

DMAP2: User Guide

DMAP2 builds on the original DMAP distribution (Stockwell, *et al.*, 2014) by providing scripts to simplify the selection of processing options and running the programs for the various steps.

Overview:

After installation of the various prerequisite software packages and genomic resources, DMAP has a series of steps:

- a. Genome preparation for `bismark` mapping (usually only needs doing once for a series of samples)
- b. Adaptor trimming of bisulphite sequence reads for each sample
- c. Mapping reads for each sample to the genome with `bismark` producing `.bam` or `.sam` files
- d. Comparing mapped data with the DMAP program `diffmeth`
- e. Relating genomic fragments or regions of interest to genomic features with the DMAP program `identgeneloc`

DMAP2 groups these steps into a number of scripts each implementing one or more operations. If prior work has already performed some steps (genome preparation, mapping) then those can be omitted and appropriate information put in the basic parameter file. While the mapping scripts are based on using the `bismark` bisulphite aligner, other aligners can be used (e.g. `bsmap`) in which case they can be processed from the `diffmeth` point at step (d).

Script details:

(i) Information common to all scripts is specified in a file like `dmap_basic_params.sh` which defines the following variables:

`verbose` - set to "yes" for informational messages
`path_to_dmap` - if DMAP executables are not defined on your path, leave blank if they are
`dmap_genome_file_location` - directory for the genome fasta files
`dmap_genome_fasta_files` - list of genome fasta files, one for each chromosome
`dmap_bismark_index_location` - where bismark index is, defaults to genome location
`dmap_path_to_bismark_exes` - where bismark executables are, blank if already on your path
`dmap_chr_info_file` - file for `diffmeth` to locate genome files: defaults to `dmap_chr_info.txt`
`bismark_build_options` - options for building bismark genomic index
`bismark_run_options` - options for bismark at runtime
`feature_annotation_type` - to define annotation format (EMBL, Genbank, SeqMonk, GTF or GFF3)
`annotation_file_location` - location of the annotation file(s)

`annotation_files` - list or array of the files: SeqMonk, EMBL or Genbank expect 1 file for each chromosome, GTF uses a single file

`dmap_annot_info_file` - name of the information file generated for

SeqMonk/EMBL/Genbank annotations: defaults to `dmap_annot_info.txt`

We have found SeqMonk annotation useful when the locations of TSS (Transcription Start Sites) and CpG Islands are needed, since these are contained in the SeqMonk files. It is also possible to include extra features into SeqMonk files if required. See **Prerequisites** section below for descriptions of how to obtain SeqMonk files or Gencode GTF annotation files.

(ii) **`dmap_index_build.sh`** – performs the genome preparation (step a) and further sets up details for the annotation step (e). The script takes information from `dmap_basic_params.sh`. By default, the bismark indices will be built in the same location as the genome fasta files, but this would require that you have write access to that location. If not then you can specify some other writeable directory in `dmap_bismark_index_location` in `dmap_basic_params.sh`.

Since all the mapping for an experiment would usually be done against the same genome, the index building step should only be needed once.

The annotation preparation depends on the annotation format and produces an information file relating chromosome IDs to files for EMBL, Genbank and SeqMonk formats. No action is needed for GTF/GFF3 formats.

Once the file `dmap_basic_params.sh` has been set up for your environment the script should be run with a command like:

```
./dmap_index_build.sh dmap_basic_params.sh
```

noting that this will take some time for the bismark genome generation process. It may be worth using Linux/Unix tools like `nohup` to keep such jobs running in the background, especially if you need to disconnect from the machine running this operation.

DMAP expects genome fasta files to be one file per chromosome. In the event that multiple chromosomes or contigs are present in one file, the script `split_fasta.awk` in Appendix VI will separate them into individual files, each named by its fasta ID by default. Script documentation is in the file.

(iii) **`map_bisulphite_reads.sh`** performs the steps of adaptor trimming (b) and bismark mapping (c). Parameters specific to each sample are in a file like `sample_params.sh` (example in Appendix II) where `dmap_run_type` is set to RRBS or WGBS, needed here since RRBS library preparation adds a CG dinucleotide to the end of each MspI fragment and these require removal during adaptor trimming. In order to avoid issues with overrepresentation of C's central to short reads with paired end mapping, we routinely used read 1 only. See Appendix I for other possible options.

