

TMAP: the flow sequence mapping program ¹

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Contents

Table of Contents	iv
Preface	v
1 Basic Usage	1
1.1 Program Organization	1
1.2 Common Mapping Options	1
1.2.1 Global Options	2
1.2.2 Flowspace Options	5
1.2.3 Pairing Options	6
1.3 tmap index	7
1.3.1 Usage	8
1.3.2 IUPAC Ambiguity Codes	8
1.4 tmap map1	9
1.4.1 Usage	9
1.5 tmap map2	10
1.5.1 Usage	11
1.6 tmap map3	11
1.6.1 Usage	11
1.7 tmap map4	12
1.7.1 Usage	12
1.8 tmap mapvsw	13
1.8.1 Usage	13
1.9 tmap mapall	13
1.9.1 Overview	14
1.9.2 Usage	14
1.10 tmap server	15
1.10.1 Usage	15

2	File Formats	17
2.1	SAM Alignment Format	17
2.1.1	SAM Header Fields	17
2.1.2	SAM Record Optional Tags	18
3	Commonly Asked Questions	21
3.1	How are IUPAC ambiguity codes handled in the reference/target FASTA? .	21
3.2	How are insertions and deletions scored?	21
3.3	How are insertions and deletions justified?	21
3.4	Will the output of TMAP be the same if run twice?	22
3.5	How does TMAP soft clip bases in the read?	23
3.6	What is flow space re-alignment	24
4	Appendix	27
	References	27

Preface

This document is meant to serve as a guide for the practical use of TMAP. It includes explanations of all command-line options for each command and binary in TMAP to give an idea of basic usage. Input and output file formats are also detailed. The default options of the various programs are designed intelligently to adapt to the various program uses, but in some cases the options may need to be customized.

This document does not try to explain the underlying algorithms or data-structures used in TMAP. Without proper understanding of the underlying algorithms, it is difficult to use this very flexible program knowledgeably to obtain your desired results. Please see [chapter 3](#) for common questions and their answers.

If you have anything that you would be useful to add to this guide, feel free to relay the addition to the TMAP developers. This includes but is not limited to bugs, typos, and explanations. Please see <http://ioncommunity.iontorrent.com> for more details.

Enjoy!

Chapter 1

Basic Usage

1.1 Program Organization

TMAP consists of a set of utilities, combined into one command line program called `tmap`. Each utility is specified by a unique name or command. Specifying TMAP without a command will give a list of commands, while specifying TMAP with a command but no options will give a list of options for that command.

There are two steps in mapping with TMAP. The first step is to build an index of the reference genome onto which we map. This index needs only to be built once for each genome, performing much of the mapping work upfront. The second step is to map the reads to the reference genome using this index.

TMAP is implemented as a command-line program. It accepts many command-line options to customize and tune the mapping algorithm. The key commands are organized into one binary program called `tmap`. To access each command, we use `tmap <name>`, where `<name>` is the name of the command we wish to execute.

1.2 Common Mapping Options

Some common options exist across some or all of the mapping commands (`map1`, `map2`, `map3`, `map4`, `mapvsw` and `mapall`). These options will be discussed here to avoid duplication.

The TMAP mapping commands can accept their input file from the standard input stream. The output of these commands also write to the standard output stream. This facilitates the use of these TMAP commands in a pipe-and-filter model.

1.2.1 Global Options

`-f, --fn-fasta FILE`

Specifies the file name of the reference genome in FASTA format. The maximum number of bases in the reference genome can be 4294967295, due to the use of 32-bit values to represent each position in the reference.

`-r, --fn-reads FILE`

Specifies the file name of the reads to mapped. The input reads can be in FASTA, FASTQ, or SFF format. If an SFF is input, only reads that have a matching key sequence will be mapped.

`-i, --reads-format STRING`

Specifies the file format of the reads file. Without this option, the format is auto-recognized by the file extension. The valid inputs are `fa` or `fasta` for FASTA, `fq` or `fastq` for FASTQ, or `sff` for SFF. If compiled with SAMTools support, the option also supports `sam` for SAM, or `bam` for BAM.

`-s, --fn-sam FILE`

Specifies the file name of the output file SAM format. If this is not specified, the output will be written to stdout.

`-A, --score-match INT`

Specifies the match score. This number must always be positive.

`-M, --pen-mismatch INT`

Specifies the mismatch penalty. This number must always be positive.

