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```
%{
Yonatan Carver
ECES 352 - Lab 3

%}
clear; clc; close all
```



```
%{
1) 31 results for "hbb"
3) 280 results for "hbb AND human[orgn]"
%}
```

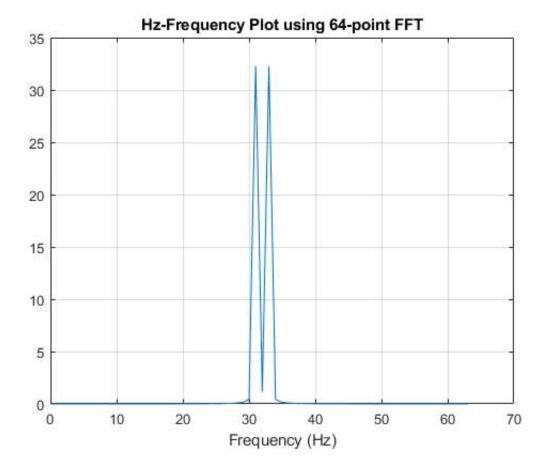

(1)

```
N = 64;
fft_x = abs(fftshift(fft(x, N))); % N-point FFT
```

(3)

range = 0 : 1 : 63;

```
range = linspace(0, 63, N);
figure
plot(range, fft_x)
grid on
xlabel('Frequency (Hz)')
title(['Hz-Frequency Plot using ', num2str(N), '-point FFT'])
% [yy, xx] = findpeaks(fft_x);
% frequency spikes @ f = 26 & 38
```

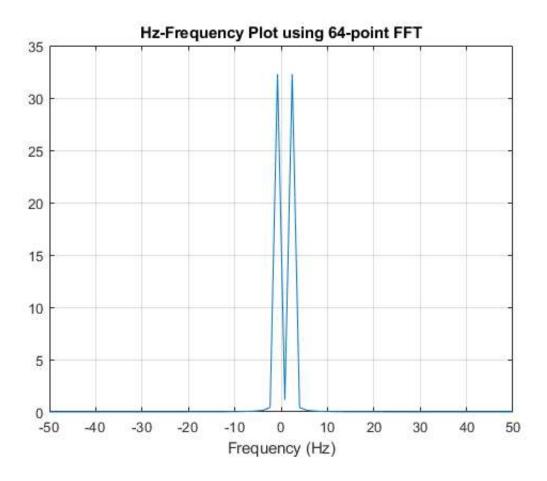


(4)

```
range = linspace(-50, 50, N); % plot from -50 to 50 (100Hz sampling rate)

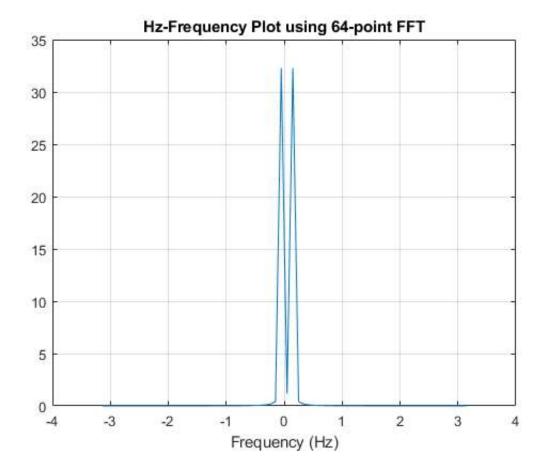
figure
plot(range, fft_x)
grid on
```

```
xlabel('Frequency (Hz)')
title(['Hz-Frequency Plot using ', num2str(N), '-point FFT'])
% frequency spikes @ -8.7302 & 10.3175
```

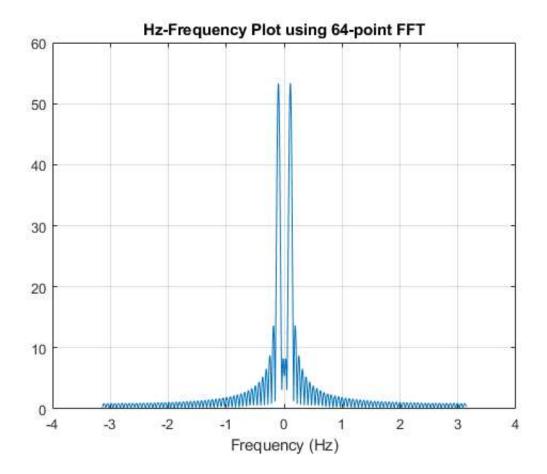


(6)