`map_bisulphite_reads.sh` creates and uses an adaptor file `contam.fa` for the adaptor trimming step, it also makes a file `<sample_name>_bismark_cmd.sh` to execute bismark, where `<sample_name>` is based on the name in `dmap_sample_files` in the `sample_params.sh` file for the sample.

For example: the original untrimmed read file `mysample1_R1.fastq` would be adaptor trimmed to `mysample1_R1_at.fastq` then mapped using the command in `mysample1_R1_bismark_cmd.sh` to produce output files like `mysample1_R1_at_bismark_bt2.bam` and `mysample1_R1_at_bismark_bt2_SE_report.txt`.

Running the mapping for each sample would be done with a command like:

```
./map_bisulphite_reads.sh dmap_basic_params.sh sample_params.sh
```

again noting that some time will be needed for each sample. The use of `nohup` or other utilities to run these jobs in the background is useful.

(iv) `dmap_run_diffmeth.sh` takes mapped data from `bismark` and performs the required differential methylation analysis (step d) with parameters from a file which specifies the type of analysis, the input files and the `diffmeth` output file. An example `diffmeth` parameter file is in Appendix III and, in this case, is for a 3 group Anova run. The important variables are:

`dmap_run_type` - defines RRBS or WGBS and should match the setting for the `map_bisulphite_reads.sh` step
`diffmeth_run_type` - defines the analysis type (Anova, chisquare, pairwise CpGlist or binlist)
`file_list1`, `file_list2` and `file_list3` each contains a list (array) of bam files for each group: the Anova analysis here requires 3 groups
`diffmeth_out_name` sets the name of the output file.

Analyses needing only a single file or group of files (`diffmeth_run_type` in `chisquare`, `CpGlist` or `binlist`) will only need a single file in `file_list1`. Pairwise runs need two file names in that list.

Anova runs also need `anova_group_number` and `anova_detail` to be defined as well as additional lists of file names in `file_list2`, etc.: one such list for each group. If the variable `anova_fold_difference` is set to "yes" a column of fold methylation differences will be added to the output.

Further control of `diffmeth` parameters is *via* the variables:

`diffmeth_fragment_min` - minimum length of RRBS fragments
`diffmeth_fragment_max` - maximum length of RRBS fragments
`wgbs_window` - window length for WGBS runs
`min_cpg_hits` - minimum number of hits for a CpG to be included in analysis
`min_valid_cpgs` - number of CpGs meeting `min_cpg_hits` for a fragment/region to be included
`min_sample_count` - minimum number of samples fulfilling `min_cpg_hits` and `min_valid_cpgs` for a fragment/region to be included
`map_init_CpG_to_prev` - for 3' matching reads, the initial C is counted in the adjacent RRBS fragment
`pr_threshold` - reject fragments/regions with Pr above this threshold

For SAM mapping files it may be necessary to define `max_sam_readlength` if the reads exceed 150bp. This is not required for BAM files.

Run the `diffmeth` script with a command like:

```
./dmap_run_diffmeth.sh dmap_basic_params.sh my_diffmeth_params.sh
```

noting that the execution time can range from 20 minutes to some hours, depending on the amount of data and the computer performance. `diffmeth` runs will typically require some 5Gb of RAM or more depending on the genome size, the number of samples and the coverage.

(v) `dmap_run_genloc.sh` implements step e on the output from `diffmeth` runs or other appropriately formatted input in which `identgeneloc` looks upstream (in the sense of the feature) for the nearest feature of interest to each region/fragment. The minimum input is 3 columns (chromosome ID, start, stop) but more usually the other columns generated by `diffmeth` would be included.