`-O, --pen-gap-open INT`

Specifies the gap open penalty. This number must always be positive.

See [section 3.2](#) for more information how insertions and deletions are scored.

`-E, --pen-gap-extension INT`

Specifies the gap extension penalty. This number must always be positive. See [section 3.2](#) for more information how insertions and deletions are scored.

`-w, --band,width INT`

Specifies the band width for local alignment. This number must always be positive.

`-g, --softclip-type INT`

Specifies that the type of soft-clipping to perform.

0. soft-clip both the left and right portions of the read.
1. soft-clip only the left portion of the read.
2. soft-clip only the right portion of the read.
3. do not soft-clip any portion of the read

`-W, --duplicate-window INT`

Specifies to remove duplicate mappings that occur within this bp window.

`-B, --max-seed-band INT`

Specifies the window in bases in which to group seeds.

`-U, --no-unroll-banding`

Specifies to not unroll the grouped seeds from banding if multiple alignments are found

`-T, --score-thres INT`

Specifies the number of multiples of the match score (`-A`) for the minimum scoring threshold.

`-q, --reads-queue-size INT`

Specifies the number of reads to cache or load into memory at one time.

`-n, --num-threads INT`

Specifies the number of threads to run.

`-a, --aln-output-mode INT`

Specifies the output filter for the mappings.

0. returns the mapping with the best score only if all other mappings had worse score, otherwise the read is unmapped.
1. returns the mapping with the best score. If more than one mapping has this score, a random mapping with this score is returned.
2. returns all the mappings with the best score.
3. returns all the mappings, regardless of score.

Reads that have no mapping are returned as unmapped reads.

`-R, --sam-read-group STRING`

Specifies the RG (read group) line to use in the SAM file (with tab separators). The “PG”, “FO”, and “KS” tags should not be specified in this string; they will be added by tmap. Alternatively, multiple `-R` options can also be used to populate the RG line. For example, `-R CN:SEQUENCING_CENTER -R PG:TMAP -R PL:IONTORRENT` will populate the CN, PG, and PL tags in the RG record.

`-j, --input-bz2, -z, --input-gz`

Specifies that the input is bzip2 (`-j`) or gzip (`-z`) compressed. This is auto-recognized if the input file name has the extension `.bz2` for bzip2 and `.gz` for gzip.

`-J, --output-bz2, -Z, --output-gz`

Specifies that the output should be bzip2 (`-J`) or gzip (`-Z`) compressed.

`-k, --shared-memory-key INT`

Specifies the shared memory key if the reference index has been loaded into shared memory.

`-v, --verbose`

Specifies to print verbose progress messages, otherwise progress messages will be suppressed.

`-h, --help`

Specifies to print a help message, listing all available options.

`--min-seq-length INT`

Specifies the minimum sequence length to consider (inclusive). This is applied to each algorithm independently. Therefore, when using `mapall`, we could specify a unique sequence length range (using `---min-seq-length` and `---max-seq-length`) each algorithm.

`--max-seq-length INT`

Specifies the maximum sequence length to consider (inclusive). This is applied to each algorithm independently. Therefore, when using `mapall`, we could specify a unique sequence length range (using `---min-seq-length` and `---max-seq-length`) each algorithm.

1.2.2 Flowspace Options

These options control how TMAP uses and outputs information about the flow order from the sequencing machine. Flow space re-alignment will be performed when the flow order is specified (`-F, --flow-order`). See [section 3.6](#) for a description of flow space re-alignment.

`-X, --pen-flow-error INT`

Specifies the flow score penalty. This number must always be positive.

`-F, --flow-order STRING`

Specifies the flow order ([ACGT]4+ or “file” to obtain it from the input file.

`-K, --key-sequence STRING`

Specifies the key sequence ([ACGT]4+ or “file” to obtain it from the input file.

`-y, --softclip-key`

Specifies to soft clip only the last base of the key sequence.

`-Y, --sam-sff-tags`

Specifies that SFF specific tags should be added to the output SAM file. See [section 2.1](#) for more informaton.

`-N, --ignore-flowgram`

Specifies to not use the flowgram, otherwise uses the flowgram for flowspace re-alignment when available.

`-G, --remove-sff-clipping`

Specifies to not remove bases and qualities based on the adapter and quality clipping fields found in the an input SFF file.

