The *Vmatch* large scale sequence analysis software

A Manual

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1 Introduction

This document describes Vmatch, a versatile software tool for efficiently solving large scale sequence matching tasks. Vmatch subsumes the software tool REPuter [7] but is much more general, with a very flexible user interface, and improved space and time requirement. Here are the most important features of Vmatch.

Persistent index

Usually, in a large scale matching problem, extensive portions of the sequences under consideration are static, i.e. they do not change much over time. Therefore it makes sense to preprocess this static data to extract information from it and to store this in a structured manner, allowing efficient searches. Vmatch does exactly this: it preprocesses a set of sequences into an index structure. This is stored as a collection of several files constituting the persistent index. The index efficiently represents all substrings of the preprocessed sequences and, unlike many other sequence comparison tools, allows matching tasks to be solved in time, independent of the size of the index. Different matching tasks require different parts of the index, but only the required parts of the index are accessed during the matching process.

Alphabet independency

Most software tools for sequence analysis are restricted to DNA and/or protein sequences. In contrast, Vmatch can process sequences over any user defined alphabet not larger than 250 symbols. Vmatch fully implements the concept of symbol mappings, denoting alphabet transformations. These allow the user to specify that different characters in the input sequences should be considered identical in the matching process. This feature is used to group similar amino acids, for example.

Versatility

Vmatch allows a multitude of different matching tasks to be solved using the persistent index. Every matching task is basically characterized by (1) the kind of sequences to be matched, (2) the kind of matches sought, (3) additional constraints on the matches, and (4) the kind of post-processing to be done with the matches.

In the standard case, Vmatch matches sequences over the same alphabet. Additionally, DNA sequences can be matched against a protein sequence index in all six reading frames. Finally, DNA sequences can be transformed in all six reading frames and compared against itself.

Where appropriate, Vmatch can compute the following kinds of matches, using state-of-the-art algorithms:
1 INTRODUCTION

- maximal repeats using the algorithm of [2].
- branching tandem repeats using the algorithm of [1],
- supermaximal repeats using the algorithm of [2].
- maximal substring matches using the algorithm of [6].
- maximal unique substring matches using the algorithm of [6].
- complete matches using the algorithms of [8] and [9]

To compute degenerate substring matches or degenerate repeats, each kind of match (with the exception of tandem repeats and complete matches) can be taken as an exact seed and extended by either of two different strategies:

- the maximum error extension strategy, as described in [7] for repeat detection,
- the greedy extension strategy of [12].

Matches can be selected according to their length, their E-value, their identity value, or match score.

In the standard case, a match is displayed as an alignment including positional information. Alternatively, a match can directly be postprocessed in different ways:

- inverse output, i.e. reporting of substrings not covered by a match.
- masking of substrings covered by a match.
- clustering of sequences according to the matches found.
- chaining of matches, i.e. finding optimal subsets of matches which do not cross, using the algorithms described in [3].
- clustering of matches according to pairwise sequence similarities computed by the dynamic programming algorithm of [10].
- clustering of matches according to the positions where they occur, following the approach of [11].
Efficient algorithms and data structures

*Vmatch* is based on enhanced suffix arrays described in [2]. This data structure has been shown to be as powerful as suffix trees, with the advantage of a reduced space requirement and reduced processing time. Careful implementation of the algorithms and data structures incorporated in *Vmatch* have led to exceedingly fast and robust software, allowing very large sequence sets to be processed quickly. The 32-bit version of *Vmatch* can process up to 400 million symbols, if enough memory is available. For large server class machines (e.g. SUN-Sparc/Solaris, Intel Xeon/Linux, Compaq-Alpha/Tru64) *Vmatch* is available as a 64 bit version, enabling gigabytes of sequences to be processed.

Flexible input format

The most common formats for input sequences (Fasta, Genbank, EMBL, and SWISSPROT) are accepted. The user does not have to specify the input format. It is automatically recognized. All input files can contain an arbitrary number of sequences. Gzipped compressed inputs are accepted.

Customized output and match selection

*Vmatch*’s output can be parsed by other programs easily. Furthermore, several options allow for its customization. XML output is available and new output formats can easily be incorporated without changing *Vmatch*’s program code. Certain matches can easily be selected by user defined criteria, without intermediate output and subsequent parsing.

The parts of Vmatch

Up until now we have referred to *Vmatch* as a collection of programs. In the following we use the same name, *vmatch* (in typewriter font), for the most important program in this collection. Besides *vmatch*, there are the following programs available:

1. *mkvtree* constructs the persistent index and stores it on files.
2. *mkdna6idx* constructs an index for a DNA sequence after translating this in all six reading frames.
3. *vseqinfo* delivers information about indexed database sequences.
4. *vstree2tex* outputs a representation of the index in LATEX-format. It can be used, for example, for educational or debugging purposes.
5. *vseqselect* selects indexed sequences satisfying specific criteria.
6. vsubseqselect selects substrings of a specified length range from an index.

7. vmigrate.sh converts an index from big endian to little endian architectures, or vice versa.

8. vmatchselect sort and selects matches delivered by vmatch.

9. chain2dim computes optimal chains of matches from files in Vmatch-format.

10. matchcluster computes clusters of matches from files in Vmatch-format.

Figure 1 depicts the data flow in the different tools described in this manual.

Overview

The rest of this document is organized as follows: In Section 2 we describe mkvtree, a program which constructs the index. For a DNA sequence we can construct a six-frame index using the programming mkdna6idx, described in section 3. Section 4 is devoted to the program vseqinfo which delivers information about the indexed database sequences. In Section 5 we describe the program vstree2tex which outputs the tables of a virtual suffix tree in \LaTeX-format. Section 6 is devoted to the program vseqselect which allows to select sequences from an index satisfying specific criteria. Similarly, the program vsubseqselect described in Section 7 selects substrings of a specified length range from an index. Section 8 describes the shell-script vmigrate.sh to convert an index from big endian (e.g. SUN/Sparc or SGI/IRIX) to little endian (e.g. Linux or Compaq-Alpha) architectures, or vice versa. In Section 9 we explain how to use the program vmatch which allows to solve a variety of matching tasks on the index. Section 10 is devoted to the program vmatchselect, which allows to sort and select matches delivered by vmatch. Each of these sections is completed with several examples showing how to apply the different programs to solve the different matching tasks. In the course of this, the output format is explained.

In Appendix A we define some basic notions. Appendix B explains the format of the symbol mapping. Appendix C goes into detail about the X-drop extension strategy. Substring Specifications are described in Appendix D. Appendix E is devoted to the concept of selection functions. Finally Appendix F explains the tables the index consists of.

2 mkvtree: Making Virtual Suffix Trees

The program mkvtree constructs an index for a given set of sequences. These are given as a list of input files. The sequences are referred to as database sequences. They can be over any given alphabet. The alphabet can be the DNA alphabet, or the protein alphabet, or any other alphabet consisting of printable characters. An alphabet is specified by a file storing a symbol mapping. The index consists of several files, the index files. Each such file stores a different table. The
Figure 1: The Dataflow in Vmatch. The programs are shown in ellipses. The inputs are the database and the query sequences and the alphabet transformation, represented by rectangles. At the center of all computations the persistent index is shown. All other rectangles represent the different kinds of output.
user specifies which tables (i.e. which part of the index) is written to a file, using one of eight output options, or a single option specifying that all tables are written to file. Appendix F gives a comprehensive description of the different tables. We recommend the user to only use the output options -tis, -ois, -suf, -stil, -bwt, -bck, and -lcp. Option -skp is only necessary for approximate pattern matching.

If an error occurs, the program exits with error code 1. Otherwise, the exit code is 0. We support the following formats for the input files. They are recognized according to the first non-white space symbol in the file.

**multiple FASTA format** If the file begins with the symbol >, then this file is considered to be a file in multiple FASTA format (i.e. it contains one or more sequences). Each line starting with the symbol > contains the description of the sequence following it. Each line not starting with the symbol > contains the sequence. Empty lines are allowed and ignored when reading the input.

**multiple EMBL/SWISSPROT format** If the file begins with the string ID, then this file is considered to be a file in multiple EMBL format (i.e. containing one or more sequences, each in EMBL-format). The information contained in the ID and DE-lines is taken as the description of the corresponding sequence. The EMBL format is identical to the SWISSPROT format (w.r.t. the information we need to extract from such entries). So one can also use files in multiple SWISSPROT format as input.

**multiple GENBANK format** If the file begins with the string LOCUS, then this file is considered to be a file in multiple GENBANK format (i.e. containing one or more entries in GENBANK-format). The information contained in the LOCUS and the DEFINITION-lines is taken as the description of the corresponding sequence.

**plain format** If the file does not begin with the symbol > or the strings ID or LOCUS, then the file is taken verbatim. That is, the entire file is considered to be the input sequence (white spaces are not ignored).

There is no special option necessary to tell the program the sequence format. It automatically detects the appropriate format, according to the rules given above. If none of the above rules apply, then the program cannot recognize the input format and exits with error code 1. In such a case please check you input files for if they are conform with the input formats above. Another good solution is to use a more versatile sequence format transformation programs (e.g. readseq) to first generate multiple FASTA files and then feed this into mkvtree.

Today many files containing sequence files are provided compressed by the program gzip. To simplify the use of these files, mkvtree also accepts gzipped input files. These files must have the ending .gz. The gzipped formatted files are gunzipped internally and then processed as any other file.
2.1 The Options of mkvtree

In this subsection, we describe the program mkvtree by explaining its options. An option is always beginning with the dash symbol. We write an option in *typewriter* mode. Arguments to options are written in typewriter mode if they are to be used verbatim. We use *italics* for an option argument if it is to be replaced by a some value. Table 1 gives an overview of the option available in mkvtree.

The program is called as follows:

```
mkvtree options
```

And here is a description of the options:

- **-db** *dbfiles*
  Specify a non empty list of database files separated by white spaces. Each database file contains sequences in one of the formats as specified above. However, the format of all files has to be identical. The sequence must consist of characters over the alphabet as specified by the options `-dna`, `-protein`, or `-smap`, see below. White spaces are ignored. This option is mandatory.

- **-smap** *mapfile*
  Specify the file storing the symbol mapping. If the given *mapfile* cannot be found in the directory where mkvtree is run, then all directories specified by the environment variable `MKVTREESMAPDIR` are searched. If defined correctly, this contains a list of directory paths
separated by colons (`:`). For example, if one uses the csh or the tcsh, the definition of the environment-variable could look like this:

```
$ setenv MKVTREESMAPDIR "$HOME/vstree:/usr/vstree/TRANS"
```

For the bash or the sh the definition could look like

```
$ MKVTREESMAPDIR="$HOME/vstree:/usr/vstree/TRANS"
$ export MKVTREESMAPDIR
```

Then, if `mapfile` is not available in the current directory, `mkvtree` searches for `mapfile` in the two given directories. It scans the directory-list from left to right. As soon as it has found the file it stops. If the file cannot be found an error message is reported and the program exits with error code 1. See Appendix B for a more detailed explanation of the format of the symbol mapping file.

\texttt{-dna}

This option is equivalent to the option \texttt{-smap mapfile} where `mapfile` stores exactly the following 5 lines:

\begin{verbatim}
aA
cC
gG
tTuU
nsywrkvbdhmNSYWRKVBDHM
\end{verbatim}

This specifies an alphabet of size 4 with additional wildcard symbols appearing in the fifth line. See Appendix B for a more detailed explanation of the format of the symbol mapping file.

\texttt{-protein}

This option is equivalent to the option \texttt{-smap mapfile} where `mapfile` stores exactly the following 21 lines:

\begin{verbatim}
L
V
I
F
K
R
E
D
A
G
S
T
N
\end{verbatim}
This specifies an alphabet of size 20 with additional wildcard symbols on the last line. See Appendix B for a more detailed explanation of the format of the symbol mapping file.

-indexname filepath
Specify filepath to be the name of the index, later referred to by indexname. This option is mandatory, if more than one database file is given and if additionally at least one file comprising the index is stored (i.e. if any of the output options is used). If no file from the index is stored, then this option is not allowed. If there is only one database file, and this option is not given, then indexname is the basename of the given filepath, i.e. the filename stripped by the directory path where it is stored. The filepath can be a complete path.

-pl [prefixlength]
Start the sorting of the suffixes with an initial step that sort all suffixes according to their prefixes of length prefixlength. The argument prefixlength is optional. Hence it is denoted in square brackets. If the argument is omitted, then the value for prefixlength is automatically determined. More precisely, it is \( \lceil \log_{k} \frac{n}{8} \rceil \), where \( n \) is the total length of the database sequences and \( k \) is the alphabet size (\( k = 4 \) for DNA, \( k = 20 \) for proteins, and \( k = \) one less than the number of lines in the corresponding symbol mapping file). Table 2 shows the choices for the prefixlength for DNA and protein sequences, given different ranges of values for \( n \). The maximum value for prefixlength is \( \lceil \log_{k} \frac{4n}{8} \rceil \). If you want to choose the prefix length manually, then we recommend prefixlength \( \in [4, 13] \) for DNA sequences, and prefixlength \( \in [2, 5] \) for protein sequences, depending on the total length of the input sequences, see Table 2. If this option is not specified, then the value for prefixlength is 0, in which case no initial bucket sort is performed. Note that in case you want to match a query against the index, the least length for the search must be at least prefixlength.

-tis
Store the table tistab in file indexname.tis.

-ois
Store the table oistab in file indexname.ois.

-suf
Store the table suftab in file indexname.suf.

-stil
Store the reduced version of the inverse suffix table stitab in file indexname.stil.
Table 2: The value of \textit{prefixlength} automatically determined for different ranges for DNA and proteins and different input sizes.

<table>
<thead>
<tr>
<th>DNA range of ( n )</th>
<th>prefixlength</th>
</tr>
</thead>
<tbody>
<tr>
<td>513 – 2,048</td>
<td>4</td>
</tr>
<tr>
<td>2,049 – 8,192</td>
<td>5</td>
</tr>
<tr>
<td>8,193 – 32,768</td>
<td>6</td>
</tr>
<tr>
<td>32,769 – 131,072</td>
<td>7</td>
</tr>
<tr>
<td>131,073 – 524,288</td>
<td>8</td>
</tr>
<tr>
<td>524,289 – 2,097,152</td>
<td>9</td>
</tr>
<tr>
<td>2,097,153 – 8,388,608</td>
<td>10</td>
</tr>
<tr>
<td>8,388,609 – 33,554,432</td>
<td>11</td>
</tr>
<tr>
<td>33,554,433 – 134,217,728</td>
<td>12</td>
</tr>
<tr>
<td>134,217,729 – 536,870,912</td>
<td>13</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Protein range of ( n )</th>
<th>prefixlength</th>
</tr>
</thead>
<tbody>
<tr>
<td>161 – 3,200</td>
<td>2</td>
</tr>
<tr>
<td>3,201 – 64,000</td>
<td>3</td>
</tr>
<tr>
<td>64,001 – 1,280,000</td>
<td>4</td>
</tr>
<tr>
<td>1,280,001 – 25,600,000</td>
<td>5</td>
</tr>
</tbody>
</table>

-\texttt{bwt}

Store the table \texttt{bwttab} and the integer \( i \) satisfying \( suftab[i] = 0 \) in file \texttt{indexname.bwt}.

-\texttt{bck}

Store the table \texttt{bcktab} in file \texttt{indexname.bck}.

-\texttt{lcp}

Store the tables \texttt{lcptab} and \texttt{llvtab} in the files \texttt{indexname.lcp} and \texttt{indexname.llv}.

-\texttt{skp}

Store the table \texttt{skptab} in file \texttt{indexname.skp}.

-\texttt{allout}

This option is equivalent to the combination of the output options -\texttt{tis}, -\texttt{ois}, -\texttt{suf}, -\texttt{stil}, -\texttt{bwt}, -\texttt{bck}, -\texttt{lcp}, and -\texttt{skp}.

-\texttt{v}

Be verbose, that is, give reports about the different steps as well as the resource requirements of the computation. This option is recommended.

-\texttt{version}

Show the version of the Vmatch version, the program is part of. Also report the compilation date and the compilation options.

-\texttt{help}

Show a summary of all options and terminate with exit code 0.
Note the following when combining options of `mkvtree`:

- If any of the output options `-tis`, `-ois`, `-suf`, `-bwt`, `-lcp`, `-bck`, `-skp`, or `-allout` is used, then tables `prjtab`, `ssptab`, `destab`, and `sdstab` are stored in the files `indexname.prj`, `indexname.ssp`, `indexname.des`, and `indexname.sds`.

- Option `-allout` cannot be combined with any other output option.

- Option `-bck` requires that the option `-pl` is used.

- Option `-skp` requires that the options `-lcp` and `-suf` are used.

- The options `-smap`, `-dna`, and `-protein` pairwise exclude each other. If the input files are in FASTA-, EMBL-, SWISSPROT-, or GENBANK-format, then one of these three options is required. If the input files are not in FASTA-, EMBL-, SWISSPROT-, or GENBANK-format, then none of these options is allowed, and the symbol transformation is automatically generated satisfying the following: the symbol in the input files with the smallest ASCII code is mapped to 0, the symbol with the second smallest ASCII code is mapped to 1, etc.

- There must be at least one database file. The maximum number of database files is 256.

In case an error occurs, the error code of `mkvtree` is 1. Otherwise, it the exit code is 0.

### 2.2 Applying `mkvtree`

Suppose we have a file `atEST` in multiple FASTA format containing several EST sequences from the *Arabidopsis thaliana* genome.

```
>gi|5587835|gb|AF078689.1|AF078689 AF078689 Arabidopsis thaliana
AGTGGCTACGGCGGCGGTGGCGGCGATGGAAGTAGACGGATGGTGGTAGGAATGAAAGGCTAGAAGCGGC
GGGAGAAGTATGTGGATAAGAAATAACAAAACTGAGGGGATCATGAAGTTCTTGTTATATTATAGTTTT
CAATCTGAATTTCAATTCAATTCCGCCGCTCGCCTTTTCTCTCCTCCGCTTTTCCGTCTCTCGGGATCTGCCGC
GGGGACGACCTTGTGATGATTATAGCTCTGGAAGGTCATCACAAGGATATAAAAAAAAAAAAAAAAAAAA
>gi|4714049|dbj|C99932.1|C99932 C99932 Arabidopsis thaliana
ATGRAYAMTCAAAAATGGAATMTAGGTTTTCCATATCTCTCTCGCTTTTCTCTTCATCATCTCTTCCAGCTG
...
```

The size of the file is about one megabyte:

```
$ ls -l atEST
-rw-r----- 1 kurtz users 999815 Nov 6 21:37 atEST
```

We want to construct an index using `atEST` as a database file. Since the sequences in `atEST` are DNA sequences, we use the option `-dna` when constructing the index. We add the option `-pl`, since we want to construct table `bcktab`. Moreover, we specify option `-allout` to create all files comprising the index, and `-v` to obtain information about what the program is doing, and the resources it requires.
$ mkvtree -db atEST -dna -pl -allout -v
reading file "atEST"
maximal value for argument of option -pl is 9, recommended value is 8
total length of sequences: 772376 (including 1951 separators)
alphabet of size 5: acgtn
creating file "atEST.tis"
creating file "atEST.ois"
creating file "atEST.des"
creating file "atEST.sds"
creating file "atEST.lcp"
initializing data structures
sorting suffixes according to prefix of length 8
sorting all buckets
creating file "atEST.llv"
creating file "atEST.bck"
creating file "atEST.suf"
creating file "atEST.stil"
creating file "atEST.bwt"
creating file "atEST.prj"
creating file "atEST.all"
creating file "atEST.skp"
overall space peak: main=5.34 MB (7.25 bytes/symbol), secondary=1.69 MB

The previous call of mkvtree has constructed several index files:

$ ls -l atEST.*
-rw-r----- 1 kurtz gistaff 37 2005-02-01 01:43 atEST.all
-rw-r----- 1 kurtz gistaff 524288 2005-02-01 01:43 atEST.bck
-rw-r----- 1 kurtz gistaff 772381 2005-02-01 01:43 atEST.bwt
-rw-r----- 1 kurtz gistaff 215445 2005-02-01 01:43 atEST.des
-rw-r----- 1 kurtz gistaff 772377 2005-02-01 01:43 atEST.lcp
-rw-r----- 1 kurtz gistaff 18536 2005-02-01 01:43 atEST.llv
-rw-r----- 1 kurtz gistaff 772376 2005-02-01 01:43 atEST.ois
-rw-r----- 1 kurtz gistaff 179 2005-02-01 01:43 atEST.prj
-rw-r----- 1 kurtz gistaff 7812 2005-02-01 01:43 atEST.sds
-rw-r----- 1 kurtz gistaff 3089508 2005-02-01 01:43 atEST.skp
-rw-r----- 1 kurtz gistaff 7804 2005-02-01 01:43 atEST.ssp
-rw-r----- 1 kurtz gistaff 7812 2005-02-01 01:43 atEST.suf
-rw-r----- 1 kurtz gistaff 772376 2005-02-01 01:43 atEST.tis

Currently, none of the programs described here requires the table with the skip-values. Hence the file atEST.skp can be safely removed. You can of course directly specify which files you want to generate by giving the appropriate output options.