`dmap_run_genloc.sh` requires a parameter file (example in Appendix IV) to specify the source of annotation data and to control other aspects of the run. Variables are:

`genloc_input_file` - name of 3 or more column input file
`genloc_output_file` - name of output file, default is `stdout`, but more usefully set to a named file
`genloc_column_number` - number of columns of input file to include, leave as 0 for automatically including all columns
`internal_to_genes` - control whether regions can be located internal to genes, a value of `internal` will permit this, `exon_intron` will further relate regions to intron/exon boundaries
`show_TSS` - for SeqMonk annotation, show nearest upstream transcription start sites (TSS)
`TSS_limit` - if non-zero will limit how far TSS will be scanned from regions.
`show_CpGi` - for SeqMonk annotation, show nearest upstream CpG island
`show_CpGi_internal` - allow CpG islands to be internal to regions
`show_TSS_CpGi_ranges` - show ranges for CpG islands and TSS
`genloc_seqmonk_biotypes` - for SeqMonk annotation restrict features of interest to specific biotypes (e.g. `protein_coding`): multiple values accepted
`genloc_GTF_attributes` - for GTF annotation, use the specified attributes, e.g. `transcript_name`, `transcript_id`, `gene_name`; multiples allowed
`genloc_GTF_featuretypes` - for GTF annotation, specify the desired features as in column 3 of the GTF file, e.g. `gene`, multiples allowed.

Run the `identgeneloc` script with a command like:

```
./dmap_run_genloc.sh dmap_basic_params.sh geneloc_params.sh
```

`identgeneloc` runs will typically require some 350Mb of memory, depending on the size of the annotation file(s) and would take some 12 minutes or so for input files of 500,000 regions.

Prerequisites:

In order to run DMAP the following prerequisites should be obtained and installed.

(a) Genomes

DMAP requires genomes in FASTA format with each chromosome in an individual file. Various sources exist, but one of the most convenient is <https://www.ensembl.org/info/data/ftp/index.html> following the 'DNA (FASTA)' link. Three different versions of each chromosome are offered: `.dna.`, `.dna.rm.` and `.dna.sm.` respectively being complete genome, genome with repeat regions masked as 'N' and genome with repeats in lower case. The first is probably the most useful. Files can be downloaded by selecting each individually with the browser or as a group from the command line (for Homo sapiens) using a command like:

```
curl -O ftp.ensembl.org/pub/release-105/fasta/homo_sapiens/dna/Homo_sapiens.GRCh38.dna.chromosome.[1-22].fa.gz
```

followed by

```
curl -O ftp.ensembl.org/pub/release-105/fasta/homo_sapiens/dna/Homo_sapiens.GRCh38.dna.chromosome.[X-Y].fa.gz
```

noting that each command is complete on one line. The files are gzip compressed and, when put in an appropriate directory, can be uncompressed with a command like:

```
gunzip Homo_sapiens.GRCh38.dna.chromosome.*.fa.gz.
```

In `dmap_basic_params.sh` set `dmap_genome_file_location` to the directory where these files have been decompressed and fill `dmap_genome_fasta_files` with the chromosome file names.

(b) Annotation files in one of the following formats:

(i) SeqMonk - obtained through the SeqMonk mapped sequence data analyser, available from

<https://www.bioinformatics.babraham.ac.uk/projects/seqmonk/>

by starting the application and choosing `File > New Project` then 'Import Genome From Server' and selecting then downloading the genome of choice. This should create a directory `seqmonk_genomes`, probably in your top level directory, containing the organism and genome version which contain a series of files `1.dat`, `2.dat`, ... - these contain the annotations for each chromosome. Set the location of these files in `dmap_basic_params.sh` as `annotation_file_location` and fill `annotation_files` with the list of chromosome files. `feature_annotation_type` is set to SeqMonk.

SeqMonk annotation has proven particularly useful when transcription start sites and CpG islands are needed and allows the choice of specific biotypes for the features of interest. Further, it is possible to append other feature information (e.g. enhancers) to each chromosome: Appendix V contains examples of added features.