1.2.3 Pairing Options

These options control how TMAP uses and outputs information about the paired end or mate pair reads. Pairing will be performed when exactly two input read files are given using the `-r` option twice.

`-Q, --pairing`

Specifies the pairing orientation. This will override `-S` and `-P`.

- 0. no pairing is to be performed
- 1. 1 - mate pairs (`-S 0 -P 1`)
- 2. 2 - paired end (`-S 1 -P 0`)

`-S, --strandedness`

Specifies the strand orientation between the pairs.

- 0. the reads should be mapped to the same strand
- 1. the reads should be mapped to opposite strands

`-P, --positioning`

Specifies that position orientation between the pairs.

0. the first read of the pair (A) is before the second read of the pair (B).
1. the second read of the pair (B) is before the first read of the pair (A).

`-b, --ins-size-mean FLOAT`

Specifies the expected mean insert size.

`-c, --ins-size-std FLOAT`

Specifies the expected insert size standard deviations.

`-d, --ins-size-std-max-num FLOAT`

Specifies the maximum number of standard deviations for a pair to be proper. See [section 2.1](#) for more informaton about proper pairs.

`-L, --read-rescue`

Specifies to perform read rescuing during pairing.

`-l, --read-rescue-std-num FLOAT`

Specifies the number of insert size standard deviations to search around the inferred rescue position.

1.3 tmap index

The **index** command creates a compact version of the reference genome and associated index. The index is stored as a compressed suffix array using the FM-index and BWT transform ([Ferragina and Manzini \(2000\)](#); [Burrows and Wheeler \(1994\)](#)). A hash into this index accelerates this lookups of DNA sequences in this index. In fact, a second additional index of the reference genome is created that indexes the reverse (but not complimented) reference genome. This second index further speeds up the search time.

See [section 1.2](#) for common options that are in use in this command.

1.3.1 Usage

`-o INT`

Specifies the occurrence interval size o , storing only every o th occurrence interval. This must be one, or a multiple of 16.

`-w INT`

Specifies the k-mer size (the number of bases) to hash. The size of the hash give the k-mer size k in bytes is:

$$= 2 \sum_{n=1}^k 4^n = 2 \left(\frac{1 - 4^{k+1}}{1 - 4} - 1 \right)$$

using the Taylor series.

`-i INT`

Specifies the suffix array interval size i , storing only every i th suffix array interval. This must be one, or a multiple of two.

`-a STRING`

Specifies the BWT construction algorithm (`bwtsw` or `is`). The `bwtsw` algorithm is for genomes larger than or equal to 10Mb, and the `is` algorithm is for genomes smaller than 10Mb. This will be auto-recognized during index creation if this option is omitted.

`-H`

Specifies to not validate the BWT hash.

`--version`

Specifies to print the index format that will be created by TMAP and exit. Format strings are of the form `tmap-f<n>`, where `<n>` will only increase as new formats are created. Format strings will be the same for all compatible indices.

1.3.2 IUPAC Ambiguity Codes

Please see [section 3.1](#) for more details.

1.4 tmap map1

The **map1** is a command to quickly map short sequences to a reference genome by intelligently enumerating errors. This algorithm is not well suited for longer reads (< 150bp) and based off of the BWA short-read algorithm ([Li and Durbin \(2009\)](#)).

See [section 1.2](#) for common options that are in use in this command.

1.4.1 Usage

`--seed-length INT`

Specifies the primary seed (k-mer) length; reads must be of at least this length.

`--seed-max-diff INT`

Specifies the maximum number of edits allowed in the primary seed. This includes both mismatches and each base in an insertion or deletion.

`--seed2-length INT`

Specifies the secondary seed (k-mer) length; this number of bases will be examined before extending all mappings.

`--max-diff NUM`

Specified the maximum number of edits or false-negative probability assuming the maximum error rate (`--max-error-rate`). This former includes both mismatches and each base in an insertion or deletion. The latter false-negative probability is the probability of not considering the correct mapping if it exists.