Here is an example constructing an index for a complete genome, the pathogen Ecoli O157:H7. This is stored in the file EcoliO157H7. Now we explicitely ask for the construction of a subset of the tables comprising the index, using the options -stil, -bwt, -bck, -suf, -lcp, -tis, and -ois.
$ mkvtree -db EcoliO157H7 -v -pl -stil -bwt -dna -bck -suf -lcp -tis -ois -skp
reading file "EcoliO157H7"
maximal value for argument of option -pl is 10, recommended value is 9
total length of sequences: 5471376
alphabet "aAcCcGgTtUuNsNnYyWwRrKkVvDdHhMm" (size 32) mapped to "acgtn" (size 5)
create file "EcoliO157H7.tis"
create file "EcoliO157H7.ois"
create file "EcoliO157H7.des"
create file "EcoliO157H7.sds"
create file "EcoliO157H7.lcp"
initializing data structures
sorting suffixes according to prefix of length 9
sorting all buckets
create file "EcoliO157H7.1lv"
create file "EcoliO157H7.bck"
create file "EcoliO157H7.suf"
create file "EcoliO157H7.stil"
create file "EcoliO157H7.bwt"
create file "EcoliO157H7.prj"
create file "EcoliO157H7.all"
create file "EcoliO157H7.skp"
overall space peak: main=32.46 MB (6.22 bytes/symbol), secondary=10.50 MB

The previous call of mkvtree has constructed several index files:

$ ls -l EcoliO157H7.*
-rw-r----- 1 kurtz gistaff      37 2005-02-01 23:18 EcoliO157H7.all
-rw-r----- 1 kurtz gistaff    2097152 2005-02-01 23:18 EcoliO157H7.bck
-rw-r----- 1 kurtz gistaff   5471381 2005-02-01 23:18 EcoliO157H7.bwt
-rw-r----- 1 kurtz gistaff      30 2005-02-01 23:18 EcoliO157H7.des
-rw-r----- 1 kurtz gistaff   5471377 2005-02-01 23:18 EcoliO157H7.1lv
-rw-r----- 1 kurtz gistaff  1031440 2005-02-01 23:18 EcoliO157H7.lcp
-rw-r----- 1 kurtz gistaff   5471376 2005-02-01 23:18 EcoliO157H7.ois
-rw-r----- 1 kurtz gistaff     186 2005-02-01 23:18 EcoliO157H7.prj
-rw-r----- 1 kurtz gistaff       8 2005-02-01 23:18 EcoliO157H7.sds
-rw-r----- 1 kurtz gistaff   5471377 2005-02-01 23:18 EcoliO157H7.stil
-rw-r----- 1 kurtz gistaff 21885508 2005-02-01 23:18 EcoliO157H7.suf
-rw-r----- 1 kurtz gistaff  5471376 2005-02-01 23:18 EcoliO157H7.tis

Of course, we can also construct an index from more than one file. Suppose we have two bacterial genomes in the files mgen.fna and mpneu.fna: The genome of *M. genitalis* and the genome of *M. pneumoniae*.

$ ls -l mgen.fna mpneu.fna
-rw-r----- 1 kurtz users 589593 Oct 23 18:53 mgen.fna
-rw-r----- 1 kurtz users 829789 Oct 23 18:53 mpneu.fna

$ mkvtree -dna -pl -bck -tis -lcp -suf -indexname bac -db mgen.fna mpneu.fna

Since there are two input files, we have to give a name bac for the index, using the option -indexname. Since we have not used the verbose option -v, the program works silently.

Now let us consider an example where we construct an index from a set of protein sequences given in SWISSPROT-format. The corresponding input file swissp.sp.gz is in gzip format.

After gunzipping the file, the contents looks like this:
$ zcat swissp.sp.gz
ID 110K_PLAKN STANDARD; PRT; 296 AA.
AC P13813;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 01-FEB-1994 (Rel. 28, Last annotation update)
DE 110 KD ANTIGEN (PK110) (FRAGMENT).
OS Plasmodium knowlesi.
OC Eukaryota; Alveolata; Apicomplexa; Haemosporida; Plasmodium.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 88039002.
RA PERLER F.B., MOON A.M., QIANG B.Q., MEDA M., DALTON M., CARD C.,
RA SCHMIDT-ULLRICH R., WALLACH D., LYNCH J., DONELSON J.E.;
RT "Cloning and characterization of an abundant Plasmodium knowlesi
RT antigen which cross reacts with Gambian sera."
DR EMBL; M19152; AAA29471.1; -.
DR PIR; A54527; A54527.
KW Malaria; Antigen; Repeat.
FT NON_TER 1 1
FT DOMAIN 131 296 13.5 X 12 AA TANDEM REPEATS OF E-E-T-Q-K-
FT T-V-E-P-E-Q-T.
SQ SEQUENCE 296 AA; 34077 MW; 666F88DF CRC32;
   FNSNMLRGSV CEEVSLMTS IDNMIEEIDF YEKEIYKGSH SGGVIKMDY DLEDDENED
   EMTEQMVEEV ADHITQDMID EVAHVQLDNH TDHMAHMEEH VHGLSGDVTQ IKEIVQKVNV
   AVEKVVKHVE TEETQKTVEP EQIEETQNTV EPEQTEETQK TVEPEQTEET QNTVEPEQIE
   ETQKTEPEQ TEEAQKTVEP EQTEETQKTV EPEQTEETQK TVEPEQTEET QKTVEPEQTE
   ETQKTEPEQ TEETQKTVEP EQTEETQKTV EPEQTEETQK TVEPEPQTQET QNTVEP
/

We use the gzipped-file like any other file. mkvtree automatically recognizes that it is in
gzipped format (due to the suffix .gz). It gunzips the file internally, and extracts the required
sequence information. Of course, any other sequence formats can also be in gzipped format.

$ mkvtree -protein -db swissp.sp.gz -pl -lcp -suf -tis -ois -bwt -bck -v
reading gzipped file "swissp.sp.gz"
maximal value for argument of option -pl is 2, recommended value is 2
total length of sequences: 4566 (including 7 separators)
alphabet of size 21: LVIFKREDAGSTNQYWPHMCX
creating file "swissp.sp.gz.tis"
creating file "swissp.sp.gz.ois"
creating file "swissp.sp.gz.des"
creating file "swissp.sp.gz.sds"
creating file "swissp.sp.gz.lcp"
initializing data structures
sorting suffixes according to prefix of length 2
sorting all buckets
creating file "swissp.sp.gz.llv"
creating file "swissp.sp.gz.bck"
creating file "swissp.sp.gz.suf"
creating file "swissp.sp.gz.bwt"
creating file "swissp.sp.gz.prj"
creating file "swissp.sp.gz.al1"
overall space peak: main=0.18 MB (40.80 bytes/symbol), secondary=0.00 MB

The description of the sequences is extracted from the ID-field and the DE-field of the SWISS-PROT-entry. For the first sequence, stored in the input file swissp.sp.gz, mkvtree extracts the following description:

110K_PLAKN 110 KD ANTIGEN (PK110) (FRAGMENT).

We now consider a Fasta-File with two protein sequences:

```
$ cat Proteinseq
>QPTFLLPPGEASYFNNIVKRRQTNMFILNNYYFHSKRIRTIEELAEMYLDQVRG
>QGPLFLLPPGEASYFNNIVKRRQTNMVFNYYLHSKRLRTFEELAEMYLDQVRG
```

We produce an index from this file, applying the symbol mapping TransProt11:

```
$ mkvtree -db Proteinseq -smap TransProt11 -pl -allout -v
```

We we produce an index from this file, applying the symbol mapping TransProt11:

```
$ mkvtree -db Proteinseq -smap TransProt11 -pl -allout -v
```

Note that the fourth line of the output reports that the input alphabet is the usual amino acid alphabet including some wildcards, while the transformed alphabet is of size 12.

3 mkdna6idx: generating a six frame translation index

mkdna6idx is very similar to mkvtree. While mkvtree can handle sequences over arbitrary alphabets, mkdna6idx requires DNA-sequences as input. It generates two indices, namely:
A flat index indexname for the given DNA sequences. It mainly consists of the two files indexname.tis and indexname.ois. This index is mainly used for output purpose.

An index indexname.6fr for the given DNA sequences translated in all six reading frames. This is used for computing the matches.

These two indices allow vmatch to compute matches on the protein level.

The program is called as follows:

mkdna6idx options

And here is a description of the options:

-db dbfiles
Specify a non empty list of database files separated by white spaces. Each database file contains sequences in one of the following formats: Fasta, Genbank, EMBL, and SWISS-SPROT. The user does not have to specify the input format. However, the format of all files has to be identical. The sequence must consist of characters over the alphabet a, c, g, t, or u (in lower or upper case), or wildcards n, s, y, w, r, k, v, b, d, h, m. White spaces in the input files are ignored. This option is mandatory. This option is identical with the same option of the program mkvtree.

-smap mapfile
Specify the file storing the symbol mapping applied to the amino acid symbols after the six frame translation. If the given mapfile cannot be found in the directory where mkvtree is run, then all directories specified by the environment variable MKVREETREEMAPDIR are searched. If defined correctly, this contains a list of directory paths separated by colons (`:`). For example, if one uses the csh or the tcsh, the definition of the environment-variable could look like this:

setenv MKVREETREEMAPDIR "$HOME/vstree:/usr/vstree/TRANS"

For the bash or the sh the definition could look like

MKVREETREEMAPDIR="$HOME/vstree:/usr/vstree/TRANS"
export MKVREETREEMAPDIR

Then, if mapfile is not available in the current directory, mkvtree searches for mapfile in the two given directories. It scans the directory-list from left to right. As soon as it has found the file it stops. If the file cannot be found, an error message is reported and the program exits with error code 1. See Appendix B for a more detailed explanation of the format of the symbol mapping file.
Specify the number of the codon translation table used for the six frame translation. \( t \) must be a number in the range \([1, 23]\) except for \(7, 8, 17, 18, 19\) and \(20\). Table 3 gives the possible numbers and their names. The codon translation tables, their numbers, and their names were taken from ftp://ftp.ncbi.nih.gov/entrez/misc/data/gc.prt. If this option is not used, then the standard codon translation table (number 1) is used.

Specify \( \text{filepath} \) to be the name of the index, later referred to by \( \text{indexname} \). This option is mandatory, if more than one database file is given and if additionally at least one file comprising the index is stored (i.e. if any of the output options is used). If no file from the index is stored, then this option is not allowed. If there is only one database file, and this option is not given, then \( \text{indexname} \) is the basename of the given \( \text{filepath} \), i.e. the filename stripped by the directory path where it is stored. The \( \text{filepath} \) can be a complete path.

Store the table \( \text{tistab} \) in file \( \text{indexname.6fr.tis} \).

Store the table \( \text{oistab} \) in file \( \text{indexname.6fr.ois} \).

Be verbose, that is, give reports about the different steps as well as the resource requirements of the computation. This option is recommended.

Show the version of the program. Also report the compilation date and the compilation options.

Show a summary of all options and terminate.

An example application of \( \text{mkdna6idx} \) is given in section 9.9.4.

## 4 vseqinfo: Obtaining Sequence Information

vseqinfo echoes for each database sequence its length and its description. The program has no options. It takes exactly one argument, namely the index name. The output goes to standard out.

### 4.1 Applying vseqinfo

We can obtain the information about the database sequences in \( \text{atEST} \).
1  Standard
2  Vertebrate Mitochondrial
3  Yeast Mitochondrial
4  Mold Mitochondrial; Protozoan Mitochondrial; Coelenterate Mitochondrial; Mycoplasma; Spiroplasma
5  Invertebrate Mitochondrial
6  Ciliate Nuclear; Dasycladacean Nuclear; Hexamita Nuclear
9  Echinoderm Mitochondrial
10  Euplotid Nuclear
11  Bacterial
12  Alternative Yeast Nuclear
13  Ascidian Mitochondrial
14  Flatworm Mitochondrial
15  Blepharisma Macronuclear
16  Chlorophycean Mitochondrial
21  Trematode Mitochondrial
22  Scenedesmus Obliquus Mitochondrial
23  Thraustochytrium Mitochondrial

Table 3: The possible codon translation table numbers and table names.

$ vseqinfo atEST
0 275 gi|5587835|gb|AF078689.1|AF078689 AF078689 Arabidopsis thaliana lib
1 852 gi|4714049|dbj|C99932.1|C99932 C99932 Arabidopsis thaliana library
2 719 gi|4714048|dbj|C99931.1|C99931 C99931 Arabidopsis thaliana library
3 746 gi|4714047|dbj|C99930.1|C99930 C99930 Arabidopsis thaliana library
4 668 gi|4714046|dbj|C99929.1|C99929 C99929 Arabidopsis thaliana library
5 800 gi|4714045|dbj|C99928.1|C99928 C99928 Arabidopsis thaliana library

5 vstree2tex: Pretty Printing a Virtual Tree

The program vstree2tex produces a representation of a virtual suffix tree in \LaTeX-format and outputs it to standard out.

Note that vstree2tex should only be used for very small indexes since it produces large output files. Suppose the total length of all sequences in the index is $n$. If the option $-s$ is not used, then the output size of vstree2tex is about $10n$ bytes per option. (plus some constant number of bytes for the header and the footer of the \LaTeX-file). If the option $-s$ is used, then the size of the output is proportional to $n^2$. The program is mainly designed for debugging a program based on the index and for educational purposes.

The program is called as follows:

vstree2tex [options] indexname
And here is a description of the options:

- **s**
  Output the suffixes. The \( i \)th separator symbol in a multiple sequence is shown as \( i \).

- **tis**
  Output table \( tistab \). The \( i \)th separator symbol in a multiple sequence is shown as \( i \).

- **ois**
  Output table \( oistab \). The \( i \)th separator symbol in a multiple sequence is shown as \( i \).

- **suf**
  Output table \( suftab \).

- **stil**
  Output table \( stitab1 \).

- **bwt**
  Output table \( bwttab \). The \( i \)th separator symbol in a multiple sequence is shown as \( i \).

- **bck**
  Output table \( bcktab \).

- **lcp**
  Output table \( lcptab \).

- **skp**
  Output table \( skptab \).

- **help**
  Show a summary of all options and terminate with exit code 0.

## 5.1 Applying \texttt{vstree2tex}

Suppose a file with the following contents:

```plaintext
> a short sequence
acaacata
tat
```

Let us construct the corresponding index:

```bash
$ mkvtree -dna -db smallseq.fna -pl -allout
```

Now run \texttt{vstree2tex} to produce the table in \LaTeX\-format:
Table 4: The output of vstree2tex after applying \LaTeX{} to it.

<table>
<thead>
<tr>
<th>i</th>
<th>oistab</th>
<th>suftab</th>
<th>lctab</th>
<th>bwttab</th>
<th>S_{suftab}[i]</th>
<th>w</th>
<th>bcktab[\phi(w)]</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>a</td>
<td>2</td>
<td></td>
<td></td>
<td>c aaacatat$</td>
<td>a</td>
<td>(0, 5)</td>
</tr>
<tr>
<td>1</td>
<td>c</td>
<td>3</td>
<td>2</td>
<td>a</td>
<td>a aaacatat$</td>
<td>c</td>
<td>(6, 7)</td>
</tr>
<tr>
<td>2</td>
<td>a</td>
<td>0</td>
<td>1</td>
<td></td>
<td>acaaacatat$</td>
<td>g</td>
<td>(1, 0)</td>
</tr>
<tr>
<td>3</td>
<td>a</td>
<td>4</td>
<td>3</td>
<td>a</td>
<td>aacatat$</td>
<td>t</td>
<td>(8, 9)</td>
</tr>
<tr>
<td>4</td>
<td>a</td>
<td>6</td>
<td>1</td>
<td>c</td>
<td>atat$</td>
<td></td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>c</td>
<td>8</td>
<td>2</td>
<td></td>
<td>t at$</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6</td>
<td>a</td>
<td>1</td>
<td>0</td>
<td>a</td>
<td>caaacatat$</td>
<td></td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>t</td>
<td>5</td>
<td>2</td>
<td>a</td>
<td>catat$</td>
<td></td>
<td></td>
</tr>
<tr>
<td>8</td>
<td>a</td>
<td>7</td>
<td>0</td>
<td>a</td>
<td>tat$</td>
<td></td>
<td></td>
</tr>
<tr>
<td>9</td>
<td>t</td>
<td>9</td>
<td>1</td>
<td>a</td>
<td>t$</td>
<td></td>
<td></td>
</tr>
<tr>
<td>10</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

$ \text{vstree2tex -ois -suf -bwt -bck -lcp -s smallseq.fna > tmp.tex}$

Then the command

$ \text{latex tmp.tex}$

produces Table 4.

6 vseqselect: Selecting Sequences

The program vseqselect selects sequences from a given index and prints them on standard out.

The program is called as follows:

vseqselect [options] indexname

And here is a description of the options:

- \textbf{-minlength} $\alpha$
  Select a sequence only if it is not shorter than $\alpha$.

- \textbf{-maxlength} $\omega$
  Select a sequence only if it is not longer than $\omega$.

- \textbf{-seqnum} filename
  Select the sequences with the numbers given line by line in file filename.
6 VSEQSELECT

-randomnum k
  Randomly select k sequences from the index.

-randomlength ℓ
  Randomly select sequences from the index until the total length of the selected sequences is ℓ or larger.

-help
  Show a summary of all options and terminate with exit code 0.

The options -randomnum, -randomlength, and -seqnum cannot be pairwise combined. However, each of these options can be combined with the option -minlength, and -maxlength. This allows the selection of sequences by random or by numbers specified in a file, satisfying additional constraints.

6.1 Applying vseqselect

Let us start with an example which randomly selects 3 sequences of length between 100 and 200 from the index atEST.

$ vseqselect -randomnum 3 -minlength 100 -maxlength 200 atEST
>gi|2764371|gb|T46124.1|T46124 9387 Lambda-PRL2 Arabidopsis thaliana cDNA clone 135
CCAAGCTCACATCCGATGCAGCCTTGGACCCCGCTGGTCTTGTGCGCTGTGGCTCACGTCTTTNCCCTTTNNGTTGGTGTTTCCATAGCCGNCAACATCTCCGGCGGACACCTTAACCCCGCCGTNACACTTCGGTCTCGCCGTCGGTGGCAACATCACAG
>gi|3868949|dbj|AB015100.1|AB015100 95411 Lambda-PRL2 Arabidopsis thaliana cDNA clone 136
TTTTTGGATGATGACAAGGTCTTAGCTTTGGTGGACATAACTCCCCAGGGGTAACTGGGCTCTCTAAGGGAAAAAGCACAATCGACATCTTGGGCCACCTACTCTACACTGCCTAACTTTTCTCAAAACAAAAAGGCCTAGCA
>gi|2764374|gb|T46197.1|T46197 9460 Lambda-PRL2 Arabidopsis thaliana cDNA clone 138
AATTTTTTTTTTNTTTCAGCAAAAAAGCAAAAAGGACTATACAGGTGATGGACATTCATGAC'TNCCTAAACCTAAGAAAGGTGGATTTTNCCCAAGAAAAGGCGGGAGTNCTCAAGGTGGTAACACNGGTAAATTGAGAACGTGAGTTAAGTNGGAAGAAGAAAAAGACAGAGANC
AAAGCTCTGTTTCT

For each selected sequence which appears in the output, lines of the following form are output on standard error:

# 0: select sequence with number 1579 (length 166)
# 1: select sequence with number 161 (length 186)
# 2: select sequence with number 1576 (length 194)

The first number in each line is always the consecutive number of the selected sequences. Alternatively we can randomly select sequences of some total length:

$ vseqselect -randomlength 300 -minlength 30 -maxlength 99 atEST
>gi|4713981|dbj|C99855.1|C99855 C99855 Arabidopsis thaliana library (Motohashi R) A
Now suppose we have stored the following sequence numbers in a file called myseqnum:

69
51
109
127

Then the program call

\$ vseqselect -seqnum myseqnum atEST

produces the same result as the previous call. A final example shows how to search for all sequences whose description matches the pattern T46124.1:

\$ vseqselect -matchdesc T46124.1 atEST

7  **vsubseqselect**: Selecting Subsequences

The program **vsubseqselect** selects subsequences from a given index and prints them on standard out, either line by line or in FASTA format. The selection can either be random or according to position ranges specified by the user.

The program is called as follows:

vsubseqselect [options] indexname

And here is a description of the options:

- **-minlength** $\alpha$
  Select subsequences from the index whose minimum length is $\alpha$.

- **-maxlength** $\omega$
  Select subsequences from the index whose maximum length is $\omega$. 
7 VSUBSEQSELECT

-snum k
Select k sequences from the index in the specified length range (as given by the options -minlength and -maxlength).

-range k l
Report the substring in the index starting at position k and ending at position l.

-seq l n r
Report the substring in the index of length l in sequence number n starting at the relative position r.

-help
Show a summary of all options and terminate with exit code 0.

The three options -minlength, -maxlength, and -snum can only be used in combination with each other. No other option can be combined with any of these options. The combination of these options delivers sequences s_0, s_1, ..., s_{k-1}, such that for all i ∈ [0, k – 1] the following holds: If i is even, then s_i occurs in the index. If i is odd, then the reverse of s_i occurs in the index. Options -range and -seq exclude each other. Hence there are basically three choices of options:

- If the options -minlength, -maxlength, and -snum with argument k is used, then k subsequences are randomly selected, all of which occur in the given index. The lengths of the selected sequences are evenly distributed over the interval [α, ω]. Besides the sequences, the distribution of the length ranges of the selected subsequences is reported.
- If the option -range is used with two arguments k and l, then the substring starting at position k and ending at position l is shown.
- If the option -seq is used with three arguments l, n, and r, then the substring of length n in sequence number l starting at the relative position r is shown.

7.1 Applying vsubseqselect

The following command shows how to randomly select 5 subsequences of length between 30 and 40 from the index atEST:

$ vsubseqselect -minlength 30 -maxlength 40 -snum 5 atEST
agttacttttcctctttttctgtgttggaag
tctctacctagcatactcttcctcaatc
cctttattttcctcttcattaatgctttccgtagt
ataagaattgaactactgattagtt
gttctttgtgttgcattcgatacttttcctc
# 1 subsequences of length 32
# 1 subsequences of length 33
# 2 subsequences of length 34
# 1 subsequences of length 38
Now suppose we want to select the first 30 residues of index swissp.sp.gz:

$ vsubseqselect -range 0 29 swissp.sp.gz
>110K_PLAKN 110 KD ANTIGEN (PK110) (FRAGMENT). swissp.sp.gz [0,29]
FNSNMLRGSVCEEDVSLMTSIDNMIEEIDF

8 vmigrate.sh: Migrating an Index

Some of the files comprising the virtual suffix tree contain integers in binary format. Such a binary format is not portable to a different computer architecture in general, since the byte order for 2-byte or 4-byte integers may differ. However, in some cases the user wants to construct a virtual suffix tree on a machine with large memory (e.g. a SUN-Sparc Server) and then use the virtual suffix tree on a machine with much less memory (e.g. a Linux-Laptop). For such a case the user should copy all files to the Laptop and then call vmigrate.sh indexname. This transforms the index with name indexname. That is, it reverses the byte order for all files containing 4-byte integers. A second call vmigrate.sh indexname results in the original files, i.e. it reverses the first transformation. The shell-script vmigrate.sh calls a program vendian which performs the transformation for the appropriate files. It is part of the virtual suffix tree distribution. vmigrate.sh takes the index as an argument and does not have any options.

8.1 Applying vmigrate.sh

Suppose we have created a very large index vlsi using mkvtree on a large, say SUN-server. This index cannot directly be used on our Linux-machine. So we convert it as follows:

$ vmigrate.sh vlsi
# migrate the indexfile vlsi.bck llv suf sds ssp skp
Converting vlsi.bck
Converting vlsi.llv
Converting vlsi.suf
Converting vlsi.sds
Converting vlsi.ssp
Converting vlsi.skp
Done.

