

```

IAYNYRHGQPIPRVEYTEEEKQTWGTVFRTLKALYKTHACYEHNHIFPLL 213
IAYNYRHGQPIPRVEYTEEERKTWGTVFRTLKALYKTHACYEHNHIFPLL 213
IAYNYRHGQPIPRVEYMEEEEKKTWGTVFRTLKSLYKTHACYEYNHIFPLL 213
IAYNYRHGQPIPRVEYTEEEKKTWGTVFRTLKSLYKTHACYEHNHIFPLL 212
.....DFTLPQPLDRYSAEDHATWATLYQRQCKLLPGRACDEFMEGL... 67
QTLRPDFTMEQPVHRYTAADHATWRTL YDRQEALLPGRACDEFLLQGL... 83
....PDWTIDQGWEYTYQAEDHWITLYERQTDMLHGRACDEFMRGL... 58
.....DNGFIHYPETEHQVWNITLITRQLKVIIEGRACQEYLDGI... 50
.VCRADFTVAQDYD.YSDEEQAVWRTLCDRQTKLTRKLAHHSYLDGV... 65
***** * * * ****!* ** * * * * * * * * * * * * * * * *

```

X non-conserved  
X  $\geq 50\%$  conserved

## 4 Functions for Multiple Sequence Alignment in More Detail

The example in Section 3 above simply called the function `msa()` without any additional arguments. We mentioned already that, in this case, ClustalW is called with default parameters. We can also explicitly request ClustalW or one of the two other algorithms ClustalOmega or MUSCLE:

```

myClustalWAlignment <- msa(mySequences, "ClustalW")

## use default substitution matrix

myClustalWAlignment

## CLUSTAL 2.1
##
## Call:
##   msa(mySequences, "ClustalW")
##
## MsaAAMultipleAlignment with 9 rows and 456 columns
##   aln                                     names
## [1] MAAVLENGVLSRKLSDF...SINSEVGILCNALQKIKS PH4H_Rattus_norve...
## [2] MAAVLENGVLSRKLSDF...SINSEVGILCHALQKIKS PH4H_Mus_musculus
## [3] MSTAVLENPGLGRKLSDF...SINSEIGILCSALQKIK- PH4H_Homo_sapiens
## [4] MSALVLESRALGRKLSDF...SISSEVEILCSALQKIK- PH4H_Bos_taurus
## [5] -----...GWADTEDV----- PH4H_Chromobacter...
## [6] -----...GWADTADI----- PH4H_Ralstonia_so...
## [7] -----...AYATAGGRLAGAAAG--- PH4H_Caulobacter_...
## [8] -----...----- PH4H_Pseudomonas_...
## [9] -----...----- PH4H_Rhizobium_loti
## Con -----...???????IL??A???--- Consensus

myClustalOmegaAlignment <- msa(mySequences, "ClustalOmega")

```

```

## using Gonnet

myClustalOmegaAlignment

## ClustalOmega 1.2.0
##
## Call:
##   msa(mySequences, "ClustalOmega")
##
## MsaAAMultipleAlignment with 9 rows and 467 columns
##   aln                                     names
## [1] MSALVLESRALGRKLSDF...SISSEVEILCSALQKLK- PH4H_Bos_taurus
## [2] MSTAVLENPGLGRKLSDF...SINSEIGILCSALQKIK- PH4H_Homo_sapiens
## [3] MAAVLENGVLSRKLSD...SINSEVILCNALQKIKS PH4H_Rattus_norve...
## [4] MAAVLENGVLSRKLSD...SINSEVILCNALQKIKS PH4H_Mus_musculus
## [5] -----...----- PH4H_Pseudomonas_...
## [6] -----...----- PH4H_Rhizobium_loti
## [7] -----...LAGAAAG----- PH4H_Caulobacter_...
## [8] -----...V----- PH4H_Chromobacter...
## [9] -----...I----- PH4H_Ralstonia_so...
## Con -----...???????----- Consensus

myMuscleAlignment <- msa(mySequences, "Muscle")
myMuscleAlignment

## MUSCLE 3.8.31
##
## Call:
##   msa(mySequences, "Muscle")
##
## MsaAAMultipleAlignment with 9 rows and 460 columns
##   aln                                     names
## [1] MAAVLENGVLSRKLSD...SINSEVILCNALQKIKS PH4H_Rattus_norve...
## [2] MAAVLENGVLSRKLSD...SINSEVILCNALQKIKS PH4H_Mus_musculus
## [3] MSTAVLENPGLGRKLSDF...SINSEIGILCSALQKIK- PH4H_Homo_sapiens
## [4] MSALVLESRALGRKLSDF...SISSEVEILCSALQKLK- PH4H_Bos_taurus
## [5] -----...----- PH4H_Pseudomonas_...
## [6] -----...----- PH4H_Rhizobium_loti
## [7] -----...AYATAGGRLAGAAAG--- PH4H_Caulobacter_...
## [8] -----MNDRADF...QGWADTEDV----- PH4H_Chromobacter...
## [9] MAIATPTSAAPTAPAGF...EGWADTADI----- PH4H_Ralstonia_so...
## Con M????????????????DF...????????L??A???--- Consensus

```

Please note that the call `msa(mySequences, "ClustalW", ...)` is just a shortcut for the call `msaClustalW(mySequences, ...)`, analogously for `msaClustalOmega()` and `msaMuscle()`.

In other words, `msa()` is nothing else but a wrapper function that provides a unified interface to the three functions `msaClustalW()`, `msaClustalOmega()`, and `msaMuscle()`.

All three functions `msaClustalW()`, `msaClustalOmega()`, and `msaMuscle()` have the same parameters: The input sequences are passed as argument `inputSeqs`, and all functions have the following arguments: `cluster`, `gapOpening`, `gapExtension`, `maxiters`, `substitutionMatrix`, `order`, `type`, and `verbose`. The ways these parameters are interpreted, are largely analogous, although there are some differences, also in terms of default values. See the subsections below and the man page of the three functions for more details. All of the three functions `msaClustalW()`, `msaClustalOmega()`, and `msaMuscle()`, however, are not restricted to the parameters mentioned above. All three have a `'...'` argument through which several other algorithm-specific parameters can be passed on to the underlying library. The following subsections provide an overview of which parameters are supported by each of the three algorithms.

#### 4.1 ClustalW-Specific Parameters

The original implementation of ClustalW offers a lot of parameters for customizing the way a multiple sequence alignment is computed. Through the `'...'` argument, `msaClustalW()` provides an interface to make use of most these parameters (see the documentation of ClustalW<sup>2</sup> for a comprehensive overview). Currently, the following restrictions and caveats apply:

- The parameters `infile`, `clustering`, `gapOpen`, `gapExt`, `numiters`, `matrix`, and `outorder` have been renamed to the standardized argument names `inputSeqs`, `cluster`, `gapOpening`, `gapExtension`, `maxiters`, `substitutionMatrix`, and `order` in order to provide a consistent interface for all three multiple sequence alignment algorithms.
- Boolean flags must be passed as logical values, e.g. `verbose=TRUE`.
- The parameter `quiet` has been replaced by `verbose` (with the exact opposite meaning).
- The following parameters are (currently) not supported: `bootstrap`, `check`, `fullhelp`, `interactive`, `maxseqLen`, `options`, and `tree`.
- For the parameter `output`, only the choice `"clustal"` is available.

#### 4.2 ClustalOmega-Specific Parameters

In the same way as ClustalW, the original implementation of ClustalOmega also offers a lot of parameters for customizing the way a multiple sequence alignment is computed. Through the `'...'` argument, `msaClustalOmega()` provides an interface to make use of most these parameters (see the documentation of ClustalOmega<sup>3</sup> for a comprehensive overview). Currently, the following restrictions and caveats apply:

- The parameters `infile`, `cluster-size`, `iterations`, and `output-order` have been renamed to the argument names `inputSeqs`, `cluster`, `maxiters`, and `order` in order to provide a consistent interface for all three multiple sequence alignment algorithms.

<sup>2</sup>[http://www.clustal.org/download/clustalw\\_help.txt](http://www.clustal.org/download/clustalw_help.txt)

<sup>3</sup><http://www.clustal.org/omega/README>

- ClustalOmega does not allow for setting custom gap penalties. Therefore, setting the parameters `gapOpening` and `gapExtension` currently has no effect and will lead to a warning. These arguments are only defined for future extensions and consistency with the other algorithms available in `msa`.
- ClustalOmega only allows for choosing substitution matrices from a pre-defined set of names, namely "BLOSUM30", "BLOSUM40", "BLOSUM50", "BLOSUM65", "BLOSUM80", and "Gonnet". This is a new feature — the original ClustalOmega implementation does not allow for using any custom substitution matrix. However, since these are all amino acid substitution matrices, ClustalOmega is still hardly useful for multiple alignments of nucleotide sequences.
- Boolean flags must be passed as logical values, e.g. `verbose=TRUE`.
- The following parameters are (currently) not supported: `maxSeqLength` and `help`.
- For the parameter `outFmt`, only the choice "clustal" is available.

### 4.3 MUSCLE-Specific Parameters

Finally, also MUSCLE offers a lot of parameters for customizing the way a multiple sequence alignment is computed. Through the `'...'` argument, `msaMuscle()` provides an interface to make use of most these parameters (see the documentation of MUSCLE<sup>4</sup> for a comprehensive overview). Currently, the following restrictions and caveats apply:

- The parameters `in`, `gapOpen`, `gapExtend`, `matrix`, and `seqtype` have been renamed to `inputSeqs`, `gapOpening`, `gapExtension`, `substitutionMatrix` and `type` in order to provide a consistent interface for all three multiple sequence alignment algorithms.
- Boolean flags must be passed as logical values, e.g. `verbose=TRUE`.
- The parameter `quiet` has been replaced by `verbose` (with the exact opposite meaning).
- The following parameters are currently not supported: `clw`, `clwstrict`, `fastaout`, `group`, `html`, `in1`, `in2`, `log`, `loga`, `msaout`, `msf`, `out`, `phyi`, `phyiout`, `phys`, `physout`, `refine`, `refinew`, `scorefile`, `spscore`, `stable`, `termgaps4`, `termgapsfull`, `termgapshalf`, `termgapshalflonger`, `tree1`, `tree2`, `usetree`, `weight1`, and `weight2`.