`--max-error-rate NUM`

Specified the assumed maximum per-base error rate. This classifies both mismatches and each base in an insertion or deletion as errors

`--max-mismatches NUM`

Specifies the maximum number of mismatches allowed, or the fraction of mismatches with respect to the read length. When the secondary seed is enabled, this parameter is relative to the secondary seed length, not the read length;

`--max-gap-opens NUM`

Specifies the maximum number of indels allowed in the mapping, or the fraction of indels with respect to the read length. The number of indels is equal to the number of gap opens. When the secondary seed is enabled, this parameter is relative to the secondary seed length, not the read length;

`--max-gap-extensions NUM`

Specifies the maximum number of indel extensions allowed in the mapping, or the fraction of indel extensions with respect to the read length. The number of indel extensions is equal to the number of gap extensions. When the secondary seed is enabled, this parameter is relative to the secondary seed length, not the read length;

`--max-cals-deletion INT`

Specifies the maximum number of candidate alignment locations (CALs) to allow a deletion to be extended.

`--indel-ends-bound INT`

Specifies to disallow indels within this number of bases from the ends of the read.

`--max-best-cals INT`

Specifies to stop searching for mappings when this many best scoring mappings have been found.

`--max-nodes INT`

Specifies the maximum number of alignment nodes in memory in the implicate prefix trie traversal before the search is stopped.

1.5 tmap map2

The **map2** is a command to quickly map long sequences to a reference genome. This algorithm is well suited for longer reads (≥ 150 bp) and based off of the BWA long-read algorithm (Li and Durbin (2010)).

See [section 1.2](#) for common options that are in use in this command.

1.5.1 Usage

`--length-coef` FLOAT

Specifies the coefficient to adjust the mapping score threshold based on the read length. Given a l -long read, the threshold for a mapping to be retained is $a * \max\{T, c * \log(l)\}$, where a is the match score ($-A$) and T is the minimum score threshold ($-T$).

`--max-seed-intv` INT

Specifies the maximum seed interval size (mappings) to retain at during extension of the seed.

`--z-best` INT

Specifies the maximum number of top-scoring nodes to keep on each iteration.

`--seeds-rev` INT

Specifies the maximum seeding interval size for extending a mapping.

`--narrow-rmdup`

Specifies to remove duplicate seeds for narrow suffix array.

1.6 tmap map3

The **map3** is a command to map sequences to a reference genome. This algorithm is well suited for longer reads (≥ 150 bp) and a simplification of the SSAHA long-read algorithm (Ning *et al.* (2001)).

See [section 1.2](#) for common options that are in use in this command.

1.6.1 Usage

`--hit-frac` FLOAT

Specifies the fraction of seed positions that are under the maximum (`--max-seed-hits`).

`--seed-step` INT

Specifies the number of bases to increase the seed while repetitive (-1 to disable).

`--seed-length INT`

Specifies the k -mer length to seed candidate alignment locations (CALs). With a value of `-1` will set the seed length to $(\text{ceiling}(\log_4(R)) + 2)$ where R is the reference genome length.

`--max-seed-hits INT`

Specifies the maximum number of CALs allowed to be returned by a seed before it is ignored.

`--hp-diff INT`

Specifies the number of bases to enumerate for a single homopolymer within the seed.

`--fwd-search`

Specifies to perform a forward search instead of a reverse search.

`--skip-seed-frac FLOAT`

Specifies the fraction of a seed to skip when a lookup succeeds.

1.7 tmap map4

The **map4** is a command to map sequences to a reference genome. This algorithm is well suited for longer reads ($\geq 100\text{bp}$) and is a variation of the super-maximal exact matching (SMEM) algorithm proposed by Heng Li. SMEMs are equivalent to returning all longest common substrings between the read and reference.

See [section 1.2](#) for common options that are in use in this command.

1.7.1 Usage

`--hit-frac FLOAT`

Specifies the fraction of seed positions that are under the maximum (`--max-seed-hits`).

`--max-iwidth INT`

Specifies the maximum interval size to accept a hit.

`--min-seed-length INT`

Specifies the minimum number of bases to use to seed. With a value of -1 will set the seed length to $(\text{ceiling}(\log_4(R)) + 2)$ where R is the reference genome length.

`--max-seed-length INT`

Specifies the maximum number of bases to use to seed.

1.8 tmap mapvsw

The **mapvsw** is a command to map sequences to a reference genome. This algorithm performs the full Smith Waterman algorithm alignment of the read to the reference genome, using SSE2 (vectorized) programming instructions. This algorithm should not be used for large number of reads or large genomes, and instead should be used for debugging and investigating small numbers of reads..

See [section 1.2](#) for common options that are in use in this command.

