

- ✓ Aug 21: Comprehensive Exam 10:30am
- ✓ Aug 26: make new list of dates for goals
- ✓ Sep 7: Write Up methods for clustering testing and add to substitutions paper
- Sep 9: Gene expression data for the inversions project
- Sep 14: Have all clustering testing complete for all bacteria
- Sep 14: Compile notes from comps papers into one document
- Sep 30: New intro for Substitution paper
- Sep 15-28: Apply for NSERC (if applicable)
- Oct 3: NSERC Due
- Oct 5-12: Apply for Mac Scholarships and Awards
- Oct 31: Write out methods for gene expression paper
- Sep 9: Think about/compile list of inversions in *E. coli* for new paper
- Nov 15: Think about how to better look at the COG data
- Nov 25: Complete any extra analysis needed for Substitution paper
- Dec 4: Mac Scholarships and Awards Due
- Dec 1: Write out COG methods
- Dec 15: Gather papers for COG paper intro
- Dec 15: Implement COG stuff

Last Week

I was working on getting a good chunk of the clustering testing done. There is still a bit more to complete but it should be done by the end of next week!

I also wrote out detailed methods for the clustering testing and added it to the substitutions paper write-up. I also working on finding more gene expression data for the inversions and gene expression project.

This Week

I would like to continue to work on the clustering testing. In between having this finished I would like to keep looking for gene expression data for *Escherichia coli* and have this finished by the end of the week.

Next Week

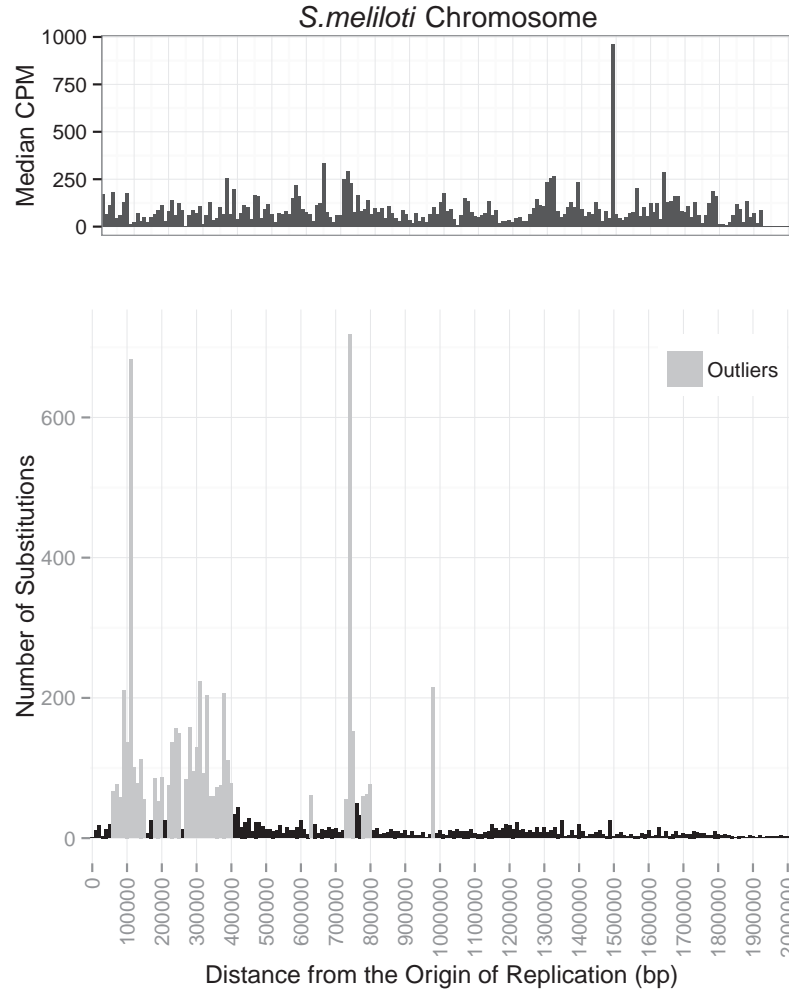
I would like to compile all my notes from comps into one file so that I can use this to update my substitutions paper intro. I would like to complete the remainder of the clustering testing.

