Appendix E: Sample Code – Matlab

Programs

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// Beginnning of Program: DataCompaction_MDWT.m //
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/* Matlab Program for Algorithm to perform Data Reduction
using Wavelet Transforms */
   Program to transform data using Haar Wavelet
Transforms on already reduced nucleotide sequence using
2-bit Binary Indicator %%
tStart=tic;
lookup seqn = cast(csvread('SRR000675.txt'), 'uint8'); %%
Sample Read of SRR000675
%lookup seqn = cast(csvread('Cnt46 EC.txt'), 'uint8'); %%
Contig 46 of E.Coli K12 Strain
%lookup seqn = cast(csvread('ch7 HS 1.txt'), 'uint8'); %%
Chr7 of Homo Sapien
wavelettype = 'haar';
maxdec = 4;
tStart wavelets = tic;
   for declev = 1:maxdec
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```
pdeclev = sprintf('dec %d', declev);
        dec
mdwtdec('r',lookup_seqn,declev,wavelettype);
    end;
tElapsed_wavelets = toc(tStart_wavelets);
    s_len4(declev) = cast(zeros(), 'uint16');
    for count = 1:declev
        s_{len4}(count) = cast(length(dec.cd{1,4})),
'uint16');
    end;
     rec = mdwtrec(dec);
     sizedec4 = dec.cd{1,4};
%%% Draw Plots %%%
    subplotm=3;
    subplotn=2;
    subplotp=0;
     subplotp=subplotp+1;
     subplot (subplotm, subplotn, subplotp);
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plot(lookup seqn); title('original seqn');
     xlabel('time'); ylabel('scale');
     subplotp=subplotp+1;
     subplot (subplotm, subplotn, subplotp);
     plot(rec); title('reconstructed seqn');
     xlabel('time'); ylabel('scale');
    for declev = 1:maxdec
     pdeclev = sprintf('level %d decomposition', declev);
     subplotp=subplotp+1;
     subplot (subplotm, subplotn, subplotp);
     plot(dec.cd{1, declev});
     title(pdeclev); xlabel('time'); ylabel('scale');
    end;
  tElapsed = toc(tStart);
// End of Matlab
Program:CompressSequence_DataCompaction_DWT.m //
//// Beginning of Program SortedDuplicateReads.m ////
%% Program to Compress (& Decompress) the sequences using
%% functions like mdwtdec & mdwtrec and
%% find the duplicate reads.
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%% The Transformed signals are compared after sorting
them based on lengths and comparing only
%% those sequences whose length are same
tStart=tic;
tStart dup = tic;
tStart wavelets = tic;
%% Metagenomics SRA data used in the paper, for finding
duplicate reads
%% To test the program for various data set, uncomment
the following list
%% of filenames one after the other and run the program
in Matlab R2009a release.
%% Various results can be checked from workspace of
matlab
%Afasta_file_reads = 'SRR000907.fna';
%Afasta file reads = 'SRR001669.fna';
%Afasta file reads = 'SRR001670.fna';
%Afasta file reads = 'SRR077225.fna';
%Afasta file reads = 'SRR000905.fna';
%Afasta_file_reads = 'SRR000906.fna';
%Afasta file reads = 'SRR000675.fna';
%Afasta file reads = 'SRR001663.fna';
Afasta file reads = 'SRR065619.fna'; % Small fasta file
of bacillus, to be used for testing the program & quick
results
[header, sequence] = fastaread(Afasta file reads);
[rows columns] = size(sequence);
seq len=columns;
wavelettype = 'haar';
total base count = 0;
for count = 1:seq len
        dbs1 = sequence(count);
        ldbs1 = cast(length(dbs1), 'uint16');
%% To generate frequency string of the given sequence
using, Electron-Ion Interaction Pseudo Potential of
Nucleotides
        fqdbs1 = eiip submitted(dbs1);
        lseq{count} = ldbs1;
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```
total base count = cast(total base count,
'uint32') + cast(ldbs1, 'uint32');
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%% Compression using multiple transforms:
mdwtdec/mdwtrec %%
        maxdec = 4;
%% Performing each level of transform and storing in an
array of decompositions
        for declev = 1:maxdec
            pdeclev = sprintf('dec %d', declev);
            dec(count) =
mdwtdec('r',fqdbs1,declev,wavelettype);
        end;
%% Performing reconstruction after all 4 levels of
decomposition is completed &
%% plotting it to verify with the original sequence's
plot for exact
%% reconstruction - a proof that wavelet transform based
data reduction is
%% appropriate for doing comparisions for recognizing
duplicate reads
         rec = mdwtrec(dec(count));
         recseq{count} = rec;
end;
tElapsed wavelets = toc(tStart wavelets);
%% To find the total length of decomposed sequence after
each level of transform is performed. %%
    s len4(seq len) = cast(zeros(), 'uint16');
    for count = 1:seq len
        s len4(count) = cast(length(dec(count).cd{1,4}),
'uint16');
    end;
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   This block identifies the duplicate reads, by
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checking those reads
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%% whose decomposed length is same, and if the lengths
are same then, only
%% compare decomposed values to each other for
similarity check.
