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## NAME

cutter.pl

## SYNOPSIS

```
% perl cutter.pl <Lmer> <#bp per chunk> <#bp overlap between chunks > <last chunk must be this %
chunk size> e.g. % perl cutter.pl 4 10000 3000 0.5
```

means chunks of 10000bp, each subsequent chunk will start 3000bp downstream, and ensure that the last chunk is at least 50% of the present chunk size; this means the last chunk(file) is in the size range of: [ 50% of chunk size .. 1.5x chunk size)

## DESCRIPTION

### SUMMARY

Cutter.pl ("Script #1") is the second of a suite of scripts designed to assist in the analysis of DNA. This particular script breaks a large DNA sequence down into several smaller chunks of user-determined size. The script then shifts downstream by a user-determined amount (note that we recommend chunk size and shift size should be the same) and repeats the process until it reaches the end of the sequence. If the last chunk is large enough in proportion to the previous chunk (again, this is determined by the user) it will be treated as its own separate chunk; otherwise it will be added on to the previous chunk. This process can be repeated multiple times for several files as needed.

The "chunks" produced by this script can be passed to motifCounts.pl ("Script #2") for analysis.

### INPUT

Run Cutter.pl using FNA files extracted from your database using the script extract\_ALL\_chr.pl.

### OUTPUT

Cutter.pl produces several FNA files containing discrete "chunks" of DNA for use with Script #2 (motifCounts.pl). These smaller chunks are much more manageable (and therefore more preferable) than using entire genomes at a time.

## AUTHORS

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## MODIFICATION HISTORY

### June 1, 2010 (nkf) --

Fixed some potential problems with regex across platforms.

### May 31, 2010 (nkf) --

Improved the internal documentation. Wrote some of the readme documentation.

Fixed a bug that caused "tailing" chunks to get deleted instead of added to the end of the previous chunk.

Modified file I/O code to implement cross-platform functionality.

**Jan 08, 2009 (mdl) --**

added separate output directory for each chromosome  
for example: in ~split\_texts/data\_all\_chr/  
Agrobacterium\_tumefaciens\_C58\_Cereon\_\_chr01/  
Agrobacterium\_tumefaciens\_C58\_Cereon\_\_chr02/

**Nov 18, 2008 (mdl) --**

changed to bp rather than #words

**October 01, 2008 (cn) --**

reworked for DNA

**June 17, 2008 (acj) --**

changed output file names to include genre and manuscript and text numbers

reworked algorithm to work work with chunk sizes and shift sizes that are  
not multiples of each other

**June 16, 2008 (acj) --**

finished commenting and wrote pod

fixed the last chunk problem

**June 12, 2008 (acj) --**

wrote and debugged script (for anglo-saxon files =:o )

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