| Position Difference | <i>E. coli</i> Chromosome | <i>B. subtilis</i> Chromosome | <i>Streptomyces</i> Chromosome | <i>S. meliloti</i> Chromosome | <i>S. meliloti</i> pSymA | <i>S. meliloti</i> pSymB |
|---------------------|-----------------------------|-------------------------------|--------------------------------|-------------------------------|----------------------------|----------------------------|
| 1bp | -1.394 $\times 10^{-7}$ *** | -2.538 $\times 10^{-8}$ *** | 1.736 $\times 10^{-8}$ ** | -1.541 $\times 10^{-6}$ ** | -9.130 $\times 10^{-7}$ ** | 2.488 $\times 10^{-7}$ *** |
| 10bp | -1.394 $\times 10^{-7}$ *** | -2.518 $\times 10^{-8}$ *** | -4.484 $\times 10^{-9}$ *** | -1.627 $\times 10^{-6}$ *** | -9.13 $\times 10^{-7}$ *** | 3.487 $\times 10^{-7}$ *** |
| 100bp | -1.764 $\times 10^{-7}$ *** | -1.417 $\times 10^{-8}$ *** | | | | |

Table 1: Logistic regression analysis of the number of substitutions along the genome of the respective bacteria replicons to test position differences. All results are marked with significance codes as followed: $< 0.001 = '***'$, $0.001 < 0.01 = '**'$, $0.01 < 0.05 = '*'$, $0.05 < 0.1 = '.'$, $> 0.1 = ''$. Logistic regression was calculated after the positions in the genome were determined to be the same at each position difference listed in the first column.

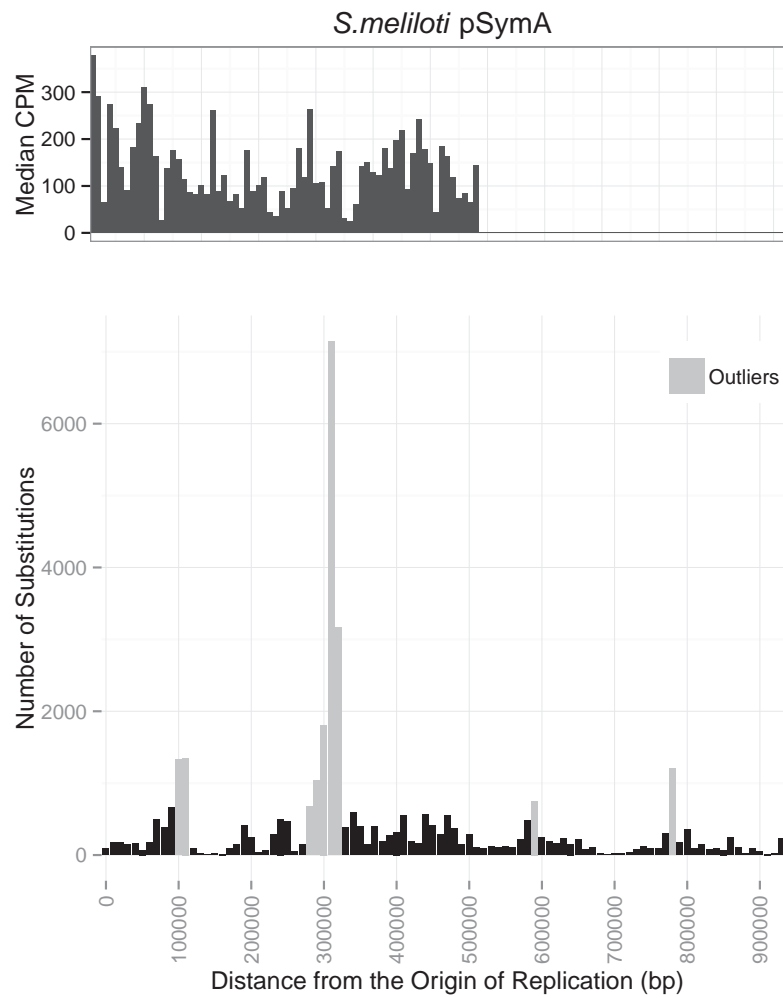
| Bacteria and Replicon | Coefficient Estimate | Standard Error | P-value |
|--------------------------------|------------------------|-----------------------|-----------------------|
| <i>E. coli</i> Chromosome | -6.41 $\times 10^{-5}$ | 1.65 $\times 10^{-5}$ | 1.1 $\times 10^{-4}$ |
| <i>B. subtilis</i> Chromosome | -9.9 $\times 10^{-5}$ | 2.18 $\times 10^{-5}$ | 6 $\times 10^{-6}$ |
| <i>Streptomyces</i> Chromosome | -1.5 $\times 10^{-6}$ | 1.4 $\times 10^{-7}$ | $< 2 \times 10^{-16}$ |
| <i>S. meliloti</i> Chromosome | 3.19 $\times 10^{-5}$ | 3.57 $\times 10^{-5}$ | 3.7 $\times 10^{-1}$ |
| <i>S. meliloti</i> pSymA | -5.36 $\times 10^{-5}$ | 6.34 $\times 10^{-4}$ | 9.33 $\times 10^{-1}$ |
| <i>S. meliloti</i> pSymB | 5.05 $\times 10^{-4}$ | 2.6 $\times 10^{-4}$ | 5.3 $\times 10^{-2}$ |

