

X Jan 6: Write up methods for COG paper

✓ May 7: Revise summer goals if accepted for Chicago Conference

✓ May 11: Find gene expression papers for bacteria and specific bacteria, printed

May 25: Read above papers and make notes (one a day?)

May 25: Think about/compile list of inversions in *E. coli* for new paper

June 8: Gather gene expression data for the above mentioned *E. coli* strains

June 12-29: Have first draft of ISMB presentation done (and present for the lab)/ prepare for conference questions

July 5: Have final edits for ISMB presentation finished

July 6-13: ISB Chicago Conference

July 20: Have date booked for Comps

July 16 - August 31: Prepare for Comps

## Last Week

I spent last week reading 1-2 papers per day about gene expression and other related topics to compile notes for the gene expression paper. I have also began to think about the gene expression and inversion paper and how to obtain data for that.

## This Week

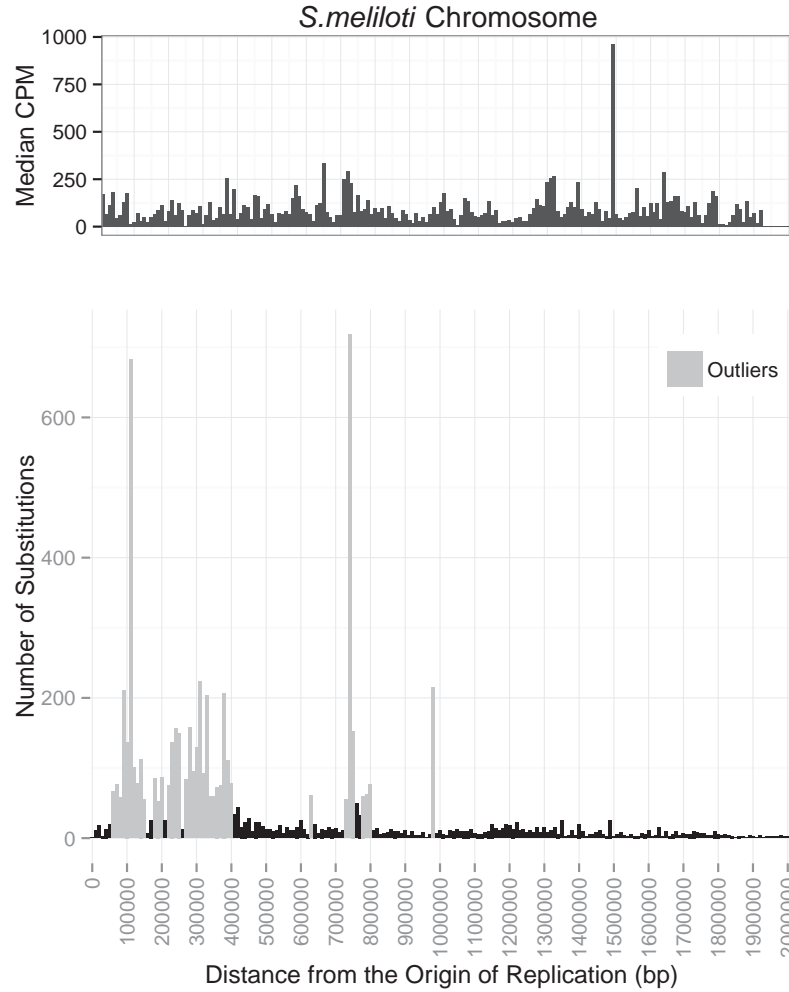
I will continue to be reading 1-2 papers per day as mentioned above and making notes. I would also like to run some tests on the ancestral position reconstruction code you sent me.

## Next Week

I plan on continuing to run test on the above mentioned code and gathering gene expression data for the inversion paper.