The conversion of the byte order is only done for the files containing integers. The size of each converted file remains unchanged.

9 vmatch: Solving Matching Tasks

The program vmatch allows to solve a multitude of different matching tasks over an index constructed by mkvtree. Each matching task is solved by a combination of options specifying
• the input,
• the kind of matches sought,
• additional constraints on the matches,
• the direction of the matches (in case of DNA),
• the kind of postprocessing to be done,
• the output mode and output format.

Additionally, if there is more than one algorithm to solve a certain matching task, vmatch allows to specify which algorithm is to be used.

vmatch allows to compute the following kinds of matches:

(1) match all substrings of the database sequences against itself. The matches can be one of the following kinds:

   (a) branching tandem repeats, i.e. repeats where the two instances of the repeat occur at consecutive positions
   (b) maximal repeats, i.e. pairs of maximal substrings occurring more than once in the database sequences
   (c) supermaximal repeats, i.e. pairs of maximal substrings occurring more than once in the database sequences, but not in any other maximal repeat

(2) match a set of query sequences (given in an extra query file) against the index. The matches can be one of the following kinds:

   (a) maximal substring matches, i.e. the substrings of the query sequences matching substrings of the database sequences. All matches exceeding some minimum length, extended maximally to the left and to the right, are reported.
   (b) maximal unique matches, i.e. the substrings of the query sequences matching substrings of the database sequences. A match is reported if it is unique in the database sequences as well as in the query sequences.
   (c) complete matches, i.e. a query sequence must completely match (i.e. from the first character to the last character) a substring of the database sequences.

For all these match kinds, the matches themselves can be direct or palindromic (i.e. on the reverse strand, in case of DNA sequences). If required, DNA sequences are translated into six reading frames and the matches are computed on the protein level, and reported on the DNA level. Besides exact matches, also degenerate matches with a maximal number of errors (insertions, deletions, and mismatches) are supported. Moreover, degenerate matches can be derived from
exact matches by extending these using a greedy extension strategy. This does not apply to complete matches.

For all different match kinds, the matches delivered by vmatch can be selected according to their E-value, their identity value, or their match score.

In the default case, a match is reported as a formatted row of numbers, containing its lengths, the positions where it occurs, the E-value, the number of errors it contains, the match score, and the identity value. Optionally, an alignment of the sequences that are involved in the match can be reported.

An important feature of vmatch is the capability of directly postprocessing the matches found in the following ways:

(i) inverse output, i.e. report substrings of the database sequences or the query sequences not covered by a match

(ii) masking substrings of the database sequences or the query sequences covered by a match

(iii) clustering of a set of database sequences according to the matches found between these sequences. The output of this option can be a representation of the clusters, or a set of sequences each being representative for a cluster.

(iv) chaining of a set of matches, i.e. finding optimal subsets of all matches which do not cross

(v) clustering of matches according to the pairwise similarities on the sequences involved in the match.

(vi) clustering of matches according to the positions where they occur.

Finally, to accommodate many more kinds of user defined post processing tasks, vmatch provides the concept of selection functions. These provide an open interface which allow arbitrary on-the-fly postprocessing of the matches without output and parsing of the matches. For more details on this concept, see Appendix E.

Since the different kinds of matches can be postprocessed in different ways, using different direction of matches, or additional match constraints, or output modes, or output formats, the number of possible combinations of options is very large. Ignoring trivial variations of options (to e.g. modify the output format), we have generated 1386 different combinations of options. In Section 9.9 we can only give 43 examples for a few number of combinations of options.

vmatch is called as follows.

vmatch options indexname

The last argument of vmatch is always the indexname. This can be a complete path. The output of vmatch goes to standard output. vmatch currently has 39 options, described on about 9 pages of this manual. To allow a new user, to get acquainted with the program, we have divided
the options into different categories. An overview of the option categories with a short one-line description of each option is given in table 5.

9.1 Input Options

-\texttt{queryfiles}
  Specify the query files containing the queries to match against the indexed database sequences. The names of the query files are separated by white spaces. Each query file can be in \textsc{fasta}-, \textsc{embl}-, \textsc{swissprot}-, or \textsc{genbank} format. The format of all files has to be identical. If none of the four formats is used, then the program exits with error code 1. Each file can be in gzipped compressed format, in which case the file must have the ending .gz.

-\texttt{dnasprot} \texttt{t [mapfile]}
  Match six frame translation of query sequences against the protein sequence index. First argument \texttt{t} specifies the number of the codon translation table used for the six frame translation of the query. \texttt{t} must be a number in the range [1,23] except for 7, 8, 17, 18, 19 and 20. Table 3 gives the possible numbers and their names. The codon translation tables were taken from the website ftp://ftp.ncbi.nih.gov/entrez/misc/data/gc.prt. The second optional argument is a filename containing a symbol mapping for DNA sequences. This is used for reading the query sequences. If the second argument is missing, then the standard DNA symbolmap is used, as if option \texttt{-dna} was used for \texttt{mkvtree}. When matching the six frame translation of the query against the index, the symbol mapping is used that was supplied to \texttt{mkvtree}, when computing the index.

9.2 Matchkind Options

-\texttt{tandem}
  Compute branching tandem repeats for an index which only contains database sequences. This option requires the use of the option \texttt{-l} to specify the minimum length of the branching tandem repeats to be reported. A tandem repeat is a sequence that occurs more than once in the database sequence at consecutive positions. For a precise definition of the notion branching repeat, see the end of Appendix A.

-\texttt{supermax}
  Compute supermaximal repeats for an index which only contains database files. This option requires the use of option \texttt{-l} to specify the minimum length of the supermaximal repeats to be reported. A repeat is a sequence that occurs more than once in the database sequence. A repeat is supermaximal if it never occurs as a substring of any other maximal repeat. Suppose that a maximal repeat of length \(l\) occurs at positions \(i_0, i_1, \ldots, i_{k-1}\). Then we report all pairs of positions \(i_q\) and \(i_p\) such that \(q \in [0,k-1], p \in [0,k-1]\), and \(i_q < i_p\), together with the usual information as reported for the other kinds of repeats.
### Table 5: Overview of the `vmatch`-Options sorted by Categories

<table>
<thead>
<tr>
<th>Category</th>
<th>Example Options</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Input parameter</strong></td>
<td>-q, -dnavsprot</td>
</tr>
<tr>
<td><strong>Kind of matches</strong></td>
<td>-tandem, -supermax, -mum, -complete</td>
</tr>
<tr>
<td><strong>Postprocessing of matches</strong></td>
<td>-dbnomatch, -qnomatch, -dbmaskmatch, -qmaskmatch, -dbcluster, -nonredundant, -pp, -selfun</td>
</tr>
<tr>
<td><strong>Algorithms</strong></td>
<td>-online, -qspeedup</td>
</tr>
<tr>
<td><strong>Direction of matches</strong></td>
<td>-d, -p</td>
</tr>
<tr>
<td><strong>Match constraints</strong></td>
<td>-l, -h, -e, -allmax, -seedlength, -hxdrop, -exdrop, -leastscore, -value, -identity</td>
</tr>
<tr>
<td><strong>Output modes</strong></td>
<td>-sort, -best, -i</td>
</tr>
<tr>
<td><strong>Output formats</strong></td>
<td>-s, -showdesc, -f, -absolute, -nodist, -noevalue, -noscore, -noidentity, -v, -version, -help, -help+</td>
</tr>
<tr>
<td><strong>Miscellaneous</strong></td>
<td>-v, -version, -help, -help+</td>
</tr>
</tbody>
</table>

**Input parameter**
- `q` specify files containing queries to be matched
- `dnavsprot` perform six frame translation of DNA

**Kind of matches**
- `tandem` compute right branching tandem repeats
- `supermax` compute supermaximal matches
- `mum` compute maximal unique matches
- `complete` specify that query sequences must match completely

**Postprocessing of matches**
- `dbnomatch` mask all database substrings containing a match
- `qnomatch` show all query substrings not containing a match
- `dbmaskmatch` mask all database substrings containing a match
- `qmaskmatch` mask all query substrings containing a match
- `dbcluster` cluster the database sequences
- `nonredundant` generate file with non-redundant set of sequences;
- `pp` generic postprocessing of matches
- `selfun` specify shared object file containing selection function

**Algorithms**
- `online` run algorithms online without using the index
- `qspeedup` specify speedup level when matching queries

**Direction of matches**
- `d` compute direct matches (default)
- `p` compute palindromic (i.e. reverse complemented matches)

**Match constraints**
- `l` specify that match must have the given length
- `h` specify the allowed hamming distance > 0
- `e` specify the allowed edit distance > 0
- `allmax` show all maximal matches in the order of their computation
- `seedlength` specify the seed length
- `hxdrop` specify the xdrop value for hamming distance extension
- `exdrop` specify the xdrop value for edit distance extension
- `leastscore` specify the minimum score of a match
- `value` specify the maximum E-value of a match
- `identity` specify minimum identity of match in range [1..100%]

**Output modes**
- `sort` sort the matches, additional argument is mode
- `best` show the best matches (those with smallest E-values)
- `i` give information about number of different matches

**Output formats**
- `s` show the alignment of matching sequences
- `showdesc` show sequence description of match
- `f` show filename where match occurs
- `absolute` show absolute positions
- `nodist` do not show distance of match
- `noevalue` do not show E-value of match
- `noscore` do not show score of match
- `noidentity` do not show identity of match

**Miscellaneous**
- `v` verbose mode
- `version` show the version of the Vmatch package
- `help` show basic options
- `help+` show all options
-mum [cand]
Compute maximal substring matches which are unique, both in the database sequences and in the query sequences. Such matches are called maximal unique matches. If the optional argument cand is used, then MUM-candidates are computed, i.e. maximal substring matches that are unique in the database but not necessarily in the query sequence. This option requires the use of the option -l to specify the minimum length of the matches to be reported. Additionally, the option -q is required.

-complete [remred]
Match each query sequence completely against the indexed database sequences. In this case, a query sequences is also called pattern. This option only works in conjunction with option -q. The pattern can be a sequence of characters (sequence pattern) over the same alphabet as the database. If this option is used together with the options -online and -e, then an additional argument remred removes redundant matches from the output.

If none of the matchkind options is given, then vmatch searches for maximal substring matches.

9.3 Postprocessing Options

-dbnomatch r [flag]
Compute substrings of length at least r in the database sequences which are not part of a match computed according to the remaining options. This works as follows: instead of showing the match in the usual way, the positions covered by a match are marked and finally only those substrings which are not marked are output if their length is at least r. The optional argument flag can be any of the following values:

- keepleft means that the left instance of the match is not marked.
- keepright means that the right instance of the match is not marked.
- keepleftifsamesequence means that the left instance of the match is not marked whenever the right instance of the match occurs in the same sequence.
- keeprightifsamesequence means that the right instance of the match is not marked whenever the left instance of the match occurs in the same sequence.

Note that the left instance of a match is always the part of the match in the indexed database sequences. If the index is compared to itself, then right instance of a match is also in the indexed database sequences. If query sequences are matched against the index, then the right instance of a match is in the query sequences. Note that positions that are covered by a match but that are not marked due to the use of one of these flags, may still be missing in the output, because they may be marked by the left or the right instance of a different match. This option was called -allunique in a previous version of this program when comparing an index against itself.
-qnomatch $r$
Compute substrings of length at least $r$ in the query sequences which are not part of a
match computed according to the remaining options. The query files containing the query
sequences then have been specified with the option -q either given to mkvtree or to
vmatch.

-dbmaskmatch (tolower toupper X) [flag]
Compute substring matches but do not output them. Instead, output the complete database
sequence in FASTA format and mask the matching substrings. One of the two keywords
tolower or toupper or a single character $X$ must be used as an argument. It determines
how the matches are masked:

- If the argument to this option is tolower, then all characters of the matching sub-
  strings are transformed from upper case to lower case. This requires that all charac-
  ters in the database sequences are in upper case.
- If the argument to this option is toupper, then all characters of the matching sub-
  strings are transformed from lower case to upper case. This requires that all charac-
  ters in the database sequences are in lower case.
- If the argument to this option is a single visible character, say $X$, then all characters
  of the matching substrings are replaced by $X$.

The optional argument flag can be any of the following values:

keepleft
keepright
keepleftifsamesequence
keeprightifsamesequence

If flag is used, then it has the same semantics as for option -dbnomatch. After showing
all masked sequences, and extra line reports the number of symbols masked. This line
is shown on stderr. If the environment variable VMATCHCOMMENTTOSTDOUT is set to the
value on, then the line is shown on stdout.

-qmaskmatch (tolower toupper X)
Compute substring matches but do not output them. Instead, output the complete query
sequence in FASTA format and mask the matching substrings. This option requires the
option -q specifying query files. The additional argument specifies how the matches are
masked. The semantics is as in option -dbmaskmatch.

-dbcluster $p_{small}$ $p_{large}$ [filenameprefix] (minclsize, maxclsize)
Cluster the database sequences according to the matches found in a self comparison of the
index. $p_{small}$ and $p_{large}$ are integers in the range $[0, 100]$. Initially each database sequence
is put into its own cluster. Then the matches are computed, but they are not shown. In-
stead, they are evaluated to form single linkage clusters. In particular, consider a match
between two database sequences, where the smaller of these sequences is of length $\ell_{\text{small}}$ and the larger of these sequences is of length $\ell_{\text{large}}$. Whenever both match instances cover at least $p_{\text{small}}\%$ of the smaller sequence and $p_{\text{large}}\%$ of the larger sequence (or equivalently when the smaller of both match instances is of length $(\ell_{\text{small}} \cdot p_{\text{small}})/100$ and the larger is of length $(\ell_{\text{large}} \cdot p_{\text{large}})/100$), then the clusters containing these two sequences are joined. After computing all matches, the clusters are output. The given cluster numbers are consecutive numbers beginning with 0. Suppose that the argument `filenameprefix` is specified. Then additional information about the clusters is reported:

- For each cluster, say with number $i$, a file `filenameprefix.s.i.match` is generated storing all matches which were used to form cluster $i$. $s$ is the size of the cluster with number $i$. The matches are reported in the same format as described in Section 9.9, and the format can be modified by the corresponding output options. However, note that the option `-s` does not give you the alignment in the `.match`-file. To obtain this, one can apply the program `vmatchselect` to the match-file. `vmatchselect -s` gives the alignment for each sequence involved in a match.

- If the option `-s` (see below) is specified, then for each cluster, say with number $i$, a file named `filenameprefix.s.i.fna` is generated. $s$ is the size of the cluster, and the file stores all sequences contained in cluster $i$ in multiple FASTA format. This includes the description of the sequence prefixed by the sequence number.

If the third optional argument is used, then there may be a fourth optional argument of the form $(\text{minclsize}, \text{maxclsize})$. minclsize and maxclsize are integers satisfying the following: $\text{minclsize} \geq 1$ and $\text{minclsize} \leq \text{maxclsize}$ or $\text{maxclsize} = 0$. If $\text{maxclsize} = 0$, then this is interpreted as $\infty$. If minclsize and maxclsize are specified, then the `.match` and `.fna` file for a cluster are only output if the cluster has at least minclsize and at most maxclsize elements. The singleton sequences of the index, i.e. those sequences not contained in any cluster are output in a single file named `filenameprefix.single.fna` if minclsize = 1 and if the option `-s` is used. Note that in some Unix-shells the symbols ( and ) have special meaning. So it may be necessary to quote the argument $(\text{minclsize}, \text{maxclsize})$, so that the shell does not evaluate it. For example, single quotes work for the C-Shell and the Bourne-Shell.

Note that most of the options determining the matches to search for, can be combined with the option `-dbcluster`. The corresponding holds for the options determining the output format of the matches. Note also that the option `-s` has a slightly different meaning when used in combination with `-dbcluster`. It produces the additional `.fna` files containing the sequences of a cluster.

`-nonredundant filename`

Output a set of non-redundant sequences stored in the index. The output is in FASTA format and goes to file `filename`. The option only works in combination with option `-dbcluster`. The non-redundant sequences are determined in the following way: All sequences are clustered according to the given options. For each cluster we output the longest sequence from that cluster, including its description, to obtain a FASTA formatted file.
-pp chain chain-parameter

Specify that matches should be chained. chain-parameter is a sequence of the following keywords (each optionally followed by some value):

<table>
<thead>
<tr>
<th>Keyword</th>
<th>meaning</th>
</tr>
</thead>
<tbody>
<tr>
<td>global</td>
<td>global chaining</td>
</tr>
<tr>
<td>local</td>
<td>local chaining</td>
</tr>
<tr>
<td>wf</td>
<td>specify weight factor</td>
</tr>
<tr>
<td>maxgap</td>
<td>specify maximal gap width</td>
</tr>
<tr>
<td>outprefix</td>
<td>specify prefix of output files</td>
</tr>
</tbody>
</table>

When prepending the symbol - to a keyword, one obtains an option of the separate program chain2dim. All strings in chain-parameters that are not keywords, are considered to be arguments of the preceeding option. For example,

-pp chain local 10p wf 1.8 maxgap 10

is translated into the options

-locat 10p -wf 1.8 -maxgap 10

of the program chain2dim. This program is part of the Vmatch-distribution and its options are described in a separate manual. We refer to this chain2dim-manual for a comprehensive description of these options and the meaning of the five keywords in turn. Note that chain2dim reads the matches from a match file before it does the chaining. In contrast vmatch computes matches, stores them internally, and then performs chaining.

-pp matchcluster matchcluster-parameter

Specify that matches should be clustered. matchcluster-parameter is a sequence of the following keywords (each optionally followed by some value):

<table>
<thead>
<tr>
<th>Keyword</th>
<th>meaning</th>
</tr>
</thead>
<tbody>
<tr>
<td>erate</td>
<td>specify maximum error rate for similarity clustering</td>
</tr>
<tr>
<td>gapsize</td>
<td>specify maximum gapsize for gap clustering</td>
</tr>
<tr>
<td>overlap</td>
<td>specify minimum percentage of overlap for overlap clustering</td>
</tr>
<tr>
<td>outprefix</td>
<td>specify prefix of output files</td>
</tr>
</tbody>
</table>

When prepending the symbol - to a keyword, one obtains an option of the separate program matchcluster. All strings in matchcluster-parameter that are not keywords, are considered to be arguments of the preceeding option. For example,

-pp matchcluster erate 35 outprefix clout
is translated into the options

\[-erate\ 35\ -outprefix\ clout\]

of the program \textit{matchcluster}. This program is part of the \textit{Vmatch}-distribution and its options are described in a separate manual. We refer to this \textit{matchcluster}-manual for a comprehensive description of these options and the meaning of the four keywords in turn. Note that \textit{matchcluster} reads the matches from a match file before it computes clusters. In contrast \textit{vmatch} computes matches, stores them internally, and then performs the clustering step.

\textbf{\texttt{-selfun \textit{filename} [extraargs]}}

Access file \textit{filename}, which must be a shared object file containing a selection function bundle. If the access to such a selection function bundle is not possible, then the program terminates with error code 1. If \textit{filename} is not an absolute path (i.e., it does not begin with a forward slash (‘/’), then the file is searched for in the colon-separated list of directories specified by the environment variable \texttt{LD_LIBRARY_PATH}. For details on selection functions see Section \texttt{E}. \textit{filename} must have the appropriate extension. This depends on the particular platform. For most Unix-platforms the extension \texttt{.so} is valid. For HP-UX, the valid file extension is \texttt{.sl}. Option \texttt{-filename} can be followed by an optional list \texttt{extraargs} of extra arguments (not beginning with the symbol ‘-’). These are ignored by \textit{vmatch} and \textit{vmatchselect}. Since they are stored in the argument vector passed to the function \texttt{selectmatchHeader} (see Section \texttt{E}), they can be evaluated in the selection function bundle.

\section*{9.4 Algorithm Options}

\textbf{\texttt{-online}}

Run online algorithms where possible. These do not use the index, except for the original and the transformed input sequences, and are therefore more space efficient. However, the online algorithms usually run not as fast as the index based algorithms. We currently provide this option for three different matching tasks:

- match a substring of the query sequences against an index with or without allowing for errors.
- match each query sequence completely against an index with or without allowing for errors

This option only works together with option \texttt{-q}.

\textbf{\texttt{-qspeedup \textit{i}}}

\texttt{vmatch} provides two algorithms to compute maximal exact substring matches of a data base and a query. We may later add more algorithms. The output of the algorithms are
identical. You can select each of these algorithms, termed Algorithm $i$ for $i \in \{0, 2\}$ by the appropriate choice of $i$:

- Algorithm 0 is the most space efficient algorithm. If there are not many long exact matches between the query sequences and the database sequences, then the algorithm is about as fast as Algorithm 2.
- Algorithm 1 is no longer available.
- Algorithm 2 requires $n$ bytes more space than Algorithm 0. If there are many long exact matches between the query sequences and the database sequences, then this Algorithm is faster than Algorithm 0.

These algorithms are applied only if you specify query files using the option $-q$, or if you are searching for palindromic matches in a self comparison of the index. The default algorithm is Algorithm 2.

### 9.5 Direction Options

$-d$

Report direct matches. This option can be combined with option $-p$.

$-p$

Report palindromic matches in case one is searching DNA sequences. This option can be combined with option $-d$. If neither option $-d$ nor $-p$ is used, then only direct matches are reported.