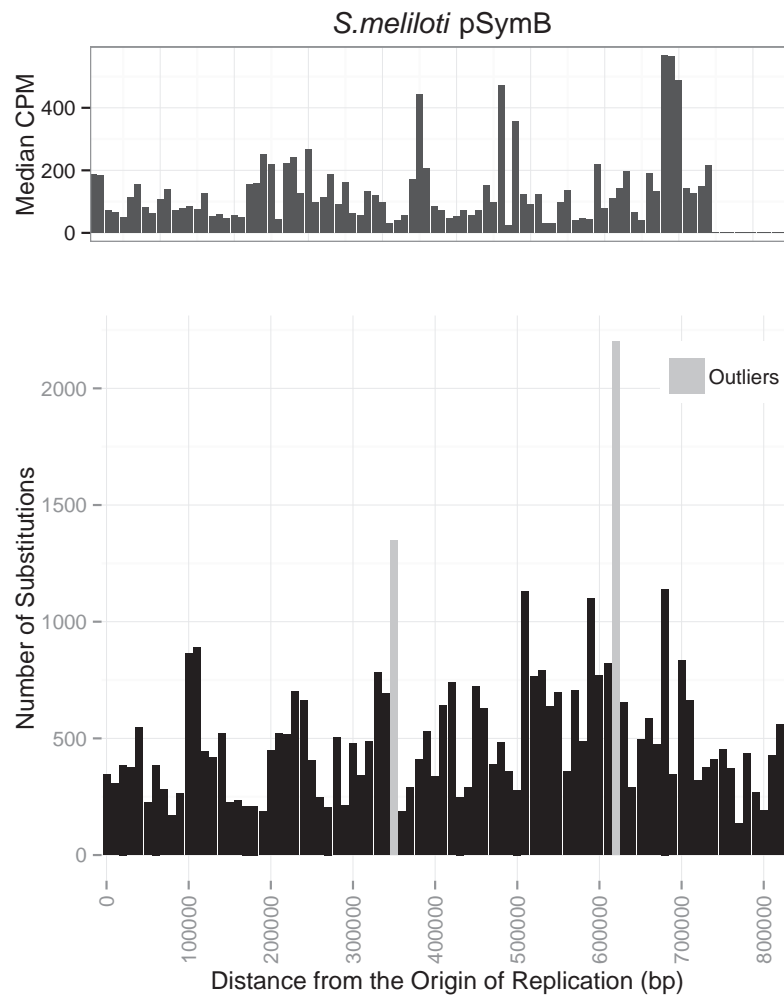
Table 2: Linear regression analysis of the median counts per million expression data along the genome of the respective bacteria replicons. Grey coloured boxes indicate statistically significant results at the 0.5 significance level. Linear regression was calculated after the origin of replication was moved to the beginning of the genome and all subsequent positions were scaled around the origin accounting for bidirectionality of replication.

