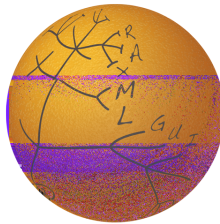


raxmlGUI

version 1.5

manual



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Contents

Table of Contents	2
Introduction	3
Requirements	3
Preferences	4
Using the CIPRES Science Gateway	4
Loading a file	5
Analysis	6
Analysis settings	6
Additional Analyses	7
Excluding sites	7
Partitioning the matrix	7
Outgroup selection	9
Setting a topological constraint	9
Output files	9
Utilities	10
Keyboard shortcuts	10
Contacts	10
References	13
Appendix	14
A A working PHYLIP example	14
B The format of the Partition file	14
C Selected papers that cite raxmlGUI	15
D References for Amino Acid substitution models	23

Introduction

RaxmlGUI is a python application, which provides a user friendly front-end for RAxML (STAMATAKIS 2014a) for Maximum Likelihood based phylogenetic analyses. The GUI interacts with RAxML executables, which are incorporated in the package, or the CIPRES Science Gateway. It enables the user to select input files, set the parameters and run ML analyses locally or in the cloud, such as phylogenetic reconstructions or ancestral state reconstructions with only a few mouse clicks. A number of options and functions are automated (e. g. checking for identical sequences, or gap-only characters) and simplified (e. g. model and outgroup selection, excluding sites, setting topological constraints and partitioning a matrix). Some features extend the usage of RAxML, e. g. assembling concatenated datasets with automatic partitioning, and providing analyses pipelines e. g. bootstrapping followed by computing a consensus tree, or a fast tree search followed by branch lengths estimation and computing of SH-like support values.

The GUI is meant to simplify the usage of RAxML, nevertheless it is strongly recommended to get familiar with the RAxML manual (STAMATAKIS 2014b) and the “hands-on session”¹ on the Exelixis page, to be aware of contents and intent of input and output files.

Requirements

- RaxmlGUI runs under Mac, Windows and Linux operating systems. The GUI automatically determines the operating system when started for the first time, and selects the respective RAxML executable.
- **Python 2.5.x or higher is required**, but please note that **Python 3.x.x is not supported**. **Under Windows make sure** that the python folder is added to your **PATH variable** (in recent python versions this is possible via a checkbox during installation). An explanation on how to do that manually you can find [here](#).
- Additionally you can install the python library [DendroPy](#) (SUKUMARAN & HOLDER 2010) to be able to import from and export to NEXUS files (MADDISON et al. 1997).
- **Furthermore, you can** set up an account for the [CIPRES REST API](#) (MILLER et al. 2015), to **run RAxML analyses on the CIPRES Science Gateway** (MILLER et al. 2011) instead of using your local RAxML executable, if you install an additional python library. For further explanations see below.
- **The application can be launched** with a double click on the file “raxmlGUI.py” under Windows (during the installation of Python it should be set as the default application for the “.py” extension). Under Unix (Mac OS and Linux) the program is launched by browsing to its directory via shell, and typing “./raxmlGUI.py” or “python raxmlGUI.py”. A Mac OSX application is also provided, that can be run with a double click.
- **Avoid special characters (like diacritics) and punctuation other than dots (“.”) and underscore (“_”) in the (path) names of the raxmlGUI folder and all input files!**