### 9.6 Matchconstraint Options

$-l \ [\text{mingapsize}] \ [\text{maxgapsize}]$

Specify the length value $\ell$. This must be a positive integer. Only matches of length at least $\ell$ are reported. The optional arguments $\text{mingapsize}$ and $\text{maxgapsize}$ can be specified when searching repeats. $\text{mingapsize}$ and $\text{maxgapsize}$ are (possibly negative) integers specifying the minimum and the maximum gap size between the two instances of a repeat. If $\text{maxgapsize}$ is specified, then also $\text{mingapsize}$ must be specified. If $\text{maxgapsize}$ is not specified, then no upper bound on the gap size is specified. Both optional arguments can be negative specifying that the repeat instances may overlap. For an explanation of the effect of these optional arguments, consider a repeat specified by four integers $(l, i, r, j)$. $l$ is the length of first instance of the repeat. $r$ is the length of second instance of the repeat. $i$ is the start position of the left instance, and $j$ is the start position of the right instance of the repeat. Note that $i < j$. If $i + l - 1 > j$, then the gapsize $g$ of the repeat is $-(i + l - j)$. If $i + l - 1 \leq j$, then the gapsize $g$ of the repeat is $j - (i + l - 1)$. The repeat is accepted if $g \geq \text{mingapsize}$ and $g \leq \text{maxgapsize}$, provided $\text{maxgapsize}$ is defined.
Specify the allowed hamming distance $> 0$. This is the hamming distance option, which only allows mismatches in matches. If this option is used together with option $-l$, then substring matches are sought such that the two instances of the match contain at most $k$ mismatches. Here the mandatory argument $k$ is a positive integer. If this option is used together with option $-\text{complete}$, then complete matches are sought, and the mandatory argument $k$ specifies the number of mismatches allowed in the match. $k$ can have three different forms:

- If $k$ is a positive integer, then this specifies the absolute number of mismatches allowed.
- If $k$ is of the form $i p$, where $i$ is a positive integer, then up to $\frac{i}{100} \cdot m$ mismatches are allowed in a match. Here $m$ is the length of the current query to be searched. Thus the character $p$ specifies the mismatch rate as a percentage of the query length. This mode is called the percentage search mode.
- If $k$ is of the form $i b$, where $i$ is a positive integer, then in a first search phase, the minimum number, say $q$, of mismatches is determined, which still delivers at least one match for the given query. Then all matches with exactly $q$ mismatches are reported. The range of values for $q$ to try in the first phase can be controlled by the positive integer. That is, $q$ must be smaller than or equal to $\frac{i}{100} \cdot m$. This mode is called the best search mode.

Specify the allowed edit distance $> 0$. This is the edit distance option, which allows differences, i.e. mismatches, insertions, and deletions in matches. If this option is used together with option $-l$, then substring matches are sought such that the two instances of the match contain at most $k$ differences. Here the mandatory argument $k$ is a positive integer. If this option is used together with option $-\text{complete}$, then complete matches are sought, and the mandatory argument $k$ specifies the number of differences allowed in the match. $k$ can have three different forms:

- If $k$ is a positive integer, then this specifies the absolute number of differences allowed.
- If $k$ is of the form $i p$, where $i$ is a positive integer, then up to $\frac{i}{100} \cdot m$ differences are allowed in a match. Here $m$ is the length of the current query to be searched. Thus the character $p$ specifies the difference rate as a percentage of the query length. This mode is called the percentage search mode.
- If $k$ is of the form $i b$, where $i$ is a positive integer, then in a first search phase, the minimum number, say $q$, of differences is determined, which still delivers at least one match for the given query. Then all matches with exactly $q$ differences are reported. The range of values for $q$ to try in the first phase can be controlled by the positive integer. That is, $q$ must be smaller than or equal to $\frac{i}{100} \cdot m$. This mode is called the best search mode.
Be very careful when you choose the parameter $k$ and do not use the option \texttt{-complete}. The running time of the program is very dependent on $k$. More precisely, it grows faster than $k^2l$. A reasonable value for $k$ is in the range $[1,10]$ (as was restricted in earlier program versions).

\texttt{-allmax}

Report all maximal matches, in the order in which they are found. We do not recommend to use this option. It is mainly used for compatibility with the program \texttt{REPuter}.

\texttt{-seedlength m}

Set the length of the exact seeds. This option is only allowed together with one of the options \texttt{-e}, \texttt{-h}, \texttt{-exdrop}, or \texttt{-hxdrop}. If options \texttt{-e} or \texttt{-h} is used, then the seed length is the maximum of $\lfloor \ell/(k+1) \rfloor$ and $m$, where the argument $m$ is a positive integer, $k$ is the argument for options \texttt{-e} or \texttt{-h}, respectively, and $\ell$ is the argument for option \texttt{-l}. If options \texttt{-exdrop} or \texttt{-hxdrop} is used, then the seed length is $m$. The user should carefully consider the choice for $m$ if using either option \texttt{-e} or \texttt{-h}: if $m$ is larger than $\lfloor \ell/(k+1) \rfloor$, \texttt{vmatch} may miss some matches of length at least $\ell$ with the specified number $k$ of errors. If the options \texttt{-exdrop} or \texttt{-hxdrop} is used, but not \texttt{-seedlength}, then the default value for the seed length is 30.

\texttt{-hxdrop X}

Specify the Xdrop-score $X$ when extending a seed in both directions allowing only for matches and mismatches. The argument $X$ must be a positive integer smaller or equal to 255. Matches are scored 2 and mismatches are scored $-1$. The extension process stops as soon as the extension involving matches and mismatches has a score smaller than $T - X$ where $T$ is the largest score seen so far. This option requires that one of the options \texttt{-l}, \texttt{-leastscore}, \texttt{-evalue} or \texttt{-identity} is used to restrict the matches. The minimum length of the seeds is specified by the argument to option \texttt{-seedlength}. If this option is not used, then the default value for the seed length is 30.

\texttt{-exdrop X}

Specify the Xdrop-score $X$ when extending a seed in both directions allowing for matches, mismatches, insertions, and deletions. The argument $X$ must be a positive integer smaller or equal to 255. Matches are scored 2, mismatches are scored $-1$, and indels are scored $-2$. The extension procedure is further explained in Appendix C. This option requires that one of the options \texttt{-l}, \texttt{-leastscore}, \texttt{-evalue} or \texttt{-identity} is used. The minimum length of the seeds is specified by the argument to option \texttt{-seedlength}. If this option is not used, then the default value for the seed length is 30.

\texttt{-leastscore ls}

Specify the least score $ls$ a match must have to be reported. That is, if an optimal alignment of the matching sequences has score smaller than the positive integer $ls$, then it is not reported. The score of the alignment is computed from the match/mismatch/indel scores as defined above.
-evalue \( e \)
Specify the maximum E-value \( e \) a match must have to be reported. That is, if a match has an E-value larger than the floating point value \( e \), then it is not reported. The argument \( e \) to this option must either be 0.0 or \( \geq 1.0 \cdot 10^{-300} \).

-identity \( q \)
Specify the minimum identity value \( q \) a match must have to be reported. The identity value of a match is defined by \( 100 \cdot (1 - \frac{d}{l}) \) where \( l \) is the maximum of the lengths of the substrings \( s_1 \) and \( s_2 \) involved in the match and \( d \) is the number of mismatches and indels in the optimal alignment of \( s_1 \) and \( s_2 \). Thus, if the identity value of the match is strictly smaller than the positive integer \( q \), then it is not reported.

## 9.7 Output Options

-sort \( \text{mode} \)
sort the reported matches according to the given mode. The following modes are available:
- \( la \): sort in ascending order of length
- \( ld \): sort in descending order of length
- \( ia \): sort in ascending order of first position
- \( id \): sort in descending order of first position
- \( ja \): sort in ascending order of second position
- \( jd \): sort in descending order of second position
- \( ea \): sort in ascending order of E-value
- \( ed \): sort in descending order of E-value
- \( sa \): sort in ascending order of score
- \( sd \): sort in descending order of score
- \( ida \): sort in ascending order of identity
- \( idd \): sort in descending order of identity

If used for \texttt{vmatch}, then this option requires to also use the option -best.

-best \( m \)
Report at most the best \( m \) matches. \( m \) is a positive integer. The matches are reported according their E-value, their length, and their start position (in this order). That is, if two matches are compared, the most important value is their E-value. The match with the smaller E-value is “better”. If both matches have the same E-value, then the longer match is “better”. If both matches have the same E-value and are of the same length, then their first start position matters, and the matches with the smaller start position is “better”. The reported matches satisfy the given length and error constraints.

-i
Do not show matches, but a distribution (preview information) about the length of the different matches found.
Additionally report an alignment of the two instances of the match. The optional integer argument \( q \) specifies the linewidth. That is, the alignments or matching substrings are formatted to \( q \) symbols per line. If \( q \) is not specified, then the default linewidth is 60. For details see Section 9.9. The optional argument \( \text{flag} \) can be any of the following five values:

- **abbrev** means that the matching substrings of an exact match or a hamming distance match are shown in abbreviated form. That is, the exact matching substring is shown verbatim, while for a hamming distance match a mismatching pair of characters, say \( a \) in the left instance of the match and \( c \) in the right instance of the match, are shown as \([ac]\). The output of an edit distance match is not affected by this flag.

- **abbreviub** has the same meaning as **abbrev**, except that each pair of different symbols in a match containing only mismatches is shown as a single IUB-character, according to the following table:

<table>
<thead>
<tr>
<th></th>
<th>A</th>
<th>C</th>
<th>G</th>
<th>T</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>m</td>
<td>r</td>
<td>w</td>
<td></td>
</tr>
<tr>
<td>C</td>
<td>m</td>
<td>s</td>
<td>y</td>
<td></td>
</tr>
<tr>
<td>G</td>
<td>r</td>
<td>s</td>
<td>k</td>
<td></td>
</tr>
<tr>
<td>T</td>
<td>w</td>
<td>y</td>
<td>k</td>
<td></td>
</tr>
</tbody>
</table>

Note that a mismatch of a wildcard character, say \( n \), with a symbol \( a \) is still reported as \([na]\) or \([an]\) since there is no IUB-character for denoting the set consisting of \( a \) and \( n \). The output of an edit distance match is not affected by this flag.

- **leftseq** means that only the sequence information of the left instance of a match is reported.

- **rightseq** means that only the sequence information of the right instance of a match is reported.

- **xml** means that the entire output is in XML-format, see an example in section 9.9.2.

**-showdesc \( m \)**

Instead of the sequence number, show the descriptions of the sequences. The mandatory argument \( m \) can have three forms:

- If \( m \) is a positive integer, then up to the first \( m \) symbols of the descriptions of the sequences are shown.

- If \( m \) is 0, then the descriptions of the sequences is shown up to the first white space character. If there is no such character, then the complete description is shown.

- If \( m \) is of the form \((\text{skipprefix}, \text{maxlength})\) where \( \text{skipprefix} \) and \( \text{maxlength} \) are non-negative integers, then the initial \( \text{skipprefix} \) symbols of the sequence descriptions are skipped. Moreover, if \( \text{maxlength} > 0 \), then up to \( \text{maxlength} \) remaining symbols are shown. If \( \text{maxlength} \) is 0, then the remaining symbols of the description are shown.
up to first white space character. If there is no such white space character, then the remaining part of the description is shown.

White spaces in the descriptions are replaced by the symbol _, so that each description is shown in a single column of the output. This simplifies parsing of the output file.

-f

Additionally report the filename the match instance is contained in.

-absolute

Show the absolute positions instead of pairs of sequence numbers and relative positions.

-nodist

Do not show the distance value of a match.

-noevalue

Do not report the E-value of a match.

-noscore

Do not report the score of a match.

-noidentity

Do not report the identity value of a match.

The previous four options reduce the size of the output considerably, and in a lot of cases also the running time of the program. This is especially true when comparing an index against itself. Then the matches are computed so fast that the generation of the output takes a considerable share of the total running time. Note however, that the options -noevalue and -nodist produce output that cannot parsed by the program vmatchselect.

9.8 Miscellaneous Options

-v

Be verbose, that is, give reports about the different steps as well as the resource requirements of the computation. Additionally, it produces an initial explanation of the format in which the matches are shown.

-help

Show a summary of all basic options and terminate with exit code 0.

-help+

Show a summary of all options and terminate with exit code 0.

Note the following when combining options of vmatch:
1. Option `-seedlength` can only be used in combination with exactly one of the options `-h`, `-e`, `-hxdrop`, and `-exdrop`.

2. Option `-online` only works together with option `-q`.

3. Option `-complete` only works together with option `-q`.

4. Option `-mum` only works together with option `-l`.

5. Option `-super` only works together with option `-l`.

6. Option `-tandem` only works together with option `-l`.

7. Option `-allmax` only works together with either option `-h` or option `-e`.

8. Option `-nonredundant` only works together with option `-dbcluster`.

9. Option `-qspeedup` only works together with either option `-query` or option `-p`.

10. Options `-hxdrop` and `-exdrop` can only be used in combination with at least one of the options `-l`, `-leastscore`, `-identity`, or `-evalue`.

11. If option `-complete` is not used, then either option `-l` or `-exdrop` or `-hxdrop` must be used.

12. If option `-complete` is not used, then options `-e` or `-h` require to use option `-l`.

Table 6 shows all 102 combinations of options that cannot be pairwise combined.

If the options `-best`, `-allmax`, and `-complete` are not used, then for each seed a best match, i.e. one with a minimum E-value is output. The matches are enumerated in the order as they are found. There is no limit on the number of matches reported.

### 9.9 Applying `vmatch`

#### 9.9.1 Self Comparison

Using the index `atEST` generated by `mkvtree` (see Section 2.2), we perform a self comparison of the database sequences, searching for exact duplicates of length at least 350 in `atEST`:

```bash
$ vmatch -v -l 350 atEST
# args=-v -l 350 atEST
# matches are reported in the following way
# l(S) n(S) r(S) t l(S) n(S) r(S) d e s i
# where:
#  l = length
#  n = sequence number
#  r = relative position
#  t = type (D-direct, P-palindromic)
```
Table 6: Options of \texttt{vmatch} that cannot be combined with each other. The entries are meant to be symmetric. That is, if option \textit{a} cannot be combined with option \textit{b}, then option \textit{b} cannot be combined with option \textit{a}.

<table>
<thead>
<tr>
<th>option</th>
<th>cannot be combined with option</th>
</tr>
</thead>
<tbody>
<tr>
<td>-dnavsprot</td>
<td>-supermax -tandem -dbcluster -nonredundant</td>
</tr>
<tr>
<td>-online</td>
<td>-supermax -tandem -dbcluster -nonredundant</td>
</tr>
<tr>
<td>-l</td>
<td>-complete</td>
</tr>
<tr>
<td>-q</td>
<td>-supermax -tandem -dbcluster -nonredundant</td>
</tr>
<tr>
<td>-complete</td>
<td>-dbcluster -nonredundant -allmax -mum -supermax</td>
</tr>
<tr>
<td></td>
<td>-tandem -seedlength -hxdrop -exdrop -qspeedup</td>
</tr>
<tr>
<td>-mum</td>
<td>-supermax -tandem</td>
</tr>
<tr>
<td>-supermax</td>
<td>-tandem -p -qspeedup</td>
</tr>
<tr>
<td>-tandem</td>
<td>-allmax -h -e -hxdrop -exdrop -qnomatch -qmaskmatch</td>
</tr>
<tr>
<td></td>
<td>-p -qsspeedup -dbcluster -nonredundant</td>
</tr>
<tr>
<td>-i</td>
<td>-dbcluster -nonredundant -pp -sort -showdesc -absolute</td>
</tr>
<tr>
<td></td>
<td>-f -dbnomatch -dbmaskmatch -qmaskmatch -qnomatch</td>
</tr>
<tr>
<td></td>
<td>-nodist -novalue -noscore -noidentity -s</td>
</tr>
<tr>
<td>-h</td>
<td>-e -hxdrop -exdrop</td>
</tr>
<tr>
<td>-e</td>
<td>-hxdrop -exdrop</td>
</tr>
<tr>
<td>-hxdrop</td>
<td>-exdrop</td>
</tr>
<tr>
<td>-allmax</td>
<td>-best -sort -hxdrop -exdrop -leastscore -evalue</td>
</tr>
<tr>
<td></td>
<td>-identity</td>
</tr>
<tr>
<td>-showdesc</td>
<td>-absolute</td>
</tr>
<tr>
<td>-dbnomatch</td>
<td>-nodist -novalue -noscore -noidentity -dbcluster</td>
</tr>
<tr>
<td></td>
<td>-nonredundant -dbmaskmatch -qmaskmatch -qnomatch</td>
</tr>
<tr>
<td></td>
<td>-pp</td>
</tr>
<tr>
<td>-qnomatch</td>
<td>-nodist -novalue -noscore -noidentity -dbcluster</td>
</tr>
<tr>
<td></td>
<td>-nonredundant -pp</td>
</tr>
<tr>
<td>-dbmaskmatch</td>
<td>-nodist -novalue -noscore -noidentity -dbcluster</td>
</tr>
<tr>
<td></td>
<td>-nonredundant -qmaskmatch -pp</td>
</tr>
<tr>
<td>-qmaskmatch</td>
<td>-nodist -novalue -noscore -noidentity -dbcluster</td>
</tr>
<tr>
<td></td>
<td>-nonredundant -pp</td>
</tr>
<tr>
<td>-pp</td>
<td>-sort</td>
</tr>
</tbody>
</table>
In all examples, the symbol $\$ is the prompt, after which user input consisting of the called program plus the arguments are shown. In the line beginning with # args=, the arguments of \texttt{vmatch} are echoed. Additionally, the verbose option gives us the remaining lines marked by the symbol #. They explain, in abbreviated form, the meaning of the different columns. Each of the remaining lines, if not marked by the symbol # in the first column, reports a match by showing the following items from left to right:

1. The length of the left instance of the match.
2. The number, say \(i\), of the input sequence, the left instance of the match occurs in. The input sequence numbers are counted from 0.
3. The relative position of the left instance of the match in sequence \(i\).
4. A character \(D\) for direct matches and a character \(P\) for palindromic matches.
5. The length of the right instance of the match.
6. The number, say \(j\), of the input sequence, the right instance of the match occurs in. The input sequence numbers are counted from 0. If the right instance occurs in a query sequence, then \(j\) is relative to the first query sequence.
7. The relative position of the right instance of the match in sequence \(j\).
8. The distance of the match. An exact match has distance 0. A \(k\)-mismatch match with \(k > 0\) mismatches has distance \(-k\). A \(k\)-differences match with \(k > 0\) differences has distance \(k\).
9. The E-value of the match.
(10) The score of the match. The score is computed from the optimal alignment of the left and the right instance of the match. Matching characters in the alignment are scored 2, mismatching characters are scored $-1$, and indels (i.e. insertions and deletions) are scored $-2$.

(11) The identity value $100 \cdot (1 - \frac{d}{\max(l_1, l_2)})$ of the match where $l_1$ and $l_2$ are the lengths of the match instances $s_1$ and $s_2$ and $d$ is the number of mismatches and indels in the optimal alignment of $s_1$ and $s_2$.

Now let us search for palindromic matches with a minimum length of 200, but allowing for up to one mismatch. Instead of a sequence number we output the corresponding sequence description using the option `-showdesc 10`. We additionally report the alignments of the different matches. This alignment is formatted to 60 characters per line. The matching sequences (on the Sbjct-lines) only contain a few mismatches. Each such mismatch between, say character $c_1$ in the left instance of the match (upper line) and character $c_2$ in the right instance of the match (lower line), is emphasized by the symbol `!`.

```
$ vmatch -p -l 200 -h 1 -showdesc 10 -s 60 atEST
# args=-p -l 200 -h 1 -showdesc 10 -s 60 atEST
218 gi|3719113  0 P 218 gi|3450463  75 -1 6.18e-118 -433 99.54
Sbjct: TGAATTGCAAGTCATATAAAAAGACCCAAAAAAATATGAAAAGAGGAGAAAAGAGCAG 60
Sbjct: TGAATTGCAAGTCATATAAAAAGACCCAAAAAAATATGAAAAGAGGAGAAAAGAGCAG 135
Sbjct: AAGCAAAATATTGGAGATCACTAAGATGCGCATGCTGTCGACATGGCATCAGACAC 120
Sbjct: AAGCAAAATATTGGAGATCACTAAGATGCGCATGCTGTCGACATGGCATCAGACAC 195
Sbjct: GCTGAGCACCGCTGGGTGATCTTCCCTCATGTCATCATAAGCCTCTTGTGTCGGTGGC 180
Sbjct: GCTGAGCACCGCTGGGTGATCTTCCCTCATGTCATCATAAGCCTCTTGTGTCGGTGGC 255
Sbjct: CCTTCCTTTTACCTCATCCATCTCAATGGTCATCAGC 218
! Sbjct: CCTTCCTTTTACCTCATCCATCTCAATGGTCATCAGC 293
209 gi|3719089  0 P 209 gi|3450056 231 0 2.48e-115 418 100.00
Sbjct: AAAAGTTTTGAAAACCTTCTCTATACATACATATCCTCCGAGATGGTGTTACTAATCCAACG 60
Sbjct: AAAAGTTTTGAAAACCTTCTCTATACATACATATCCTCCGAGATGGTGTTACTAATCCAACG 291
Sbjct: AAATATAAAAAATTTAACAACAAATTGTATTGTATCATCATCTTTCTCATAGCTTGTCGAGTCATATCAACCAAA 120
Sbjct: AAATATAAAAAATTTAACAACAAATTGTATTGTATCATCATCTTTCTCATAGCTTGTCGAGTCATATCAACCAAA 351
Sbjct: TCCCACTCCGCTCCGCGAAAATACCGAGAAATCGACAGCGAAAACGAGGATCTCCAGG 180
Sbjct: TCCCACTCCGCTCCGCGAAAATACCGAGAAATCGACAGCGAAAACGAGGATCTCCAGG 411
Sbjct: CACAAAAACACAAGCGAGCGAGAGCA 209
Sbjct: CACAAAAACACAAGCGAGCGAGAGCA 440
```

We only show the initial first 10 characters of the description. Note that this time we did not use the verbose option. The output contains one match with a single mismatch and one exact match. Instead of an alignment, we can also report both sequences involved in the match:
Here each mismatch between, say $c_1$ in the left instance of the match and $c_2$ in the right instance of the match, is reported as $[c_1c_2]$. Also note, that we use option $-\text{showdesc} \ (3,7)$ to skip the first 3 characters of the description and to show only the first 7 characters from the beginning of the rest. As an alternative we can use option $-s \ \text{abbreviub}$ to report a mismatch of $c_1$ and $c_2$ as a single IUB character (provided both $c_1$ and $c_2$ are not wildcard symbols).

Instead of showing the two matching sequences at once, we can report the left or the right instance of the the match, using the argument $\text{leftseq}$ or $\text{rightseq}$. This also works for inexact matches:

Now consider an example where we combine the search for direct and palindromic matches. At most 4 differences between the matches are allowed.
Note that, except for option \(-p\), all previously used options for \texttt{vmatch} can also be used for protein sequences. Here is an example involving the index \texttt{Proteinseq}, we constructed earlier:

\begin{verbatim}
$ vmatch -s -l 50 -exdrop 2 Proteinseq
# args=-s -l 50 -exdrop 2 Proteinseq
57 0 1 D 56 1 2 3 2.36e-47 104 94.74
Sbjct: PTFFLLPPGEGGAESYFNNIVKR-RTQMFILNYYFHSKNRIRTIKELAEMYLDQVGRG 58
    !! ! === = = =
Sbjct: PTLFLLPPGEGG--SYFNNIVKRLQRTNMVFNYYLYHSSRKLRTFEELAEMYLDQVGRG 58
\end{verbatim}

Note that, in the alignment, the line between the matching sequences contains the symbols \(!\) and \(=\). As usual, the symbol \(!\) emphasizes mismatches, or insertions, or deletions in the corresponding alignment column. The symbol \(!\) emphasizes columns with different characters, that are equivalent according to the given symbol mapping. For example, the symbols \(F\) and \(L\) in column 3 are equivalent according to the symbol map \texttt{TransProt11}.