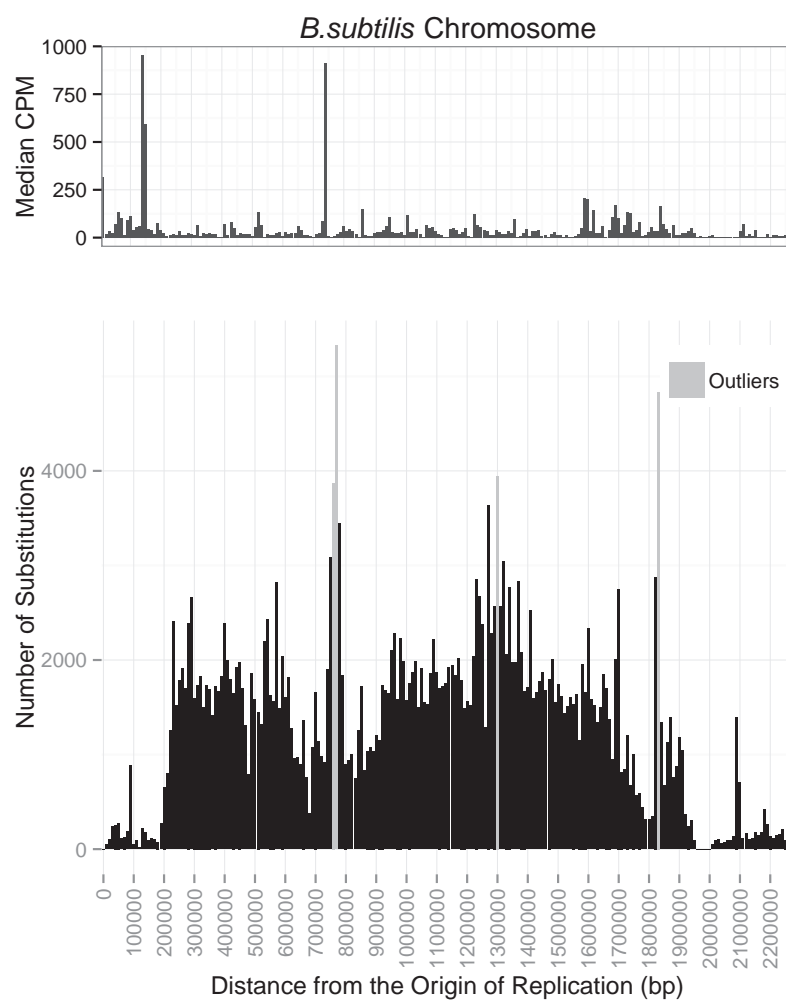


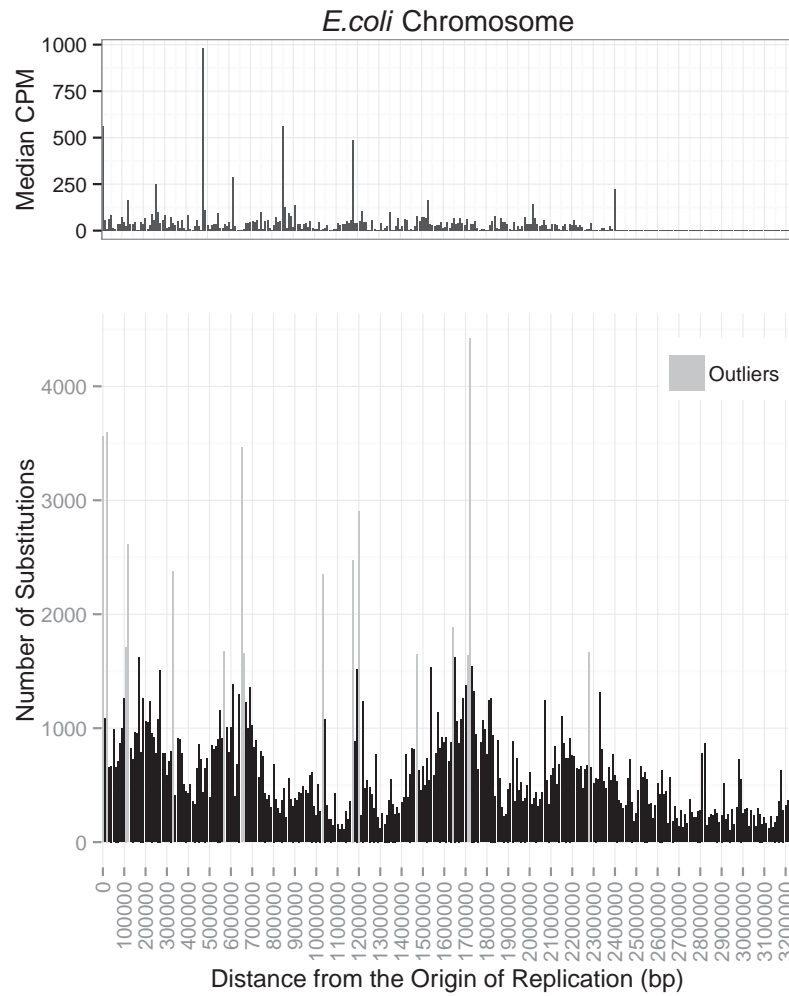
| Bacteria and Replicon | Coefficient Estimate | Standard Error | P-value |
|--------------------------------|-------------------------|-------------------------|-------------------------|
| <i>E. coli</i> Chromosome | -1.394×10^{-7} | 2.425×10^{-9} | $< 2 \times 10^{-16}$ |
| <i>B. subtilis</i> Chromosome | -1.265×10^{-8} | 1.562×10^{-9} | 5.430×10^{-16} |
| <i>Streptomyces</i> Chromosome | 1.736×10^{-8} | 7.231×10^{-10} | $< 2 \times 10^{-16}$ |
| <i>S. meliloti</i> Chromosome | -1.541×10^{-6} | 3.042×10^{-8} | $< 2 \times 10^{-16}$ |