(ii) GTF - obtained from <https://www.gencodegenes.org/> for mouse or human, by choosing the organism and genome version, then downloading the comprehensive gene annotation file as GTF which will download a file with a name like: `gencode.v39.annotation.gtf.gz` which can be moved to an appropriate location where it can be uncompressed for DMAP with:

```
gunzip gencode.v39.annotation.gtf.gz
```

producing `gencode.v39.annotation.gtf`. Set the location in `dmap_basic_params.sh` as `annotation_file_location` and `annotation_files` to the filename. `feature_annotation_type` is set to GTF.

(iii) GFF3 - obtained as for GTF, with similar basic parameter settings.

(iv) EMBL - obtained from <https://www.ensembl.org/info/data/ftp/index.html> as 'Annotated sequence (EMBL)' or consider using utilities like `ftp` or `curl` to transfer the files at the command line, with a command (for Homo sapiens) like:

```
curl -O ftp.ensembl.org/pub/release-105/embl/homo_sapiens/Homo_sapiens.GRCh38.105.chromosome.[1-22].dat.gz
```

(all complete on 1 line). X and Y chromosomes will need [1-22] replaced by [X-Y].

The retrieved files will need decompressing with a command like:

```
gunzip Homo_sapiens.GRCh38.105.chromosome.*.dat.gz
```

Set the location of these files in `dmap_basic_params.sh` as `annotation_file_location` and fill `annotation_files` with the list of chromosome files. `feature_annotation_type` is set to EMBL.

Note that EMBL (and Genbank) feature tables obtained from ENSEMBL only return ENSEMBL gene IDs with DMAP.

(v) Genbank - available from NCBI, though hard to find there and comes with awkward chromosome IDs. Better obtained similarly to EMBL from Ensembl as 'Annotated sequence (GenBank)' or at the command line with something like:

```
curl -O ftp.ensembl.org/pub/release-105/genbank/homo_sapiens/Homo_sapiens.GRCh38.105.chromosome.[1-22].dat.gz
```

as above. `feature_annotation_type` is set to Genbank.

For SeqMonk, EMBL and Genbank annotation formats the list of file names and chromosome IDs are written to the file set in `dmap_annot_info_file`, which defaults to `dmap_annot_info.txt`.

Note that Genbank (and EMBL) feature tables obtained from ENSEMBL only return ENSEMBL gene IDs with DMAP.

(c) **Bismark**: while other bisulphite aligners (e.g. BSMAP at <https://code.google.com/archive/p/bimap/>) will work effectively with DMAP, we have generally found Bismark to be a good performer. Bismark is available from <https://github.com/FelixKrueger/Bismark> or from <https://www.bioinformatics.babraham.ac.uk/projects/bismark/> and is installed as per instructions, which amounts to putting the perl executables into an appropriate directory: either system ones (/usr/local/bin - system privilege required) or your own top level bin. The variable `bismark_run_options` in `dmap_basic_params.sh` can be set to reflect your needs: we have found that '-N 1' restricts the permitted seed mismatches to 1 (defaults to 2) works well for our data and gives significantly faster mapping. The option '--bowtie2' is not now necessary for recent bismark versions, but remains for completeness.

(d) **bowtie2**: recent versions of Bismark require the aligner bowtie2. If not already installed in your systems it is available from <https://github.com/BenLangmead/bowtie2> or <https://sourceforge.net/projects/bowtie-bio/files/bowtie2/>, the latter providing precompiled binaries which can be unzipped and installed in an appropriate directory.

(e) **samtools**: required by Bismark in order to generate bam rather than sam files. (bam files are about 1/3 the size of the corresponding sam files.) If not already installed, it is available from <https://sourceforge.net/projects/samtools/> as source code which will need unpacking and compiling using instructions like:

```
bzip2 -dc samtools-1.14.tar.bz2 | tar -xvf -
cd samtools-1.14
./configure
make
make install
```

The latter instruction may require system privilege (e.g. sudo). Alternatively you can copy the executable to a location on your PATH with something like:

```
cp samtools ~/bin/
```

(f) **DMAP**: available as source code from <https://github.com/peterstockwell/DMAP> either by downloading as a zip archive from 'Code->Download ZIP' and using unzip or from the command line with:

```
git clone https://github.com/peterstockwell/DMAP
```

producing either DMAP-master or DMAP which contains instructions for building in README.md.