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    tStart dup actual = tic;
    flag = cast(zeros(size(lseq)), 'logical');
    i = 0;
    for row = 1: (seq len-1)
        if (flag(row) == 1) %% To check if the row is
already identified as a duplicate of earlier row
            continue;
        else
        j = 1;
        for col = (row + 1):seq len
            if ( seq len < 255)
                cast size = 'uint8';
            elseif (seq len >= 255 && seq len < 65535)
                cast size = 'uint16';
            elseif ( seq len >= 65535 )
                cast size = 'uint32';
            end;
             if (s_{en4}(row) == s_{en4}(col))
                if (dec(1, row).cd\{1, 4\} ==
dec(1,col).cd\{1, 4\})
                     if (j == 1)
                        i = i + 1;
                        dup read(i,j) =
cast(row, cast size);
                         j = j + 1;
                         dup read(i,j) =
cast(col, cast size);
                         j = j + 1;
                         flag(row) = 1; %% To set the
status, if the row has identified any duplicate
                         flag(col) = 1; %% To set the
status to already compared read, if the row is already
identified as a duplicate of earlier row do not check for
its duplicate again
                     else
                         dup read(i,j) =
cast(col,cast size);
                         j = j + 1;
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```
flag(col) = 1;
                    end;
                end;
            end;
        end;
        end;
    end;
tElapsed dup actual = toc(tStart dup actual);
tElapsed dup = toc(tStart_dup);
tStart stats = tic;
%% The following block of code generates the statistical
and more organized and user-friendly output. It is not
compulsory for the code to identify duplicate reads.
%% Generating the Unique list of duplicate reads, unlike
above block which
%% is generating the entire list. This block is based on
the output of the above block.
%% So, do not remove or alter the above block
%% It is also Calculating the total number of duplicate
reads and
%% its percentage ie ratio of duplicate reads to the
total number of reads
%% And the redundant total no. of reads and total number
of bases
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%% To find the total max no. of duplicates
    [dup rows, dup cols] = size(dup read);
i=1;
j=1;
total unique dup read = 0;
total dup read count = 0;
total unique dup read count bases = 0;
total redundant dup read count bases = 0;
    for i=1:dup rows
        if (dup read(i, 1) \sim 0)
            total unique dup read = total unique dup read
+1;
        for j=2:dup_cols
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```
if (dup read(i,j) \sim 0)
                read no = cast(dup read(i,1), cast_size);
                dup read details(i,1) =
cast(read no, cast size); %% finds the read no. of a
unique duplicate read
                dup read details(i,2) =
cast(lseq{1,read no},cast size); %% finds the no. of bp
in a read
                dup read details(i,3) =
cast(j,cast size);
                    %% finds total no. of copies of
this duplicate read
                dup read details(i,4) =
cast((dup read details(i,2) * (dup read details(i,3) - 1)
),cast_size); %% finds total redundant bases of this read
            end;
        end;
        total dup read count = total dup read count +
dup read details(i,3);
        total unique dup read count bases =
total_unique_dup_read_count_bases +
cast(dup read details(i,2),'double');
        total redundant dup read count bases =
total redundant dup read count bases +
cast(dup read details(i,4),'double');
        end;
    end;
    total redundant dup read = total dup read count -
total unique dup read;
    total percentage redundant read =
(cast(total redundant dup read, 'double') * 100)/
seq len;
    total percentage redundant bases =
(cast(total_redundant_dup_read_count_bases,'double') *
100)/ cast(total base count, 'double');
    total dup read count bases =
cast(total unique dup read count bases, 'double') +
cast(total_redundant_dup_read_count_bases, 'double');
    total percentage dup read bases =
(cast(total_dup_read_count_bases, 'double') * 100) /
cast(total_base count, 'double');
    total percentage dup read =
(cast(total_dup_read count, 'double') * 100)/ seq len;
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tElapsed_stats = toc(tStart_stats);
tElapsed = toc(tStart);
```

//// End of Program : SortedDuplicateReads.m ////

FindZeroIndex.m

/// Beginning of Program: FindZeroIndex.m /// /* Matlab Function to find Indices of Zeroes in the given signal. This function finds the positions or indices of zero values in a given signal, so that it can be used to detect Short Tandem Repeat Regions in a given DNA sequence. */ %% Function to find indices of zeros in a given signal %% function [findzeroindx d_rle] = findzeroindex(signal, thresholdval) extendsignal = [1, signal, 1]; convextendsignal = conv(extendsignal, ones(1,thresholdval)); tempsignal = double(convextendsignal == 0); tempconv = conv(tempsignal, ones(1,thresholdval)); findzeroindx = find(tempconv) - thresholdval; d rle = runlengthencoding(tempsignal == 0); end;

//// End of Program: FindZeroIndex.m ////