| <i>S. meliloti</i> pSymA | -9.130×10^{-7} | 1.975×10^{-8} | $< 2 \times 10^{-16}$ |
| <i>S. meliloti</i> pSymB | 2.488×10^{-7} | 1.964×10^{-8} | $< 2 \times 10^{-16}$ |

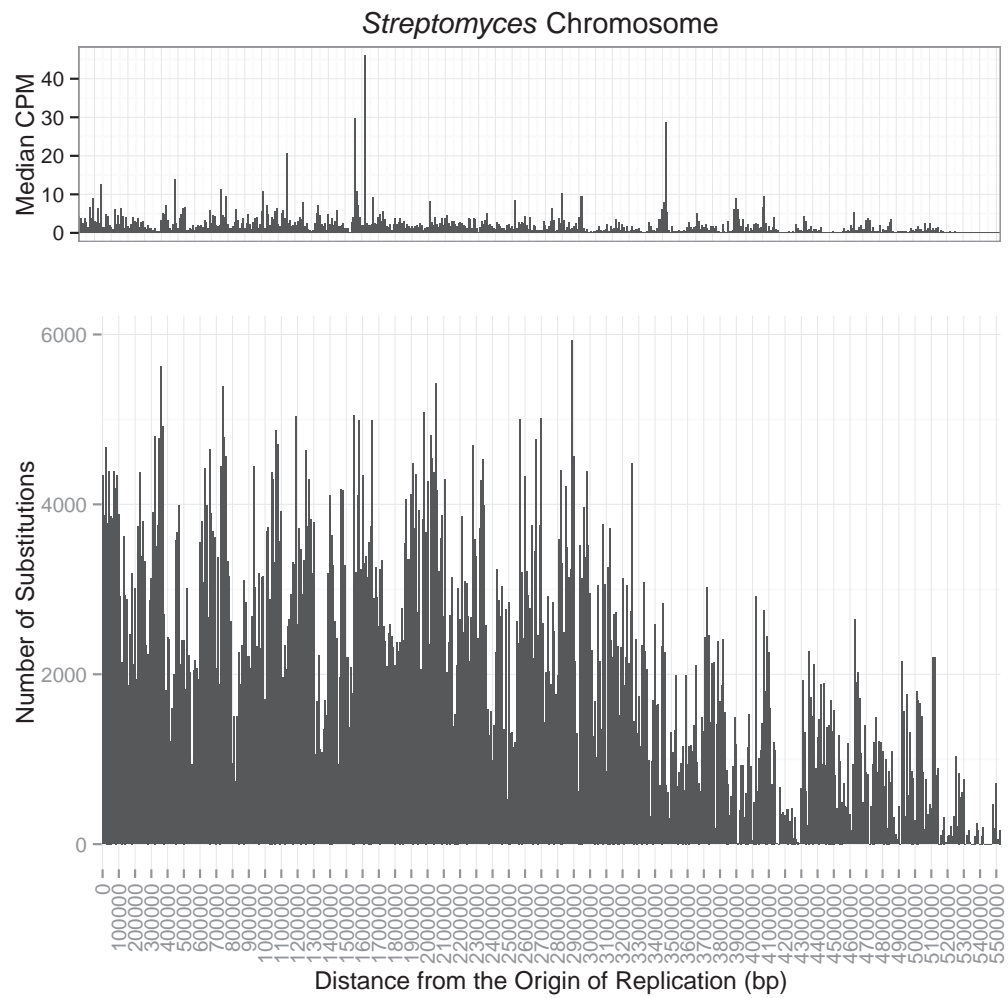
Table 3: Logistic regression analysis of the number of substitutions along the genome of the respective bacteria replicons. Grey coloured boxes indicate a negative logistic regression coefficient estimate. All results are statistically significant. Logistic regression was calculated after the origin of replication was moved to the beginning of the genome and all subsequent positions were scaled around the origin accounting for bidirectionality of replication.