¹http://sco.h-its.org/exelixis/web/software/raxml/hands_on.html

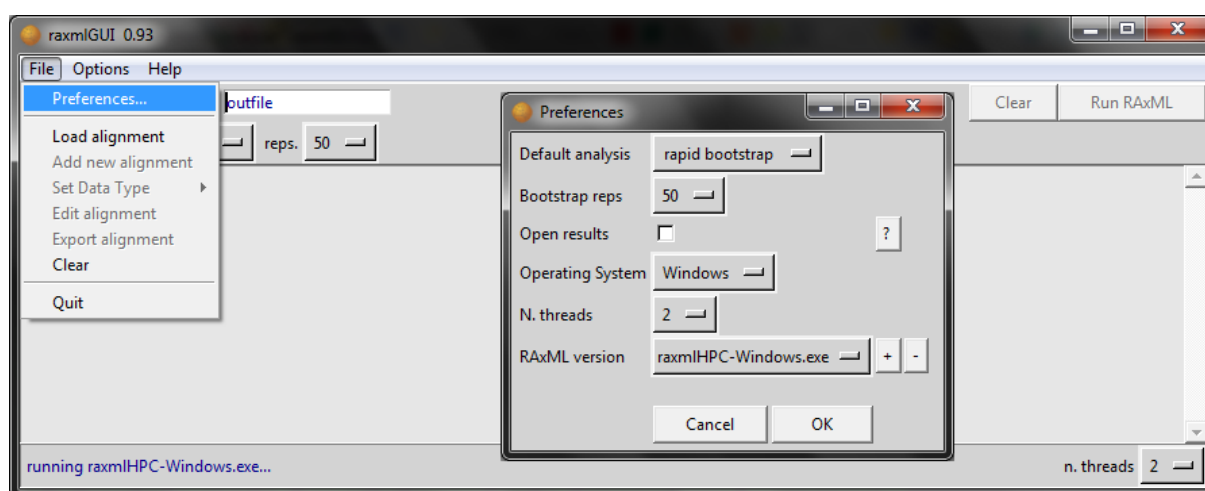


Figure 1: The preferences panel

Preferences

The preferences panel provides the possibilities to change default settings and select a RAxML executable. As default you can choose among a “normal” (Windows) or a SSE3 (Mac OSX; BERGER & STAMATAKIS 2010) and a multithread version (OTT et al. 2007). If you choose the multithread version the number of threads used by RAxML can be set. You can use the “+” button to add additional versions of RAxML (e. g. newer binaries² or those compiled for special needs). The chosen binary is then automatically copied to the application contents folder “/raxml”. Old or unused versions can be removed with “-” (WARNING: These executable files will be erased).

If the box “Open results” is checked, the resulting tree (e.g. best-scoring ML tree with bootstrap values) will be opened with the default application for tree files³ (.tre extension), if you have set one.

Please note that changes made in the preferences panel become effective only after re-starting the program.

Using the CIPRES Science Gateway ***NEW in version 1.5!***

You can opt to run your analysis (so far only working for the default analysis type: “3. ML + rapid bootstrap”, see below) on the CIPRES Science Gateway (MILLER et al. 2011) via the REST API (MILLER et al. 2015). You will need to install the necessary python toolkit, kindly provided by Terri Schwartz from the SDSC and included in the package. You will be guided through the installation when accessing the CIPRES options. Alternatively you can also install the toolkit manually by navigating to the folder “/raxmlgui/setup_python_cipres” in a command line window (using the “cd” command) and typing “python setup.py install” (without the quotes). After restarting raxmlGUI the CIPRES options should be available.

To configure the CIPRES option within raxmlGUI you can use the Utility “Configure CIPRES ID” and type in your log in information. If you don’t have an account, yet, you can set one up using the button “Sign Up” or via the web page of the [CIPRES REST API](#). **NOTE that the log in (user name and password) information entered will be saved in a plain text file**

²You can find the latest source code on Alexis Stamatakis’ [git repository](#).

³Widely used tree visualizers are [Dendroscope](#) (HUSON et al. 2007), [FigTree](#) (RAMBAUT 2006), and [Treeview](#) (PAGE 1996). For a comprehensive list of tree visualizing and editing software see http://bioinfo.unice.fr/biodiv/Tree_editors.html.

called “pypres.conf” in your home folder. Especially on public computers, you should erase the file or its contents after use, which can be done from within the same utility in raxmlGUI.

You can set a maximum time for your analysis (at the bottom of the main raxmlGUI window “CIPRES CPU time”), after which the potentially unfinished job would be aborted. The maximum time is 168 hours (= 7 days). The server will give priority to shorter jobs.

Once the CIPRES REST service is configured in raxmlGUI launching an analysis on the CIPRES servers is very easy. You can load an alignment and set substitution model, partitions etc. as in a standard raxmlGUI analysis (more details below) and select an option in the “CIPRES action” button. There are currently two options to start the analysis:

1. “Run CIPRES-RAXML” will launch the analysis in a new Terminal window that will show the progress of the run. Note that this window should not be closed until the end of the run
2. “Run CIPRES-RAXML in background” will send the analysis to the CIPRES servers as a background process. This means that the analysis will continue even after you close raxmlGUI. You will be notified when the analysis is done by email (the same address you used to register to the CIPRES services).

The “CIPRES action” button also provides options to check the current status of your job(s) running on the CIPRES servers and to download the results to a directory of your choice. The results of each job will be saved to folders named after the job’s IDs.

Depending on where you live, you have a certain CPU time available per year (currently for 50 000 CPU hours for US citizens, 30 000 CPU hours for others. Note that parallel jobs count multiplied by the number of processes). If you exceed this time you won’t be able to submit jobs any more.