## 5 Printing Multiple Sequence Alignments

As already shown above, multiple sequence alignments can be shown in plain text format on the R console using the `print()` function (which is implicitly called if just the object name is entered on the R console). This function allows for multiple customizations, such as, enabling/disabling to display a consensus sequence, printing the entire alignment or only a subset, enabling/disabling to display sequence names, and adjusting the width allocated for sequence names. For more information, the reader is referred to the help page of the `print` function:

<sup>4</sup><http://www.drive5.com/muscle/muscle.html>

```
help("print,MsaDNAMultipleAlignment-method")
```

We only provide some examples here:

```
print(myFirstAlignment)
```

```
## CLUSTAL 2.1
##
## Call:
##   msa(mySequences)
##
## MsaAAMultipleAlignment with 9 rows and 456 columns
##   aln                               names
## [1] MAAVLENGVLSRKLSDF...SINSEVGILCNALQKIKS PH4H_Rattus_norve...
## [2] MAAVLENGVLSRKLSDF...SINSEVGILCHALQKIKS PH4H_Mus_musculus
## [3] MSTAVLENPGLGRKLSDF...SINSEIGILCSALQKIK- PH4H_Homo_sapiens
## [4] MSALVLESRALGRKLSDF...SISSEVEILCSALQKLK- PH4H_Bos_taurus
## [5] -----...GWADTEDV----- PH4H_Chromobacter...
## [6] -----...GWADTADI----- PH4H_Ralstonia_so...
## [7] -----...AYATAGGRLAGAAAG--- PH4H_Caulobacter_...
## [8] -----...----- PH4H_Pseudomonas_...
## [9] -----...----- PH4H_Rhizobium_loti
## Con -----...???????IL??A???--- Consensus
```

```
print(myFirstAlignment, show="complete")
```

```
##
## MsaAAMultipleAlignment with 9 rows and 456 columns
##   aln (1..39)                               names
## [1] MAAVLENGVLSRKLSDFGQETSYIEDNSNQNGAISLIF PH4H_Rattus_norve...
## [2] MAAVLENGVLSRKLSDFGQETSYIEDNSNQNGAVSLIF PH4H_Mus_musculus
## [3] MSTAVLENPGLGRKLSDFGQETSYIEDNCNQNNGAISLIF PH4H_Homo_sapiens
## [4] MSALVLESRALGRKLSDFGQETSYIEGNSDQN-AVSLIF PH4H_Bos_taurus
## [5] ----- PH4H_Chromobacter...
## [6] ----- PH4H_Ralstonia_so...
## [7] ----- PH4H_Caulobacter_...
## [8] ----- PH4H_Pseudomonas_...
## [9] ----- PH4H_Rhizobium_loti
## Con ----- Consensus
##
##   aln (40..78)                               names
## [1] SLKEEVGALAKVLRRLFEEENDINLTHIESRPSRLNKDEYE PH4H_Rattus_norve...
## [2] SLKEEVGALAKVLRRLFEEENEINLTHIESRPSRLNKDEYE PH4H_Mus_musculus
## [3] SLKEEVGALAKVLRRLFEEENDVNLTHIESRPSRLKKDEYE PH4H_Homo_sapiens
## [4] SLKEEVGALARVLRRLFEEENDINLTHIESRPSRLRKDEYE PH4H_Bos_taurus
```

```

## [5] ----- PH4H_Chromobacter...
## [6] ----- PH4H_Ralstonia_so...
## [7] ----- PH4H_Caulobacter_...
## [8] ----- PH4H_Pseudomonas_...
## [9] ----- PH4H_Rhizobium_loti
## Con ----- Consensus
##
##      aln (79..117)                                names
## [1] FFTYLDKRTKPVLSGIKSLRNDIGATVHELSDKEKNT PH4H_Rattus_norve...
## [2] FFTYLDKRSKPVLGSIKSLRNDIGATVHELSDKEKNT PH4H_Mus_musculus
## [3] FFTHLDRSLPALTNIKILRHDIGATVHELSDKKKDT PH4H_Homo_sapiens
## [4] FFTNLDQRSVPALANI KILRHDIGATVHELSDKKKDT PH4H_Bos_taurus
## [5] ----- PH4H_Chromobacter...
## [6] ----- PH4H_Ralstonia_so...
## [7] ----- PH4H_Caulobacter_...
## [8] ----- PH4H_Pseudomonas_...
## [9] ----- PH4H_Rhizobium_loti
## Con ----- Consensus
##
##      aln (118..156)                                names
## [1] VPWFPRTIQELDRFANQILSYGAELDADHPGFKDPVYRA PH4H_Rattus_norve...
## [2] VPWFPRTIQELDRFANQILSYGAELDADHPGFKDPVYRA PH4H_Mus_musculus
## [3] VPWFPRTIQELDRFANQILSYGAELDADHPGFKDPVYRA PH4H_Homo_sapiens
## [4] VPWFPRTIQELDNFANQVLSYGAELDADHPGFKDPVYRA PH4H_Bos_taurus
## [5] -----MNDRADFVVPD-----ITTRKNVG PH4H_Chromobacter...
## [6] -----MAIATPTSAAPTPAPAGFTGTLTDKLEQ PH4H_Ralstonia_so...
## [7] -----MSG-----DGLSNG PH4H_Caulobacter_...
## [8] -----MKTQY PH4H_Pseudomonas_...
## [9] -----MSVAEYAR-----DCAAQG PH4H_Rhizobium_loti
## Con -----?????????Y?????D?????????D????? Consensus
##
##      aln (157..195)                                names
## [1] RRKQFADIAYNRHHGQPIPRVEYTEEEKQWGTWVFRSLK PH4H_Rattus_norve...
## [2] RRKQFADIAYNRHHGQPIPRVEYTEEEKQWGTWVFRSLK PH4H_Mus_musculus
## [3] RRKQFADIAYNRHHGQPIPRVEYMEEEKQWGTWVFKSLK PH4H_Homo_sapiens
## [4] RRKQFADIAYNRHHGQPIPRVEYTEEEKQWGTWVFRSLK PH4H_Bos_taurus
## [5] LSHDAN-----DFTLPQPLDRYSAEDHATWATLYQRQC PH4H_Chromobacter...
## [6] FAEGLDGQTLRPDFTMEQPVHRYTAADHATWRTLYDRQE PH4H_Ralstonia_so...
## [7] PPPGAR-----PDWTIDQGWETYTQAEDVWITLYERQT PH4H_Caulobacter_...
## [8] VARQPD-----DNGFIHYPETEHQVWNTLITRQL PH4H_Pseudomonas_...
## [9] LRGDYS--VCRADFTVAQDYD--YSDEEQAVWRTLCDRQT PH4H_Rhizobium_loti
## Con ?R?Q?????????????P?P???YTEEE??TW?TL??RQ? Consensus
##
##      aln (196..234)                                names
## [1] ALYKTHACYEHNHIFPILLEKYCGFREDNIPQLEDVVSQFL PH4H_Rattus_norve...
## [2] ALYKTHACYEHNHIFPILLEKYCGFREDNIPQLEDVVSQFL PH4H_Mus_musculus

```



```

## [3] SLYKTHACYEYNHIFPILLEKYCGFHEDNIPQLEDVSQFL PH4H_Homo_sapiens
## [4] SLYKTHACYEYNHIFPILLEKYCGFREDNIPQLEEVSQFL PH4H_Bos_taurus
## [5] KLLPGRACDEFMEGL-----ERLEVDADRVPDFNKLNQKL PH4H_Chromobacter...
## [6] ALLPGRACDEFQGL-----STLGMSREGVPSFDRLNETL PH4H_Ralstonia_so...
## [7] DMLHGRACDEFMRGL----DALDLHRSGIPDFARINEEL PH4H_Caulobacter_...
## [8] KVIEGRACQEYLDGI----EQLGLPHERIPQLDEINRVL PH4H_Pseudomonas_...
## [9] KLTRKLAHHSYLDGV----EKLGL-LDRIPDFEDVSTKL PH4H_Rhizobium_loti
## Con ?L????AC?E???G?----??LG???D?IPQLE?VSQ?L Consensus
##
##      aln (235..273)                                names
## [1] QTCTGFRLRPVAGLLSSRDFLGGLAFRVFHCTQYIRHGS PH4H_Rattus_norve...
## [2] QTCTGFRLRPVAGLLSSRDFLGGLAFRVFHCTQYIRHGS PH4H_Mus_musculus
## [3] QTCTGFRLRPVAGLLSSRDFLGGLAFRVFHCTQYIRHGS PH4H_Homo_sapiens
## [4] QSCTGFRLRPVAGLLSSRDFLGGLAFRVFHCTQYIRHGS PH4H_Bos_taurus
## [5] MAATGWKIVAVPGLIPDDVFFEHLANRRFPVTWWLREPH PH4H_Chromobacter...
## [6] MRATGWQIVAVPGLVPDEVFFEHLANRRFPASWWMRRPD PH4H_Ralstonia_so...
## [7] KRLTGWTVVAVPGLVPDDVFFDHLANRRFPAGQFIRKPH PH4H_Caulobacter_...
## [8] QATTGWRVARVPALIPFQTFELLASQQFPVATFIRTPE PH4H_Pseudomonas_...
## [9] RKLTGWEIIAVPGLIPAAPFFDHLANRRFPVTNWLRTQ PH4H_Rhizobium_loti
## Con Q???TGWR???VPGL?P???FF??LA?R?FP?TQ?IR??? Consensus
##
##      aln (274..312)                                names
## [1] KPMYTPEPDICHELLGHVPLFSDRSFAQFSQEIG-LASL PH4H_Rattus_norve...
## [2] KPMYTPEPDICHELLGHVPLFSDRSFAQFSQEIG-LASL PH4H_Mus_musculus
## [3] KPMYTPEPDICHELLGHVPLFSDRSFAQFSQEIG-LASL PH4H_Homo_sapiens
## [4] KPMYTPEPDICHELLGHVPLFSDRSFAQFSQEIG-LASL PH4H_Bos_taurus
## [5] QLDYLQEPDVFHDLFGHVPLLINPVFADYLEAYGKGGVK PH4H_Chromobacter...
## [6] QLDYLQEPDGFHDIFGHVPLLINPVFADYMQAYGQGGLK PH4H_Ralstonia_so...
## [7] ELDYLQEPDIFHDVFGHVPLMTDPVFADYMQAYGEGGRR PH4H_Caulobacter_...
## [8] ELDYLQEPDIFHEIFGHCPLLTNPWFAEFTHTYKGLGLK PH4H_Pseudomonas_...
## [9] ELDYIVEPDMFHDFFGHVPLSQPVFADFMQMYGKKAGD PH4H_Rhizobium_loti
## Con ?LDY??EPDIFHELFGHVPLSDP?FA?F?Q?YG?LA?? Consensus
##
##      aln (313..351)                                names
## [1] GAPDEYIEKLATIIYWFTVEFGLCKEG-DSIKAYGAGLLS PH4H_Rattus_norve...
## [2] GAPDEYIEKLATIIYWFTVEFGLCKEG-DSIKAYGAGLLS PH4H_Mus_musculus
## [3] GAPDEYIEKLATIIYWFTVEFGLCKQG-DSIKAYGAGLLS PH4H_Homo_sapiens
## [4] GAPDEYIEKLATIIYWFTVEFGLCKQG-DSIKAYGAGLLS PH4H_Bos_taurus
## [5] AKALGALPMLARLYWYTVEFGLINTP-AGMRIYGAGILS PH4H_Chromobacter...
## [6] AARLGALDMLARLYWYTVEFGLIRTP-AGLRIYGAGIVS PH4H_Ralstonia_so...
## [7] ALGLGRLANLARLYWYTVEFGLMNTP-AGLRIYGAGIVS PH4H_Caulobacter_...
## [8] ASKE-ERVFLARLYWMTIEFGLVETD-QGKRIYGGGILS PH4H_Pseudomonas_...
## [9] IIALGGDEMITRLYWYTAEYGLVQEAGQPLKAFGAGLMS PH4H_Rhizobium_loti
## Con ?A?????E?LARLYW?TVEFGL????-???KAYGAGLLS Consensus
##
##      aln (352..390)                                names