| Origin Location | <i>E. coli</i> Chromosome | <i>B. subtilis</i> Chromosome | <i>Streptomyces</i> Chromosome | <i>S. meliloti</i> Chromosome | <i>S. meliloti</i> pSymA | <i>S. meliloti</i> pSymB |
|-------------------|----------------------------|-------------------------------|--------------------------------|-------------------------------|----------------------------|----------------------------|
| Moved 100kb Left | $-1.445 \times 10^{-7***}$ | $4.374 \times 10^{-9*}$ | $6.909 \times 10^{-9***}$ | $-1.316 \times 10^{-6***}$ | $-1.058 \times 10^{-6***}$ | $-2.009 \times 10^{-7***}$ |
| Moved 90kb Left | $-1.544 \times 10^{-7***}$ | $-1.036 \times 10^{-7***}$ | $5.677 \times 10^{-9***}$ | $-1.32 \times 10^{-6***}$ | $-1.246 \times 10^{-6***}$ | $-1.357 \times 10^{-7***}$ |
| Moved 80kb Left | $-1.65 \times 10^{-7***}$ | $-1.072 \times 10^{-7***}$ | $8.11 \times 10^{-9***}$ | $-1.338 \times 10^{-6***}$ | $-1.398 \times 10^{-6***}$ | $-6.57 \times 10^{-8***}$ |
| Moved 70kb Left | $-1.667 \times 10^{-7***}$ | $-1.102 \times 10^{-7***}$ | $6.716 \times 10^{-9***}$ | $-1.363 \times 10^{-6***}$ | $-1.405 \times 10^{-6***}$ | 9.83×10^{-8} |
| Moved 60kb Left | $-1.64 \times 10^{-7***}$ | $-1.19 \times 10^{-7***}$ | $8.7 \times 10^{-9***}$ | $-1.324 \times 10^{-6***}$ | $-1.394 \times 10^{-6***}$ | $1.129 \times 10^{-7***}$ |
| Moved 50kb Left | $-1.446 \times 10^{-7***}$ | $-1.211 \times 10^{-7***}$ | $1.045 \times 10^{-8***}$ | $-1.36 \times 10^{-6***}$ | $-1.403 \times 10^{-6***}$ | $1.521 \times 10^{-7***}$ |
| Moved 40kb Left | $-1.4 \times 10^{-7***}$ | $-1.299 \times 10^{-7***}$ | $1.214 \times 10^{-8***}$ | $-1.255 \times 10^{-6***}$ | $-1.422 \times 10^{-6***}$ | $1.543 \times 10^{-7***}$ |
| Moved 30kb Left | $-1.498 \times 10^{-7***}$ | $-1.292 \times 10^{-7***}$ | $1.24 \times 10^{-8***}$ | $-1.26 \times 10^{-6***}$ | $-1.392 \times 10^{-6***}$ | $1.63 \times 10^{-7***}$ |
| Moved 20kb Left | $-1.51 \times 10^{-7***}$ | $-1.1 \times 10^{-7***}$ | $1.395 \times 10^{-8***}$ | $-1.525 \times 10^{-6***}$ | $-1.412 \times 10^{-6***}$ | $1.603 \times 10^{-7***}$ |
| Moved 10kb Left | $-1.262 \times 10^{-7***}$ | -2.602×10^{-9} | $1.563 \times 10^{-8***}$ | $-1.599 \times 10^{-6***}$ | $-9.499 \times 10^{-7***}$ | $2.973 \times 10^{-7***}$ |