The source code should compile with any reasonable C compiler, but note that the library zlib is needed for full functionality with .bam files. The main executables needed are `difmeth` and `identgeneloc` which should be copied to an appropriate directory on your PATH or set `path_to_dmap` in `dmap_basic_params.sh` to the actual location.

References:

Stockwell, P.A., Chatterjee, A., Rodger, E.J. and Morison, I.M. "DMAP: Differential Methylation Analysis Package for RRBS and WGBS data" *Bioinformatics* (2014) DOI: 10.1093/bioinformatics/btu126.

Krueger, F., and Andrews, S.R., Bismark: a flexible aligner and methylation caller for Bisulfite-Seq applications, *Bioinformatics*, (2011), <https://doi.org/10.1093/bioinformatics/btr167>

Appendix I: Options for mapping paired end reads.

The authors of `bismark` (Felix Krueger and Simon Andrews, 2011) advise that the use of paired end reads with RRBS fragments will result in over-representation of the overlapping portion of short fragments, given that the 40-220 bp size selection will generate many reads below the usual read lengths presently available. A possible solution is to use tools like `FLASH` (<http://www.cbcb.umd.edu/software/flash>) to generate longer reads where paired end reads clearly overlap. This process generates 3 fastq output files: longer overlapped reads, R1 non-overlapped and R2 non-overlapped. Since `bismark` is only configured to accept either single ended files (or a list of them) or paired end files for R1 & R2 (or a list for each), a strategy would be for two `bismark` runs, one single-ended for joined reads, the other paired-ended for unjoined reads, then use `samtools merge` to combine the results before continuing with `DMAP` analysis. Our experience with this was that it did not improve the mapping efficiency sufficiently to justify the extra work, hence our decision to work with single-ended data.

Appendix II: An example sample parameter file for mapping:

```
# sample_params.sh: parameters to describe a single bisulphite
# sequence sample, including the raw fastq file(s) and any
# trimming that might be justified by the read qualities.

# identify if this was RRBS or WGBS: this determines a number
# of downstream operations, including adaptor trimming and
# differential methylation calculations. Values are RRBS or WGBS.

dmap_run_type="RRBS";
# dmap_run_type="WGBS";

# name of adaptor-trimmed output dir:

adtrimmed_out_dir="./";

# sample file name(s): we usually only map read 1, so
# read 2 name would omitted.

dmap_sample_files=("CRC-P-10_1.fastq.gz" "CRC-P-10_2.fastq.gz");

# hard trim length, leave 0 for no trimming.

dmap_sample_trim_length=0;

# minimum read length for retention: usually set this to 20

dmap_min_read_length=20;

# mapping output directory

bismark_out_dir="bismark_out/";
```