```

```

## [1] SFGELQYCLSD-KPKLLPLELEKTACQEYSVTEFQPLY PH4H_Rattus_norve...
## [2] SFGELQYCLSD-KPKLLPLELEKTACQEYTVTEFQPLY PH4H_Mus_musculus
## [3] SFGELQYCLSE-KPKLLPLELEKTAIQNYTVTEFQPLY PH4H_Homo_sapiens
## [4] SFGELQYCLSD-KPKLLPLELEKTAVQEYTITEFQPLY PH4H_Bos_taurus
## [5] SKSESIYCLDSASPNRVGFDMRIMNTRYRIDTFQKTYF PH4H_Chromobacter...
## [6] SKSESVYALDSASPNRIGFDVHRIMRTRYRIDTFQKTYF PH4H_Ralstonia_so...
## [7] SRTESIFALDDPSPNRIGFDLERVMRTLYRIDDFQVYF PH4H_Caulobacter_...
## [8] SPKETVYSLSD-EPLHQAFNPLEAMRTPYRIDILQPLYF PH4H_Pseudomonas_...
## [9] SFTELQFAVEGKDAHHVPFDLETVMRTGYEIDKFQRAYF PH4H_Rhizobium_loti
## Con SF?ELQYCLSD-?P???PF?LE??M?T?Y?ID?FQPLYF Consensus
##
##      aln (391..429)                names
## [1] VAESFSDAKEKVRTFAATIPRPFVRYDPYTQRVEVLDN PH4H_Rattus_norve...
## [2] VAESFNDAKEKVRTFAATIPRPFVRYDPYTQRVEVLDN PH4H_Mus_musculus
## [3] VAESFNDAKEKVRNFAATIPRPFVRYDPYTQRIEVLDN PH4H_Homo_sapiens
## [4] VAESFNDAKEKVRNFAATIPRPFVHYDPYTQRIEVLDN PH4H_Bos_taurus
## [5] VIDSFQKLFDATA-PDFAPLYLQLADAQPWGAGDVAPDD PH4H_Chromobacter...
## [6] VIDSFQKLFDATA-PDFTPLYEALGTLPTFGAGDVVDGD PH4H_Ralstonia_so...
## [7] VIDSIQTLQEVTL-RDFGAIYERLASVSDIGVAEIVPGD PH4H_Caulobacter_...
## [8] VLPDLKRLFQLAQ-EDIMALVHEAMRLG-LHAPLFPPKQ PH4H_Pseudomonas_...
## [9] VLPSFDALRDAFQTADFEAIVARRKDQKALDPATV---- PH4H_Rhizobium_loti
## Con V??SF??L?E??R??D?T????????????P??????V?D? Consensus
##
##      aln (430..456)                names
## [1] TQQLKILADSINSEVGILCNALQKIKS PH4H_Rattus_norve...
## [2] TQQLKILADSINSEVGILCHALQKIKS PH4H_Mus_musculus
## [3] TQQLKILADSINSEIGILCSALQKIK- PH4H_Homo_sapiens
## [4] TQQLKILADSISSVEILCSALQKIK- PH4H_Bos_taurus
## [5] LVLNAGDRQGWADTEDV----- PH4H_Chromobacter...
## [6] AVLNAGTREGWADTADI----- PH4H_Ralstonia_so...
## [7] AVLTRGT-QAYATAGGRLAGAAAAG--- PH4H_Caulobacter_...
## [8] AA----- PH4H_Pseudomonas_...
## [9] ----- PH4H_Rhizobium_loti
## Con ?????????????????IL??A???--- Consensus

print(myFirstAlignment, showConsensus=FALSE, halfNrow=3)

## CLUSTAL 2.1
##
## Call:
##   msa(mySequences)
##
## MsaAAMultipleAlignment with 9 rows and 456 columns
##      aln                names
## [1] MAAVLENGVLSRKLSD...SINSEVGILCNALQKIKS PH4H_Rattus_norve...
## [2] MAAVLENGVLSRKLSD...SINSEVGILCHALQKIKS PH4H_Mus_musculus

```

```

## [3] MSTAVLENPGLGRKLSDF...SINSEIGILCSALQKIK- PH4H_Homo_sapiens
## ... ..
## [7] -----...AYATAGGRLAGAAAG--- PH4H_Caulobacter_...
## [8] -----...----- PH4H_Pseudomonas_...
## [9] -----...----- PH4H_Rhizobium_loti

print(myFirstAlignment, showNames=FALSE, show="complete")

##
## MsaAAMultipleAlignment with 9 rows and 456 columns
##     aln (1..60)
## [1] MAAVLENGVLSRKLSDFGQETSYIEDNSNQNGAISLIFSLKEEVGALAKVLRRLFEEENDI
## [2] MAAVLENGVLSRKLSDFGQETSYIEDNSNQNGAVSLIFSLKEEVGALAKVLRRLFEEENEI
## [3] MSTAVLENPGLGRKLSDFGQETSYIEDNCNQNNGAISLIFSLKEEVGALAKVLRRLFEEENDV
## [4] MSALVLESRALGRKLSDFGQETSYIEGNSDQN-AVSLIFSLKEEVGALARVLRRLFEEENDI
## [5] -----
## [6] -----
## [7] -----
## [8] -----
## [9] -----
## Con -----
##
##     aln (61..120)
## [1] NLTHIESRPSRLNKDEYEFFTYLDKRTKPVLSGSIKSLRNDIGATVHELSDRDKKNTVPW
## [2] NLTHIESRPSRLNKDEYEFFTYLDKRSKPVLSGSIKSLRNDIGATVHELSDRDKKNTVPW
## [3] NLTHIESRPSRLKKDEYEFFTHLDRSLPALTNIKILRHDIGATVHELSDRDKKNTVPW
## [4] NLTHIESRPSRLRKDEYEFFTNLDQRSVPALANI KILRHDIGATVHELSDRDKKNTVPW
## [5] -----
## [6] -----
## [7] -----
## [8] -----
## [9] -----
## Con -----
##
##     aln (121..180)
## [1] FPRTIQELDRFANQILSYGAELDADHPGFKDPVYRARRKQFADIAYNRHHGQPIPRVEYTT
## [2] FPRTIQELDRFANQILSYGAELDADHPGFKDPVYRARRKQFADIAYNRHHGQPIPRVEYTT
## [3] FPRTIQELDRFANQILSYGAELDADHPGFKDPVYRARRKQFADIAYNRHHGQPIPRVEYMT
## [4] FPRTIQELDNFANQVLSYGAELDADHPGFKDPVYRARRKQFADIAYNRHHGQPIPRVEYTT
## [5] -----MNDRADFVVPD-----ITTRKNVGLSHDAN-----DFTLPQPLDRYS
## [6] -----MAIATPTSAAPTPAPAGFTGTLTDKLRQFAEGLDGQTLRPDFTMEQPVHRYT
## [7] -----MSG-----DGLSNGPPPGAR-----PDWTIDQGWETYT
## [8] -----MKTQYVARQPD-----DNGFIHYP
## [9] -----MSVAEYAR-----DCAAQGLRGDYS--VCRADFTVAQDYD-YS
## Con -----?????????Y?????D?????????D?????????R?Q?????????????P?P????YT
##