| Moved 10kb Right | $-1.305 \times 10^{-7***}$ | $-2.045 \times 10^{-8***}$ | $1.578 \times 10^{-8***}$ | $1.614 \times 10^{-6***}$ | $-1.026 \times 10^{-6***}$ | $3.505 \times 10^{-7***}$ |
| Moved 20kb Right | $-1.454 \times 10^{-7***}$ | $-1.006 \times 10^{-7***}$ | $1.903 \times 10^{-8***}$ | $-1.634 \times 10^{-6***}$ | $-1.475 \times 10^{-6***}$ | $1.649 \times 10^{-7***}$ |
| Moved 30kb Right | $-1.548 \times 10^{-7***}$ | $-8.596 \times 10^{-8***}$ | $2.046 \times 10^{-8***}$ | $-1.698 \times 10^{-6***}$ | $-1.417 \times 10^{-6***}$ | $1.526 \times 10^{-7***}$ |
| Moved 40kb Right | $-1.632 \times 10^{-7***}$ | $-8.378 \times 10^{-8***}$ | $2.125 \times 10^{-8***}$ | $-1.719 \times 10^{-6***}$ | $-1.367 \times 10^{-6***}$ | $1.589 \times 10^{-7***}$ |
| Moved 50kb Right | $-1.856 \times 10^{-7***}$ | $-7.879 \times 10^{-8***}$ | $1.957 \times 10^{-8***}$ | $-1.735 \times 10^{-6***}$ | $-1.277 \times 10^{-6***}$ | $1.654 \times 10^{-7***}$ |
| Moved 60kb Right | $-1.91 \times 10^{-7***}$ | $-6.98 \times 10^{-8***}$ | $1.974 \times 10^{-8***}$ | $-1.788 \times 10^{-6***}$ | $-1.169 \times 10^{-6***}$ | $1.645 \times 10^{-7***}$ |
| Moved 70kb Right | $-1.892 \times 10^{-7***}$ | $-6.634 \times 10^{-8***}$ | $1.934 \times 10^{-8***}$ | $-1.854 \times 10^{-6***}$ | $-1.059 \times 10^{-6***}$ | $1.843 \times 10^{-7***}$ |
| Moved 80kb Right | $-1.879 \times 10^{-7**}$ | $-5.814 \times 10^{-8***}$ | $2.313 \times 10^{-8***}$ | $-1.891 \times 10^{-6***}$ | $-9.07 \times 10^{-7***}$ | $1.90 \times 10^{-7***}$ |
| Moved 90kb Right | $-1.862 \times 10^{-7***}$ | $-4.314 \times 10^{-8***}$ | $2.304 \times 10^{-8***}$ | $-1.865 \times 10^{-6***}$ | $-7.171 \times 10^{-7***}$ | $2.415 \times 10^{-7***}$ |
| Moved 100kb Right | $-1.799 \times 10^{-7***}$ | $-2.597 \times 10^{-8***}$ | $1.945 \times 10^{-8***}$ | $-1.525 \times 10^{-6***}$ | $-6.572 \times 10^{-7***}$ | $3.095 \times 10^{-7***}$ |

Table 4: Logistic regression analysis of the number of substitutions along the genome of the respective bacteria replicons. All results are marked with significance codes as followed: $< 0.001 = '***'$, $0.001 < 0.01 = '**'$, $0.01 < 0.05 = '*'$, $0.05 < 0.1 = '.'$, $> 0.1 = ''$. Logistic regression was calculated after the origin of replication was moved to the beginning of the genome and all subsequent positions were scaled around the origin accounting for bidirectionality of replication.