Consider some more examples on the computation of maximal repeats for the \textit{Ecoli O157:H7} genome, for which we have earlier constructed an index. To determine the appropriate parameter set for computing matches, we recommend to start with the option \(-i\). This reports the distribution of the length of the matches, but not the match positions or matching sequences.

\begin{verbatim}
$ vmatch -i -l 12 EcoliO157H7
# args=-i -l 12 EcoliO157H7
# all 1507204
# 12 1066546
# 13 301935
# 14 88735
# 15 27097
# 16 8845
# 17 3753
# 18 1693
# 19 1088
# 20 895
# 21 476
# 22 537
# 23 528
# 24 323
# 25 200
# 26 517
# 27 136
# 28 180
# 29 278
\end{verbatim}

This output shows the distribution of the length of all matches of length at least 12. The line starting with the keyword \texttt{all} reports the total number of the maximal repeats of length \(\geq 12\). Then each line of the form

\begin{verbatim}
# \ell k
\end{verbatim}
VMATCH

reports that there are \(k\) direct repeats of length exactly \(\ell\). The output (of which only the first 20 lines are shown), reports that there are more 1 million exact repeats of length 12, and 301935 exact repeats of length 13. We conclude that the minimum length of 12 or 13 is probably not selective enough to compute repeats in the Ecoli O157:H7 genome. A minimum length value of 15 or larger seems to be more appropriate.

We can also quickly get an overview of the number of very long repeats. For example, let us compute the number of matches of length at least 4000 with exact seeds of length 30 (the default seed length) and identity of 85% or more.

```
$ vmatch -i -exdrop 5 -l 4000 -identity 85 EcoliO157H7
# args=-i -exdrop 5 -l 4000 -identity 85 EcoliO157H7
# all 223
# 4113 3
# 4289 29
# 4884 36
# 5446 39
# 5480 19
# 6147 16
# 7494 21
# 7753 46
# 83654 14
```

Instead of restricting the length of the matches, we can restrict the E-value. In the following example, we compute all matches with seed length 20 and E-value smaller than \(10^{-100}\):

```
$ vmatch -seedlength 20 -exdrop 5 -evalue 10e-100 EcoliO157H7
# args=-seedlength 20 -exdrop 5 -evalue 10e-100 EcoliO157H7
2793 0 1654897 D 2812 0 1725693 249 0.00e+00 4858 91.15
1840 0 1280857 D 1848 0 1725693 89 0.00e+00 3421 95.18
7753 0 1272934 D 7773 0 1647113 496 0.00e+00 14038 93.62
2013 0 4746246 D 2017 0 4842428 60 0.00e+00 3850 97.03
3212 0 1267826 D 3216 0 1862616 82 0.00e+00 6182 97.45
2294 0 1641528 D 2278 0 1862438 297 0.00e+00 3681 87.05
2439 0 2688397 D 2440 0 2909349 108 0.00e+00 4555 95.57
395 0 2688268 D 396 0 3492312 123 0.00e+00 422 68.94
502 0 1639778 D 513 0 1860288 94 1.20e-117 733 81.68
451 0 1251809 D 449 0 1628124 79 3.46e-107 663 82.48
6147 0 1287968 D 6141 0 1917801 828 0.00e+00 9804 86.53
665 0 922807 D 670 0 1290526 247 0.00e+00 594 63.13
```

Now suppose we want to compute only the best 10 palindromic repeats, ordered in descending order of their length. We also want to allow a few errors in the repeats.

```
$ vmatch -p -l 1000 -best 10 -exdrop 1 -sort ld EcoliO157H7
# args=-p -l 1000 -best 10 -exdrop 1 -sort ld EcoliO157H7
9441 0 1904945 P 9441 0 2124427 12 0.00e+00 18846 99.87
3873 0 1335481 P 3873 0 2949532 33 0.00e+00 7647 99.15
```
The following example shows how to restrict the computation to supermaximal repeats (which only works for direct repeats):

```
$ vmatch -supermax -l 2000 EcoliO157H7
# args=-supermax -l 2000 EcoliO157H7
5031 0 1106367 D 5031 0 1501974 0 0.00e+00 10062 100.00
22545 0 1066894 D 22545 0 1462501 0 0.00e+00 45090 100.00
4256 0 1125607 D 4256 0 1521213 0 0.00e+00 8512 100.00
2433 0 1143845 D 2433 0 1539451 0 0.00e+00 4866 100.00
2430 0 345369 D 2430 0 4597386 0 0.00e+00 4860 100.00
8316 0 1058577 D 8316 0 1454184 0 0.00e+00 16632 100.00
4923 0 1120684 D 4923 0 1516291 0 0.00e+00 9846 100.00
5904 0 1147797 D 5904 0 1510386 0 0.00e+00 11808 100.00
2473 0 1112305 D 2473 0 1507913 0 0.00e+00 4946 100.00
2162 0 1129864 D 2162 0 1525470 0 0.00e+00 4324 100.00
9438 0 1132027 D 9438 0 1527633 0 0.00e+00 18876 100.00
6002 0 1100364 D 6002 0 1495971 0 0.00e+00 12004 100.00
2583 0 1896937 D 2583 0 2708503 0 0.00e+00 5166 100.00
10923 0 1089440 D 10923 0 1485047 0 0.00e+00 21846 100.00
```

Note that 14 supermaximal repeats are reported. Since there are 19 maximal repeats of length \( \geq 2000 \) (see above), there are five maximal repeats which are not supermaximal. We can further restrict the set of maximal repeats by only computing branching tandem repeats. We decide to only report tandem repeats of length at least 50.

```
$ vmatch -tandem -l 50 EcoliO157H7
# args=-tandem -l 50 EcoliO157H7
62 0 2658616 D 62 0 2658678 0 3.96e-25 124 100.00
282 0 1293997 D 282 0 1294279 0 1.39e-157 564 100.00
141 0 1292479 D 141 0 1294420 0 1.08e-72 282 100.00
162 0 2925000 D 162 0 2925162 0 2.46e-85 324 100.00
57 0 2099948 D 57 0 2100005 0 4.05e-22 114 100.00
87 0 670299 D 87 0 670386 0 3.52e-40 174 100.00
92 0 391492 D 92 0 391584 0 3.43e-43 184 100.00
141 0 4600881 D 141 0 4601022 0 1.08e-72 282 100.00
```

Note that tandem repeats can only be computed on the forward strand. Sometimes the option `-tandem` is too restrictive, and one wants to allow short gaps between the match instances. This can be done by supplying two extra arguments for the option `-l`. 

Note that tandem repeats can only be computed on the forward strand. Sometimes the option `-tandem` is too restrictive, and one wants to allow short gaps between the match instances. This can be done by supplying two extra arguments for the option `-l`. 

9.9.2 Matching Queries against an Index

In the previous applications we had always compared the index against itself. This section shows applications where query sequences are involved. The corresponding query files containing the query sequences are specified via the option `-q`.

**Computing Substring Matches:** Let us match a DNA sequence in file $U89959$ against $atEST$ to find all direct substring matches of length at least 250 with at most 6 differences:

```
$ vmatch -l 250 -e 6 -q U89959 -s 70 atEST
# args=-l 250 -e 6 -q U89959 -s 70 atEST

270 1265 63  D 269 0 72484 6 1.81e-135 521 97.78
Sbjct: AGTATGGGAAGCCCTGTCTCAGCCTNNGCTTCCAGCNCNTCTCCTCCTCCACATCNCNCTTAAAAACTCCACCT 133
! !!! ! !
Query: agtatgggaag-cctgtctcagccacggcttccagcctctcctccacatattaataactccacct 72553

Sbjct: TGGAAGATTATTAGGAGAATGAGAGCGACACGCTCTGTGCTTTCTTTCCTTATGATCCAGCTCTTCCACGC 203
Query: tggaagatttttaggagaatgagagcgacacgctctgtgctttcttcccacattagaacgccacgccagctctctccacgc 72623

Sbjct: ACAAATGAACTATGAAACATATATAAAGCGCACACATATATTTATATGCATATCAGCTTTTGTGATTATG 273
Query: acaaatgaaaactaattatagaaacgagacacacattatataattatgcatatacaagcttttgggtgtattatg 72693

Sbjct: GTATTGATAGTGTTAGTAAAGGCACACATATATTATATGCATATCAGCTTTTGTGATTATG 333
Query: gtattgatagtgattataaagctcgtgtgactatggtattaaaaagaagtataaatcctctctctct 72753
```

So by additionally using the option `-q` we tell `vmatch` to not do a self comparison, but to match all query sequences against $atEST$. The right instance of the match is always in $U89959$, and this file only contains one sequence. Hence the second reported sequence number (in column 6 of the output) is always 0. Moreover, we have used the option `-s 70` to report an optimal alignment between the two match instances. The sequences in the alignment are marked by the keywords `Sbjct` and `Query`. The `Sbjct`-line always shows the the left instance of the match (i.e. the part in the indexed database sequence). The `Query`-line always shows the the right instance of the match (i.e. the part in the query sequence). Alignment columns with insertions, deletions or mismatches are marked by the symbol `!`. The last position of the alignment in every row is shown on the right. If the match instances are longer than the width of a line, it is split into different lines.
The output format is designed to contain all necessary information about the found matches. The kind of information shown on each line can be distinguished according to the first character of the line:

- If the first character of a line is the symbol #, then the rest is a comment. It, for example, shows the arguments of the program. The user can also specify a comment line beginning with

  # smallheading=

  or with

  # largeheading=

  followed by some user defined heading. This can be considered as an annotation, which may be displayed in a graphical user interface visualizing the output of vmatch.

- If the first non-white space character is a digit, then this line shows the match record.

- If the first non-white space character is neither the symbol # nor a digit, then the line shows the match instances, or an alignment of the match instances. In the latter case the line begins with Sbjct or Query.

vmatch can also produce XML-output format. This is not designed for the human eye, but easy to parse by standard XML-parsers. If we replace the parameter 70 of option -s by the keyword xml and add the option -showdesc, then we obtain the following XML output:

```bash
$ vmatch -l 250 -e 6 -q U89959 -s xml -showdesc 0 atEST
<?xml version="1.0"?>
<!DOCTYPE Vmatchoutput PUBLIC "-//VMATCH//VMATCH Vmatchoutput/EN" "Vmatchoutput.dtd"
<Vmatchoutput>
  <Vmatchglobalparams>
    <Vmatchversion>1.0</Vmatchversion>
    <Vmatchindex>atEST</Vmatchindex>
    <Vmatchquery>U89959</Vmatchquery>
    <Vmatchnumofdbseq>1952</Vmatchnumofdbseq>
    <Vmatchdatabaselength>772375</Vmatchdatabaselength>
    <Vmatchnumofqueryseq>1</Vmatchnumofqueryseq>
    <Vmatchquerylength>106972</Vmatchquerylength>
    <Vmatchalphabet>
      <Vmatchalphabetdomainsize>32</Vmatchalphabetdomainsize>
      <Vmatchalphabetmapsize>5</Vmatchalphabetmapsize>
      <Vmatchalphabetmappedwildcards>22</Vmatchalphabetmappedwildcards>
      <Vmatchalphabetundefsymbol>253</Vmatchalphabetundefsymbol>
      <Vmatchalphabetdomain>aAcGcTtUuNsywzrKvbhMNSYWRKVDHM</Vmatchalphabetdomain>
      <Vmatchalphabetverbosechar>acgtn</Vmatchalphabetverbosechar>
    </Vmatchalphabet>
  </Vmatchglobalparams>
  <Vmatchiterationmatches>
    <Match>
      <Vmatchmatchidnumber>0</Vmatchmatchidnumber>
      <Vmatchlength1>270</Vmatchlength1>
      <Vmatchseqnum1>1265</Vmatchseqnum1>
      <Vmatchdescription1>gi|3449674|gb|AI099935.1|AI099935</Vmatchdescription1>
      <Vmatchrelpos1>63</Vmatchrelpos1>
      <Vmatchflag>D</Vmatchflag>
    </Match>
  </Vmatchiterationmatches>
</Vmatchoutput>
```
The XML-output starts with some header giving the kind of output a name, in this case it is called *Vmatchoutput*. The output of a single *vmatch* run is shown between `<Vmatchoutput>` and `</Vmatchoutput>`. In the first part of the output some global parameters are defined, like the name of the index and the query (if any). Their sizes and the numbers of sequences they contain are also given. The alphabet of the given sequences is followed a list of matches. For each match, its ID-number is given, and all important parameters, all tagged such that they are easily identified by an XML-parser to process them by other tools. Since the option `-showdesc` is given, the sequence number and the sequence description is reported. The match is completed by an alignment in form of a sequence of edit operations. There are four kinds of edit operations, namely match, deletion, insertion, and mismatch. For each of these operations a positive number is specified which gives the number of character involved in the operation, i.e. the number of characters matched, or deleted, etc.

**Computing Maximal Unique Matches:** In some applications it is helpful to restrict the set of maximal substring matches to those which only occur exactly once in the database and once in the query. That is, one wants to restrict to maximal unique matches. Using the option `-mum`, we can compute these. In the following example, we only output the best 10 maximal unique matches (out of 28828).
There are 32236 maximal substring matches of length 30 or longer between the two Ecoli genomes. The number of MUM-candidates of this length is 28828, and the number of maximal maximal unique matches is 28409.

**Computing Complete Matches:** Now suppose, we have a set of strings in a file ORFs:

```
$ cat ORFs
>orf1
CTCTTCCAGT
>orf1
GTCTAGGTTT
```

We want to match each of these strings completely against the index. We can use the option `-complete`. We want to allow direct matches as well as palindromic matches. So we combine option `-d` and `-p`. Option `-s` additionally reports the matching sequences. For palindromic matches the reverse complement of the matching sequence is shown:

```
$ vmatch -complete -d -p -q ORFs -s -showdesc 10 atEST
# args=-complete -d -p -q ORFs -s -showdesc 10 atEST
 10 gi|2764223 272 D 10 orf1 0 0 4.14e-01 20 100.00
  Sbjct: CTCTTCCAGT 282
  Query: CTCTTCCAGT

  10 gi|4714046 26 D 10 orf1 0 0 4.14e-01 20 100.00
  Sbjct: GTCTAGGTTT 36
  Query: GTCTAGGTTT

  10 gi|4714003 10 D 10 orf1 0 0 4.14e-01 20 100.00
  Sbjct: GTCTAGGTTT 20
  Query: GTCTAGGTTT

  10 gi|3449668 225 P 10 orf1 0 0 4.14e-01 20 100.00
  Sbjct: AAACCTAGAC 235
  Query: AAACCTAGAC
```

Consider another example where we have two patterns in a file called Patterns:

```
$ cat Patterns
>pattern 1

```
Both patterns do not occur in the \textit{Ecoli O157:H7} genome, as can easily be verified:

\begin{verbatim}
$ vmatch -complete -d -q Patterns EcoliO157H7
# all 145
# 14 2
# 15 9
# 16 18
# 17 23
# 18 35
# 19 22
# 20 20
# 21 9
# 22 7
\end{verbatim}

Thus we obtain 145. Far more than we want to see. Maybe the pattern occurs with less than 4 respectively 5 differences in the genome. We can use the best search mode, to find the best matches, i.e. those with the smallest number of differences:

\begin{verbatim}
$ vmatch -complete -e 25b -q Patterns -s EcoliO157H7
# all 145
# 14 2
# 15 9
# 16 18
# 17 23
# 18 35
# 19 22
# 20 20
# 21 9
# 22 7
\end{verbatim}

That is, for the first pattern we have found a match with three differences, and for the second pattern a match with four differences.
9.9.3 Matching a DNA sequence against a Protein Index

Using the option \(-dnavsprot\), we can match a DNA query sequence against an index built from a protein sequence. The DNA sequence is translated in all six reading frames. The resulting sequence of aminoacids is compared to the protein sequence. The position and length of the match referring to the query are shown with respect to the original DNA sequence. Hence a match of length \(l\) in the indexed protein database sequence is reported as a match of length \(3l\) on the codon level. Consider a file \texttt{Codonseq.fna}:

\[
\text{\$ cat Codonseq.fna}
\]

\[
> \\
\text{TACTTCAACACCGATTACGACAAACAAACATGGTAGTATTCAACAACACTAC} \\
\text{TACTTACACTCAAAACGATTACGAACAAACATGTTCATATTAAACAACACTAC}
\]

We match this file against the index \texttt{Proteinseq}, which was constructed earlier, using the symbol mapping \texttt{TransProt11}:

\[
\text{\$ vmatch -q Codonseq.fna -dnavsprot 1 -e 1 -l 19 -s Proteinseq} \\
\text{# args=-q Codonseq.fna -dnavsprot 1 -e 1 -l 19 -s Proteinseq} \\
\text{28 1 15 F 84 0 0 0 2.71e-84 112 100.00} \\
\text{Sbjct: YFNNIVKRLRQTNNMVFPNYYLHSKRLR} \\
\text{Query: YFNNIVKRLRQTNNMVFPNYYLHSKRLR} \\
\]

\[
\text{27 0 16 F 84 0 0 1 9.03e-81 108 98.81} \\
\text{Sbjct: YFNNIVKR-RQTMFILNNYFHSKIR} \\
\text{Query: YFNNIVKRLRQTNNMVFPNYYLHSKRLR} \\
\]

Note that the symbol mapping \texttt{TransProt11} is inherited from \texttt{Proteinseq}, when comparing the translated codons of the six reading frames of \texttt{Codonseq.fna} with the protein sequence. Also note that the fourth column of the position line shows the symbol \(F\). This means that the right match instance (in the query sequence) refers to one of the three reading frames on the forward strand. In case the right match instance refers to the one of the three reading frames on the reverse strand, the symbol \(G\) would be shown.

9.9.4 Matching a DNA sequence on the Protein Level

To match a DNA sequence on the Protein level, we construct a six frame translation index, using the program \texttt{mkdna6idx}. We do this for the file \texttt{Codonseq.fna} of the previous section.

\[
\text{\$ mkdna6idx -db Codonseq.fna -indexname Codonseq -tis -ois -v} \\
\text{reading file "Codonseq.fna"} \\
\text{total length of sequences: 102} \\
\text{create file "Codonseq.tis"} \\
\text{create file "Codonseq.ois"} \\
\text{create file "Codonseq.des"}
\]
create file "Codonseq.sds"
create file "Codonseq.prj"
create file "Codonseq.al1"
maximal value for argument of option -pl is 1, recommended value is 1
total length of sequences: 205 (including 5 separators)
alphabet "LVIFKREDAGSTNQYWPHMCXUBZ" (size 26) mapped to "LVIFKREDAGSTNQYWPHMCX" (size 21)
create file "Codonseq.6fr.tis"
create file "Codonseq.6fr.ois"
create file "Codonseq.6fr.des"
create file "Codonseq.6fr.sds"
create file "Codonseq.6fr.lcp"
initializing data structures
sorting suffixes according to prefix of length 1
sorting all buckets
create file "Codonseq.6fr.llv"
create file "Codonseq.6fr.suf"
create file "Codonseq.6fr.bwt"
create file "Codonseq.6fr.prj"
create file "Codonseq.6fr.al1"

# overall space peak: main=0.13 MB, secondary=0.00 MB

The generated index has a name ending with suffix .6fr. The additional plain text index named Codonseq is used for transforming and showing the matches. We want to compute exact matches of length at least 12 on the protein level:

```
$ vmatch -l 12 -s Codonseq.6fr
# args=-l 12 -s Codonseq.6fr
12 0 20 F 12 0 71 0 1.15e-12 24 100.00
Sbjct: ACAAACAAACAT 32
  !
Sbjct: ACGAACAAACAT 83

12 0 70 I 12 0 19 0 1.15e-12 24 100.00
Sbjct: TGTTTGTTCGTA 82
  !
Sbjct: TGTTTGTTTGTC 31
```

As usual, the symbol ! shows alignment columns with mismatches. However, the mismatching bases still allow a match on the corresponding translated codon. Note that the fourth column of the position line shows the symbols F and I. These symbols describe if the match instances are on the three reading frames of the forward strand or of the three reading frames on the reverse strand. Since all combinations are possible, there are two additional symbols G and H besides F and I. The following table shows which symbol stands for which combination:

<table>
<thead>
<tr>
<th>symbol</th>
<th>left match instance</th>
<th>right match instance</th>
</tr>
</thead>
<tbody>
<tr>
<td>F</td>
<td>forward strand reading frame</td>
<td>forward strand reading frame</td>
</tr>
<tr>
<td>G</td>
<td>forward strand reading frame</td>
<td>reverse strand reading frame</td>
</tr>
<tr>
<td>H</td>
<td>reverse strand reading frame</td>
<td>forward strand reading frame</td>
</tr>
<tr>
<td>I</td>
<td>reverse strand reading frame</td>
<td>reverse strand reading frame</td>
</tr>
</tbody>
</table>
9.9.5 Computing Regions not containing a Match

There are applications, where it is not important to know where the matches are. Instead, one is interested in knowing those regions of the database sequences or of the query sequences where no match occurs. For these kinds of applications, `vmatch` provides the options `-dbnomatch` and `-qnomatch`. Consider, for example, a comparison of the index against itself. The corresponding matches can be called repeats, since they occur more than once in the same sequence, or sets of sequences. Those parts of the sequence not covered by a repeat can then be considered unique substrings. To compute these, we can use the option `-dbnomatch` which requires an additional argument specifying the least length of the unique substrings to be reported. For example, the following program call reports all substrings of length at least 700 which do not contain a direct repeat of length 25 with at most 1 difference:

```
$ vmatch -dbnomatch 700 -l 25 -e 1 atEST
# args=-dbnomatch 700 -l 25 -e 1 atEST
>10 0 815
>26 0 1019
>32 0 1039
>77 191 742
>304 0 725
```

Each line beginning with the symbol `>` reports the sequence number, the relative start position, and the length of a unique substring (i.e. a substring not covered by a repeat instance) in the sequence under consideration. As usual, the output format can be modified by the options `-f`, `-showdesc`, `-absolute`, and `-s`. For example, if we additionally use the option `-s`, then also the sequence content of the unique substrings is reported. This gives an output file in FASTA-format. Additionally, the `>`-line shows the character distribution of the sequence reported.