Loading a file

With the button “Load alignment” you can load a data set as input for the RAxML analysis, which must be in PHYLIP format⁴. A FASTA or NEXUS file (MADDISON et al. 1997) can be loaded with the menu option “Import FASTA/NEXUS file”. The latter will be converted to a PHYLIP file for which you are asked to specify a path and file name. For converting NEXUS to PHYLIP the additional python library DendroPy (SUKUMARAN & HOLDER 2010) is needed. In case it is not yet installed, you will be guided through the installation process⁵. **Please note, that blanks and special characters in taxon names in the imported file will cause errors, as they are not allowed in PHYLIP format**, please rename your taxa according to PHYLIP limitations⁶.

The data type of the loaded file is determined automatically. If the file contains only one data type (i.e. nucleotides, amino acid, binary, or multistate), the alignment will be checked for readability (through the RAxML option “-f c”). A warning appears in case identical sequences and/or gap-only characters are detected: you can choose to run the analyses on either the original or the reduced data set. Note that if you want to exclude sites, partition the matrix and/or add additional data sets to a combined alignment, you should retain the

⁴For an example see Appendix A

⁵If you agree to install the library, the latest source code from Jeet Sukumaran’s [git repository](#) is downloaded as a zip folder. You will need to unzip this folder and pass its path to raxmlGUI, when it asks for it (point the installer to the folder which contains the file “setup.py”). You can follow the installation process in the terminal/console window. You will need to restart raxmlGUI to make use of DendroPy.

⁶Prohibited characters in PHYLIP are: , ') (: ;] [

original file, since through this option taxa might be removed and/or column numbers might change.

Analysis

Analysis settings

Seven different main analyses can be carried out through raxmlGUI:

1. “**Fast tree search**” – very fast, superficial tree search (RAxML option “-f E”) followed by optional computations of branch lengths (“-f e”) and SH-like support values (“-f J”; SHIMODAIRA & HASEGAWA 1999)⁷. The analysis result is comparable to FastTree (PRICE et al. 2010) outputs, but is expected to yield better likelihood scores.
2. “**ML search**” – Maximum likelihood reconstruction using the rapid hill-climbing algorithm (“-f d”; STAMATAKIS et al. 2007), optionally followed by the computation of SH-like support values (see 1.), which will be plotted on the single best-scoring tree⁷. To combine the resulting trees of independent ML searches in one file, check the box “combined output”.
3. “**ML + rapid bootstrap**” (default) – Rapid bootstrap analysis and search for a best-scoring Maximum Likelihood tree (equivalent to 2.). The number of ML searches is equal to 20% of the BS replicates (“-f a”; STAMATAKIS et al. 2008). The bootstrap values are reported on the ML tree.
4. “**ML + slow bootstrap**” – Slow bootstrap analysis (“-b”), followed by a ML search (2.). The bootstrap support values are drawn on the most likely tree (“-f b”).
5. “**Bootstrap + consensus**” – Rapid bootstrap analysis (“-x”) and a subsequent majority rule consensus tree calculation from all bootstrap trees⁷ (“-J MR”).
6. “**Ancestral states**” – Compute marginal ancestral states based on a user provided rooted tree and a character matrix (“-f A”).
7. “**Pairwise distances**” – Compute distances for all taxa pairs in the data set (“-f x”). As default a MP starting tree will be calculated, alternatively you can provide a user defined tree. This function is only available for GAMMA models.

Depending on the kind of alignment loaded you can choose the substitution model (GTR, BIN, MULTI, or PROT) with GAMMA[I] (YANG 1994), or CAT[I] (STAMATAKIS 2006) rate heterogeneity, or without rate heterogeneity (“-V”). For large data sets it is possible to select the RAxML option “-F” from the menu to reduce the used memory (works best in combination with CAT[I]).

If the file contains **amino acid** data, you can specify the substitution model⁸ you want to apply, and whether base frequencies should be determined empirically (note that the GTR substitution models always rely on empirical frequencies). If the data type in the file is **multistate** you can choose between GTR, Ordered, and MK substitution model.

⁷Such trees can be converted to a FigTree compatible format using the menu option “Convert to FigTree”.

⁸Available substitution models for amino acids are: DAYHOFF, DCMUT, JTT, MTREV, WAG, RTREV, CPREV, VT, BLOSUM62, MTMAM, LG, MTART, MTZOA, PMB, HIVB, HIVW, JTTDCMUT, FLU, GTR and GTR_UNLINKED. For the references of these see Appendix D.