```
range = linspace(-pi, pi, N);
figure
plot(range, fft_x)
grid on
xlabel('Frequency (Hz)')
title(['Hz-Frequency Plot using ', num2str(N), '-point FFT'])
% frequency spikes @ -0.5484 & 0.6483
```



(8)




```
clear; clc; close all
hbb = genbankread('hbb_region_chr11.gb'); % load *.gb file as a struct
cds_indices = hbb.CDS.indices; % get range of indices of CDS
```

(1)

There are 5 CDS regions in the sequence length(hbb.CDS)

(2)

The length of hbb.CDS(1).translation is 147 length(hbb.CDS(1).translation)

(3)

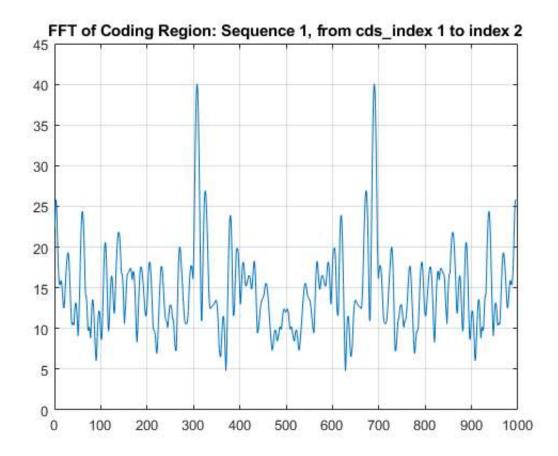
```
% use function "get_sequence_ft"
% function sequence_FT = get_sequence_ft(hbb, sequence_number)

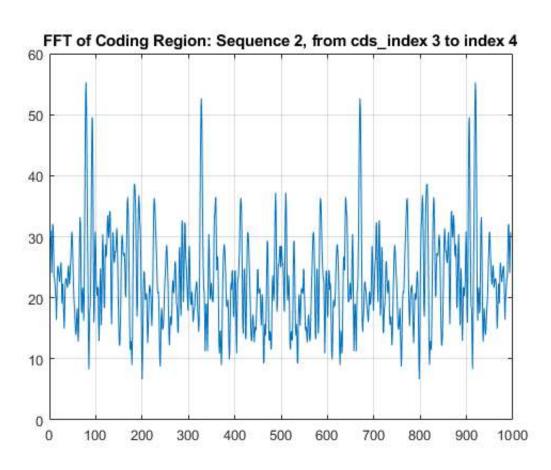
N = 1024;

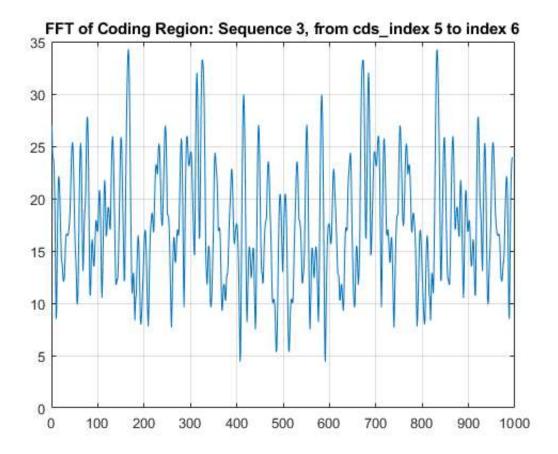
sequence1_ft = get_sequence_ft(hbb, 1, N);
sequence2_ft = get_sequence_ft(hbb, 2, N);
sequence3_ft = get_sequence_ft(hbb, 3, N);