The option `-qnomatch` can be used when comparing a set of query sequences against the database sequences. For example, the following program call reports all regions of length 10000 in the genome of *Ecoli K12* which do not contain any exact match to a substring of length 18 or longer in the *Ecoli O157:H7* genome.

```
$ vmatch -qnomatch 10000 -q EcoliK12 -l 18 EcoliO157H7
# args=-qnomatch 10000 -q EcoliK12 -l 18 EcoliO157H7
>0 1385319 10800
>0 1398241 10711
>0 1570168 20414
>0 1599821 12375
>0 2755646 11820
```

Each line beginning with the symbol `>` reports the starting position and the length of a substring in the query sequences which is not common to both genomes under consideration. If we additionally use the option `-s`, then also the sequences in the query which are not common with the database sequences are reported.
Alternatively, we can output the substrings in the *Ecoli O157:H7* genome not occurring in *Ecoli K12*. These could be good candidates to explain the pathogenicity of *Ecoli O157:H7*.

```
$ vmatch -dbnomatch 10000 -q EcoliK12 -l 18 EcoliO157H7
# args=-dbnomatch 10000 -q EcoliK12 -l 18 EcoliO157H7
>0 664171 11617
>0 1087635 12158
>0 1105699 10152
>0 1117238 14153
>0 1278595 13158
>0 1434907 13857
>0 1483242 12158
>0 1501306 10152
>0 1512845 14152
>0 1652802 11515
>0 1718335 11241
>0 1956644 11322
>0 2030357 12870
>0 2111045 12614
>0 2934852 12334
>0 3206647 12308
>0 3729821 15836
>0 4603137 15160
>0 4618316 13631
>0 5319491 10210
>0 5339473 11560
```

### 9.9.6 Masking Matches

The options `-dbmaskmatch` and `-qmaskmatch` are very similar to the `nomatch`-options used in the previous section. While the `nomatch`-options output those parts of the sequences not covered by a match, the `maskmatch`-options allows to mask the matches in the database sequences or in the query sequences. For example, the following program call masks the left instance of substring matches of length \( \geq 20 \) when comparing the index `swissp.sp.gz` against itself. X is used as a masking character.

```
$ vmatch -l 20 -s 90 -dbmaskmatch X swissp.sp.gz
>110K_PLAKN 110 KD ANTIGEN (PK110) (FRAGMENT).
FNSNMLRGSVCEEDVSVSMISVM rangenm11d23s49vsmkshsgV1KMDYDLEDENDEDEDEQMQVEEVADHITQOMIDEEVAHVDN
TDHMHEMEEIEGILGSDGTQIKEIVKQVNAVAVKHIYETETQKTVPEEIETQOTQXXXXXNXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX
ETQKTVPEEQTEEAXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX
XXXXXXXXXXXXXXXXXPTQETQOTNTVEP
```

Of course, the masking options can also be used when matching a query against an index. Then there are two choices. One can either mask the database sequences or the query sequences. For example, the following application masks both instances of the substring matches of length 70 or longer between *Ecoli O157:H7* and *Ecoli K12*. That is, the matches in the database sequence represented by the index `EcoliO157H7` are masked by converting the original lower case characters to upper case characters.
It is easily visible that there are several matches in the first few hundred bases of \textit{EcoliO157H7}. To mask the substrings of the query sequence \textit{Ecoli K12} covered by a match, we replace option \texttt{-dbmaskmatch} by \texttt{-qmaskmatch}. We also use the flag tolower, since the query sequence consists of upper case characters.

Clustering Indexed Sequences

Clustering a set of sequences according to matches occurring in the sequences is an important task. It is done, for example, when clustering ESTs or protein sequences. The following command computes clusters of all database sequences in \textit{atEST}, based on all matches of length at least 200 with at most 10 differences. The matches must cover at least 30\% of the smaller of the two matching sequences and 50\% of the larger of the two sequences.

\begin{verbatim}
$ vmatch -dbcluster 30 50 -p -d -l 200 -e 10 atEST
# args=-dbcluster 30 50 -p -d -l 200 -e 10 atEST
# 152 clusters
# 389 elements out of 1952 (19.93\%) are in clusters
# 1563 elements out of 1952 (80.07\%) are singlets
# 124 clusters of size 2
# 18 clusters of size 3
# 4 clusters of size 4
# 2 clusters of size 5
# 1 cluster of size 6
\end{verbatim}
# 1 cluster of size 7
# 1 cluster of size 8
# 1 cluster of size 40
0: 1646 1862
1: 3 39 6
2: 1764 1896 1745 1767 1882

We only show the first 3 clusters of size at least two. For example, cluster 2 contains the sequences with numbers 1764, 1896, 1745, 1767, 1882.

Instead of the sequence numbers, we can output the descriptions, using the option -showdesc 0.

$ vmatch -dbcluster 30 50 -p -d -l 200 -e 10 -showdesc 0 atEST
# args=-dbcluster 30 50 -p -d -l 200 -e 10 -showdesc 0 atEST
# 152 clusters
# 389 elements out of 1952 (19.93%) are in clusters
# 1563 elements out of 1952 (80.07%) are singlets
# 124 clusters of size 2
# 18 clusters of size 3
# 4 clusters of size 4
# 2 clusters of size 5
# 1 cluster of size 6
# 1 cluster of size 7
# 1 cluster of size 8
# 1 cluster of size 40
0:
  gi|2764304|gb|R65326.1|R65326
  gi|2764088|gb|R30410.1|R30410
1:
  gi|4714047|dbj|C99930.1|C99930
  gi|4714011|dbj|C99887.1|C99887
  gi|4714044|dbj|C99927.1|C99927
2:
  gi|2764186|gb|R64887.1|R64887
  gi|2764054|gb|R30268.1|R30268

To output more information about the clustered sequences we supply the option -dbcluster with a third argument CLatEST and add the option -s.

$ vmatch -dbcluster 30 50 CLatEST -s -p -d -l 200 -e 10 atEST
# args=-dbcluster 30 50 CLatEST -s -p -d -l 200 -e 10 atEST
# 152 clusters
# 389 elements out of 1952 (19.93%) are in clusters
# 1563 elements out of 1952 (80.07%) are singlets
# 124 clusters of size 2
# 18 clusters of size 3
# 4 clusters of size 4
# 2 clusters of size 5
# 1 cluster of size 6
This generates files CLatEST.s.i.match and CLatEST.s.i.fna containing the matches and the clustered sequences for cluster \( i \). \( s \) is the number of sequences in the cluster. For example,

```
$ cat CLatEST.5.2.match
228 1882 70 D 229 1896 73 8 4.36e-106
218 1767 90 D 219 1896 83 7 1.85e-102
218 1745 90 D 219 1896 83 7 1.85e-102
336 1764 74 D 336 1882 63 7 1.39e-171
330 1764 74 D 334 1767 73 10 2.86e-163
310 1745 73 D 308 1764 74 9 1.92e-151
...```

shows the matches responsible for the cluster 2 of size 5. Furthermore,

```
$ cat CLatEST.5.2.fna
>gi|2764186|gb|R64887.1|R64887
Lambda-PRL2 Arabidopsis thaliana cDNA
ACAAGAATCTCAACACTCTTAAACACTCACTTTAAGCCTTNNAAAACACTCAATC
ATGAATCAGCAACATCGCAGGCGCTAGTCTGACAAAGACCCGATTGCTAAAGAAGGGA
ACCAAGCTAGACCTTCCACCGCAGGACCTCCGAGAGCAGGCGAGATCCAGCAGCTG
GACCTTGCAGCCGAGGACACACTGGAATTCAGTGCGCTTCTGAGCTGCTGACAGTGG
ATGCATCGAGACCTTGTTCGAGGATCCAGTCTGAGCAGGGAGAGTTCTACAGGCGT
GCGCTGCACTGCTGCCACCACACTAATGANGCTTCTTTTAATCAAAATGTAATATGTA
ATAAAAAGTGTATGTGGCTACTATGTTAGGCTCAGTNCTNTTATTACTACTCTCTCAGT
ATGGTGGATGAATGGTTATTGACACCTTCTCTCCTCTCCCTATATAAATTNAGACCTT
GCAAAGTTT
>gi|2764054|gb|R30268.1|R30268
Lambda-PRL2 Arabidopsis thaliana cDNA
CTCAACACTCTTAAACACTCACTTTAAGCCTTNNAAAACACTCAATC
ATGAATCAGCAACATCGCAGGCGCTAGTCTGACAAAGACCCGATTGCTAAAGAAGGGA
ACCAAGCTAGACCTTCCACCGCAGGACCTCCGAGAGCAGGCGAGATCCAGCAGCTG
GACCTTGCAGCCGAGGACACACTGGAATTCAGTGCGCTTCTGAGCTGCTGACAGTGG
ATGCATCGAGACCTTGTTCGAGGATCCAGTCTGAGCAGGGAGAGTTCTACAGGCGT
GCGCTGCACTGCTGCCACCACACTAATGANGCTTCTTTTAATCAAAATGTAATATGTA
ATAAAAAGTGTATGTGGCTACTATGTTAGGCTCAGTNCTNTTATTACTACTCTCTCAGT
ATGGTGGATGAATGGTTATTGACACCTTCTCTCCTCTCCCTATATAAATTNAGACCTT
GCAAAGTTT
...```

shows the sequences of this cluster.

### 9.9.8 Computing a non-redundant set of sequences

A colleague recently sent a list of 8124 sequences for different virus strains of the Mouth and Foot Disease. The goal was to produce a multiple alignment of the sequences. Given the GI-numbers of the sequences, it was easy to download the sequences from Genbank resulting in 126 files. The lengths of the sequences is in the range between 7733 and 8280. The total length of all 8124 sequences was. A simple call to mkvtree with all 126 files given as arguments to option `-db`, creates an index for all sequences. The name of the index is mfdindex. Since the construction of the index mfdindex takes longer than expected, there seemed to be long common substrings in the sequences. This can easily be verified by clustering the matches:

```
vmatch -l 7000 -seedlength 1000 -exdrop 2 -dbcluster 100 100 mfdindex
```
This means that the sequences are clustered based on matches of length 7000 or longer with seeds of length 1000 or longer, allowing an Xdrop threshold of 2. The parameters 100 100 to option -dbcluster specify that two sequences are clustered only if a match covers the sequence completely. That is, the sequences in each cluster are pairwise identical.

We obtain 110 clusters and 1 one singlet. That is 8213 sequences are in cluster of size at least two, and one sequence is singleton. So to compute an optimal alignment, it makes sense to only continue with the singlet and exactly one sequence from each cluster. These 111 sequences can be extracted by adding the option -nonredundant to the previous call of vmatch. The argument of this option is the name of the file to store the nonredundant sequences:

```
vmatch -l 7000 -seedlength 1000 -exdrop 2 -dbcluster 100 100 -v -nonredundant nrset mfdindex
```

Having computed the non-redundant set, we can, for example, continue with computing a multiple alignment using the software MGA [5].

### 9.9.9 Chaining Matches

We only give two examples of using the option -pp chain. We compare the E.coli genomes Ecoli K12 and Ecoli O157:H7 and compute all 659 matches of length 500 or longer between these sequences. The first example shows how to compute a global chain. The optimal global chain contains 597 matches (i.e. 91% of all matches), of which we only show the first and the last seven.

```
$ vmatch -l 500 -q EcoliK12 -pp chain global EcoliO157H7
# args=-l 500 -q EcoliK12 -pp chain global EcoliO157H7
# chain 0: length 596 score 835932
512 0 13736 D 512 0 13719 0 0.00e+00 1024 100.00
645 0 59313 D 645 0 54337 0 0.00e+00 1290 100.00
1501 0 59959 D 1501 0 54983 0 0.00e+00 3002 100.00
895 0 61917 D 895 0 56941 0 0.00e+00 1790 100.00
515 0 65876 D 515 0 60943 0 0.00e+00 1030 100.00
502 0 67856 D 502 0 62923 0 0.00e+00 1004 100.00
671 0 5448270 D 671 0 4615599 0 0.00e+00 1342 100.00
915 0 5449694 D 915 0 4617023 0 0.00e+00 1830 100.00
577 0 5450982 D 577 0 4618311 0 0.00e+00 1154 100.00
597 0 5455239 D 597 0 4623611 0 0.00e+00 1194 100.00
553 0 5457015 D 553 0 4625387 0 0.00e+00 1106 100.00
871 0 5457972 D 871 0 4626344 0 0.00e+00 1742 100.00
828 0 5469153 D 828 0 4637525 0 0.00e+00 1656 100.00
```

To find spots of high local similarity, we instead perform local chaining. We compute the local chains with the best and the second best scores: We obtain two chains. The highest scoring chain has score 23118 and is of length 10. The second highest scoring chain has score 12654 and is of length 7:
\$ vmatch -l 500 -q EcoliK12 -pp chain local 2b EcoliO157H7
# args=-l 500 -q EcoliK12 -pp chain local 2b EcoliO157H7
# chain 0: length 7 score 12654
  749 0 193444 D 749 0 189957 0 0.00e+00 1498 100.00
  592 0 194194 D 592 0 190707 0 0.00e+00 1184 100.00
  1210 0 194787 D 1210 0 191300 0 0.00e+00 2420 100.00
  1065 0 196408 D 1065 0 193063 0 0.00e+00 2130 100.00
  505 0 197510 D 505 0 194165 0 0.00e+00 1010 100.00
  2632 0 198385 D 2632 0 195040 0 0.00e+00 5264 100.00
  589 0 201135 D 589 0 197790 0 0.00e+00 1178 100.00
# chain 1: length 10 score 23118
  2153 0 4184202 D 2153 0 3437201 0 0.00e+00 4306 100.00
  1845 0 4186356 D 1845 0 3439355 0 0.00e+00 3690 100.00
  575 0 4188373 D 575 0 3441372 0 0.00e+00 1150 100.00
  537 0 4188949 D 537 0 3441948 0 0.00e+00 1074 100.00
  2567 0 4189568 D 2567 0 3442567 0 0.00e+00 5134 100.00
  798 0 4192136 D 798 0 3445135 0 0.00e+00 1596 100.00
  833 0 4192935 D 833 0 3445934 0 0.00e+00 1666 100.00
  952 0 4194011 D 952 0 3447010 0 0.00e+00 1904 100.00
  1552 0 4195622 D 1552 0 3448621 0 0.00e+00 3104 100.00
  917 0 4197175 D 917 0 3450174 0 0.00e+00 1834 100.00

For examples showing the effect of the different chaining options, see the corresponding manual for the program \textit{chain2dim}.

\subsection*{9.9.10 Clustering Matches}

Suppose we have constructed an index for yeast chromosome III:

\$ mkvtree -dna -db ychrIII.fna -v -tis -ois -bwt -suf -lcp
reading file "ychrIII.fna"
total length of sequences: 315339
create file "ychrIII.fna.tis"
create file "ychrIII.fna.ois"
create file "ychrIII.fna.des"
create file "ychrIII.fna.sds"
create file "ychrIII.fna.lcp"
initializing data structures
sorting suffixes
create file "ychrIII.fna.llv"
create file "ychrIII.fna.suf"
create file "ychrIII.fna.bwt"
create file "ychrIII.fna.prj"
create file "ychrIII.fna.all"
overall space peak: main=2.82 MB (9.38 bytes/symbol), secondary=0.31 MB

Now compute all repeats of length $\geq 100$ in yeast chromosome III, and cluster these matches by similarity, using an error rate of 35. The cluster files produced have the prefix \textit{clout}:
$ vmatch -l 100 -pp matchcluster erate 35 outprefix clout ychrIII.fna
# args=-l 100 -pp matchcluster erate 35 outprefix clout ychrIII.fna
# cluster 16 matches
# create cluster 0 of size 2
# create cluster 1 of size 3
# create cluster 2 of size 2

Three files were generated by this program call:

$ ls -l clout.*
-rw-r----- 1 kurtz gistaff 285 2005-02-21 16:41 clout.2.0.match
-rw-r----- 1 kurtz gistaff 289 2005-02-21 16:41 clout.2.2.match
-rw-r----- 1 kurtz gistaff 491 2005-02-21 16:41 clout.3.1.match

Consider the last file clout.3.1.match representing cluster 1 with three matches:

$ cat clout.3.1.match
# args=-l 100 ychrIII.fna
# id 7
280 0 83954 D 280 0 84469 0 7.41e-159 560 100.00
# id 8
239 0 83954 D 239 0 90099 0 3.58e-134 478 100.00
# id 12
286 0 84422 D 286 0 90052 0 1.81e-162 572 100.00
# linked 8 and 12 with edit distance 47 (error rate 19.67%)
# linked 7 and 12 with edit distance 88 (error rate 31.43%)
# linked 7 and 8 with edit distance 41 (error rate 17.15%)

Cluster 1 thus contains the matches with identification numbers 7, 8, and 12. For example, match 7 and 12 achieve a distance of 88, which corresponds to an error rate of 31.43%, well below the maximum error rate of 35%. Note that the generated files are in a format that they can be read by vmatchselect.

Instead of clustering by similarity, we can cluster by gap size, allowing gaps of size up to 1000.

$ vmatch -l 100 -pp matchcluster gapsize 1000 outprefix clout ychrIII.fna
# args=-l 100 -pp matchcluster gapsize 1000 outprefix clout ychrIII.fna
# cluster 16 matches
# create cluster 0 of size 9
# create cluster 1 of size 4

Consider the file clout.4.1.match representing cluster 1 with four matches: It contains the matches with identification numbers 14, 7, 8, and 12. For example, match 8 and 7 have a gap of 276 = 84469 – 83954 – 239 which is well below the maximum gap size of 1000.
In a third run we cluster the matches by overlap, allowing overlaps of minimum 10%.

Consider the file `clout.3.2.match` representing cluster 2 with three matches: It contains the matches with identification numbers 0, 6, and 1. For example, match 0 and 6, overlap by \(82 = 199591 + 203 - 199712\) positions, which is 40.39% of the length 203 of the longer match.

9.9.11 Selection Functions

The concept of selection functions (bundles) is extensively explained in Appendix E. The following function bundle (only consisting of the function `selectmatch`), defined in a file `sel392.c`, accepts matches of length at most 392.
#include <string.h>
#include "select.h"

int selectmatch(Alphabet *alpha,
                Multiseq *virtualmultiseq,
                Multiseq *querymultiseq,
                StoreMatch *storematch)
{
    if(storematch->Storelength1 <= 392)
    {
        return 1; /* accept */
    } else
    {
        return 0; /* reject */
    }
}

To compile the appropriate shared object file under Linux or a Compaq-Alpha/True64 system, we use the gcc compiler with the following options:

$ gcc -Wall -Werror -O3 -shared sel392.c -o sel392.so

On a SUN-Sparc/Solaris computer, the option -shared is replaced by the option -G. Note that in case you are using the 64-bit version of the program you have to add the compiler option -m64. For statically linked executables of vmatch and vmatchselect selection functions only work under very restrictive circumstances: The machine which runs vmatch or vmatchselect must have the same versions of the libraries as the machine on which the statically linked binaries were compiled.

Now we can select the matches using the shared object sel392.so. This rejects the two matches of length 517 and 402, see Section 9.9.1, above.

$ vmatch -l 350 -selfun sel392.so atEST
   388   79  236   D  388  1495  180   0  4.22e-223  776  100.00
   392  229   0   D  392  270   0   0  1.65e-225  784  100.00
   369  736  141   D  369  1151  85   0  1.16e-211  738  100.00
   369  902  54   D  369  1151  87   0  1.16e-211  738  100.00
   367  736  143   D  367  902  54   0  1.86e-210  734  100.00

In the usual case you want to call the program vmatch or vmatchselect in any directory. If you are using option -selfun, you have to specify the path to the directory where the shared object can be found. This is done by defining the environment variable LD_LIBRARY_PATH. For the csh or the tcsh you can, for example, define it as follows:

$ setenv LD_LIBRARY_PATH ".":"/usr/vstree/SELECT"

The corresponding definition for the bash or the sh is

$ LD_LIBRARY_PATH="."":"/usr/vstree/SELECT"
$ export LD_LIBRARY_PATH
This means that the shared objects are found in the current directory or in the directory

/usr/vstree/SELECT

The above example is a very simple selection function. The distribution of the programs described in this manual comes with a subdirectory SELECT containing more well-documented selection functions for diverse tasks.

10  **vmatchselect**: Sorting and Selecting Matches

**vmatchselect** allows to select interesting matches from the output of **vmatch** as specified by user-defined criteria. It delivers matches of chosen length, degeneracy or significance into further analysis routines. **vmatchselect** removes from the input all those matches that are contained in another match. To do this efficiently, the matches are sorted by the position in the database sequence, and hence in the order in which the matches are output, unless the user specifies otherwise. Moreover, the sequences of the virtual suffix tree for which the match file was produced can be clustered according to the matches. The input for **vmatchselect** is a file produced by **vmatch**, called a *match file*.

**vmatchselect** is called as follows:

```
vmatchselect [options] matchfile
```

The output of **vmatchselect** goes to standard output and is sorted in ascending order of the positions of the left instance of a matches. Two matches where the left instance occurs at the same position, are sorted in descending order of their length. Two matches of the same length where the left instance occurs in the same position, are sorted in ascending order of the position of the right instance of the match.

**vmatchselect** provides a subset of the options of **vmatch**, namely the following options:

<table>
<thead>
<tr>
<th>Option</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>-l</td>
<td>-leastscore</td>
</tr>
<tr>
<td>-best</td>
<td>-value</td>
</tr>
<tr>
<td>-identity</td>
<td>-identity</td>
</tr>
<tr>
<td>-showdesc</td>
<td></td>
</tr>
<tr>
<td>-f</td>
<td>-nolead</td>
</tr>
<tr>
<td>-nodist</td>
<td>-nolead</td>
</tr>
<tr>
<td>-noidentity</td>
<td></td>
</tr>
<tr>
<td>-s</td>
<td>-score</td>
</tr>
<tr>
<td>-absolute</td>
<td></td>
</tr>
<tr>
<td>-v</td>
<td>-help</td>
</tr>
<tr>
<td>-nonredundant</td>
<td></td>
</tr>
</tbody>
</table>

The main difference to **vmatch** is that **vmatchselect** gets the matches from a *match file*, while **vmatch** computes the matches from scratch. Therefore options specifying the index and/or the query sequences to be matched, as well as options specifying how to match are not available in **vmatchselect**. The options of **vmatchselect** have the same meaning as in the program **vmatch**. Thus, for a description, see above.