1.8.1 Usage

There are no algorithm specific options.

1.9 tmap mapall

The **mapall** is a command to quickly map short sequences to a reference genome. This command combines available mapping algorithms for fast and sensitive alignment. The algorithm follows a multi-stage approach, with a set of algorithms and associated settings for each stage. If there are no mappings for a read by applying the algorithms in the i th stage that pass filters, then the algorithms in the $i + 1$ th stage are applied. For example, a set of algorithms to quickly align near-perfect reads may be used in the first stage, while a set of sensitive algorithms may be used to map difficult reads in the second stage

It is recommended that **mapvsw** is not used for any large scale mapping project. Please see [section 1.8](#) for more details.

See [section 1.2](#) for common options that are in use in this command.

1.9.1 Overview

First, a set of global options is specified that will be used for the algorithms to be applied at all stages. Global options should not be given in the options for a specific stage or algorithm. Next, a stage will be specified and its associate options that will be applied across algorithms in that stage. Stage-specific options should not be given in the options for any algorithm in that stage. The stage name should be specified by “stage%d”, where “%d” is the stage number (one-based). Finally, the algorithms and their associated options will be specified for the given stage.

An example would be:

```
tmap mapall -f ref.fasta -r reads.fastq -g 1 -M 3 stage1 --stage-keep-all
map1 --seed-length 12 --seed-max-diff 4 stage2 map2 --z-best 5 map3
--max-seed-hits 10.
```

In this case, the global options `-f ref.fasta -r reads.fastq -g 1 -M 3` will be applied to all stages and algorithms. Therefore, `map1` algorithm will be applied with the options `-g 1 -M 3 --seed-length 12 --seed-max-diff 4` in the first stage. If no mapping is found for a read, the `map2` algorithm with the options `-g 1 -M 3 --z-best 5` and the `map3` algorithm with the options `-g 1 -M 3 --max-seed-hits 10` will be applied in the second stage. Notice how the global options `-g 1 -M 3` are applied to all the algorithms where applicable. It is possible to have the same algorithm run in two different stages.

The recommended values are pre-set, and the following command line is recommended:

```
tmap mapall -f <in.fasta> -r <in.fastq> -n <num.threads> -v map1 map2
map3 > <out.sam>.
```

1.9.2 Usage

The following options apply to a single stage, and control how mappings are handled between stages.

```
--stage-score-thres INT
```

Specifies the number of multiples of the match score (`-A`) for the minimum scoring threshold for the current stage. An alignment is filtered in the given stage if it has an alignment score less than this threshold, and there are more stages to process.

`--stage-mapq-thres INT`

Specifies the mapping quality threshold for the current stage. An alignment is filtered in the current stage if it has a mapping quality less than this threshold, and there are more stages to process.

`--stage-keep-all`

Specifies to keep mappings from the current stage for the next stage. If this option is given, the mappings from the previous stage are added to any mappings from the subsequent stage as candidates.

`--stage-seed-freq-cutoff FLOAT`

Specifies the minimum frequency a seed must occur in order to be considered for mapping.

1.10 tmap server

The **server** command loads the reference genome index and data into shared memory. This lets other mapping instances avoid having to load the index upon each execution. This program will try to fail gracefully, detaching shared memory upon exiting.

See [section 1.2](#) for common options that are in use in this command.

1.10.1 Usage

`-c STRING`

Specifies the command for the server (`start`, `stop`, `kill`). The `start` command loads data into shared memory, and waits for a `ctrl-c` or `SIGINT` signal. The `stop` command will signal a currently running server to stop and detach shared memory. The `kill` command will forcibly detach the shared memory segment and signal a currently running server to exit. The latter command is especially useful for killing zombied processes and detaching lost shared memory segments.

`-k INT`

Specifies the shared memory key for this server. The shared memory segment will be identified by this key.

-a

Specifies to load all reference genome data structures into memory.

-r

Specifies to load the forward packed reference sequence.

-R

Specifies to load the reverse packed reference sequence.

-b

Specifies to load the forward BWT sequence.

-B

Specifies to load the reverse BWT sequence.

-s

Specifies to load the forward suffix array.

-S

Specifies to load the reverse suffix array.

Chapter 2

File Formats

2.1 SAM Alignment Format

TMAP produces mappings in the SAM format (Li *et al.* (2009)). Optional tags are used to store information about mappings useful for downstream processing.