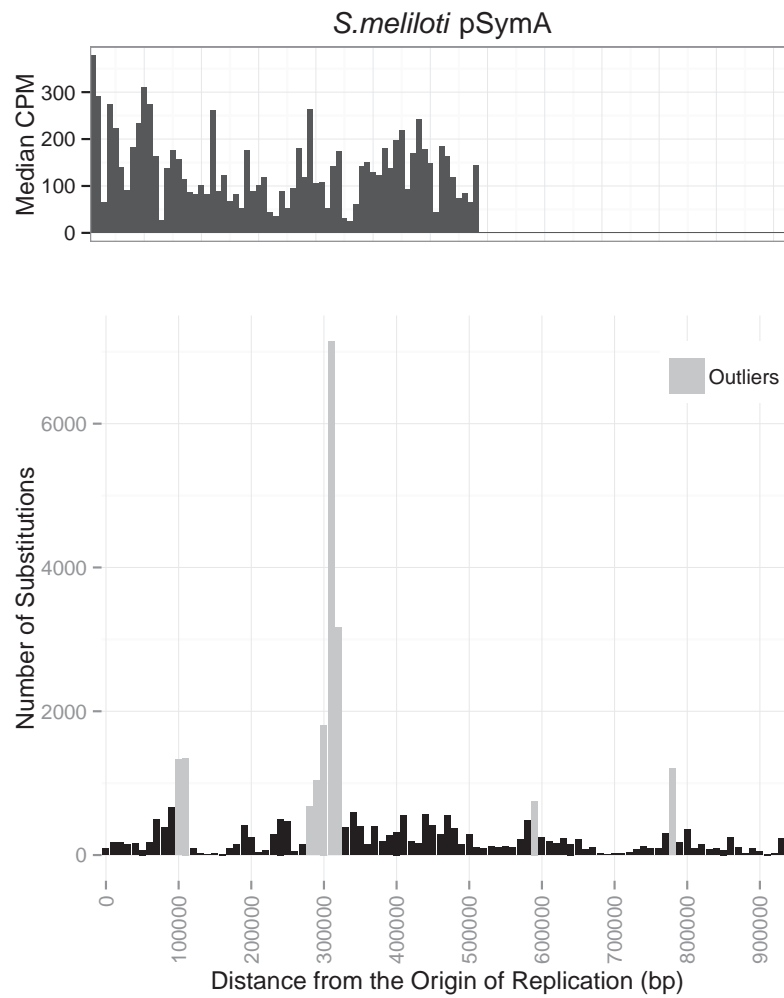
Bacteria and Replicon	Coefficient Estimate	Standard Error	P-value
<i>E. coli</i> Chromosome	$-6.41 \times 10^{-5}$	$1.65 \times 10^{-5}$	$1.1 \times 10^{-4}$
<i>B. subtilis</i> Chromosome	$-9.9 \times 10^{-5}$	$2.18 \times 10^{-5}$	$6 \times 10^{-6}$
<i>Streptomyces</i> Chromosome	$-1.5 \times 10^{-6}$	$1.4 \times 10^{-7}$	$< 2 \times 10^{-16}$
<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	$5.05 \times 10^{-4}$	$2.6 \times 10^{-4}$	$5.3 \times 10^{-2}$