```

```

##      a1n (181..240)
## [1] EEEKQWGTVFRTLKALYKTHACYEHNHIFPPLLEKYCGFREDNIPQLEDVVSQFLQTCTGF
## [2] EEERKTWGTVFRTLKALYKTHACYEHNHIFPPLLEKYCGFREDNIPQLEDVVSQFLQTCTGF
## [3] EEEKKTWGTVFKTLKSLYKTHACYEYHNHIFPPLLEKYCGFHEDNIPQLEDVVSQFLQTCTGF
## [4] EEEKKTWGTVFRTLKSLYKTHACYEHNHIFPPLLEKYCGFREDNIPQLEEVVSQFLQSCTGF
## [5] AEDHATWATLYQRQCKLLPGRACDEFMEGL----ERLEVDADRVPDFNKLNLQKLMAATGW
## [6] AADHATWRTLYDRQEALLPGRACDEFQLGGL----STLGMSREGVPSFDRNLNETLMRATGW
## [7] QAEHDVWITLYERQTDMLHGRACDEFMRGL----DALDLHRSGIPDFARINEELKRLTGW
## [8] ETEHQVWNTLITRQLKVIETRACQEYLDGI----EQLGLPHERIPQLDEINRVLQATTGW
## [9] DEEQAVWRTLCDRQTKLTRKLAHHSYLDGV----EKLGL-LDRIPDFEDVSTKLRKLTGW
## Con EEE??TW?TL??RQ??L????AC?E???G?----??LG???D?IPQLE?VSQ?LQ??TGW
##
##      a1n (241..300)
## [1] RLRPVAGLLSSRDFLGGLAFRVFHCTQYIRHGSKPMYTPEPDICHELLGHVPLFSDRSFA
## [2] RLRPVAGLLSSRDFLGGLAFRVFHCTQYIRHGSKPMYTPEPDICHELLGHVPLFSDRSFA
## [3] RLRPVAGLLSSRDFLGGLAFRVFHCTQYIRHGSKPMYTPEPDICHELLGHVPLFSDRSFA
## [4] RLRPVAGLLSSRDFLGGLAFRVFHCTQYIRHGSKPMYTPEPDICHELLGHVPLFSDRSFA
## [5] KIVAVPGLIPDDVFFEHLANRRFPVTWWLREPHQLDYLQEPDVFHDLFGHVPLLINPVFA
## [6] QIVAVPGLVPDEVFFEHLANRRFPASWMMRRPDQDYLQEPDGFHDIFGHVPLLINPVFA
## [7] TVVAVPGLVPDDVFFDHLANRRFPAGQFIRKPHELDYLQEPDIFHDVFGHVPMMLTDPVFA
## [8] RVARVPALIPFQTFEELLASQQFPVATFIRTPEELDYLQEPDIFHEIFGHCPLLTNPWFA
## [9] EIIAVPGLIPAAPFFDHLANRRFPVTNWLRTQELDYIVEPDMFHDFFGHVPLSQPVFA
## Con R???VPGL?P???FF??LA?R?FP?TQ?IR????LDY??EPDIFHELFGHVPLSDP?FA
##
##      a1n (301..360)
## [1] QFSQEIG-LASLGAPDEYIEKLATIIYWFTVEFGLCKEG-DSIKAYGAGLLSSFGEQYCL
## [2] QFSQEIG-LASLGAPDEYIEKLATIIYWFTVEFGLCKEG-DSIKAYGAGLLSSFGEQYCL
## [3] QFSQEIG-LASLGAPDEYIEKLATIIYWFTVEFGLCKQG-DSIKAYGAGLLSSFGEQYCL
## [4] QFSQEIG-LASLGAPDEYIEKLATIIYWFTVEFGLCKQG-DSIKAYGAGLLSSFGEQYCL
## [5] DYLEAYGKGGVKAKALGALPMLARLYWYTVEFGLINTP-AGMRIYGAGILSSKSESIYCL
## [6] DYMQAYGQGGKKAARLGALDMLARLYWYTVEFGLIRTP-AGLRIYGAGIVSSKSESVYAL
## [7] DYMQAYGEGRRALGLGRLANLARLYWYTVEFGLMNTP-AGLRIYGAGIVSSRTESIFAL
## [8] EFTHTYKGLGLKASKE-ERVFLARLYWMTIEFGLVETD-QGKRIYGGGILSSPKETVYSL
## [9] DFMQMYGKKAGDIIALGGDEMITRLYWYTAEYGLVQEAGQPLKAFGAGLMSSTELQFAV
## Con ?F?Q?YG?LA???A?????E?LARLYW?TVEFGL????-???KAYGAGLLSSF?ELQYCL
##
##      a1n (361..420)
## [1] SD-KPKLLPLELEKTACQEYSVTEFQPLYYYVAESFSDAKEKVRTFAATIPRPFVRYDPY
## [2] SD-KPKLLPLELEKTACQEYTVTEFQPLYYYVAESFNDAKEKVRTFAATIPRPFVRYDPY
## [3] SE-KPKLLPLELEKTAIQNYTVTEFQPLYYYVAESFNDAKEKVRNFAATIPRPFVRYDPY
## [4] SD-KPKLLPLELEKTAVQEYTIITEFQPLYYYVAESFNDAKEKVRNFAATIPRPFVHYDPY
## [5] DSASPNRVGFDLMRIMNTRYRIDTFQKTYFVIDSFQKLFDATA-PDFAPLYLQLADAPW
## [6] DSASPNRIGFDVHRIMRTRYRIDTFQKTYFVIDSFQKLFDATA-PDFTPLYEALGTLPTF
## [7] DDPSPNRIGFDLERVMRTLYRIDDFQQVYFVIDSIQTLQEVTL-RDFGAIYERLASVSDI
## [8] SD-EPLHQAFNPLEAMRTPYRIDILQPLYFVLPDLKRLFQLAQ-EDIMALVHEAMRLG-L
## [9] EGKDAHHPFDLETVMRTGYEIDKFRAYFVLPSPFDALRDAFQTADFEAIVARRKDKAL

```

```
## Con SD-?P???PF?LE??M?T?Y?ID?FQPLYFV??SF??L?E??R??D?T????????????P?
##
##      aln (421..456)
## [1] TQRVEVDNTQQLKILADSINSEVGILCNALQKIKS
## [2] TQRVEVDNTQQLKILADSINSEVGILCHALQKIKS
## [3] TQRIEVDNTQQLKILADSINSEIGILCSALQKIK-
## [4] TQRIEVDNTQQLKILADSISSVEILCSALQKIK-
## [5] GAGDVAPDDLVLNAGDRQGWADTEDV-----
## [6] GAGDVVDGDAVLNAGTREGWADTADI-----
## [7] GVAEIVPGDAVLTRGT-QAYATAGGRLAGAAAG---
## [8] HAPLFPPKQAA-----
## [9] DPATV-----
## Con ?????V?D????????????????????IL??A???
```

## 6 Processing Multiple Alignments

### 6.1 Methods Inherited From Biostrings

The classes defined by the `msa` package for storing multiple alignment results have been derived from the corresponding classes defined by the `Biostrings` package. Therefore, all methods for processing multiple alignments are available and work without any practical limitation. In this section, we highlight some of these.

The classes used for storing multiple alignments allow for defining masks on sequences and sequence positions via their row and column mask slots. They can be set by `rowmask()` and `colmask()` functions which serve both as setter and getter functions. To set row or column masks, an `IRanges` object must be supplied:

```
myMaskedAlignment <- myFirstAlignment
colM <- IRanges(start=1, end=100)
colmask(myMaskedAlignment) <- colM
myMaskedAlignment

## CLUSTAL 2.1
##
## Call:
##   msa(mySequences)
##
## MsaAAMultipleAlignment with 9 rows and 456 columns
##      aln                                     names
## [1] #####...SINSEVGILCNALQKIKS PH4H_Rattus_norve...
## [2] #####...SINSEVGILCHALQKIKS PH4H_Mus_musculus
## [3] #####...SINSEIGILCSALQKIK- PH4H_Homo_sapiens
## [4] #####...SISSEVEILCSALQKIK- PH4H_Bos_taurus
## [5] #####...GWADTEDV----- PH4H_Chromobacter...
```

```
## [6] #####...GWADTADI----- PH4H_Ralstonia_so...
## [7] #####...AYATAGGRLAGAAAG--- PH4H_Caulobacter_...
## [8] #####...----- PH4H_Pseudomonas_...
## [9] #####...----- PH4H_Rhizobium_loti
## Con #####...??????IL??A??- Consensus
```

The `unmasked()` allows for removing these masks, thereby casting the multiple alignment to a set of aligned Biostrings sequences (class `AAStringSet`, `DNASTringSet`, or `RNASTringSet`):

```
unmasked(myMaskedAlignment)
```

```
## AAStringSet object of length 9:
##      width seq                                names
## [1]   456 MAAVLENGVLSRKLS...SEVGILCNALQKIKS PH4H_Rattus_norve...
## [2]   456 MAAVLENGVLSRKLS...SEVGILCHALQKIKS PH4H_Mus_musculus
## [3]   456 MSTAVLENPGLGRKLS...SEIGILCSALQKIK- PH4H_Homo_sapiens
## [4]   456 MSALVLESRALGRKLS...SEVEILCSALQKIK- PH4H_Bos_taurus
## [5]   456 -----...DTEDV----- PH4H_Chromobacter...
## [6]   456 -----...DTADI----- PH4H_Ralstonia_so...
## [7]   456 -----...TAGGRLAGAAAG--- PH4H_Caulobacter_...
## [8]   456 -----...----- PH4H_Pseudomonas_...
## [9]   456 -----...----- PH4H_Rhizobium_loti
```

Consensus matrices can be computed conveniently as follows:

```
conMat <- consensusMatrix(myFirstAlignment)
dim(conMat)

## [1] 30 456

conMat[, 101:110]

##   [,1] [,2] [,3] [,4] [,5] [,6] [,7] [,8] [,9] [,10]
## A    0    0    0    4    0    0    0    0    0    0
## R    0    0    0    0    0    0    0    0    0    0
## N    0    0    0    0    0    0    0    0    0    0
## D    4    0    0    0    0    0    0    0    0    0
## C    0    0    0    0    0    0    0    0    0    0
## Q    0    0    0    0    0    0    0    0    0    0
## E    0    0    0    0    0    0    0    4    0    0
## G    0    0    4    0    0    0    0    0    0    0
## H    0    0    0    0    0    0    4    0    0    0
## I    0    4    0    0    0    0    0    0    0    0
## L    0    0    0    0    0    0    0    0    4    0
```

```
## K    0    0    0    0    0    0    0    0    0    0
## M    0    0    0    0    0    0    0    0    0    0
## F    0    0    0    0    0    0    0    0    0    0
## P    0    0    0    0    0    0    0    0    0    0
## S    0    0    0    0    0    0    0    0    0    4
## T    0    0    0    0    4    0    0    0    0    0
## W    0    0    0    0    0    0    0    0    0    0
## Y    0    0    0    0    0    0    0    0    0    0
## V    0    0    0    0    0    4    0    0    0    0
## U    0    0    0    0    0    0    0    0    0    0
## O    0    0    0    0    0    0    0    0    0    0
## B    0    0    0    0    0    0    0    0    0    0
## J    0    0    0    0    0    0    0    0    0    0
## Z    0    0    0    0    0    0    0    0    0    0
## X    0    0    0    0    0    0    0    0    0    0
## *    0    0    0    0    0    0    0    0    0    0
## -    5    5    5    5    5    5    5    5    5    5
## +    0    0    0    0    0    0    0    0    0    0
## .    0    0    0    0    0    0    0    0    0    0
```

If called on a masked alignment, `consensusMatrix()` only uses those sequences/rows that are not masked. If there are masked columns, the matrix contains NA's in those columns:

```
conMat <- consensusMatrix(myMaskedAlignment)
conMat[, 95:104]

##   [,1] [,2] [,3] [,4] [,5] [,6] [,7] [,8] [,9] [,10]
## A   NA   NA   NA   NA   NA   NA    0    0    0     4
## R   NA   NA   NA   NA   NA   NA    0    0    0     0
## N   NA   NA   NA   NA   NA   NA    0    0    0     0
## D   NA   NA   NA   NA   NA   NA    4    0    0     0
## C   NA   NA   NA   NA   NA   NA    0    0    0     0
## Q   NA   NA   NA   NA   NA   NA    0    0    0     0
## E   NA   NA   NA   NA   NA   NA    0    0    0     0
## G   NA   NA   NA   NA   NA   NA    0    0    4     0
## H   NA   NA   NA   NA   NA   NA    0    0    0     0
## I   NA   NA   NA   NA   NA   NA    0    4    0     0
## L   NA   NA   NA   NA   NA   NA    0    0    0     0
## K   NA   NA   NA   NA   NA   NA    0    0    0     0
## M   NA   NA   NA   NA   NA   NA    0    0    0     0
## F   NA   NA   NA   NA   NA   NA    0    0    0     0
## P   NA   NA   NA   NA   NA   NA    0    0    0     0
## S   NA   NA   NA   NA   NA   NA    0    0    0     0
## T   NA   NA   NA   NA   NA   NA    0    0    0     0
## W   NA   NA   NA   NA   NA   NA    0    0    0     0
## Y   NA   NA   NA   NA   NA   NA    0    0    0     0
```

```
## V  NA  NA  NA  NA  NA  NA  0  0  0  0
## U  NA  NA  NA  NA  NA  NA  0  0  0  0
## O  NA  NA  NA  NA  NA  NA  0  0  0  0
## B  NA  NA  NA  NA  NA  NA  0  0  0  0
## J  NA  NA  NA  NA  NA  NA  0  0  0  0
## Z  NA  NA  NA  NA  NA  NA  0  0  0  0
## X  NA  NA  NA  NA  NA  NA  0  0  0  0
## *  NA  NA  NA  NA  NA  NA  0  0  0  0
## -  NA  NA  NA  NA  NA  NA  5  5  5  5
## +  NA  NA  NA  NA  NA  NA  0  0  0  0
## .  NA  NA  NA  NA  NA  NA  0  0  0  0
```

Multiple alignments also inherit the `consensusString()` method from the `Biostrings` package. However, for more flexibility and consistency, we rather advise users to use the `msaConsensusSequence()` method (see below).