You can set the **number of bootstrap replicates** with the option button “reps”. You can choose a predefined number, set a user defined value, or select options of automatic “bootstopping” (PATTENGAL et al. 2010) according to different methods such as majority rule tree based criteria (RAxML options “-N autoMR” [recommended], “-N autoMRE”, and “-N autoMRE_IGN”) and the frequency-based criterion (option “-N autoFC”). If you check the box “**BS brL**” (option “-k”), branch lengths will be saved in the bootstrap trees (which increases computation time). Additionally you can select the **number of independent ML searches** in the second and the fourth option.

You can load a Newick file to provide a **starting tree** for the ML-search through the analysis menu (RAxML option “-t”).

With a nucleotide alignment you can load a file in which brackets ((), [], { }, <>) define stems and pseudo-knot regions of the sequence’s **secondary structure**. The absolute number of characters of this file must be identical to the number of sites of the alignment. Nucleotide positions within the specified regions are represented by dots (“.”).

Additional Analyses

With a file containing a set of trees (e. g. the “RAxML_bootstrap” output file from a bootstrap analysis) you can generate majority rule or strict **consensus trees** (RAxML option “-J”). Also you can create a **set of pruned trees**, which lack a set of rogue taxa, that decrease support values (PATTENGAL et al. 2011), a consensus tree from these pruned trees is automatically created.

In addition you can compute **Robinson-Foulds pairwise distances** (ROBINSON & FOULDS 1981) between trees (option “-f r”).

With an alignment loaded and a file containing one or more trees you can also compute **per site log Likelihoods** (option “-f g”). The output that can be read by CONSEL⁹ (SHIMODAIRA 2001) to calculate p-values. In addition you can compute **SH-like support values** (SHIMODAIRA & HASEGAWA 1999) on a given (best) tree (option “-f J”).

Excluding sites

In order to exclude sites you can use the interactive panel, or load a file which has to be in the RAxML standard format:

```
1-200 333-333 500-667
```

Loading a file with this content, or specifying these ranges in the panel, will produce a new alignment file (using the RAxML flag “-E”), reduced by the first 200 columns, the single site 333 and the sites 500–667. The new alignment file is automatically set as input and is again checked for identical sequences and/or gap-only characters.

Partitioning the matrix

There are three different ways of defining partitions (RAxML option “-q”):

Maybe the easiest way is to load individual partitions one after the other as separate files with the “Add Alignment” -button. RaxmlGUI will automatically set the partitions according to the determined data type. **CAUTION! RaxmlGUI will ONLY check for identical taxon counts! Make sure taxon order and names are identical in all files!**

⁹You can use the CONSEL command “seqmt --puzzle YOURFILE” to convert it to CONSEL format