Note that **vmatchselect** also allows to use the option `-dbcluster`. If **vmatchselect** is called with this option, then parses the given match file and performs single linkage clustering.
based on the matches in this file. Thus \texttt{vmatch} and \texttt{vmatchselect} allow hierarchical clustering. In a first step an initial set of matches with loose matching criteria is computed, using \texttt{vmatch}. Then one clusters these matches by calling \texttt{vmatchselect}. In a second round one applies more strict choices for the matches by using the options \texttt{-l}, \texttt{-leastscore}, \texttt{-evalue}, or \texttt{-identity}, etc. This allows stepwise refinement of clusters without much computational effort and no new index construction for the sequence of a cluster. The output of \texttt{vmatchselect} is the same as the output of \texttt{vmatch}.

11 Recent Changes

The following new features have recently been integrated into the software described in this manual.

\begin{itemize}
  \item \textbf{2003-May-08; mkvtree: } Option \texttt{-pl} can now be used without an argument. In this case, the value for \texttt{prefixlength} is determined automatically. If option \texttt{-pl} is used with an argument, then the behavior is as before. See page 9 for more details.
  \item \textbf{2003-May-15; vsubseqselect: } The previous restriction on the maximum size of the substring to be selected was removed. Otherwise, the behavior of the program did not change. See page 22 for more details.
  \item \textbf{2003-May-17; vmatch, vmatchselect: } Option \texttt{-iub} was removed. The effect of this option can be achieved by the argument \texttt{abbreviub} of option \texttt{-s}. See page 38 for more details.
  \item \textbf{2003-May-18; vmatch: } The default speedup-parameter for matching a query against an index is now 2. That is, the option \texttt{-qspeedup 2} is implicitly set. As a consequence, table \texttt{sitab1} is required for the matching, i.e. it has to be created. However, by using \texttt{-qspeedup 0}, the matching works as before with additional tables, but probably not as efficient as with option \texttt{-qspeedup 2}. See page 33 for more details.
  \item \textbf{2003-May-31; mkvtree: } \texttt{mkvtree} now scans the input files in GENBANK, EMBL or SWISSPROT format in such a way that additionally the \texttt{LOCUS}-information (in case of GENBANK) and the \texttt{ID}-information (in case of EMBL and SWISSPROT) is extracted and later reported as part of the description of the scanned sequence. See page 6 for more details.
  \item \textbf{2003-May-31; mkvtree: } \texttt{mkvtree} can now read gzipped database files. These are internally unzipped and then processed as any other input files. See page 6 for more details.
\end{itemize}
11 RECENT CHANGES

2003-May-31; vmatch: vmatch can now read gzipped query files. These are internally unzipped and then processed as any other input files. See page 6 for more details.

2003-Jun-4; vseqselect: vseqselect now gives an error message when the chosen minimum length is larger than the chosen maximum length. See page 20 for more details.

2003-Jun-4; vmatch, vmatchselect: The argument $e$ to option -evalue is now restricted to either be 0.0 or $\geq 1.0 \cdot 10^{-300}$. See page 36 for more details.

2003-Jun-15; vmatch, vmatchselect: The optional flags keepleft, keepright, and keepleftfifsamesequence of option -dbnomatch have now been complemented by the flag keeprightfifsamesequence. All four flags can now also be used for the option -dbmaskmatch. See page 29 for more details.

2003-Jun-28; vmatch, vmatchselect: The option -best has now been implemented with more efficient data structures. This gives a speedup if many matches are sought in order of their quality. See page 37 for more details.

2003-Jul-1; vsubseqselect: If option -range or option -seq is used, then the output shows the sequence description of the sequence, the substring was selected from. See page 22 for more details.

2003-Jul-2; vmatch, vmatchselect: Many new combinations of options are now possible. As a consequence, Table 6, contains considerably less entries of forbidden option combinations.

2003-Jul-12; vmatch, vmatchselect: Option -selfun can now have an arbitrary number of arguments, the first being the shared object containing the selection function bundle. All other arguments (not beginning with the symbol -) are ignored by vmatch and vmatchselect. See page 33 for more details.

2003-Jul-28; vstree2tex: Separator symbols are shown appropriately as different symbols. That is, the $i$th separator symbol is shown as number $i$. See page 18 for more details.

2003-Nov-13; all programs: Option -version added.
2004-Jun-18; vmatch, vmatchselect: Added new selection function bundle for reporting matches in CGviz-format. See the file SELECT/cgvizout.c. The selection function bundle was contributed by Katrin Wimmenauer. See page 81 for more details.

2004-Jun-25; vmatch: When using the options -dbmaskmatch or -qmaskmatch, it is reported how many characters are masked. Previously, this information was shown on stdout. Now it is shown on stderr. See page 30 for more details.

2004-Aug-20; vmatch: Option -online now works properly in all possible combinations with other options. We extensively verified that it produces the same result as when running vmatch without this option. See page 33 for more details.

2004-Aug-21; vmatch: Many combinations of options that were not possible before now work properly. This mainly concerns the options -i and -online. As a consequence there are only 102 combinations of options in vmatch excluded (see Table 6), compared to 112 in the previous program version (which had even less options).

2004-Sep-03; vmatch, vmatchselect: In the Sbjct- and Query-lines showing an alignment, a position shown on the right end is now displayed correctly. In particular, gaps are ignored when counting the positions. This leads to smaller positions and makes the output more compatible with other programs.

2004-Sep-18; vmatch, vmatchselect: Modified the interface to function selectmatch-Header. Besides the parameter used for computing matches, the function also gets access to the parameters used in the call of the program using option -selfun, see Section E. See page 79 for more details.

2004-Sep-18; vmatch, vmatchselect: Added an extra function selectmatchFinal-table to the selection function bundle. It allows to e.g. display matches stored in a user defined table. See page 33 for more details.

2004-Sep-19; vmatch, vmatchselect: Added new selection function bundle for merging overlapping matches. See the file SELECT/mergematches.c. See page 82 for more details.

2004-Oct-8; vmatch, vmatchselect: Changed the signature for the following functions of the selection function bundle: selectmatchInit, selectmatch, selectmatchWrap. These function are now called with three arguments:

- alphabet of the sequences to be compared,
• the Multiseq-structure representing the sequences in the index.
• the Multiseq-structure representing the sequences of the query.

See page 79 for more details.

2004-Oct-21; vmatch: When searching for repeats, option -l can now optionally be supplied with a second and third argument. Both arguments are integers specifying the minimum and maximum gap size of the repeat instances. See page 34 for more details.

2004-Dec-5; mkvtree: Given the indexname, mkvtree now removes all existing files comprising the index, before it generates the first index. This prevents from mixing index files from different runs of mkvtree. Furthermore, mkvtree now reports an error message, if an index file to be generated already exists.

2004-Dec-20; mkvtree: All symbol mappings for protein sequences now contain an additional wildcard symbol -, in addition to *, to represent stop codons. This feature was suggested by Volker Brendel.

2004-Dec-22; vmatch: The maximum number of errors allowed with option -h and -e has been extended from 10 to 25. See page 35 for more details

2004-Dec-22; vmatch: Computing complete matches with errors is now much more convenient: options -h and -e have been extended to also allow the percentage search mode and the best search mode. In the percentage search mode the user only specifies the percentage of errors relative to the length of the query to be searched. In the best search mode, the user specifies that matches for a given query sequence are reported only for the minimum number of errors. See page 35 for more details

2004-Dec-22; vmatch: Computing complete matches with errors is now also possible on the index, without using option -online. For many combinations of error thresholds and query lengths this leads to reduced running times at the cost of increased space requirement.

2004-Dec-29; vmatch: If the given index is for sequences over the protein alphabet and the given query sequences are over the DNA alphabet, then one can use option -dnasprot. This translates the DNA query sequences in all six reading frames. The six reading frames are matched against the protein database index, with all parameters allowed to control the kind of matches computed. The length of the matches and their positions are reported with respect to the original DNA sequence. An alignment for the matching substrings is shown on the protein level. See page 27 for more details
**2004-Dec-31; vmatch:** The recognition of symbol mappings for the DNA and the Protein alphabet has been improved. Now all symbol mappings in the subdirectory TRANS are correctly recognized. That is, all files in the directory TRANS with names containing the keyword DNA are recognized as DNA symbol mappings. All files in the directory TRANS with names containing the keyword Prot are recognized as Protein-symbol mappings. Of course, the recognition does not depend on the name of the symbol mapping files. See page 7 for more details.

**2005-Jan-18; vmatch:** When using the options -dbmaskmatch or -qmaskmatch, it is reported how many characters are masked. If the environment variable VMATCHCOMMENTTOSTDOUT is set to the value on, then this information is shown on stdout. Otherwise it is shown on stderr. See page 30 for more details.

**2005-Feb-1; vmatch:** Option -s now allows an optional argument xml. This produces XML-output.

**2005-Feb-21; vmatch:** Recall that in the vmatch-output, the line beginning with the symbol # args= shows the arguments of vmatch. The last argument of a vmatch-call is the indexname. By default, the indexname is shown with its absolute path. If the environment variable VMATCHRELATIVEINDEXPATH is set to the value on, then the path before the indexname is omitted.

**2005-Feb-25; vmatch:** The option -s abbreviub now works correctly for all mismatching characters which are not wildcards.

**2005-Feb-25; vmatch:** The option -mum now allows an optional flag cand to compute MUM-candidates.

**2005-Feb-28; vmatch:** Fixed bug in the codon translation on the reverse strand.

**2005-Feb-28; vmatch:** Added program mkdna6idx to compute a six frame translation index. vmatch can now read such an index to perform a self comparison of the indexed DNA database sequences on the protein level.

**2005-Feb-28; vmatch:** Added program chain2dim. This computes global and local optimal chains. Added corresponding option -pp chain to directly compute chains in vmatch.

**2005-Feb-28; vmatch:** Added program matchcluster. This clusters matches. Added corresponding option -pp matchcluster to directly compute compute clusters in vmatch.
2005-Mar-11; mkvtree, vmatch: Added perl script repfind.pl. This allows to emulate the program repfind from the REPuter software package.

2005-Mar-15; vmatch: In former versions of the program, the maximum number of errors allowed with option \(-h\) and \(-e\) was 10 and later 25. This fixed maximum error number is removed now. That is you can choose any number of errors. See page 35 for more details.

2005-Sep-13; vmatch: When combining option \(-hxdrop\) and \(-leastscore\), vmatch did not show a match with 0, no matter what score it had. This was a bug reported by Martin Frith, and is now fixed.

2006-Apr-05; vmatchselect: When using vmatchselect with option \(-s\) applied to matchfiles containing query matches, the program produced a segmentation fault. This bug is now removed.

2006-Apr-12; mkvtree, vmatch: The sequences file parser previously tolerated sequences containing no symbols. This caused problems in several cases. Now mkvtree and vmatch produce an error message if empty sequences are detected. They exit with exit code 1.

2007-Jun-2; vmatch: Martin Frith pointed out that option \(-supermax\) delivers repeats which are not always supermaximal. The problem is fixed now, i.e. the algorithm for computing supermaximal repeats now works correctly.

2007-Jun-2; vmatch: Martin Frith pointed out that option \(-i\) combined with option \(-supermax\) delivers incorrect results. This also holds if option \(-i\) is combined with option \(-mum\) and self comparison is performed. This problems is fixed and vmatch now works correctly for these options.

2007-Nov-2; vmatch: Volker Brendel pointed out that in some cases vmatch reports different alignments for the same protein to genome comparison, depending on whether the protein query is supplied singly or together with other targets. This problem is fixed now.

2008-Feb-16; vmatch: Anar Khan reported that Vmatch exits with an error message, when reverse complemented repeats are computed and constraints on the distance of the matches are specified. This problem is fixed now.
2010-Apr-26; vmatch: option -nonredundant now delivers the longest sequence from the corresponding cluster (instead of an unspecified representative). For every binary the distribution now contains an additional statically linked binary with the same name in the directory static/ (except for Mac OS X).

Acknowledgements

Sven Rahmann, Kirstin Weber, Volker Brendel, Alexander Sczyrba, Jomuna Choudhuri, Gordon Gremme, Marisano James, Martin Frith, Thomas Jahns, Peter Andrews, Wei Zhu, Anar Khan, and Jeremy Leipzig gave valuable hints on improving the manual and the programs described herein. Their help is much appreciated.

12 References


A Basic Notions

Throughout this section we consider a sequence $u$ of length $m$ over some alphabet $\Sigma$. The symbols are indexed from 0, i.e. $u = u_0u_1 \ldots u_{m-1}$.

If $\Sigma$ is the DNA alphabet $\{a, c, g, t\}$, then we define a function $wcc : \Sigma \rightarrow \Sigma$ by the following equations:

$$
wcc(a) = t \\
wcc(g) = c \\
wcc(c) = g \\
wcc(t) = a
$$

We extend $wcc$ to any sequence $u = u_0u_1 \ldots u_{m-1}$ over $\Sigma$ by defining

$$
wcc(u) = wcc(u_0)wcc(u_1) \ldots wcc(u_{m-1})
$$

Consider two sequences $u$ and $v$ of equal length. The Hamming distance of $u$ and $v$, denoted by $d_H(u,v)$, is the number of positions where $u$ and $v$ differ, that is,

$$
d_H(u,v) = |\{i \in [1, |u|] \mid u_i \neq v_i\}|
$$

There are three kinds of edit operations: deletions, insertions, and mismatches of single symbols. The edit distance or Levenshtein distance of $u$ and $v$, denoted by $d_E(u,v)$, is the minimum number of edit operations needed to transform $u$ into $v$.

Let $\approx$ be a binary relation on sequences, and assume that $u$ and $v$ are sequences of length $m$ and $n$. Let $l, r > 0$, $i \in [0, m-l]$, and $j \in [0, n-r]$. We define two different kinds of matches between $u$ and $v$:

- $(l, i, r, j)$ is a direct match if and only if
  
  $$
u_iu_{i+1} \ldots u_{i+l-1} \approx v_jv_{j+1} \ldots v_{j+r-1}
  $$(1)

  If $u = v$, then we additionally assume $i < j$.

- $(l, i, r, j)$ is a palindromic match if and only if
  
  $$
u_iu_{i+1} \ldots u_{i+l-1} \approx wcc(v_{j+r-1}v_{j+r-2} \ldots v_j)
  $$(2)

  If $u = v$, then we additionally assume $i \leq j$.

$(l, i, r, j)$ is a match, if it either is a direct or a palindromic match. Each match $(l, i, r, j)$ specifies two sequences:
1. the sequence \( u_i u_{i+1} \ldots u_{i+l-1} \) occurring on the left-hand side of (1) or (2). This is the left instance of \((l, i, r, j)\).

2. the sequence occurring on the right-hand side of (1) or (2). This is the right instance of \((l, i, r, j)\).

By specifying the relation \( \approx \), we obtain different types of matches:

1. If \( \approx \) is the identity on sequences, then a match \((l, i, r, j)\) is an exact match.

2. Let \( k > 0 \). If for any sequences \( x, y \) the relation \( \approx \) satisfies
   \[
   x \approx y \iff |u| = |v| \text{ and } d_H(x, y) \leq k
   \]
   then a match \((l, i, r, j)\) is a \( k \)-mismatch match.

3. Let \( k \in \mathbb{IN} \). If for any sequences \( x, y \) the relation \( \approx \) satisfies
   \[
   x \approx y \iff d_E(x, y) \leq k
   \]
   then a match \((l, i, r, j)\) is a \( k \)-differences match.

4. A match is degenerate if it is a \( k \)-mismatch or a \( k \)-differences match for some \( k \geq 0 \).

To distinguish between the different types of matches, we assign to them a distance value according to the following rules:

- An exact match has distance 0.
- A \( k \)-mismatch match with \( k > 0 \) mismatches has distance \(-k\).
- A \( k \)-differences match with \( k > 0 \) differences has distance \( k \).

A match \((l, i, r, j)\) is contained in a match \((l', i', r', j')\) if and only if \( i' \leq i \leq i+l-1 \leq i'+l' - 1 \) and \( j' \leq j \leq j+r-1 \leq j'+r'-1 \).

By requiring matches to be maximal we can reduce the number of interesting matches:

- An exact match is maximal if it is not contained in another exact match of the same kind.
- A \( k \)-mismatch match is maximal if it is not contained in another \( k \)-mismatch match of the same kind.
- A \( k \)-differences match is maximal if it is not contained in another \( k \)-differences match of the same kind.

A maximal unique match is a sequence \( w \) with the following properties:
\begin{itemize}
  \item $|w| \geq l$.
  \item $w$ occurs exactly once in $u$ and it occurs exactly once in $v$.
  \item For any character $a$ neither $wa$ nor $aw$ occurs both in $u$ and in $v$.
\end{itemize}

$w$ is represented by its length and its start positions in $u$ and in $v$.

**Example:** Let $u = gattctcgt$, $v = cctcgtgctg$, and $l = 2$. Then there are two maximal unique matches $ct$ and $tcg$, represented by $(2, 4, 1)$ and $(4, 6, 2)$.

When comparing a sequence against itself, we also call a match $(l, i, r, j)$ a repeat, if $i < j$. In other words, the two instances of a match do not occur at the same position. In this way, we do not report the trivial repeat where both instances of the match is the sequence itself.

Consider a repeat. If the left and the right instance of the repeat directly follow each other (without a gap or overlap), then the repeat is called tandem repeat. Suppose that the left instance of the tandem repeat starts at position $p$ and the repeat is of length $l$. Then the right instance starts at position $p + l$. Such a tandem repeat is branching if the characters at position $p + l$ and $p + 2l$ are distinct.

A supermaximal repeat is a maximal repeat that never occurs as a substring of any other maximal repeat.

## B Specification of a symbol mapping

`mkvtree` can be applied to sequences over any set of printable ASCII symbols. So, in particular it can be used for DNA, RNA, and for protein sequences. The user specifies in a symbol mapping the symbols the alphabet consists of (including wildcards). By grouping the symbols on different lines, the user also defines which symbols are considered to be identical, and how the symbols should be output when reporting a match. The symbol mapping implicitly defines an alphabet transformation $\phi$, which is applied to the input file. `mapsize` is the number of lines in the symbol mapping file. If a symbol mapping is given, then all symbols in the input files must occur in the alphabet. Otherwise, the program exits with error code 1.

The format of the symbol mapping file is best explained by some examples. Consider the file `TransDNA` containing the following lines:

```
aA
cC
gG
tTuU
nsywrkvdhmNSYWRKVDHM
```

These lines specify that the sequences in the input files are allowed to contain the symbols $a, c, g, t, u, n, s, y, w, r, k, v, b, d, h, m$ in either lower or upper case. Moreover, the first four lines
B SPECIFICATION OF A SYMBOL MAPPING

specify that, whenever characters are compared, the following equalities hold: $a = A$, $c = C$, $g = G$, and $t = T = u = U$. The last line specifies the wildcard symbols, which are replaced by unique symbols. As an effect, for each two (not necessary different) symbols $\alpha$ and $\beta$ appearing on the last line, we have $\alpha \neq \beta$ and $\alpha \neq a$, $\alpha \neq A$, $\ldots$, $\alpha \neq u$, $\alpha \neq U$.

The internal symbol mapping $\phi$ is as follows: the symbols on the first line are mapped to 0, the symbols of the second line to 1, etc. However, the symbols on the last line are mapped to the constant WILDCARD, which is a non-printable ASCII character. When matching symbols, each occurrence of WILDCARD is handled in such a way, that it does not match anywhere. Thus no false exact matches containing a wildcard will be delivered. However, a degenerate match may contain a WILDCARD, but this always leads to a pair of mismatching characters, or it must be deleted or inserted.

When matches are reported, the sequence output (if any) usually consists of the first symbol of each line, except if on that line there is a symbol following a white space. For example, if a match contains the symbol $\Lambda$ in the original input file, then this match is reported as $\Lambda$ in the output of vmatch. So if the user wants to output capital letters, he/she has to reverse the columns in the symbol mapping file.

Here is another example (see file TRANS/TransProt11), where different groups of amino acids are mapped to single symbols. We have also added comment lines. Each line at the beginning of the file starting with the symbol # is considered a comment line. It is ignored when parsing the symbol mapping file. Any number of comment lines is allowed, but only at the beginning of the file.

```
# 11-character alphabet as defined by Volker Brendel
LVIFJ i
KR +
ED -
AG s
ST o
NQ n
YW a
P p
H h
M m
C c
XUBZO* x
```

This specifies the protein alphabet $L, V, I, F, K, R, E, D, A, G, S, T, N, Q, Y, W, P, H, M, C$. Some extra wildcard symbols $X, B, Z, *, -$ are specified on the last line. All symbols occurring on the same line to the left of the first white space (if any) are considered to be equivalent. In each match, the symbol after the first white space will be shown for each symbol to its left. The symbols on the last line are considered to be wildcards. Again they are replaced by the symbol WILDCARD. The implicit mapping $\phi$ maps the symbol on line number $i \in [0, \text{mapsize} - 1]$ to integer $i$. In our case, $L, V, I, F$ are mapped to 0, $K, R$ are mapped to 1, etc.
C THE X-DROP EXTENSION STRATEGY

The X-Drop Extension Strategy

Consider the problem of aligning initial portions of two sequences, say $u$ and $v$ of length $m$ and $n$, respectively. Alignments are scored such that each mismatch has score $-1$, an indel (i.e. insertion or a deletion) has score $-2$, and each match has score $2$. The goal is to find a highest-scoring (i.e. optimal) alignment of some prefix of $u$ and some prefix of $v$, which are chosen to maximize the score. Here $m$ is the length of $u$ and $n$ is the length of $v$. There is a simple method to compute such an optimal alignment, but this method is inefficient, since it requires $O(mn)$ time. Zhang et. al. have proposed a method to prune the quadratic search space. This method, called greedy algorithm in the sequel, does not guarantee that the highest scoring alignment is found. However, in practice the alignments reported score very close to the optimal alignment. The greedy algorithm requires to specify a positive integer $X$ which influences the size of the evaluated search space. The smaller $X$, the smaller the evaluated search space. The greedy algorithm keeps track of the best alignment it has seen so far. Suppose this has score $T$. Whenever it extends an alignment, it checks whether the score of the extended alignment is smaller than $T - X$. If this is the case, than the extension is discarded. This effectively prunes the search space. The width of the search space expands in regions where the aligned sequence parts are different while it reduces in regions where the aligned sequences are similar. As a consequence, the greedy algorithm terminates if a similar part is followed by a region of many differences. The integer $X$ determines how much differences are tolerated.