Appendix III: Example parameter file for **diffmeth** runs

In this case an Anova run with three groups.

```
# diffmeth_run_params.sh: parameters to define differential
# methylation run type and mapped files for the DMAP diffmeth
# program.

# identify if this was RRBS or WGBS: this determines a number
# of downstream operations, including adaptor trimming and
# differential methylation calculations. Values are "RRBS" or "WGBS"
# where:
# "RRBS" compares methylation on fragments (usually MspI) in size range
# "WGBS" compares methylation on defined windows or regions
#
# this is previously specified for the adaptor trimming and mapping
step
# and obviously the values here should be consistent with those.

dmap_run_type="RRBS";
# dmap_run_type="WGBS";

# type of diffmeth run: values are one of "pairwise" "chisquare"
"Anova" "CpGlist" or "binlist"
# where:
# "pairwise" runs Fisher's Exact statistic on two samples
# "chisquare" runs a Chi Square test on multiple samples to identify
fragments/regions with greatest variation
# "Anova" run Analysis of Variance on two sets of multiple samples,
producing F statistic
# "CpGlist" lists methylation counts for each CpG
# "binlist" lists methylation information for each fragment/region

diffmeth_run_type="Anova";

case $diffmeth_run_type in
pairwise)
# "pairwise" requires two mapping output files
file_list1=(("./rundata/bismark_out/CRC-P-10_1_at_bismark_bt2_pe.bam"
"./rundata/bismark_out/NC-P-20_1_at_bismark_bt2.bam"));
;;
chisquare)
# "chisquare" requires a list of at least 2 mapping output files

file_list1=(
"./rundata/comb_D1.bam"
"./rundata/comb_D2.bam"
"./rundata/comb_D3.bam"
);
;;
Anova)
# Anova requires a group number to define how many groups are involved
# and a list of at least 2 mapping output files for each group
# a further parameter defines the amount of information returned, being
# one of: "anova_simple" "anova_medium" or "anova_full"
# where
# "anova_simple" returns F statistic and its probability
# "anova_medium" adds sample counts and greater methylated group
```

```

# "anova_full"    further adds methylation figures for each CpG in
fragment/region

anova_group_number=3;

file_list1=(
  "./rundata/comb_D1.bam"
  "./rundata/comb_D2.bam"
  "./rundata/comb_D3.bam"
  "./rundata/comb_D4.bam"
);

file_list2=(
  "./rundata/comb_D5.bam"
  "./rundata/comb_D6.bam"
  "./rundata/comb_D7.bam"
);

file_list3=(
  "./rundata/comb_D8.bam"
  "./rundata/comb_D9.bam"
  "./rundata/comb_D10.bam"
)

anova_detail="anova_full";
;;
CpGlist | binlist)
# "CpGlist" and "binlist" require a single mapping file
file_list1=("./rundata/comb_D1.bam")
;;

esac

# diffmeth requires a fragment size specification, even for WGBS
(ignored then)

diffmeth_fragment_min=40;
diffmeth_fragment_max=220;

# "WGBS" requires a window (tile) length

wgbs_window=1000;

# output file name for diffmeth

diffmeth_out_name="diffmeth_output_anova_test_I6_mod.txt";

# threshold criteria for including fragment/regions

# No. of hits per CpG for fragment/region to be valid
# default is 1

min_cpg_hits=10;

# No. of CpG of fragment/region that must meet min_cpg_hits criterion
# 0 indicates no check for this

min_valid_cpgs=0;

```

```

# Minimum No. samples reaching criteria for a fragment/region to be
included
# defaults to 2

min_sample_count=6;

# For RRBS 3' mapping reads, count initial CpG to previous fragment.

map_init_CpG_to_prev="yes";

# Probability threshold: omit fragments/regions with Pr greater.
# default (1.0) implies no filtering

#pr_threshold=1.0;

# Max read length for .SAM files (not needed for .BAM, though)
# defaults to 150bp, for 0

#max_sam_read_length=150;

```

Appendix IV: Example parameter file for `dmap_run_genloc.sh`

```

# genloc_run_params.sh: parameters to define gene location
# operation with identgeneloc. Some relevant parameters
# are already set with the basic parameters, this file
# relates to the specific settings required for a project.

# file of chromosome & coordinate data (min 3 columns) as:
# chrId start stop
#

genloc_input_file="diffmeth_output_binlist_nz.txt";

# name of output file: identgeneloc writes to stdout, we need
# to direct that somewhere, else will still go to stdout

genloc_output_file="genloc_binlist_nz_chrchk_gc.txt";

# the number of columns can be identified automatically
# putting a non-zero value here will override the automatically
# chosen value

genloc_column_number=0;

# Control whether regions internal to genes are allowed
# and whether they are to be related to exon/intron
# boundaries.

# values for the parameter internal_to_genes are:

# "" = disallow
# "internal" = allow
# "exon_intron" = relate to exon/intron boundaries

internal_to_genes="exon_intron";