2.1.1 SAM Header Fields

The HD, RG, SQ, and PG SAM header fields will be outputted. Specific details can be found below.

RG

The RG field can be populated using the `-R` option. By default the ID and PG tags are included. When specifying an RG line, do not include the PG tag as it will be populated by tmap: If the input is an SFF file and the `-Y` option is used, then a comment line will be placed in the header with the following tags: RG, FO, and KS. The RG tag in the comment is the associated read group ID. The FO and KS tags indicate the flow order and the key sequence.

PG

The PG field will include the ID, VN, and CL tags. The ID tag should always be tmap.

2.1.2 SAM Record Optional Tags

RG

This tag stores the read group identifier corresponding to the RG SAM header field.

PG

This tag stores the program group identifier corresponding to the PG SAM header field.

MD

This tag stores the MD array. The goal of this array is to follow the conventions in SAMTools (Li *et al.* (2009)). Any IUPAC code originally in the reference will be present in this field. Please see [section 3.1](#) for more details.

NM

This tag stores the edit distance from the reference sequence. Matches to any non-DNA IUPAC ambiguity code (B, D, H, K, M, N, R, S, V, W, Y) will be counted as a mismatch. Please see [section 3.1](#) for more details.

AS

This tag stores the alignment score. All IUPAC code positions are treated as mismatches. If the alignment contains an N in the reference but an A in the read, this score will be incorrect (overestimated) as the N was converted to an A in the reference. Please see [section 3.1](#) for more details.

NH

This tag stores the number of hits found during alignment. The number of alignments reported in the output is determined by `-a` and therefore fewer alignments may be present in the output than specified by the NH tag.

XM

This tag stores the number of mismatches in the mapping, or in the secondary seed if used.

XO

This tag stores the number of indels (gap opens) in the mapping, or in the secondary seed if used.

XG

This tag stores the number of indel extensions (gap extensions) in the mapping, or in the secondary seed if used.

XS

This tag stores the alignment score of next-best sub-optimal mapping.

XT

This tag stores the number of seeds supporting this mapping.

XF

This tag stores from where the mappings originated: one indicates from the forward search, two indicates from the reverse search, and three indicates from both the forward and reverse search.

XE

This tag stores the number of seeds supporting this mapping specifically for map2.

XI

This tag stores the size of the suffix interval for this mapping.

XA

This tag stores the algorithm that produced this mapping and from what stage. The format is the algorithm name, and then the zero-based stage, separated by a dash.

XZ

This tag stores the original alignment score, if an SFF was inputted.

FZ

This tag stores the flow signals when an SFF is inputted and the $-Y$ option is used. The flow-gram values are stored as a string of hexadecimal values, with each group of four hexadecimal values corresponding to one flow signal.

Chapter 3

Commonly Asked Questions

3.1 How are IUPAC ambiguity codes handled in the reference/target FASTA?

Ambiguous IUPAC codes in the reference/target FASTA will be converted to the lexicographically smallest DNA base that is not compatible to the IUPAC code to ensure minimum reference bias. For example, an IUPAC base R, which represents an A or a G, will be converted to a C. All Ns in the reference will be converted to As. Furthermore, any non-IUPAC character will be treated as an N. The ambiguity codes will only be re-considered when calculating the NM and MD SAM record optional tags.

3.2 How are insertions and deletions scored?

Given gap open and gap extension penalties O and E , a contiguous indel of length L will have a score of $O + (E * L)$.

3.3 How are insertions and deletions justified?

TMAP will attempt to justify insertions and deletions (indels) to the 5' (left-most) end of the genomic (reference) forward strand. This allows the indel to be consistently placed with respect the reference, regardless of the strand during sequencing. Neither the query sequence nor the reference sequence are changed, only the indel position in the alignment.

For example, if we have the alignment (query on top, target on the bottom):

```
CGAACATTTTTTCGTGGA
```

```

|||||-----|||
CGAACATTTT-CGTGGA

```

the insertion will be moved to its left-most equivalent position:

```

CGAACATTTTTCGTGGA
|||||-----|||
CGAACA-TTTTCGTGGA

```

Another example would be in a non-homopolymer deletion:

```

CGAACAT----CGTGGA
|||||-----|||
CGAACATACATCGTGGA

```

Justifying this deletion to the left would give:

```

CGA----CAATCGTGGA
|||-----|||
CGAACATACATCGTGGA

```

Justification will not occur if it disconnects a contiguous indel, causing a potential difference in the alignment score. The following alignment:

```

AGGTGC--TTTACAAGC
|||||---|||
AGGTGCCTTTTACAAGC

```

could be changed to:

```

AGGTG-C-TTTACAAGC
||||-|-|||
AGGTGCCTTTTACAAGC

```

This would change the alignment and decrease) the resulting score.

