

- 1) My first approach to finding the *ori* of Salmonella Enterica will start with finding the genome's minimum skew in the genome.

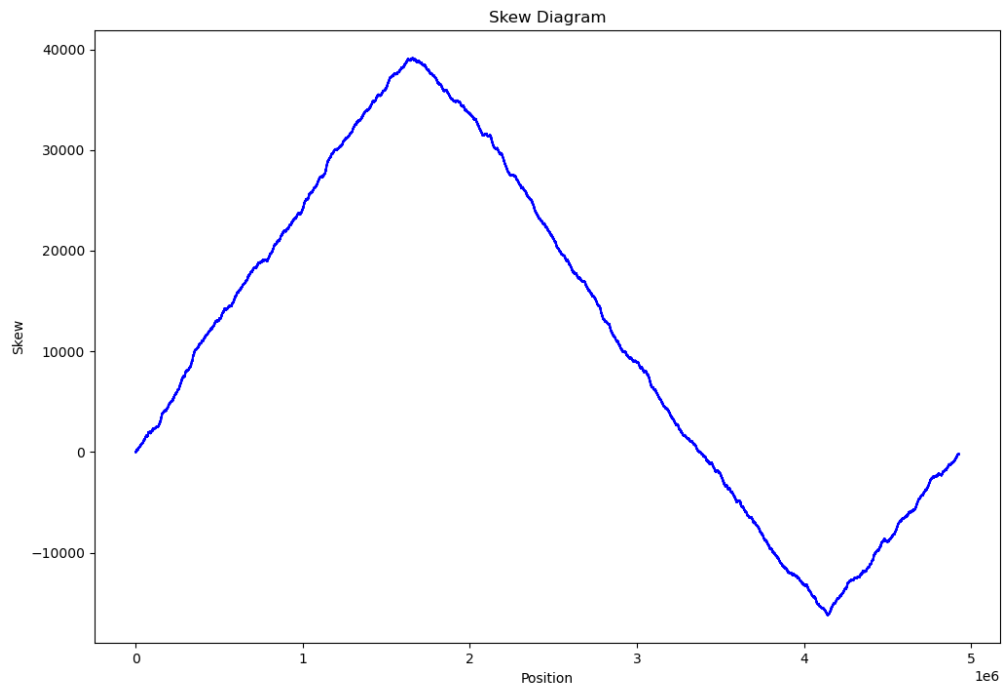
My output for the skew finding algorithm is as below:

```
hw1.py > generate_kmers
1  import matplotlib.pyplot as plt
2  import numpy as np
3
4
5  def find_minimum_skew(genome):
6      num_list = [0]
7      for i in range(len(genome)):
8          if genome[i] == 'C':
9              num_list.append(num_list[i]-1)
10         elif genome[i] == 'G':
11             num_list.append(num_list[i]+1)
12         else:
13             num_list.append(num_list[i])
14     min_value = min(num_list)
15     indices = [i for i, value in enumerate(num_list) if value == min_value]
16     print(indices)
17     return num_list
18
19
20
21 def most_frequent_kmer(text, k, d):
22     bases = ['A', 'T', 'C', 'G']
23     possible_kmers = generate_kmers(k, bases)
24     my_dict = {}
25     for kmer in possible_kmers:
26         f1 = len(approximate_occurrences_of_pattern(kmer, text, d))
27         f2 = len(approximate_occurrences_of_pattern(reverse_complement(kmer), text, d))
28         my_dict[kmer] = f1 + f2
29     max_val = max(my_dict.values())
30     result = ''
31     for key in my_dict:
32         if my_dict[key] == max_val:
33             result += ' '
34             result += key
35     print(result)
36
37
38 def generate_kmers(k, bases):
39     if k == 1:
40         return bases
41     small_kmers = generate_kmers(k-1, bases)
42     kmers = []
43     for kmer in small_kmers:
```

PROBLEMS OUTPUT DEBUG CONSOLE TERMINAL PORTS

```
gafur@gafur-HP-Spectre-x360-Convertible-13-ap0xxx:~/Documents/computational_biology$ /bin/python3 /home/gafur/Documents/computational_biology/hw1.py
[4142725, 4142727]
```

This is the skew diagram that I created for better understanding. I made the conclusion that the position of minimum skew is **4142727**.



- 2) After finding the minimum skew position, to make sure that it is the origin of replication, I checked the most frequent 9-mer with 1 mismatch and its reverse complements.
- I did this check in between positions [**4142725: 4142725+500**].
  - It turns out that the most frequent 9-mers are – **AACACGATC, AACCAGATC, GATCTGGTT, GATCGTGTT** (GATCTGGTT and GATCGTGTT are reverse complements of AACACGATC, AACCAGATC).
  - This is the screenshot of my output:

```
hw1.py > generate_kmers
55 def reverse_complement(pattern):
56     for i in range(len(pattern)-1, -1, -1):
57         if pattern[i] == 'A':
58             reverse_complement += 'T'
59         elif pattern[i] == 'C':
60             reverse_complement += 'G'
61         elif pattern[i] == 'G':
62             reverse_complement += 'C'
63         else:
64             reverse_complement += 'A'
65     return(reverse_complement)
66
67 def hamming_distance(str1, str2):
68     hamming_distance = 0
69     for i in range(len(str1)):
70         if str1[i] != str2[i]:
71             hamming_distance += 1
72     return hamming_distance
73
74 def main():
75     # Open the file in read mode and store its content in a string variable
76     with open("salmonella_enterica_sequence.fasta", "r", encoding="utf-8") as file:
77         salmonella_sequence = file.read()
78
79     skew_data = find_minimum_skew(salmonella_sequence)
80
81     x = np.arange(len(skew_data))
82     plt.figure(figsize=(12, 6))
83     plt.plot(x, skew_data, linestyle='-', marker='', color='b')
84     plt.xlabel("Position")
85     plt.ylabel("Skew")
86     plt.title("Skew Diagram")
87     plt.show()
88
89     most_frequent_kmer(salmonella_sequence[4142727:4142727+500], 9, 1)
90
91 # Ensures the script runs only when executed directly
92 if __name__ == "__main__":
93     main()
94
95
96
97
```

PROBLEMS OUTPUT DEBUG CONSOLE TERMINAL PORTS

```
gafur@gafur-HP-Spectre-x360-Convertible-13-ap0xxx:~/Documents/computational_biology$ /bin/python3 /home/gafur/Documents/computational_biology/hw1.py
[4142725, 4142727]
AACACGATC AACACGATC GATCTGGTT GATCTGGTT
gafur@gafur-HP-Spectre-x360-Convertible-13-ap0xxx:~/Documents/computational_biology$ /bin/python3 /home/gafur/Documents/computational_biology/lab2.py
```

- My algorithm starts with creating all possible 9-mers and then uses dictionary to find the most frequent ones. (I did not use the neighbor technique)
- 3) In conclusion, I identified the positions **[4142725: 4142725+500]** as DnaA box because, first, the position of minimum skew starts from there. Secondly, there are 2 most frequent 9-mers in between these positions.
  - 4) One drawback of my approach is that it requires more computation because it does not utilize d-neighborhood technique.