## 6.2 Consensus Sequences and Conservation Scores

With version 1.7.1 of `msa`, new methods have been provided that allow for the computation of consensus sequences and conservation scores. By default, the `msaConsensusSequence()` method is a wrapper around the `consensusString()` method from the `Biostrings`:

```
printSplitString <- function(x, width=getOption("width") - 1)
{
  starts <- seq(from=1, to=nchar(x), by=width)

  for (i in 1:length(starts))
    cat(substr(x, starts[i], starts[i] + width - 1), "\n")
}

printSplitString(msaConsensusSequence(myFirstAlignment))

## -----
## -----?
## ?????????Y????D????????D????????R?Q????????????P?P???YTEEE?TW?TL??
## RQ??L????AC?E???G?----??LG???D?IPQLE?VSQ?LQ??TGWR???VPGL?P????FF?
## ?LA?R?FP?TQ?IR????LDY??EPDIFHELFGHVPLSDP?FA?F?Q?YG?LA????A?????E
## ?LARLYW?TVEFGL????-???KAYGAGLLSSF?ELQYCLSD-?P???PF?LE??M?T?Y?ID?
## FQPLYFV??SF??L?E??R??D?T????????????P????????V?D????????????????IL?
## ?A????---
```

However, there is also a second method for computing consensus sequence that has been implemented in line with a consensus sequence method implemented in `TeXshade` that allows for specify an upper and a lower conservation threshold (see example below). This method can be accessed via the argument `type="upperlower"`. Additional customizations are available, too:



```
printSplitString(msaConsensusSequence(myFirstAlignment, type="upperlower",
                                     thresh=c(40, 20)))

## .....-.....
## .....
## .....d.....p..y..ee..tw.t...
## ...l...ac.e.....g...d.ip.....l...tg....v.gl....f..
## .la.r.f..t..ir.....y..epdi.h...ghvpl....fa.f.q.g.....
## .la..yw.tvefgl....-....ygag.lss..e..y.l...p.....le.....y.i..
## fq..y.v..sf.....v.....
## .....-
```

Regardless of which method is used, masks are taken into account: masked rows/sequences are neglected and masked columns are shown as “#” in the consensus sequence:

```
printSplitString(msaConsensusSequence(myMaskedAlignment, type="upperlower",
                                     thresh=c(40, 20)))

## #####
## #####.....
## .....d.....p..y..ee..tw.t...
## ...l...ac.e.....g...d.ip.....l...tg....v.gl....f..
## .la.r.f..t..ir.....y..epdi.h...ghvpl....fa.f.q.g.....
## .la..yw.tvefgl....-....ygag.lss..e..y.l...p.....le.....y.i..
## fq..y.v..sf.....v.....
## .....-
```

The main purpose of consensus sequences is to get an impression of conservation at individual positions/columns of a multiple alignment. The `msa` package also provides another means of analyzing conservation: the method `msaConservationScore()` computes sums of pairwise scores for a given substitution/scoring matrix. Thereby, conservation can also be analyzed in a more sensible way than by only taking relative frequencies of letters into account as `msaConsensusSequence()` does.

```
library(pwalign) # for the BLOSUM62 matrix

##
## Attaching package: 'pwalign'
## The following objects are masked from 'package:Biostrings':
##
## PairwiseAlignments, PairwiseAlignmentsSingleSubject,
## aligned, alignedPattern, alignedSubject,
## compareStrings, deletion, errorSubstitutionMatrices,
## indel, insertion, mismatchSummary, mismatchTable,
## nedit, nindel, nucleotideSubstitutionMatrix,
```

```
## pairwiseAlignment, pid, qualitySubstitutionMatrices,  
## stringDist, unaligned, writePairwiseAlignments  
  
data(BLOSUM62)  
msaConservationScore(myFirstAlignment, BLOSUM62)
```

```

##   R   ?   F   P   ?   T   Q   ?   I   R   ?   ?   ?
## 341 45 486 223 172 184 106 343 233 405 117 117 100
##   ?   L   D   Y   ?   ?   E   P   D   I   F   H   E
## 209 92 110 567 124 132 405 567 486 149 214 648 301
##   L   F   G   H   V   P   L   L   S   D   P   ?   F
## 186 214 486 648 249 567 249 196 108 239 175 27 486
##   A   ?   F   ?   Q   ?   Y   G   ?   L   A   ?   ?
## 324 213 387 90 286 70 199 486 -73 7 196 3 26
##   ?   A   ?   ?   ?   ?   ?   E   ?   L   A   R   L
## 124 83 76 29 49 40 71 108 71 292 261 165 244
##   Y   W   ?   T   V   E   F   G   L   ?   ?   ?   ?
## 567 891 301 405 244 405 439 486 324 173 157 129 97
##   -   ?   ?   ?   K   A   Y   G   A   G   L   L   S
## 6 92 143 140 285 124 502 486 262 486 244 217 324
##   S   F   ?   E   L   Q   Y   C   L   S   D   -   ?
## 324 52 93 405 77 81 451 261 276 171 183 -117 131
##   P   ?   ?   ?   P   F   ?   L   E   ?   ?   M   ?
## 436 104 45 137 163 214 254 175 131 175 120 149 8
##   T   ?   Y   ?   I   D   ?   F   Q   P   L   Y   F
## 165 0 567 102 288 190 73 388 405 169 108 567 382
##   V   ?   ?   S   F   ?   ?   L   ?   E   ?   ?   R
## 324 100 182 262 306 118 82 124 62 264 64 95 129
##   ?   ?   D   ?   T   ?   ?   ?   ?   ?   ?   ?   ?
## -75 -10 134 92 87 44 68 51 56 18 41 44 61
##   ?   ?   P   ?   ?   ?   ?   ?   ?   V   ?   D   ?
## 79 83 91 161 56 69 51 12 42 66 -40 78 116
##   ?   ?   ?   ?   ?   ?   ?   ?   ?   ?   ?   ?   ?
## 43 1 -40 -71 -11 -86 -70 -67 37 -8 -41 -38 -11
##   ?   ?   ?   I   L   ?   ?   A   ?   ?   ?   -   -
## -28 -87 -7 -5 -44 4 -110 -44 -84 -68 -74 -83 -55
## -
## -47

```

As the above example shows, a substitution matrix must be provided. The result is obviously a vector as long as the alignment has columns. The entries of the vector are labeled by the consensus sequence. The way the consensus sequence is computed can be customized:

```

msaConservationScore(myFirstAlignment, BLOSUM62, gapVsGap=0,
                    type="upperlower", thresh=c(40, 20))

## - - - - - - - - - - - - -
## -80 -120 -119 -134 -96 -96 -80 -96 -144 -146 -96 -120 -80
## - - - - - - - - - - - - -
## -80 -96 -96 -64 -64 -64 -80 -80 -80 -96 -48 -96 -80
## - - - - - - - - - - - - -
## -106 -64 -121 -94 -80 -64 -90 -96 -104 -96 -96 -96 -64

```

```

## - - - - - - - - - - - - - -
## -96 -96 -80 -80 -80 -96 -64 -96 -96 -96 -98 -96 -96
## - - - - - - - - - - - - - -
## -80 -96 -64 -80 -80 -64 -89 -102 -64 -96 -80 -32 -96
## - - - - - - - - - - - - - -
## -80 -96 -80 -48 -96 -80 -96 -122 -80 -64 -80 -48 -80
## - - - - - - - - - - - - - -
## -64 -64 -80 -116 -96 -64 -104 -80 -113 -146 -48 -128 -96
## - - - - - - - - - - - - - -
## -135 -112 -96 -96 -80 -144 -96 -80 -96 -64 -96 -64 -96
## - - - - - - - - - - - - - -
## -80 -96 -32 -80 -96 -96 -80 -64 -80 -112 -80 -104 -80
## - - - - - - - - - - - - - -
## -96 -48 16 -64 -48 -80 -80 -96 -80 -80 -75 -76 -129
## . . . . . . . y . . . . d
## -76 -91 8 43 -48 -63 -15 14 -49 -62 -80 -100 -10
## . . . . . . . d . . . . .
## -90 -82 -43 -58 -75 -36 -60 246 52 74 52 97 74
## . r . q . . . . . . . .
## -5 100 77 125 60 51 -66 -115 -71 -45 -28 -36 48
## . . . p . p . . Y t e e
## 95 48 -3 166 30 218 6 21 33 567 141 160 165
## e . . t W . T l . . r q .
## 325 179 71 216 891 38 405 204 208 79 165 109 42
## . l . . . . A c . e . . .
## 79 249 113 106 93 172 324 536 109 324 261 72 157
## g . - - - - . . l g . . .
## 54 156 -48 -96 -96 -80 163 60 204 267 153 -20 60
## d . i P q l e . v s q . L
## 288 141 296 567 183 196 181 145 228 236 189 20 324
## q . . T G w r . . . V p g
## 171 74 145 405 486 411 167 216 36 125 324 199 388
## L . p . . . F f . . L A .
## 324 216 199 86 86 42 486 214 125 108 324 324 92
## r . F p . t q . i R . . .
## 341 45 486 223 172 184 106 343 233 405 117 117 100
## . l d Y . . E P D i f H e
## 209 92 110 567 124 132 405 567 486 149 214 648 301
## l f G H v P l l s d p . F
## 186 214 486 648 249 567 249 196 108 239 175 27 486
## A . f . q . y G . l a . .
## 324 213 387 90 286 70 199 486 -89 7 196 3 26
## . a . . . . . e . l a r l
## 124 83 76 29 48 40 71 108 71 292 261 165 244
## Y W . T v E f G L . . . .
## 567 891 301 405 244 405 439 486 324 173 157 129 97