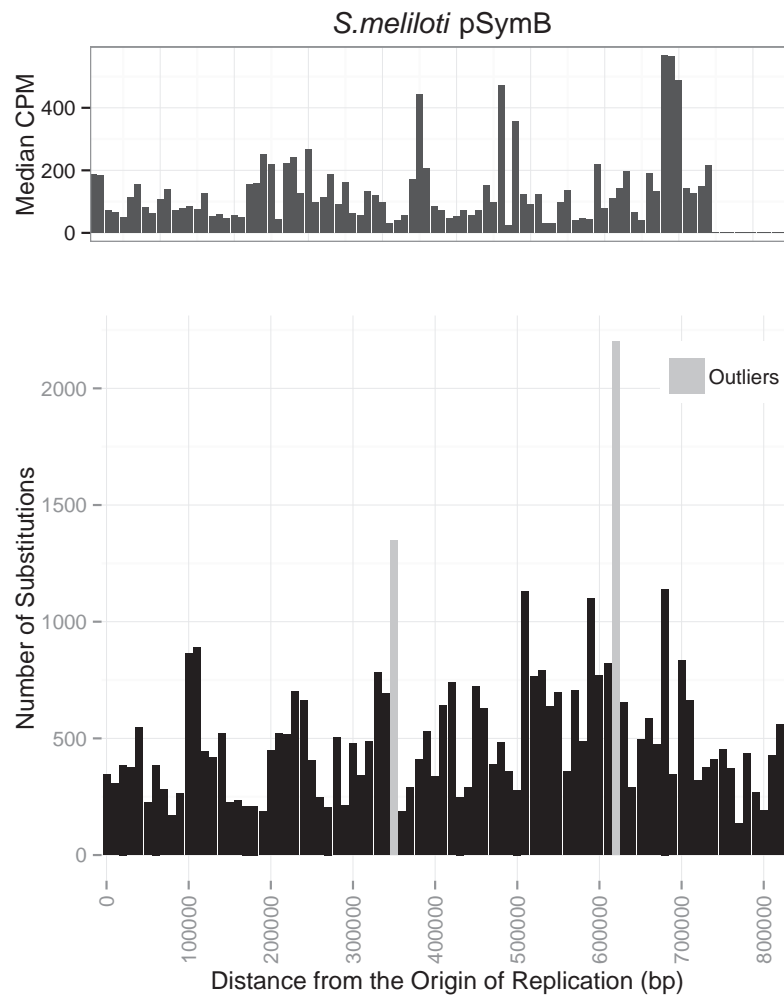
Table 1: Linear regression analysis of the median counts per million expression data along the genome of the respective bacteria replicons. Grey coloured boxes indicate statistically significant results at the 0.5 significance level. Linear regression was calculated after the origin of replication was moved to the beginning of the genome and all subsequent positions were scaled around the origin accounting for bidirectionality of replication.