```

```

# SeqMonk annotations define TSS and CpG islands
# ignored for other annotation formats

show_TSS="yes";

# limit for TSS ranges, zero=nolimit

TSS_limit=0;

show_CpGi="yes";

# CpG islands can be shown within regions: "yes" permits this

show_CpGi_internal="yes";

show_TSS_CpGi_ranges="yes";

# For SeqMonk annotations, can restrict to particular biotypes;
# valid values are: 3prime_overlapping_ncrna IG_C_gene IG_C_pseudogene
IG_D_gene IG_J_gene IG_J_pseudogene
# IG_V_gene IG_V_pseudogene Mt_rRNA Mt_tRNA Mt_tRNA_pseudogene TEC
TR_C_gene TR_D_gene TR_J_gene TR_J_pseudogene
# TR_V_gene TR_V_pseudogene ambiguous_orf antisense lincRNA miRNA
miRNA_pseudogene misc_RNA misc_RNA_pseudogene
# ncrna_host non_coding non_stop_decay nonsense_mediated_decay
polymorphic_pseudogene processed_transcript
# protein_coding pseudogene rRNA rRNA_pseudogene retained_intron
scRNA_pseudogene sense_intronic sense_overlapping
# snRNA snRNA_pseudogene snoRNA snoRNA_pseudogene tRNA_pseudogene
transcribed_unprocessed_pseudogene

# protein_coding is probably one of the most useful.

genloc_seqmonk_biotypes=(
"protein_coding"
)

# GTF & GFF3 annotations allow selection of attributes and feature
types
#
# Attributes default to "transcript_name,transcript_id,gene_name" if
# not specified. Actual attributes can vary widely for GTF format

genloc_GTF_attributes=(
"transcript_name"
"transcript_id"
"gene_name"
)

# feature types default to "exon,CDS,start_codon,stop_codon,transcript"
if
# not specified. Valid values for GTF are are any of:
#
CDS,start_codon,stop_codon,exon,intron_CNS,intron,5UTR,3UTR,CNS,inter,t
ranscript,gene
# GFF3 values are far more diverse

```

```
genloc_GTF_featuretypes=(  
"gene"  
)
```

Appendix V: Example of additional SeqMonk annotation appended to a chromosome feature file

```
FT    gene    856539..856757  
FT      /description="enhancer chr1:856539-856757"  
FT      /name="enh_856539"  
FT      /score="2"  
FT      /biotype="enhancer"  
FT    gene    858256..858648  
FT      /description="enhancer chr1:858256-858648"  
FT      /name="enh_858256"  
FT      /score="5"  
FT      /biotype="enhancer"  
FT    gene    868565..868684  
FT      /description="enhancer chr1:868565-868684"  
FT      /name="enh_868565"  
FT      /score="2"  
FT      /biotype="enhancer"
```

Appendix VI: **split_fasta.awk** script to split multi-sequence fasta files.

```
# split_fasta.awk: script to break multi-entry fasta
# files into separate files, each containing 1 entry.
# files named by the fasta IDs, with optional header
#
# Peter Stockwell: 21-Feb-2022
#
# usage:
# awk -f split_fasta.awk <multi_fasta_file>
#
# or piped, e.g.
#
# cat <multi_fasta_file> | awk -f split_fasta.awk
#
# Can define optional useful parameters in command line:
# namehdr - prefixed to each output file name, can be a directory or
# string or both (e.g. namehdr="singles_fasta/")
# extension - use as file extension, default is ".fa"
# infofile - will write table of Chr IDs and full file paths to this
# file
#
# use as: e.g.
#
# awk -f split_fasta.awk namehdr="genome_singles/" extension=".fasta"
# infofile="chr_info.txt" <multi_fasta_file>
#
# to produce chr_info.txt which is suitable for the diffmeth -G option
# If namehdr is a directory, it must already exist.
#

BEGIN {namehdr="";
extension=".fa";
outfile="";
infofile=""
cwd=ENVIRON["PWD"];
}

$0~/>/{if (outfile != "")
    close(outfile);
outfile = sprintf("%s%s%s",namehdr,substr($1,2),extension);
printf("%s\n",$0) > outfile;
if (infofile != "")
    printf("\t%s\t\t%s/%s\n",substr($1,2),cwd,outfile) > infofile;
}

$0!~/>/{
printf("%s\n",$0) > outfile;
}

END{
if (outfile != "")
    close(outfile);
if (infofile != "")
    close(infofile);
}
```