```



```

##      -      -      -      -      -      -      -      -      -      -      -      -      -
## -80  -96  -32  -80  -96  -96  -80  -64  -80 -112  -80 -104  -80
##      -      -      -      -      -      -      -      -      -      -      .      .      .
## -96  -48   16  -64  -48  -80  -80  -96  -80  -80  -75  -76 -129
##      .      .      .      .      .      .      .      y      .      .      .      .      d
## -76  -91   8   43  -48  -63  -15   14  -49  -62  -80 -100  -10
##      .      .      .      .      .      .      .      d      .      .      .      .      .
## -90  -82  -43  -58  -75  -36  -60  246   52   74   52   97   74
##      .      r      .      q      .      .      .      .      .      .      .      .      .
##  -5  100   77  125   60   51  -66 -115  -71  -45  -28  -36   48
##      .      .      .      p      .      p      .      .      .      Y      t      e      e
##  95   48   -3  166   30  218   6   21   33  567  141  160  165
##  e      .      .      t      W      .      T      l      .      .      r      q      .
## 325  179   71  216  891   38  405  204  208   79  165  109   42
##      .      l      .      .      .      .      A      c      .      e      .      .      .
##  79  249  113  106   93  172  324  536  109  324  261   72  157
##  g      .      -      -      -      -      .      .      l      g      .      .      .
##  54  156  -48  -96  -96  -80  163   60  204  267  153  -20   60
##  d      .      i      P      q      l      e      .      v      s      q      .      L
## 288  141  296  567  183  196  181  145  228  236  189   20  324
##  q      .      .      T      G      w      r      .      .      .      V      p      g
## 171   74  145  405  486  411  167  216   36  125  324  199  388
##  L      .      p      .      .      .      F      f      .      .      L      A      .
## 324  216  199   86   86   42  486  214  125  108  324  324   92
##  r      .      F      p      .      t      q      .      i      R      .      .      .
## 341   45  486  223  172  184  106  343  233  405  117  117  100
##      .      l      d      Y      .      .      E      P      D      i      f      H      e
## 209   92  110  567  124  132  405  567  486  149  214  648  301
##  l      f      G      H      v      P      l      l      s      d      p      .      F
## 186  214  486  648  249  567  249  196  108  239  175   27  486
##  A      .      f      .      q      .      y      G      .      l      a      .      .
## 324  213  387   90  286   70  199  486  -89   7  196   3   26
##      .      a      .      .      .      .      .      e      .      l      a      r      l
## 124   83   76   29   48   40   71  108   71  292  261  165  244
##  Y      W      .      T      v      E      f      G      L      .      .      .      .
## 567  891  301  405  244  405  439  486  324  173  157  129   97
##  -      .      .      .      k      a      y      G      a      G      l      l      S
## -58   92  143  140  285  124  502  486  262  486  244  217  324
##  S      f      .      E      l      q      y      c      l      s      d      -      .
## 324   52   93  405   77   81  451  261  276  171  183 -142  131
##  p      .      .      .      p      f      .      l      e      .      .      m      .
## 436  104   45  137  163  214  254  175  131  175  120  149   8
##  t      .      Y      .      i      d      .      f      Q      p      l      Y      f
## 165   0  567  102  288  190   73  388  405  169  108  567  382
##  V      .      .      s      f      .      .      l      .      e      .      .      r
## 324  100  182  262  306  118   82  124   62  264   64   95  129

```

```
##      .      .      d      .      t      .      .      .      .      .      .      .      .
## -91  -10  134   92   87   44   68   51   56   18   41   44   61
##      .      .      p      .      .      .      .      .      .      v      .      d      .
##  79   83   90  161   56   69   51   12   42   65  -41   77  115
##      .      .      .      .      .      .      .      .      .      .      .      .      .
##  42    0  -44  -75  -15  -90  -74  -76   33  -12  -45  -42  -15
##      .      .      .      i      l      .      .      a      .      .      .      -      -
## -32  -91  -11   -9  -60  -12 -126  -60 -100  -84  -90 -108  -80
##      -
## -96
```

### 6.3 Interfacing to Other Packages

There are also other sequence analysis packages that use or make use of multiple sequence alignments. The `msa` package does not directly interface to these packages in order to avoid dependencies and possible incompatibilities. However, `msa` provides a function `msaConvert()` that allows for converting multiple sequence alignment objects to other types/classes. Currently, five such conversions are available, namely to the classes `alignment` (`seqinr` package [2]), `align` (`bios2mds` package [14]), `AAbin/DNAbin` (`ape` package [10]), and `phyDat` (`phangorn` package [11]). Except for the conversion to the class `phyDat`, these conversion are performed without loading or depending on the respective packages.

In the following example, we perform a multiple alignment of Hemoglobin alpha example sequences and convert the result for later processing with the `seqinr` package:

```
hemoSeq <- readAAStringSet(system.file("examples/HemoglobinAA.fasta",
                                       package="msa"))
hemoAln <- msa(hemoSeq)

## use default substitution matrix

hemoAln

## CLUSTAL 2.1
##
## Call:
##   msa(hemoSeq)
##
## MsaAAMultipleAlignment with 17 rows and 143 columns
##      aln                                     names
## [1] -VLSPADKTNVKAAWGKV...LDKFLASVSTVLTSKYR HBA1_Homo_sapiens
## [2] MVLSPADKTNVKAAWGKV...LDKFLASVSTVLTSKYR HBA1_Pan_troglodytes
## [3] -VLSPADKSNVKAAWGKV...LDKFLASVSTVLTSKYR HBA1_Macaca_mulatta
## [4] -VLSAADKGNVKAAWGKV...LDKFLANVSTVLTSKYR HBA1_Bos_taurus
## [5] -VLSPADKTNVKGTSKI...LDKFLASVSTVLTSKYR HBA1_Tursiops_tru...
```

```
## [6] -VLSGEDKSNIKAAWGKI...LDKFLASVSTVLTSKYR HBA1_Mus_musculus
## [7] MVLSADDKTNKNCWGKI...LDKFLASVSTVLTSKYR HBA1_Rattus_norve...
## [8] -VLSATDKANVKTFWGKL...LDKFLATVATVLTSKYR HBA1_Erinaceus_eu...
## [9] -VLSAADKSNVKACWGKI...LDKFFSAVSTVLTSKYR HBA1_Felis_silves...
## [10] -VLSPADKTNKSTWDKI...LDKFFTAVSTVLTSKYR HBA1_Chrysocyon_b...
## [11] -VLSNDKTNVKATWSKV...LDKFLSNVSTVLTSKYR HBA1_Loxodonta_af...
## [12] -VLSAADKTNVKAWSKV...LDKFLALLSTVLTSKYR HBA1_Monodelphis_...
## [13] -MLTDAEKKEVTALWGKA...MDKFLSKVATVLTSKYR HBA1_Ornithorhync...
## [14] -VLSAADKNNVKGIFTKI...LDKFLCAVGTVLTAKYR HBA1_Gallus_gallus
## [15] -HLTADKKHIKAIWPSV...LDKFLVSVSNVLTSKYR HBA1_Xenopus_trop...
## [16] -VLTEEDKARVRVAWVPV...VDKFLGQISKVLASRYR HBA1_Microcephalo...
## [17] -SLSDTDKAVVKAIWAKI...VDKFFNNLALALSEKYR HBA1_Danio_rerio
## Con -VLS?ADK?NVKA?WGK?...LDKFLA?VSTVLTSKYR Consensus

hemoAln2 <- msaConvert(hemoAln, type="seqinr::alignment")
```

Now we compute a distance matrix using the `dist.alignment()` function from the `seqinr` package:

```
library(seqinr)

d <- dist.alignment(hemoAln2, "identity")
as.matrix(d)[2:5, "HBA1_Homo_sapiens", drop=FALSE]

##                HBA1_Homo_sapiens
## HBA1_Pan_troglodytes            0.000000
## HBA1_Macaca_mulatta             0.1684304
## HBA1_Bos_taurus                 0.3472281
## HBA1_Tursiops_truncatus         0.4038819
```

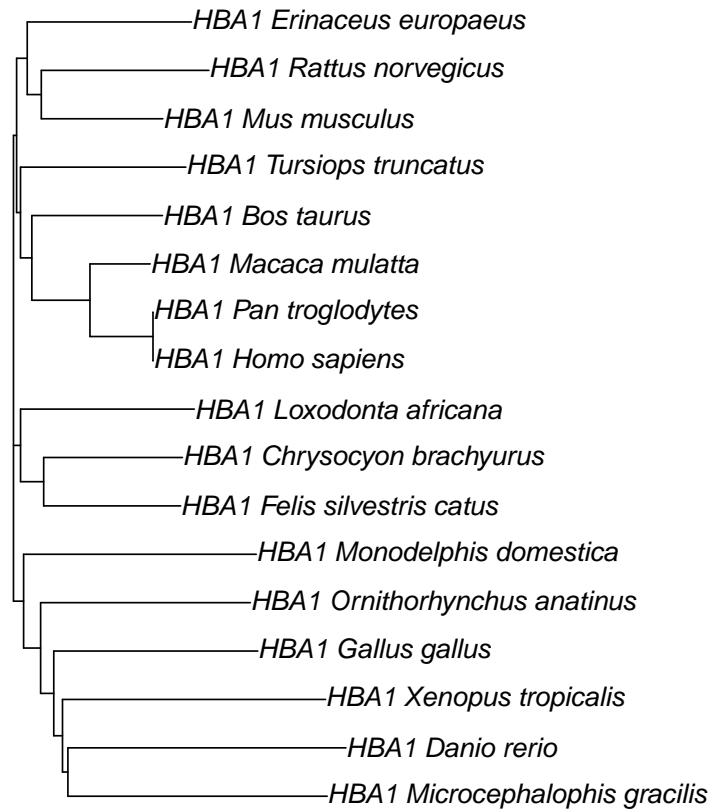
Now we can construct a phylogenetic tree with the neighbor joining algorithm using the `nj()` function from the `ape` package:

```
library(ape)

hemoTree <- nj(d)
plot(hemoTree, main="Phylogenetic Tree of Hemoglobin Alpha Sequences")
```