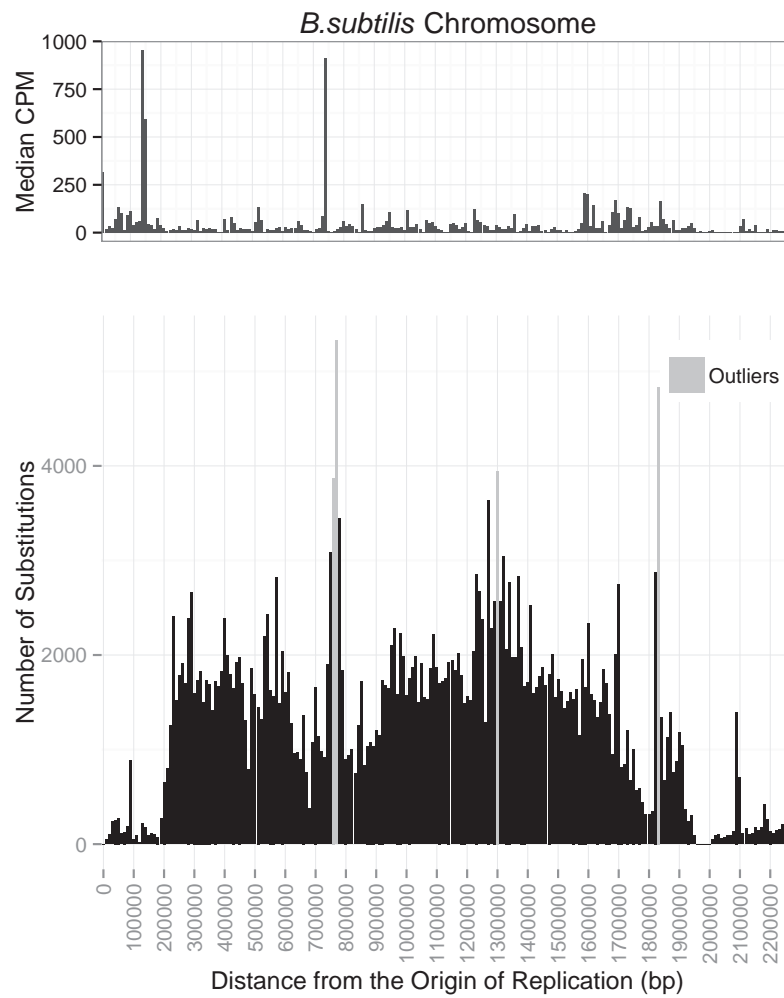


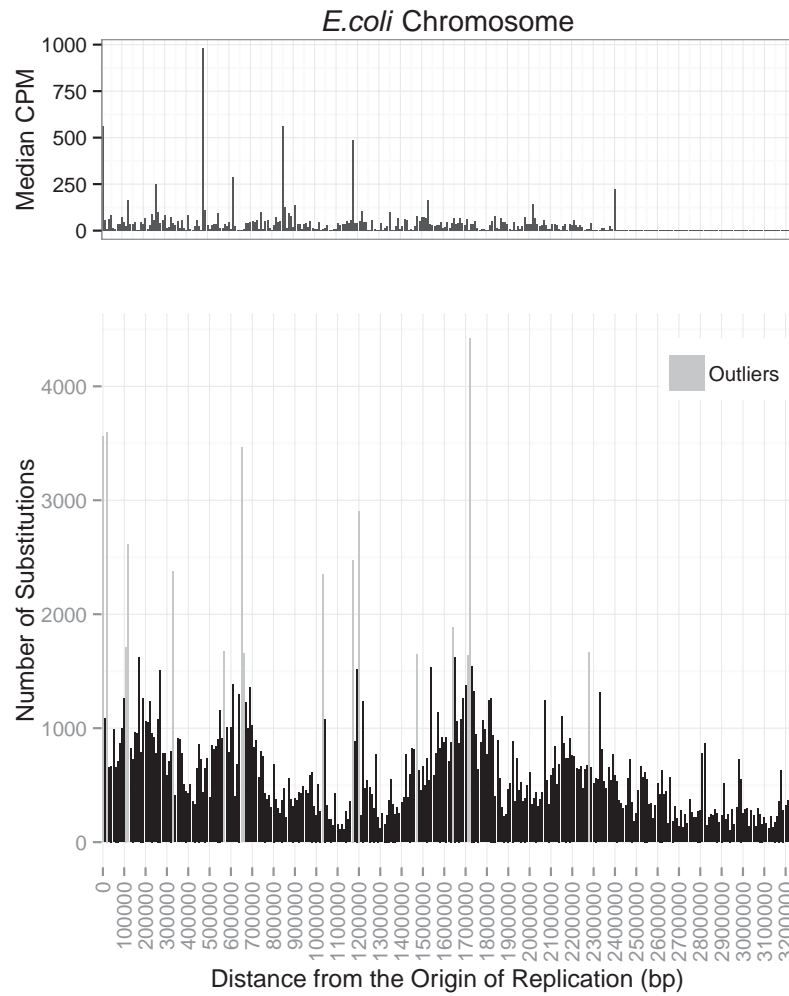
Bacteria and Replicon	Coefficient Estimate	Standard Error	P-value
<i>E. coli</i> Chromosome	$-1.394 \times 10^{-7}$	$2.425 \times 10^{-9}$	$< 2 \times 10^{-16}$
<i>B. subtilis</i> Chromosome	$-2.538 \times 10^{-8}$	$1.58 \times 10^{-9}$	$< 2 \times 10^{-16}$
<i>Streptomyces</i> Chromosome	$1.736 \times 10^{-8}$	$7.231 \times 10^{-10}$	$< 2 \times 10^{-16}$
<i>S. meliloti</i> Chromosome	$-1.541 \times 10^{-6}$	$3.042 \times 10^{-8}$	$< 2 \times 10^{-16}$
<i>S. meliloti</i> pSymA	$-9.130 \times 10^{-7}$	$1.975 \times 10^{-8}$	$< 2 \times 10^{-16}$
<i>S. meliloti</i> pSymB	$2.488 \times 10^{-7}$	$1.964 \times 10^{-8}$	$< 2 \times 10^{-16}$