figure
```

```
plot(sequence1 ft)
title('FFT of Coding Region: Sequence 1, from cds\_index 1 to index 2')
grid on
figure
plot(sequence2 ft)
title('FFT of Coding Region: Sequence 2, from cds\ index 3 to index 4')
grid on
figure
plot(sequence3 ft)
title('FFT of Coding Region: Sequence 3, from cds\ index 5 to index 6')
grid on
[y1, x1] = findpeaks(sequence1 ft);
[y2, x2] = findpeaks(sequence2 ft);
[y3, x3] = findpeaks(sequence3_ft);
응 {
signal: sequence1
a = sort(y1);
b = sort(y2);
c = sort(y3);
x1(find(y1 == a([end])))
232.4562
frequency peaks @ 308 & 690
x2(find(y2 == b([end])))
251.8894
frequency peaks @ 292 & 706
x2(find(y2 == b([end-2])))
224.3544
frequency peaks @ 92 & 906
x2(find(y2 == b([end-4])))
209.4380
frequency peaks @ 329 & 669
_____
x3(find(y3 == c([end])))
165.9027
frequency peaks @ 328 & 670
x3(find(y3 == c([end-2])))
142.8912
frequency peaks @ 164 & 834
x3(find(y3 == c([end-4])))
126.3875
frequency peaks @ 230 & 768
응 }
```





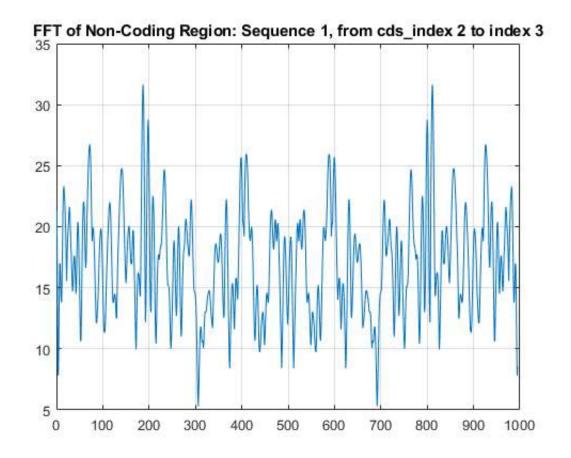


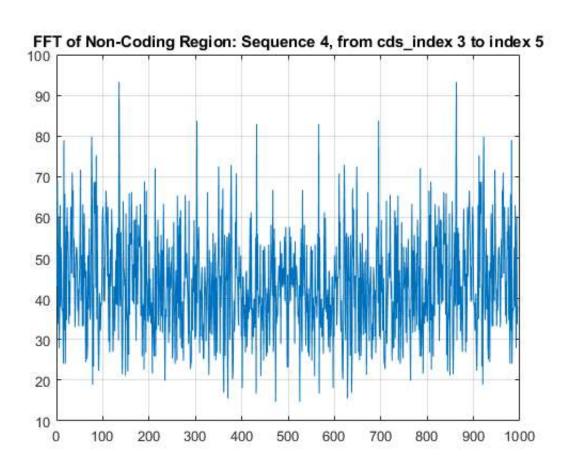
N = 1024;

```
sequence1a_ft = get_sequence_ft(hbb, '1a', N);
sequence2a_ft = get_sequence_ft(hbb, '2a', N);

figure
plot(sequence1a_ft)
title('FFT of Non-Coding Region: Sequence 1, from cds\_index 2 to index 3')
grid on

figure
plot(sequence2a_ft)
title('FFT of Non-Coding Region: Sequence 4, from cds\_index 3 to index 5')
grid on
```





```
DNA_SEQUENCE = hbb.Sequence;
WINDOW_LENGTH = 10;
NFFT = 1024;

% DNA_SEQUENCE = hbb.Sequence;
% WINDOW_LENGTH = 1000;
% NFFT = 1024;

output = threebasefreq_stft(DNA_SEQUENCE, WINDOW_LENGTH, NFFT);
plot(output)
```