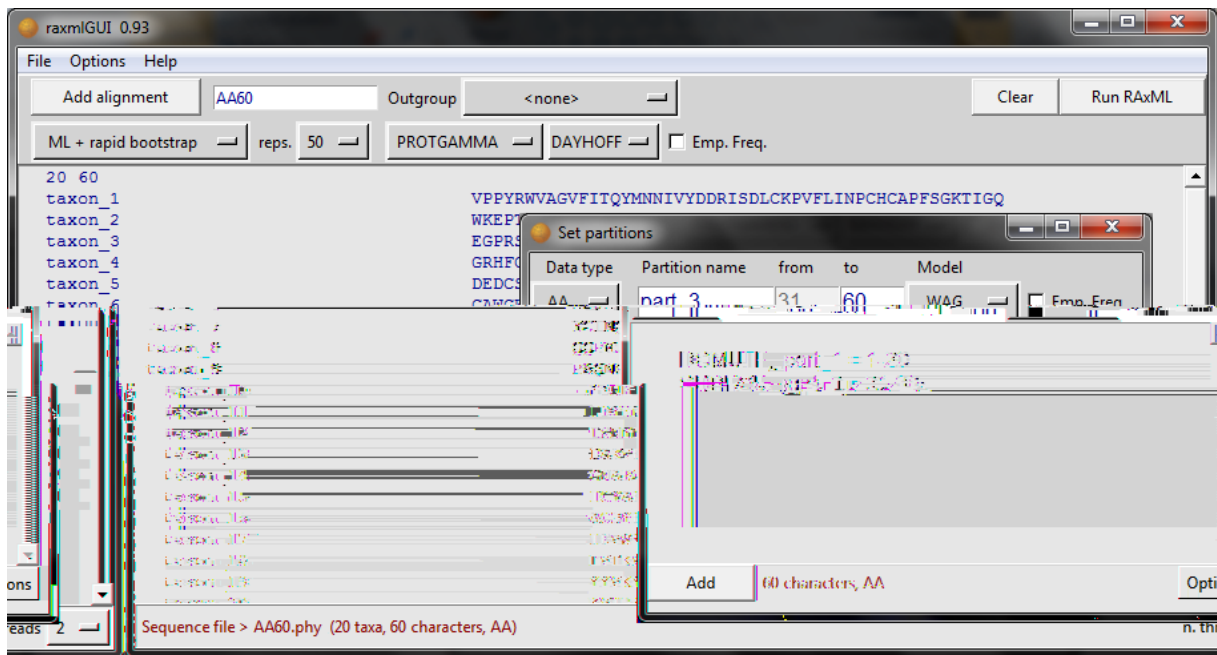


Figure 2: The partitioning panel

The second way is to load the combined data set as one and to set partitions in an interactive way with the option “Set/Edit partitions...” .

The third way is to load a file with the format specified in STAMATAKIS (2014b)¹⁰; note that in a partitioned analysis every site has to be assigned to one partition, i. e. sites must not be assigned to two different partitions, and no site is allowed to be not assigned to any partition.

In any case you can edit, delete, and export the partitions set.

If you successfully set a partitioning scheme on your data set, you will be asked before the run starts if you want to calculate the branch lengths independently for each partition (RAxML option “-M”). This will produce best-scoring likelihood trees with branch length optimized for each partition (identical topology).

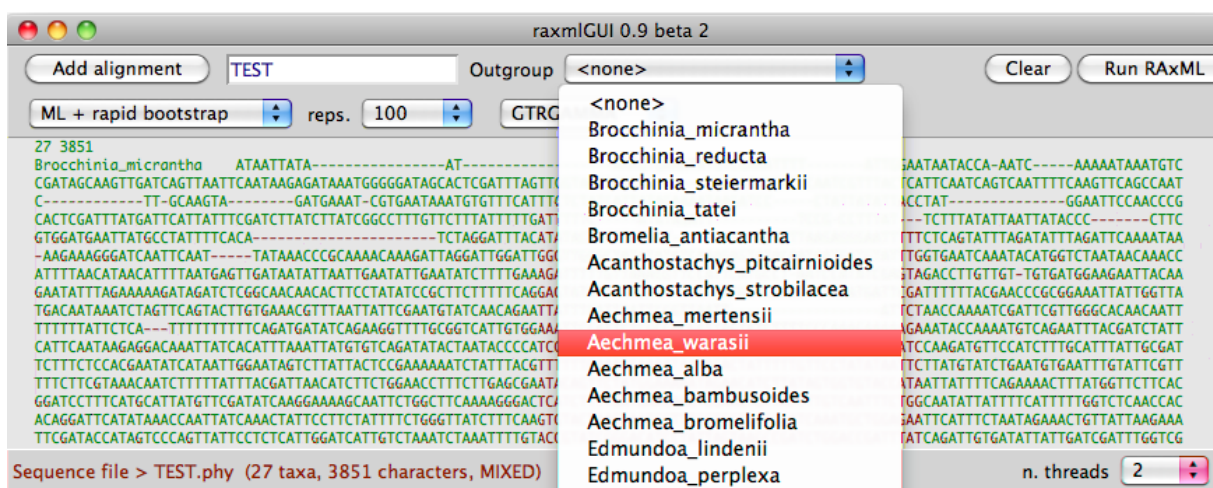


Figure 3: Outgroup selection

¹⁰See also Appendix B

Outgroup selection

A list of all taxa in the matrix is created in the program's toolbar for a quick single-taxon outgroup selection (RAxML option “-o”). If you want to specify more than one taxon to be in the outgroup, choose the option “select multiple outgroup” from the analysis menu. A window will open with a list of all taxa. You can mark more than one taxon by holding ☐ (selecting ranges) or ☐ (selecting single entries) for selecting those as outgroup. If RAxML finds the multiple outgroup not to be monophyletic, it will take the first taxon in the list as outgroup. If no outgroup is selected, the tree will be unrooted.

Setting a topological constraint

You can enforce topological constraints to your analysis through the menu option “enforce constraints”. You can define taxon groups through a panel (“define topological constraint...”) or by uploading a Newick formatted tree file, which can be binary or multifurcating. RAxML options “-r” or “-g” will be used for binary or multifurcating tree constraints, respectively.