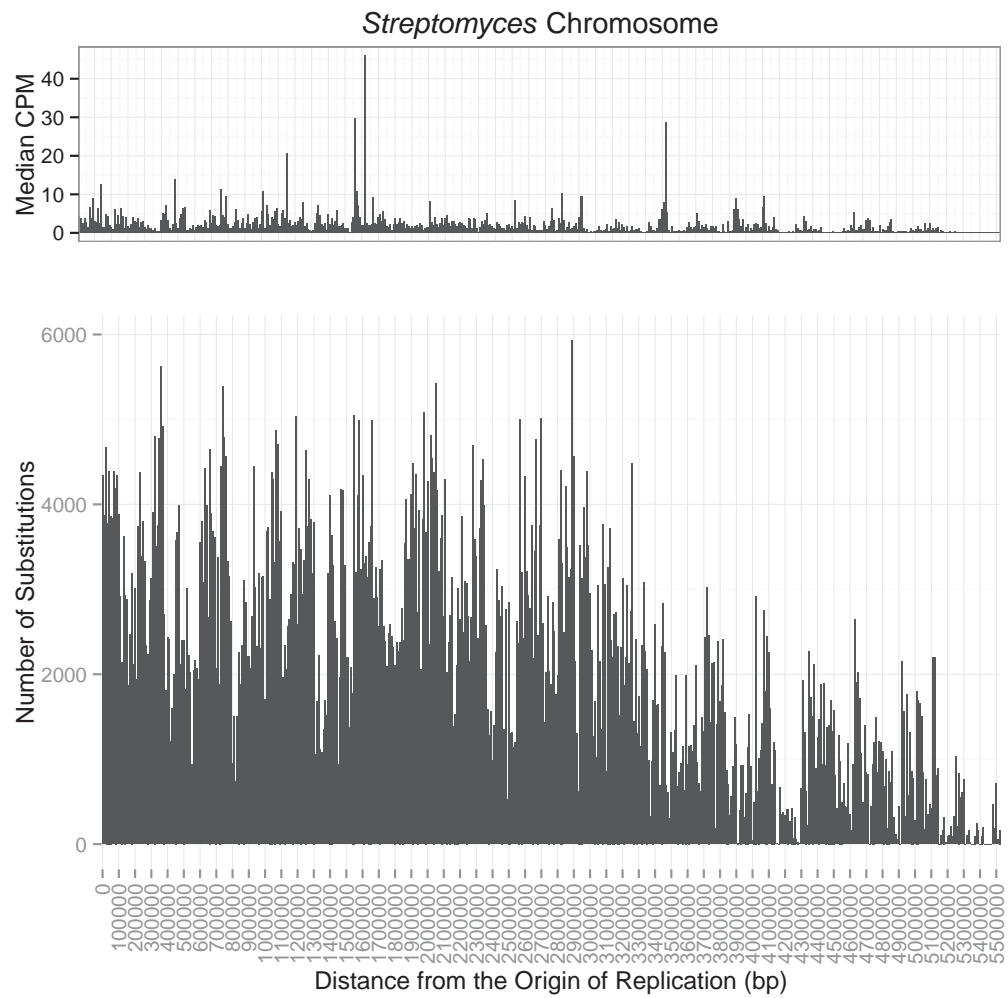
Table 2: Logistic regression analysis of the number of substitutions along the genome of the respective bacteria replicons. Grey coloured boxes indicate a negative logistic regression coefficient estimate. All results are statistically significant. Logistic regression was calculated after the origin 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	$-1.667 \times 10^{-7***}$	$-1.102 \times 10^{-7***}$	$6.716 \times 10^{-9***}$	$-1.363 \times 10^{-6***}$	$-1.405 \times 10^{-6***}$	$9.83 \times 10^{-8}$
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	$-1.262 \times 10^{-7***}$	$-2.602 \times 10^{-9}$	$1.563 \times 10^{-8***}$	$-1.599 \times 10^{-6***}$	$-9.499 \times 10^{-7***}$	$2.973 \times 10^{-7***}$
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 3: 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.