3.4 Will the output of TMAP be the same if run twice?

TMAP will always produce the same output if run twice, even when using multiple threads (-n). This is enabled by using a complementary-multiply-with-carry random number generator, one per thread. Of course, if different options are used, then the results will post likely vary, even if a different number of threads are used.

3.5 How does TMAP soft clip bases in the read?

TMAP has the ability to soft clip bases during alignment, that is to not include in the alignment bases that are present in the read. This may be useful to only map high quality portions of the read, for example if the 3' end of the read has many low quality bases called but their inclusion in the alignment results in many mismatches or insertions/deletions (indels).

The user may specify the desired type of soft-clipping using the `-g` option, with any combination of 5' and 3' soft-clipping (or neither) allowed. The clipped bases are determined from local alignment (glocal or semi-global alignment). For example, if 3' end soft-clipping is enabled, then a prefix of the read is aligned to a sub-sequence of the reference. Where the read prefix ends in the alignment all comes from the local alignment, which is heavily influenced by the scoring parameters.

For example, consider the alignment below, which aligns the full read (GATTACA) to (a subsequence of) the reference (AGATTAAC). Suppose we have scoring parameters (1,-3,-7,-2) for the match, mismatch, indel start, and indel extend operations. The alignment score would be $1+1+1+1+1-7+1 = -1$.

```

READ:  GATTACA
      |||||+|
REF:   GATTA-A

```

If instead we allow soft-clipping at the end of the read, we need only align a prefix of the read to (a subsequence of) the reference. The optimal alignment score would be 5, with the alignment shown below.

```

READ:  GATTACA
      |||||SS
REF:   GATTA

```

The details of the local alignment algorithms and their variations are well characterized in the Literature or in an Introduction to Bioinformatics text book (Jones and Pevzner (2004)). Using the local alignment and the scoring parameters to determine the soft-clipping is a powerful and flexible solution to trade off reduced alignment length for higher quality alignments via soft-clipping.

For how soft-clipping is represented in SAM files, see the SAM specification for more details (<http://samtools.sourceforge.net/SAM1.pdf>).

3.6 What is flow space re-alignment

Re-alignment in flow space is possible with TMAP, and is turned on only when the flow order option is set (-F). With normal Smith Waterman, errors in homopolymers are treated similarly to insertions and deletions (indels) that are from biological origins. Re-alignment in flow space allows TMAP to consider indels and errors in homopolymers differently than other indels, even though the resultant base space alignment may be similar.

For example, the following alignment (read on top) shows a one-base insertion of a G. This could be due to an error estimating the number of Gs from a specific flow, or a true biological difference/insertion.

```
TAGGACACCGGTCTGA
||||||| - |||||
TAGGACACC-GTCTGA
```

In the following example, we observe a mismatch/SNP difference in base space:

```
TAGGACACCGGTCTGA
||||||| |||||
TAGGACACCCGTCTGA
```

This could be due to a biological difference/SNP, or due to an under-call of the C and an overcall of the adjacent G. In the latter case, an acceptable alignment may be:

```
TAGGACACC-GGTCTGA
||||||| -- |||||
TAGGACACCC-GTCTGA
```

Finally, more complicated scenarios can arise, where flow space re-alignment may yield a more faithful alignment. In the following example, a base space alignment may be:

```
TCAT--AAATTTT
||| --|||||||
TCAATAAAATTTT
```

A suitable flow space re-alignment would be:

```
TCA-T-AAATTTT
||| -|-|||||||
TCAATAAAATTTT
```

The first alignment may be preferred in base space due to the use of affine gap penalties (see [section 3.2](#)) and the other scoring parameters. If we penalize insertions or deletions due to homopolymer differences less than other insertions or deletions, the latter alignment may be more likely. Given Ion Torrent data, the latter may be more faithful to the underlying error mode.

Chapter 4

Appendix

Bibliography

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