

Unix, Perl and BioPerl

II: Sequence Analysis with Perl

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Sequence Analysis with Perl

- Introduction
- Input/output
- Variables
- Functions
- Control structures
- Comparisons
- Sample scripts

Objectives

• write, modify, and run simple Perl scripts

 design customized and streamlined sequence manipulation and analysis pipelines with Perl scripts

Why Perl?

- Good for text processing (sequences and data)
- Easy to learn and quick to write
- Built from good parts of lots of languages/tools
- Lots of bioinformatics tools available
- Open source: free for Unix, PC, and Mac
- TMTOWTDI

A first Perl program

• Create this program and call it hey.pl

```
#!/usr/local/bin/perl -w
# The Perl "Hey" program
print "What is your name? ";
chomp ($name = <STDIN>);
print "Hey, $name, welcome to the
 Bioinformatics course.\n";
• To run: perl hey.pl
• To run: chmod +x hey.pl
         ./hey.pl
```

Perl Input/Output

- Types of input:
 - keyboard (STDIN)
 - files
- Types of output:
 - screen (STDOUT)
 - files
- Unix redirection can be very helpful
 ex: ./hey.pl > hey output.txt

Variables

Scalar variables start with \$

• Arrays (lists of scalar variables) start with @:

```
@genes = ("BMP2", "GATA-2", "Fez1");
@orfs = (395, 475, 431);
print "The ORF of $genes[0] is $orfs[0] nt.";
# The ORF of BMP2 is 395 nt.
```

Perl functions - a sample

print	tr///	closedir	open	m//
chomp	mkdir	split	close	die
length	chdir	join	chmod	rename
substr	opendir	pop	uc	use
s///	readdir	push	lc	sort

Control Structures 1

```
if (condition) # note that 0, ", and (undefined) are false
   do this; then this;...
          # optional; 'if' can be used alone; elsif also possible
else
   do this instead;
   ($exp >= 2)
                        # gene is up-regulated
  print "The gene $seq is up-regulated ($exp)";
```

Control Structures 2

```
while (condition)
  do this;
  then this;...
while ($orfLength > 100)
                          # Add to table
                      # "\t" = tab
 print "$seq\t";
 print "$orfLength\n"; # "\n" = newline
```

Control Structures 3

```
for (initialize; test; increment)
   do this;...
for ($i = 0; $i <= $#seqs; $i++)
# $#seqs = index of the last element in @seqs
   # Add elements of @seqs and @orf to table
  print "$seq[$i]\t";
  print "$orf[$i]\n";
```

Arithmetic & numeric comparisons

```
• Arithmetic operators: + - / * %
• Notation: \$i = \$i + 1; \$i += 1; \$i++;
• Comparisons: >, <, <=, >=, ==, !=
 if ($num1 != $num2)
    print "$num1 and $num2 are different";
• Note that = = is very different from =
           used as a test: if (\text{num} = 50)
           used to assign a variable: num = 50
```

String comparisons

• Choices: eq, ne

```
if ($gene1 ne $gene2)
{
    print "$gene1 and $gene2 are different";
}
else
{
    print "$gene1 and $gene2 are the same";
}
```

Multiple comparisons

```
• AND
          22
• OR
if (($exp > 2) | |
  ($exp > 1.5 \&\& $numExp > 10))
 print "Gene $gene is up-regulated";
```

Filehandles

```
To read from or write to a file in Perl, it first needs to be opened.
In general, open (filehandle, filename);
Filehandles can serve at least three purposes:
open(IN, $file);
                 # Open for input
open(OUT, ">$file"); # Open for output
open(OUT, ">>$file"); # Open for appending
Then, get data all at once @lines = <IN>;
or one line at a time
  while (<IN>) {
      $line = $ ; do stuff with this line;
      print OUT "This line: $line"; }
```

Embedding shell commands

- use backquotes (`) around shell command
- example using EMBOSS to reverse-complement: `revseq mySeq.fa mySeq_rc.fa`;
- Capture stdout from shell command if desired
- EMBOSS qualifier "-filter" prints to stdout
 \$date = `date`;
 \$rev_comp = `revseq mySeq.fa -filter`;
 print "\$date";
 print "Reverse complement:\n\$rev_comp\n";