If you use the program `vmatch` with the option `-exdrop`, then the Xdrop-alignment strategy is used when extending seeds (i.e. maximal exact matches) to the left and to the right. More precisely, if a maximal exact match $(l, i, r, j)$ of sequences $u$ and $v$ is found, then it is extended by applying the greedy algorithm twice:

- It is applied to the reverse of the two sequences $u_0u_1\ldots u_{i-1}$ and $v_0v_1\ldots v_{j-1}$
- It is applied to the two sequences $u_{i+l}u_{i+l+1}\ldots u_m$ and $v_{j+r}v_{j+r+1}\ldots v_n$

The two alignments reported by the greedy algorithm are concatenated with the seed to form an alignment of the substrings $u_{i'}\ldots u_{i''}$ and $v_{j'}\ldots v_{j''}$ where $i' \leq i$, $i + l - 1 \leq i''$, $j' \leq j$, and $j + r - 1 \leq j''$.

D Substring Specifications

There are applications where one wants to restrict the matching process to a substring of the query or to a substring of an database sequences. For example, if one already knows that the matches occur in a certain region of the database or the query, and one wants to allow less stringent matching conditions without spending time to search for matches in other regions. For these applications, `vmatch` allows two kinds of substring specifications:
E SELECTION FUNCTION BUNDLES

If only one query file with exactly one sequence, say $s$, is used as an argument to option –q, then the query file may optionally be followed by an extra argument of the form $(i, j)$. $i$ and $j$ are integers in the range $[0, |s| - 1]$ such that $i < j$ or $j = 0$. If $j = 0$, then this is interpreted as $|s| - 1$. That is, let $j' = j$ if $j > 0$ and $j' = |s| - 1$, otherwise. The pair $(i, j)$ specifies that the matching is to be restricted to the query substring $s_i \ldots s_j'$. Note that in some Unix-shells the symbols ( and ) have special meaning. Hence it may be necessary to quote the argument $(i, j)$, so that the shell does not evaluate it. For example, single quotes work for the csh, the tcsh, the sh, and the bash. The matches reported are as before, with the exception of the E-value. This is reported with respect to the shorter query substring, and thus becomes smaller. This feature does not work in connection with the option –online.

If the options –online and –q are used, then vmatch checks whether the descriptions of the query sequences begin with $>@ i j$ where $i$ and $j$ are numbers such that $0 \leq i$ and $i < j$ or $j = 0$. If this is the case for all descriptions of the query sequences, then the index is allowed to only contain one sequence, say $t$. If this is the case, then for each substring specification $>@ i j$ the matching is restricted to the substring $t_i \ldots t_j'$ where $j' = j$, if $j > 0$ and $j' = |t| - 1$, otherwise. The matches reported are as before, with the exception of the E-value. This is reported with respect to the shorter database substring, and thus becomes smaller. Note that the substring specifications are only taken into account if the option –online is used.

E Selection Function Bundles

Although vmatch allows many different ways to postprocess matches or to modify the output, the wide range of option combinations may not deliver what is required in a special application. The standard approach, used for example for BLAST, works as follows: (1) output the matches to a file, (2) parse this file with a user written program, and (3) postprocess the matches as required. However, this approach has several disadvantages:

- The user has to write a program to parse the output of the search engine. Even if the output format is designed s.t. it is easy to parse, this still requires extra effort.

- The size of the output can be huge for large scale matching problems. Due to slow I/O processing, output and parsing the matches can often take more time than computing them.

- It is often not only the values reported in the output of a typical search that is relevant for postprocessing, but also the set of sequences (with the sequence content and the corresponding sequence descriptions) subject to the matching task. Hence extra effort is required to parse the input sequences in such a way that the postprocessing program has access to the sequence information.
To circumvent these disadvantages, `vmatch` provides the concept of selection functions. These allow flexible and fast on-the-fly postprocessing of the matches computed by `vmatch`, without extra effort for output and parsing of the matches and sequence information.

Selection functions allow to access the matches and the sequences involved in the match immediately after they are computed. The selection is applied before further postprocessing or output is done, so that all other postprocessing options can be combined with selection functions.

Technically, a selection function is specified by a bundle of functions written in the programming language C. The program code implementing the selection functions is compiled independently from `vmatch`, but based on the data structures used in `vmatch`. Using the option `-selfun`, `vmatch` calls the selection functions via shared library calls (sometimes also called dynamic libraries). Since the definition of a selection function bundle requires a basic knowledge in C-programming, this concept is probably not suited for the average user. Nevertheless, we describe this concept carefully to give an idea of the flexibility it allows.

A selection function bundle consists of four functions with the following function headers:

```c
Sint selectmatchHeader(int argc,  
   char **argv,  
   int callargc,  
   char **callargv)

Sint selectmatchInit(Alphabet *alpha,  
   Multiseq *virtualmultiseq,  
   Multiseq *querymultiseq)

Sint selectmatch(Alphabet *alpha,  
   Multiseq *virtualmultiseq,  
   Multiseq *querymultiseq,  
   StoreMatch *storematch)

ArrayStoreMatch *selectmatchFinaltable(Alphabet *alpha,  
   Multiseq *virtualmultiseq,  
   Multiseq *querymultiseq);

Sint selectmatchWrap(Alphabet *alpha,  
   Multiseq *virtualmultiseq,  
   Multiseq *querymultiseq)
```

Sint is a type synonym for signed integers. The C-structures `Multiseq` and `StoreMatch` are defined in different header files. These are part of the distribution of the programs described in this manual.

Now suppose that these functions are specified in an extra file, and that they are compiled in the appropriate way (see below). Then the following holds:

- The function `selectmatchHeader` is called before the index and the query sequences (if any) are read. The first argument of the function is the number of arguments of the
corresponding call of \texttt{vmatch} and the second argument is the corresponding argument vector storing pointers

\begin{verbatim}
argv[0],...,argv[argc - 1]
\end{verbatim}

to the arguments of \texttt{vmatch}. In a call of \texttt{vmatch}, we have \texttt{callargc=argc} and \texttt{callargv=argv}. In a call of \texttt{vmatchselect}, \texttt{callargc} is the number of arguments of the program call to \texttt{vmatchselect}. \texttt{callargv} is the corresponding argument vector.

- The function \texttt{selectmatchInit} is called \textit{after} the index and the query files (if any) are read and \textit{before} the first match is processed. The indexed database sequences can be accessed via the pointer \texttt{virtualmultiseq}, while the query sequence is accessed via the pointer \texttt{querymultiseq}. The latter is \texttt{NULL} if the index is compared to itself. This function can e.g. be used to initialize internal data structures defined by the user.

- The function \texttt{selectmatch} is applied to each match referenced by \texttt{storematch}.
  
    - If \texttt{selectmatch} returns a value smaller than 0, then this is interpreted as an error, and \texttt{vmatch} or \texttt{vmatchselect} exits with error code 1.
    
    - If \texttt{selectmatch} returns 0, then this is interpreted as \texttt{false} and the match pointed to by \texttt{storematch} is not selected. That is, it is discarded.
    
    - If \texttt{selectmatch} returns 1, then this is interpreted as \texttt{true} and the match pointed to by \texttt{storematch} is selected. That is, it is output or postprocessed by \texttt{vmatch} according to the options given.

- The function \texttt{selectmatchFinaltable} is called \textit{after} the last match has been processed, but \textit{before} \texttt{selectmatchWrap} is called. If specified by the user, it must return a pointer to a memory area which is available after the call to this function. The memory area stores an array of structures of type \texttt{StoreMatch}. See the selection function bundle in \texttt{mergematches.c} for an example. \texttt{vmatch} or \texttt{vmatchselect} accesses the memory area and processes it according to the options given by the user. In the standard case, the merged matches are shown on standard out.

- The function \texttt{selectmatchWrap} is called \textit{after} the last match has been processed. It can, e.g., be used to output internal data structures which store accumulated information about the matches found.

A selection function bundle must define at least the function \texttt{selectmatch}. The other functions can be omitted in the C-file declaring the selection function bundle.

The distribution of the programs described in this manual comes with a subdirectory \texttt{SELECT} containing well-documented selection functions for diverse tasks:

\begin{verbatim}
cgvizout.c  Output matches in a format readable for the visualization program CGviz, see \cite{4}.
\end{verbatim}
\textbf{F THE TABLES COMPRISING THE INDEX}

\texttt{dbseqstat.c} counts the number of matches to different query sequences and outputs all database sequences containing at least two matches to different query sequences.

\texttt{endmatch.c} selects all matches which start with the first character of a database sequence or which end with the last character of a database sequence.

\texttt{lowcomplex.c} discards all matches where the matching sequences are of low complexity, according to a simple model based on the distribution of characters.

\texttt{mstat.c} groups matches according to their occurrence in the database sequences.

\texttt{mergematches.c} merge two matches if they overlap by a significant number of positions. This number is specified as some percentage of the length of the matches.

\texttt{rightmost.c} selects the rightmost matches in all database sequences and discards those matches which are not rightmost.

\texttt{sel392.c} selects matches which are not longer than some maximum value (see above). The latter can be defined at run time by the user.

\texttt{selsplicesite.c} selects matches where the right instance of the match (in the query) does not have AG to the right and TC to the left.

\texttt{selstartend.c} select matches where the left instance begins with the bases AC and ends with GT.

\texttt{xmlout.c} shows the numeric information of the matches (not the alignment or other sequence information) in XML-format.

Besides these files, subdirectory SELECT also contains a \texttt{makefile}, specifying how to compile a shared object for the supported platforms.

\section{The Tables Comprising the Index}

Suppose the input files contain the sequences $T_1, \ldots, T_k$ in this order. Let \# be a symbol not occurring in any $T_i$ and suppose $S = T_1 \# T_2 \# \ldots T_{k-1} \# T_k$. That is, \# is used as a separator symbol. Let $n = |S| = k - 1 + \sum_{i=1}^{k} |T_i|$. The index consists of the tables \texttt{prjtab}, \texttt{tistab}, \texttt{oistab}, \texttt{suftab}, \texttt{bwttab}, \texttt{lcptab}, \texttt{llvtab}, \texttt{skptab}, \texttt{bcktab}, \texttt{ssptab}, \texttt{destab}, \texttt{sdstab}.

These are defined as follows:

- \texttt{prjtab} contains the ”’project information’’”, i.e. some basic information about the index in human readable form. Here is an example:
<table>
<thead>
<tr>
<th>dbfile</th>
<th>atEST 999815 772376</th>
</tr>
</thead>
<tbody>
<tr>
<td>totallength</td>
<td>772376</td>
</tr>
<tr>
<td>numofsequences</td>
<td>1952</td>
</tr>
<tr>
<td>numofdbsequences</td>
<td>1952</td>
</tr>
<tr>
<td>numofquerysequences</td>
<td>0</td>
</tr>
<tr>
<td>prefixlength</td>
<td>7</td>
</tr>
<tr>
<td>largelcpvalues</td>
<td>2317</td>
</tr>
<tr>
<td>maxbranchdepth</td>
<td>517</td>
</tr>
<tr>
<td>integersize</td>
<td>32</td>
</tr>
</tbody>
</table>

This information shall be read as follows: There is only one database file atEST of length 999815. The total length of the sequences in this file plus the separator symbols between the sequences is 772376. The total length of all database sequences and their number is given in the next two lines. The construction of the index was done using an initial bucket sorting step with prefixlength set to 7. The number of lcp-values (see below) larger or equal to 255 is 2317, and the largest lcp-value is 517. The last line tells that the index has been constructed by a version of mkvtree using 32-bit integer.

- **tistab** is the transformed input sequence (thus the abbreviation *tis*). In particular it is a table of length \( n \) such that, for any \( i \in [0,n-1] \),

\[
tistab[i] = \begin{cases} 
\text{SEPARATOR} & \text{if } S_i = \# \\
\phi(S_i) & \text{otherwise}
\end{cases}
\]

Here SEPARATOR is a non-printable ASCII-character different from WILDCARD. Each entry in tistab is stored in 1 byte.

- **oistab** is the original input sequence after parsing (thus the abbreviation *ois*). In particular it is a table of length \( n \) such that, for any \( i \in [0,n-1] \),

\[
oistab[i] = \begin{cases} 
\text{SEPARATOR} & \text{if } S_i = \# \\
S_i & \text{otherwise}
\end{cases}
\]

Each entry in oistab is stored in 1 byte.

- **suftab** is a table of length \( n + 1 \). To define it, we need some preliminaries: tistab can be interpreted as a sequence, say \( T \), of length \( n \). Let \( \$ \) be a new symbol greater than any symbol in \( T \). Consider the string \( T\$ \) and let \( X_i = T_i \ldots T_{n-1}\$ \) for any \( i \in [0,n] \). That is, \( X_i \) is the suffix of \( T\$ \) starting at position \( i \). Suppose \( X_{j_0}, X_{j_1}, \ldots, X_{j_n} \) is the sequence of suffixes in ascending lexicographic order. Then table suftab of length \( n + 1 \) is defined by \( suftab[i] = j_i \) for any \( i \in [0,n] \). If \( n < 2^{32} \), then each entry in suftab is stored in 4 bytes. Otherwise, it is stored in 8 bytes.

- **stitab1** is the reduced version of the inverse suffix table. It is a table of length \( n + 1 \). To define it, we have to introduce table stitab. This is the inverse of table suftab, i.e. it
satisfies the equality $suftab[stitab[i]] = i$ for all $i \in [0, n]$. Now suppose that $stitab[i] \in [bcktab[w], bcktab[w] − 1]$ for some string $w$ of length $prefixlength$. Then

$$stitabI[i] = \begin{cases} 255 & \text{if } stitab[i] − bcktab[w] \geq 255 \\ stitab[i] − bcktab[w] & \text{otherwise} \end{cases}$$

Each entry in $stitabI$ is stored in 1 byte.

- $bwttab$ is a table of length $n + 1$ such that, for any $i \in [0, n]$,

$$bwttab[i] = \begin{cases} \text{undefined} & \text{if } suftab[i] = 0 \\ tistab[suftab[i] − 1] & \text{otherwise} \end{cases}$$

Each value in $bwttab$ is stored in 1 byte. $bwttab$ is the Burrows and Wheeler transformation.

- $lcptab$ is a table of length $n + 1$. Suppose that for any $i \in [1, n]$, $lcp(i)$ is the length of the longest common prefix of the suffix $T_{suftab[i − 1]}$ and $T_{suftab[i]}$. If $lcp(i) < 255$, then $lcptab[i] = lcp(i)$. Otherwise, $lcptab[i] = 255$ and in table $llvtab$ we store the pair $(i, lcptab[i])$. In other words, $llvtab$ stores the large lcp values. The pairs in table $llvtab$ are ordered according to their left component. $lcptab[0]$ is undefined. Each entry in $lcptab$ is stored in one byte. If $n < 2^{32}$, then each entry in $llvtab$ is stored in 8 bytes. Otherwise, it is stored in 16 bytes.

- $skptab$ is a table of length $n + 1$ such that, for any $i \in [0, n]$,

$$skptab[i] = \min(\{n\} \cup \{j \in [i + 1, n] \mid lcp(i) > lcp(j)\})$$

The abbreviation $skp$ stands for skip-value. If $n < 2^{32}$, then each entry in $skptab$ is stored in four bytes. Otherwise it is stored in 8 bytes.

- For some value $prefixlen > 0$ as specified in $prjtab$, table $bcktab$ stores the bucket boundaries. These are defined as follows: Let $r = mapsize − 1$, $l = prefixlen$, and $\Sigma = [0, r − 1]$. For any sequence $w \in \Sigma^l$, define $\kappa(w) = \Sigma_{i=0}^{l−1} r^{l−1−i} w_i$. $\kappa : \Sigma^l \to [0, r^l − 1]$ is a bijective function, mapping each $w \in \Sigma^l$ to the corresponding integer to base $r$. For each $w \in \Sigma^l$, $bcktab[\kappa(w)]$ stores a pair of integers $(p, q + 1)$ such that:

- $T_{suftab[p]}$ is the lexicographically smallest suffix of $T$ which has $w$ as a prefix
- $T_{suftab[q]}$ is the lexicographically largest suffix of $T$ which has $w$ as a prefix

If $w$ does not occur as a substring of $T$, then $p = q + 1$. $bcktab$ is of size $r^l$. If $n < 2^{32}$, then each entry in $bcktab$ is stored in 8 bytes. Otherwise, it is stored in 16 bytes.

- $ssptab$ is a table of length $k − 1$ storing, in sorted ordered, the positions in $S$ where the separator symbol # occurs. If $n < 2^{32}$, then each entry is stored in four bytes. Otherwise, it is stored in 8 bytes. The abbreviation $ssp$ stands for sequence separator positions.

- $destab$ stores the concatenated descriptions from the input files including a separator \n.
Table 7: Overview of the tables comprising the index. \( n \) is the total length of all database sequences including the \( k - 1 \) separator symbols. \( k \) is the number of database sequences. \( r \) is \( \text{mapsize} - 1 \) and \( \text{mapsize} \) is the number of lines in the symbol mapping file. The size of each entry is given in bytes, assuming that \( n < 2^{32} \). If \( n \leq 2^{32} \), then the size of each entry in table \( \text{suftab}, \text{llvtab}, \text{skptab}, \text{bcktab}, \) and \( \text{ssptab} \) is twice as large. If any of the options \(-\text{smap}, -\text{dna}, \) or \(-\text{protein} \) was used for \text{mkvtree}, then there exists a corresponding table \( \text{al1tab} \) storing the symbol mapping. Otherwise, all symbols contained in the file are stored in \( \text{al2tab} \).

<table>
<thead>
<tr>
<th>table</th>
<th>description</th>
<th>length</th>
<th>size per entry</th>
</tr>
</thead>
<tbody>
<tr>
<td>prjtab</td>
<td>project information</td>
<td>a few lines</td>
<td></td>
</tr>
<tr>
<td>tistab</td>
<td>transformed input sequences</td>
<td>( n )</td>
<td>1</td>
</tr>
<tr>
<td>oistab</td>
<td>original input sequences</td>
<td>( n )</td>
<td>1</td>
</tr>
<tr>
<td>suftab</td>
<td>suffix array</td>
<td>( n + 1 )</td>
<td>4</td>
</tr>
<tr>
<td>stitab1</td>
<td>reduced inverse suffix array</td>
<td>( n + 1 )</td>
<td>1</td>
</tr>
<tr>
<td>bwttab</td>
<td>Burrows Wheeler Transformation</td>
<td>( n + 1 )</td>
<td>1</td>
</tr>
<tr>
<td>lcptab</td>
<td>longest common prefixes (&lt;\ 255)</td>
<td>( n + 1 )</td>
<td>1</td>
</tr>
<tr>
<td>llvtab</td>
<td>longest common prefixes (\geq\ 255)</td>
<td>(</td>
<td>{i \in [1,n] \mid lcp(i) \geq 255}</td>
</tr>
<tr>
<td>skptab</td>
<td>skip values</td>
<td>( n + 1 )</td>
<td>4</td>
</tr>
<tr>
<td>bcktab</td>
<td>bucket boundaries</td>
<td>( r )</td>
<td>8</td>
</tr>
<tr>
<td>ssptab</td>
<td>sequence separator positions</td>
<td>( k - 1 )</td>
<td>4</td>
</tr>
<tr>
<td>destab</td>
<td>descriptions for sequences</td>
<td>sum of description lengths</td>
<td></td>
</tr>
<tr>
<td>sdstab</td>
<td>start pos. descriptions</td>
<td>( k + 1 )</td>
<td>4</td>
</tr>
<tr>
<td>al1tab</td>
<td>symbol mapping used</td>
<td>a few lines</td>
<td></td>
</tr>
<tr>
<td>al2tab</td>
<td>enumeration of symbols</td>
<td>( \leq 255 )</td>
<td></td>
</tr>
</tbody>
</table>

- \( \text{sdstab} \) is a table of length \( k + 1 \) storing the starting positions of the descriptions of the sequences. That is, for any \( i \in [0,k - 1] \), the description of sequence \( i \) is stored from position \( \text{sdstab}[i] \) to \( \text{sdstab}[i + 1] - 1 \). If \( n < 2^{32} \), then each entry is stored in four bytes. Otherwise, it is stored in 8 bytes.

Table 7 gives an overview of the different tables.

To satisfy the different matching tasks, different tables are required, according to the following rules:

**Self comparison:** If only the number of exact substring matches is to be reported, then \( \text{bwttab} \) and \( \text{lcptab} \). Otherwise, \( \text{lcptab}, \text{suftab}, \) and additionally:

- \( \text{bwttab} \), if direct matches are to be reported.
- \( \text{tistab} \), if degenerate matches or palindromic matches or the string content of a match is to be reported.
**•** `bcktab`, if palindromic matches are to be reported.

**query comparison:** If the matching task is to be performed online, then `tistab`. Otherwise, `lcptab`, `tistab`, `suftab`, and `bcktab`.

The following additional rules apply. In all cases, `destab`, `sdstab`, and `prjtab` are required. If there is more than one sequence in the index, then also `ssptab`. If `lcptab` is required, then also `llvtab`. Table `skptab` is currently not required for any application implemented in `vmatch`.

### G Environment Variables

There are some environment variable which allow to influence the behavior of the programs described here:

- **MKVTTREESMAPDIR**: is a colon separated list of directories to search for symbol map files.

- **VMATCHCOMMENTTOSTDOUT**: if set to the value `on`, then the line showing how much symbols are masked (for option `-dbmaskmatch` or `-qmaskmatch`) is shown on stdout. If set to `off`, then the line is shown on stderr.

- **VMATCHRELATIVEINDEXPATH**: The last argument of a `vmatch`-call is the indexname. By default, the indexname is shown with its absolute path. If this environment is set to the value `on`, then the path before the indexname is omitted.

- **LD_LIBRARY_PATH**: colon separated list of directories, where to find the shared object files used as arguments for the option `-selfun`. 
## Index

<table>
<thead>
<tr>
<th>t</th>
<th>17, 27</th>
</tr>
</thead>
<tbody>
<tr>
<td>.fna</td>
<td>31, 59</td>
</tr>
<tr>
<td>.match</td>
<td>31, 59</td>
</tr>
<tr>
<td>.single</td>
<td>31</td>
</tr>
<tr>
<td>.sl</td>
<td>33</td>
</tr>
<tr>
<td>.so</td>
<td>33</td>
</tr>
<tr>
<td>cand</td>
<td>29</td>
</tr>
<tr>
<td>chain</td>
<td>32</td>
</tr>
<tr>
<td>matchcluster</td>
<td>32</td>
</tr>
<tr>
<td>remred</td>
<td>29</td>
</tr>
</tbody>
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