### Phylogenetic Tree of Hemoglobin Alpha Sequences



The following example shows how to convert a multiple alignment object in an object of class `align` as defined by the `bios2mds` package:

```
hemoAln3 <- msaConvert(hemoAln, type="bios2mds::align")
str(hemoAln3)

## List of 17
## $ HBA1_Homo_sapiens      : chr [1:143] "-" "V" "L" "S" ...
## $ HBA1_Pan_troglodytes  : chr [1:143] "M" "V" "L" "S" ...
## $ HBA1_Macaca_mulatta   : chr [1:143] "-" "V" "L" "S" ...
## $ HBA1_Bos_taurus       : chr [1:143] "-" "V" "L" "S" ...
## $ HBA1_Tursiops_truncatus : chr [1:143] "-" "V" "L" "S" ...
## $ HBA1_Mus_musculus     : chr [1:143] "-" "V" "L" "S" ...
```

```
## $ HBA1_Rattus_norvegicus      : chr [1:143] "M" "V" "L" "S" ...
## $ HBA1_Erinaceus_europaeus   : chr [1:143] "-" "V" "L" "S" ...
## $ HBA1_Felis_silvestris_catus : chr [1:143] "-" "V" "L" "S" ...
## $ HBA1_Chrysocyon_brachyurus  : chr [1:143] "-" "V" "L" "S" ...
## $ HBA1_Loxodonta_africana    : chr [1:143] "-" "V" "L" "S" ...
## $ HBA1_Monodelphis_domestica  : chr [1:143] "-" "V" "L" "S" ...
## $ HBA1_Ornithorhynchus_anatinus : chr [1:143] "-" "M" "L" "T" ...
## $ HBA1_Gallus_gallus         : chr [1:143] "-" "V" "L" "S" ...
## $ HBA1_Xenopus_tropicalis     : chr [1:143] "-" "H" "L" "T" ...
## $ HBA1_Microcephalophis_gracilis : chr [1:143] "-" "V" "L" "T" ...
## $ HBA1_Danio_rerio           : chr [1:143] "-" "S" "L" "S" ...
## - attr(*, "class")= chr "align"
```

The conversions to the standard Biostrings classes are straightforward using standard `as()` methods and not provided by the `msaConvert()` function. The following example converts a multiple alignment object to class `BStringSet` (e.g. the `msaplot()` function from the `ggtree` package [16] accepts `BStringSet` objects):

```
hemoAln4 <- as(hemoAln, "BStringSet")
hemoAln4

## BStringSet object of length 17:
##      width seq                                names
## [1]   143 -VLSPADKTNVKAAW...KFLASVSTVLTSKYR HBA1_Homo_sapiens
## [2]   143 MVLSPADKTNVKAAW...KFLASVSTVLTSKYR HBA1_Pan_troglodytes
## [3]   143 -VLSPADKSNVKAAW...KFLASVSTVLTSKYR HBA1_Macaca_mulatta
## [4]   143 -VLSAADKGNVKAAW...KFLANVSTVLTSKYR HBA1_Bos_taurus
## [5]   143 -VLSPADKTNVKGW...KFLASVSTVLTSKYR HBA1_Tursiops_tru...
## ...   ...   ...
## [13]  143 -MLTDAEKKEVTALW...KFLSKVATVLTSKYR HBA1_Ornithorhync...
## [14]  143 -VLSAADKNNVKGIF...KFLCAVGTVLTAKYR HBA1_Gallus_gallus
## [15]  143 -HLTADDKKHIAIW...KFLVSVSNVLTSKYR HBA1_Xenopus_trop...
## [16]  143 -VLTEEDKARVRVAW...KFLGQISKVLASRYR HBA1_Microcephalo...
## [17]  143 -SLSDTDKAVVKAIW...KFFNNLALALSEKYR HBA1_Danio_rerio
```

**Note:** The `msaConvert()` function has been introduced in version 1.3.3 of the `msa` package. So, to have this function available, at least Bioconductor 3.3 is required, which requires at least R 3.3.0.

## 7 Pretty-Printing Multiple Sequence Alignments

As already mentioned above, the `msa` package offers the function `msaPrettyPrint()` which allows for pretty-printing multiple sequence alignments using the `LATEX` package `TEXshade` [1].

Which prerequisites are necessary to take full advantage of the `msaPrettyPrint()` function is described in Section 2.

The `msaPrettyPrint()` function writes a multiple sequence alignment to an alignment (`.aln`) file and then creates  $\LaTeX$  code for pretty-printing the multiple sequence alignment on the basis of the  $\LaTeX$  package `TeXshade`. Depending on the choice of the `output` argument, the function `msaPrettyPrint()` either prints a  $\LaTeX$  fragment to the R session (choice `output="asis"`) or writes a  $\LaTeX$  source file (choice `output="tex"`) that it processes to a DVI file (choice `output="dvi"`) or PDF file (choice `output="pdf"`). Note that no extra software is needed for choices `output="asis"` and `output="tex"`. For `output="dvi"` and `output="pdf"`, however, a  $\TeX/\LaTeX$  distribution must be installed in order to translate the  $\LaTeX$  source file into the desired target format (DVI or PDF).

The function `msaPrettyPrint()` allows for making the most common settings directly and conveniently via an R interface without the need to know the details of  $\LaTeX$  or `TeXshade`. In the following, we will describe some of these customizations. For all possibilities, the user is referred to the documentation of `TeXshade`.<sup>5</sup>

## 7.1 Consensus Sequence and Sequence Logo

The consensus sequence of the alignment is one of the most important results of a multiple sequence alignment. `msaPrettyPrint()` has a standard possibility to show this consensus sequence with the parameter `showConsensus`. The default value is `"bottom"`, which results in the following:

```
msaPrettyPrint(myFirstAlignment, output="asis", y=c(164, 213),
               subset=c(1:6), showNames="none", showLogo="none",
               consensusColor="ColdHot", showLegend=FALSE,
               askForOverwrite=FALSE)
```

```
IAYNYRHGQPIPRVEYTEEEKQ TWGTVFRTLKALYKTHACYEHNHIFPLL 213
IAYNYRHGQPIPRVEYTEEERKTWGTVFRTLKALYKTHACYEHNHIFPLL 213
IAYNYRHGQPIPRVEYMEEEKKTWGTVFKTLKSLYKTHACYEYNHIFPLL 213
IAYNYRHGQPIPRVEYTEEKKKTWGTVFRTLKSLYKTHACYEHNHIFPLL 212
. . . . .DFTLPQPLDRYSAEDHATWATLYQRQCKLLPGRACDEFMEGL. . . 67
QTLRPDFTMEQPVHRYTAADHATWRTLYDRQEALLPGRACDEFLLQGL. . . 83
*****!***!*****!!*!*****!*****!!*!*****
```

Consensus sequences can also be displayed on top of a multiple sequence alignment or omitted completely.

In the above example, an exclamation mark `‘!` in the consensus sequence stands for a conserved letter, i.e. a sequence positions in which all sequences agree, whereas an asterisk `‘*’` stands for positions in which there is a majority of sequences agreeing. Positions in which the sequences disagree are left blank in the consensus sequence. For a more advanced example how to customize the consensus sequence, see the example in Subsection 7.4 below.

<sup>5</sup><https://www.ctan.org/pkg/texshade>

The color scheme of the consensus sequence can be configured with the `consensusColors` parameter. Possible values are "ColdHot", "HotCold", "BlueRed", "RedBlue", "GreenRed", "RedGreen", or "Gray". The above example uses the color scheme "RedGreen".

Additionally, `msaPrettyPrint()` also offers a more sophisticated visual representation of the consensus sequence — sequence logos. Sequence logos can be displayed either on top of the multiple sequence alignment (`showLogo="top"`), below the multiple sequence alignment (`showLogo="bottom"`), or omitted at all (`showLogo="none"`):

```
msaPrettyPrint(myFirstAlignment, output="asis", y=c(164, 213),
               subset=c(1:6), showNames="none", showLogo="top",
               logoColors="rasmol", shadingMode="similar",
               showLegend=FALSE, askForOverwrite=FALSE)
```



The color scheme of the sequence logo can be configured with the `logoColors` parameter. Possible values are "chemical", "rasmol", "hydropathy", "structure", "standard area", and "accessible area". The above example uses the color scheme "rasmol".

Note that a consensus sequence and a sequence logo can be displayed together, but only on opposite sides.

Finally, a caveat: for computing consensus sequences, `msaPrettyPrint()` uses the functionality provided by `TeXshade`, therefore, the results need not match to the results of the methods described in Section 6 above.

## 7.2 Color Shading Modes

`TeXshade` offers different shading schemes for displaying the multiple sequence alignment itself. The following schemes are available: "similar", "identical", and "functional". Moreover, there are five different color schemes available for shading: "blues", "reds", "greens", "grays", or "black". The following example uses the shading mode "similar" along with the color scheme "blues":

```
msaPrettyPrint(myFirstAlignment, output="asis", y=c(164, 213),
               showNames="none", shadingMode="similar",
               shadingColors="blues", showLogo="none",
               showLegend=FALSE, askForOverwrite=FALSE)
```

```

IAYNYRHHGQPIPRVEYTEEEKQWTGTVFRTLKALYKTHACYEHNHIFPLL 213
IAYNYRHHGQPIPRVEYTEEERKTWGTVFRTLKALYKTHACYEHNHIFPLL 213
IAYNYRHHGQPIPRVEYMEEEKKTWGTVFRTLKSLYKTHACYEYHNHIFPLL 213
IAYNYRHHGQPIPRVEYTEEEKKTWGTVFRTLKSLYKTHACYEHNHIFPLL 212
.....DFTLPQPLDRYS AEDHATWATLYQRQCKLLPGRACDEFMEGL... 67
QTLRPDFTMEQPVHRYTAADHATWRTLYDRQEALLPGRACDEFLLQGL... 83
....PDWTIDQGWEYTYTQA EHDVWITLYERQTDMLHGRACDEFMRGL... 58
.....DNGFIHYPETE HQVWNTLITRQLKVI EGRACQEYLDGI... 50
.VCRADFTVAQDYD.YSDEEQAVWR TLCDRQTKLITRKL AHHSYLDGV... 65
***** * *****!***** *!*!***** *** *!******

```

If the shading modes "similar" or "identical" are used, the `shadingModeArg` argument allows for setting a similarity threshold (a numerical value between 0 and 100). For shading mode "functional", the following settings of the `shadingModeArg` argument are possible: "charge", "hydropathy", "structure", "hemical", "rasmol", "standard area", and "accessible area". The following example uses shading mode "functional" along with `shadingModeArg` set to "structure":

```

msaPrettyPrint(myFirstAlignment, output="asis", y=c(164, 213),
               showNames="none", shadingMode="functional",
               shadingModeArg="structure",
               askForOverwrite=FALSE)

```

```

IAYNYRHHGQPIPRVEYTEEEKQWTGTVFRTLKALYKTHACYEHNHIFPLL 213
IAYNYRHHGQPIPRVEYTEEERKTWGTVFRTLKALYKTHACYEHNHIFPLL 213
IAYNYRHHGQPIPRVEYMEEEKKTWGTVFRTLKSLYKTHACYEYHNHIFPLL 213
IAYNYRHHGQPIPRVEYTEEEKKTWGTVFRTLKSLYKTHACYEHNHIFPLL 212
.....DFTLPQPLDRYS AEDHATWATLYQRQCKLLPGRACDEFMEGL... 67
QTLRPDFTMEQPVHRYTAADHATWRTLYDRQEALLPGRACDEFLLQGL... 83
....PDWTIDQGWEYTYTQA EHDVWITLYERQTDMLHGRACDEFMRGL... 58
.....DNGFIHYPETE HQVWNTLITRQLKVI EGRACQEYLDGI... 50
.VCRADFTVAQDYD.YSDEEQAVWR TLCDRQTKLITRKL AHHSYLDGV... 65

```

```

X external
X ambivalent
X internal

```

In the above example, a legend is shown that specifies the meaning of the color codes with which the letters are shaded. In some of the other examples above, we have suppressed this legend with the option `showLegend=FALSE`. The default, however, is that a legend is printed underneath the multiple sequence alignment like in the previous example.