Programming issues

- What should it do and when is it "finished"?
- Who will be using/updating your software?
 - Reusability
 - Commenting
 - Error checking
- Development vs. execution time?
- Debugging tools: printing and commenting
- Beware of OBOB ("off-by-one bug")

Example: patscan_batch.pl

```
#!/usr/local/bin/perl -w
# Run patscan on all seqs in a folder
$myDir = "/home/elvis/seqs";
$patFile = "/home/elvis/polyA.pat";
chdir($myDir);
                              # Go to $myDir
opendir(DIR, $myDir);
                              # Open $myDir
foreach $seqFile (sort readdir(DIR))
if (\$seqFile =~ /\.fa\$/)
                              # if file ends in .fa
 print "Processing $seqFile\n";
 $outFile = $seqFile;
                     # Create $outFile name
 $outFile =~ s/\.fa/\.out/; # s/old/new/;
 `scan for matches $patFile < $seqFile > patscan/$outFile`;
                                                    18
```

Example: oligo analysis



sample fasta sequence:

>gi|16493450|gb|BB659629.1|BB659629
GCCTGCTTGAGTTTTGAAGTCTTGGAGCCACAGAA
AGCACTGGCCAGAGGAGGAGGTAATCACTTCTAATG
CCAGGCCTGCTGTGCAGTGCGCATGTGTGATCTCA
GTCTGCTTCTGCCCTAGCTAATGAAGGCATGGACA
ATGGAATAGCCACATGGCAGCACCGGAAAACAAGC
TTACTTCTGCAGTACACAGCCTGCTTTGCCTGATT
TCTGTCCACTGG

Basic steps for oligos.pl

Open fasta sequence

Get raw sequence

Extract oligos

Analyze oligos

Print out results

(Modify script to analyze multiple seqs)

oligos.pl: part 1

```
#!/usr/local/bin/perl -w
# Extract oligos from a sequence and analyze %GC
$seqFile = "mySeq.fa";  # input sequence
mer = 35;
                           # length of oligo to make
\$start = 5;
                           # nt to start oligos
\$end = 11;
                           # nt to stop oligos
# Get continuous sequence from sequence file
open (SEQ, $seqFile ) || die "cannot open $seqFile: $!";
@seq = \langle SEO \rangle;
                           # make array of lines
$defline = $seq[0];
                           # get defline
seq[0] = "";
                           # delete defline
$seq = join ("", @seq);
                                  # join($glue, @list)
s = x/\s*//q;
                                  # delete whitespace
```

oligos.pl: part 2

```
$seqLength = length ($seq);
print "Oligos ($mer mers) for $defline
  ($seqLength nt) and % GC content\n";
# Beware of OBOB
for (\$i = \$start - 1; \$i < \$end - 1; \$i++)
  # oligo = substr(seq, start, length);
  $ = substr($seq, $i, $mer);
  st = si + 1;
  $numGC = tr/GC//;
                                    # count GCs
  pcGC = 100 * pumGC / per;
                               # find %GC
  print "$nt\t$ \t$pcGC\n";
```

Summary

- Input/output
- Variables
- Functions (scalars and arrays)
- Control structures
- Comparisons
- Sample scripts:
 - patscan_batch.pl
 - oligos.pl

For more information, books:

- Learning Perl (Schwartz et al.) O'Reilly
- The Perl CD Bookshelf O'Reilly
- Beginning Perl for Bioinformatics Tisdall
- Using Perl to Facilitate Biological Analysis (Stein) in *Bioinformatics* (Baxevanis & Ouellette)

AND several good web sites (see course page)

Demo scripts on the web site

- hey.pl
- input and output options
- patscan_batch.pl
- rev_comp.pl
- oligos.pl
- parse_genbank.pl

Exercises

 Retrieving and aligning a list of humanmouse orthologs

 Retrieving a set of genes encoding growth factors, extracting their proximal promoters, and analyzing them.