Note that RAxML accepts only backbone constraints, which means, that unconstrained taxa can be placed at any position in the resulting tree, including within constrained clades. If you want to constrain monophyletic groups you can check the respective box in the panel, this will automatically append the remaining taxa to the set of constraints, so all constrained clades will result monophyletic.

Output files

You can set the name of the RAxML output files in the text field. By default the output file name is the same as the input file (without extension). The suffix “_red” is appended if identical sequences (or gap-only characters) are excluded from the analyses, and “_exc” is used when the “exclusion site” option is applied. If a RAxML info file with the same ID is found in the directory, you are prompted to change the output name before starting the analysis. For all types of output files, and their contents please refer to STAMATAKIS (2014b).

Utilities









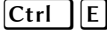

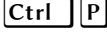


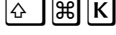
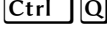



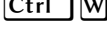

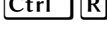

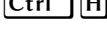

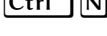


In some trees RAxML associates support values to the branches, rather than to the nodes (e.g. consensus trees, and those with SH-like support values). This format is not supported by FigTree (RAMBAUT 2006). However, if you want to use this program, you can produce a modified version of those tree files using the menu option “**Convert to FigTree format**”.

It is possible to **export your alignment in NEXUS format**. If the necessary python library DendroPy (SUKUMARAN & HOLDER 2010) is not yet installed, you will be guided through its installation.⁵

You can **inspect the RAxML command** that will be executed in the terminal before pressing “run RAxML” with the menu option “show RAxML command”. In case of pipelined analyses this can contain many commands. Further it is possible to save the command(s) to a file.

You can **export the citation** for raxmlGUI in the following formats: Text, BiBTeX, EndNote (xml), and Reference manager (RIS).

Keyboard shortcuts

Action	Short cut Win/Linux	Short cut Mac
Open alignment		
Export alignment		
Change analysis type		
Import FASTA file		
Exclude sites		
Set partitions		
Clear		
Quit raxmlGUI		
Preferences		
Close window		
Run analysis		
Open this raxmlGUI help		
Import NEXUS file		
Save changes (exclude sites/ define partitions/preferences)		

Contacts

If you find any problems/bugs or want to give us a feedback, please contact us:

raxmlgui.help@gmail.com.

If you want to get the latest news (e.g. new releases and updates), you can subscribe to our mailing list on <http://lists.sourceforge.net/lists/listinfo/raxmlgui-news>.

If you have problems or questions regarding RAxML, please have a look at the [RAxML google group](#).

If you have problems or questions regarding CIPRES, please direct them to [the cipres-rest-users forum](#).

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Appendix

A A working PHYLIP example

```
6 40
1_first_row_reads_nroftaxa_blank_nrofcharacters GTGGCGGTCATTCTCATTG
2_this_is_a_working_example_taxon_names_can_be_very_long ATTCGTGGTCATTTCGTGGTC
3_not_allowed_in_names_are_blanks_and_special_characters CGTGACATTCGTGCTTGGT
4_use_blanks_or_tabs_to_delimit_taxon_names_from_characters TCATTCGTGCGATGTCGTG
5_there_is_no_option_for_comments_like_in_Nexus_format TGTTGCGTTGGTCATCTCAG
6_for_interleaved_format_dont_repeat_the_taxon_names TTTCTTGGGCGGTCGTTCAA

TCTCATTGGCGGTCATTGGT
ATGTTGGCGGTCATTCTCTG
ATTTGGCCTCCAGGTGTGTT
GGGTCATTCATTCGTGTTGT
TGTGATTGCGGTCATTCTCG
GTTTGACTCATTGCGGTC
```

B The format of the Partition file

General format

Parttype, partname = partrange

e. g.:

```
DNA, partition1 = 1-100
BIN, partition2 = 101-200
```

A slightly more complex example

```
JTT, AAgene1 = 1-500

WAGF, AAgene2withempiricalfrequencies = 501-800

MULTI, multistatepartition3 = 801-900

BIN, binarypartition4 = 901-1000

DNA, DNAgene5codon1and2 = 1001-1500\3, 1002-1500\3
DNA, DNAgene5codon3 = 1003-1500\3

DNA, DNAgene6codon1 = 1501-2000\3
DNA, DNAgene6codon2 = 1502-2000\3
DNA, DNAgene6codon3 = 1503-2000\3

DNA, DNAsplitgene7 = 2001-2200, 2800-3000
DNA, intronofgene7 = 2201-2799
```

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