### 7.3 Subsetting

In case that not the complete multiple sequence alignment should be printed, `msaPrettyPrint()` offers two ways of sub-setting. On the one hand, the `subset` argument allows for selecting only a

subset of sequences. Not surprisingly, `subset` must be a numeric vector with indices of sequences to be selected. On the other hand, it is also possible to slice out certain positions of the multiple sequence alignment using the `y` argument. In the simplest case, `y` can be a numeric vector with two elements in ascending order which correspond to the left and right bounds between which the multiple sequence alignment should be displayed. However, it is also possible to slice out multiple windows. For this purpose, the argument `y` must be an `IRanges` object containing the starts and ends of the windows to be selected.

## 7.4 Additional Customizations

The `msaPrettyPrint()` function provides an interface to the most common functionality of `TeXshade` in a way that the user does not need to know the specific commands of `TeXshade`. `TeXshade`, however, provides a host of additional customizations many of which are not covered by the interface of the `msaPrettyPrint()` function. In order to allow users to make use of all functionality of `TeXshade`, `msaPrettyPrint()` offers the `furtherCode` argument through which users can add `LATEX` code to the `texshade` environment that is created by `msaPrettyPrint()`. Moreover, the `code` argument can be used to bypass all of `msaPrettyPrint()`'s generation of `TeXshade` code.

Here is an example how to use the `furtherCode` argument in order to customize the consensus sequence and to show a ruler on top:

```
msaPrettyPrint(myFirstAlignment, output="asis", y=c(164, 213),
               subset=c(1:6), showNames="none", showLogo="none",
               consensusColor="ColdHot", showLegend=FALSE,
               shadingMode="similar", askForOverwrite=FALSE,
               furtherCode=c("\\defconsensus{.}{lower}{upper}",
                             "\\showruler{1}{top}"))
```

```

      170          180          190          200          210
IAYNYRHGQPIPRVEYTEEEKQTWGTVFRTLKALYKTHACYEHNHIFPLL 213
IAYNYRHGQPIPRVEYTEEERKKTWGTVFRTLKALYKTHACYEHNHIFPLL 213
IAYNYRHGQPIPRVEYMEEEKKTWGTVFRTLKSLYKTHACYEYNHIFPLL 213
IAYNYRHGQPIPRVEYTEEEKKTWGTVFRTLKSLYKTHACYEHNHIFPLL 212
.....DFTLPQPLDRYS AEDHATWATLYQRQCKLLPGRACDEFMEGL... 67
QTLRPDFTMEQPVHRYTAADHATWRTLYDRQEALLPGRACDEFLLQGL... 83
iaynyrhggppiPrveYteeekktWgTvfrtlkkaLykthACyEhnhifpll
```

## 7.5 Sweave or knitr Integration

The function `msaPrettyPrint()` is particularly well-suited for pretty-printing multiple alignments in Sweave [6] or knitr [15] documents. The key is to set `output` to "asis" when calling `msaPrettyPrint()` and, at the same time, to let the R code chunk produce output that is directly included in the resulting `LATEX` document as it is. This can be accomplished with the code chunk option `results="tex"` in Sweave and with the code chunk option `results="asis"` in

knitr. Here is an example of a Sweave code chunk that displays a pretty-printed multiple sequence alignment inline:

```
<<AnyChunkName,results="tex">>=
msaPrettyPrint(myFirstAlignment, output="asis")
@
```

The same example in knitr:

```
<<AnyChunkName,results="asis">>=
msaPrettyPrint(myFirstAlignment, output="asis")
@
```

Note that, for processing the resulting  $\LaTeX$  source document, the  $\TeX$ `shade` package must be installed (see Section 2) and the  $\TeX$ `shade` package must be loaded in the preamble:

```
\usepackage{texshade}
```

## 7.6 Sequence Names

The `Biostrings` package does not impose any restrictions on the names of sequences. Consequently, `msa` also allows all possible ASCII strings as sequence (row) names in multiple alignments. As soon as `msaPrettyPrint()` is used for pretty-printing multiple sequence alignments, however, the sequence names are interpreted as plain  $\LaTeX$  source code. Consequently,  $\LaTeX$  errors may arise because of characters or words in the sequence names that  $\LaTeX$  does not or cannot interpret as plain text correctly. This particularly includes appearances of special characters and backslash characters in the sequence names.

The `msa` package offers a function `msaCheckNames()` which allows for finding and replacing potentially problematic characters in the sequence names of multiple alignment objects (see `?msaCheckNames`). However, the best solution is to check sequence names carefully and to avoid problematic sequence names from the beginning. Note, moreover, that too long sequence names will lead to less appealing outputs, so users are generally advised to consider sequence names carefully.

## 7.7 Pretty-Printing Wide Alignments

If the alignment to be printed with `msaPrettyPrint()` is wide (thousands of columns or wider),  $\LaTeX$  may terminate prematurely because of exceeded  $\TeX$  capacity. Unfortunately, this problem remains opaque to the user, since `texi2dvi()` and `texi2pdf()` do not convey much details about  $\LaTeX$  problems when typesetting a document. We recommend the following if a user encounters problems with running `msaPrettyPrint()`'s output with `texi2dvi()` and `texi2pdf()`:

1. Run `pdflatex` on the generated `.tex` file to see whether it is actually a problem with  $\TeX$  capacity.

2. If so, split the alignment into multiple chunks and run `msaPrettyPrint()` on each chunk separately.

The following example demonstrates this approach for a multiple alignment object ‘aln’:

```
chunkSize <- 300 ## how much fits on one page depends on the length of
## names and the number of sequences;
## change to what suits your needs

for (start in seq(1, ncol(aln), by=chunkSize))
{
  end <- min(start + chunkSize - 1, ncol(aln))
  alnPart <- DNAMultipleAlignment(subseq(unmasked(aln), start, end))

  msaPrettyPrint(x=alnPart, output="pdf", subset=NULL,
                 file=paste0("aln_", start, "-", end, ".pdf"))
}
```

This creates multiple PDF files all of which show one part of the alignment. Please note, however, that the numbering of columns is restarted for each chunk.

## 7.8 Further Caveats

- Note that `texi2dvi()` and `texi2pdf()` always save the resulting DVI/PDF files to the current working directory, even if the  $\LaTeX$  source file is in a different directory. That is also the reason why the temporary file is created in the current working directory in the example below.
- `TeXshade` has a wide array of functionalities. Only the most common ones have been tested for interoperability with R. So the use of the arguments `furtherCode` and `code` is the user’s own risk!

## 8 Known Issues

### Memory Leaks

The original implementations of ClustalW, ClustalOmega, and MUSCLE are stand-alone command line programs which are only run once each time a multiple sequence alignment is performed. During the development of the `msa` package, we performed memory management checks using Valgrind [8] and discovered multiple memory leaks in ClustalW and MUSCLE. These memory leaks have no effect for the command line tools, since the program is closed each time the alignment is finished. In the implementation of the `msa` package, however, these memory leaks may have an effect if the same algorithm is run multiple times.

For MUSCLE, we managed to eliminate all memory leaks by deactivating the two parameters `weight1` and `weight2`. ClustalOmega did not show any memory leaks. ClustalW indeed has



several memory leaks which are benign if the algorithm is run only a few times, but which may have more severe effects if the algorithm is run many times. ClustalOmega also has a minor memory leak, but the loss of data is so small that no major problems are to be expected except for thousands of executions of ClustalOmega.

### **ClustalOmega vs. Older GCC Versions on Linux/Unix**

We have encountered peculiar behavior of ClustalOmega if the package was built using an older GCC version: if we built the package on an x86\_64 Linux system with GCC 4.4.7, ClustalOmega built smoothly and could be executed without any errors. However, the resulting multiple sequence alignment was more than sub-optimal. We could neither determine the source of this problem nor which GCC versions show this behavior. We therefore recommend Linux/Unix users to use an up-to-date GCC version (we used 4.8.2 during package development, which worked nicely) or, in case they encounter dubious results, to update to a newer GCC version and re-install the package.

### **ClustalOmega: OpenMP Support on Mac OS**

ClustalOmega is implemented to make use of OpenMP (if available on the target platform). Due to issues on one of the Bioconductor build servers running Mac OS, we had to deactivate OpenMP generally for Mac OS platforms. If a Mac OS user wants to re-activate OpenMP, he/she should download the source package tarball, untar it, comment/uncomment the corresponding line in `msa/src/ClustalOmega/msaMakefile` (see first six lines), and build/install the package from source.

### **Build/installation issues**

Some users have reported compiler and linker errors when building `msa` from source on Linux systems. In almost all cases, these could have been tracked down to issues with the R setup on those systems (e.g. a `Rprofile.site` file that makes changes to the R environment that are not compatible with `msa`'s Makefiles).<sup>6</sup> In most cases, these issues can be avoided by installing `msa` in a “vanilla R session”, i.e. starting R with option `--vanilla` when installing `msa`.

On some systems (e.g. FreeBSD), the required Gnu make command is installed under a different name, e.g., `gmake`. The `msa` package installation can handle this under the condition that an environment variable `MAKE` is defined beforehand that contains the name of the Gnu make binary.

## **9 Future Extensions**

We envision the following changes/extensions in future versions of the package:

- Integration of more multiple sequence alignment algorithms, such as, T-Coffee [9] or DIALIGN [7]
- Support for retrieving guide trees from the multiple sequence alignment algorithms

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<sup>6</sup>See, e.g., <https://support.bioconductor.org/p/90735/>

- Interface to methods computing phylogenetic trees (e.g. as contained in the original implementation of ClustalW)
- Elimination of memory leaks described in Section 8 and re-activation of parameters that have been deactivated in order to avoid memory leaks
- More tolerant handling of custom substitution matrices (MUSCLE interface)

## 10 How to Cite This Package

If you use this package for research that is published later, you are kindly asked to cite it as follows:

U. Bodenhofer, E. Bonatesta, C. Horejš-Kainrath, and S. Hochreiter (2015). *msa*: an R package for multiple sequence alignment. *Bioinformatics* **31**(24):3997–3999. DOI: [bioinformatics/btv494](https://doi.org/10.1093/bioinformatics/btv494).

To obtain a BibTeX entries of the reference, enter the following into your R session:

```
toBibtex(citation("msa"))
```

Moreover, we insist that, any time you cite the package, you also cite the original paper in which the original algorithm has been introduced (see bibliography below).

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