

1. Project Information

Program	Microbial/CSP 2012
PMO Project	0
Seq Proj ID	1027049
Sequencing Project Name	Nanoarchaeum sp. JGI 000156CP-E21
JGI Project ID	0

2. Read Statistics

Illumina Std PE Statistics

File name	7667.5.80864.ACAGTG.fastq
Library	TGPZ
Number of reads	26,674,140
Sequencing depth [†]	800X
Read type	2x150 bp

[†] A genome size of 5.0 Mbp was assumed in this calculation.

3. Read QC Results

The following are the results of reads screened against contaminants. Pairs of matching reads were removed from the dataset.

Illumina Std PE Read Filter Statistics

Description	Num Reads	Pct Reads
Input	26,674,140	100
Contam removed	98	0.0
Artifact removed	1,110,670	4.2
Total removed	6,674,140	25.0
Total remaining	20,000,000	75.0

List of Contaminants Removed

Description	Num Reads	Pct Reads
gi 357579577 Canis.lupus_familiaris_chr3	54	0.00
human_chr2	50	0.00
gi 357579535 Canis.lupus_familiaris_chr20	32	0.00
gi 357579571 Canis.lupus_familiaris_chr5	8	0.00
human_chr7	4	0.00
human_chr8	4	0.00
human_chr11	2	0.00

human_chr19	2	0.00
human_chr4	2	0.00

The following are the results of reads screened against potential reagent and process contaminants but were not removed from the dataset.

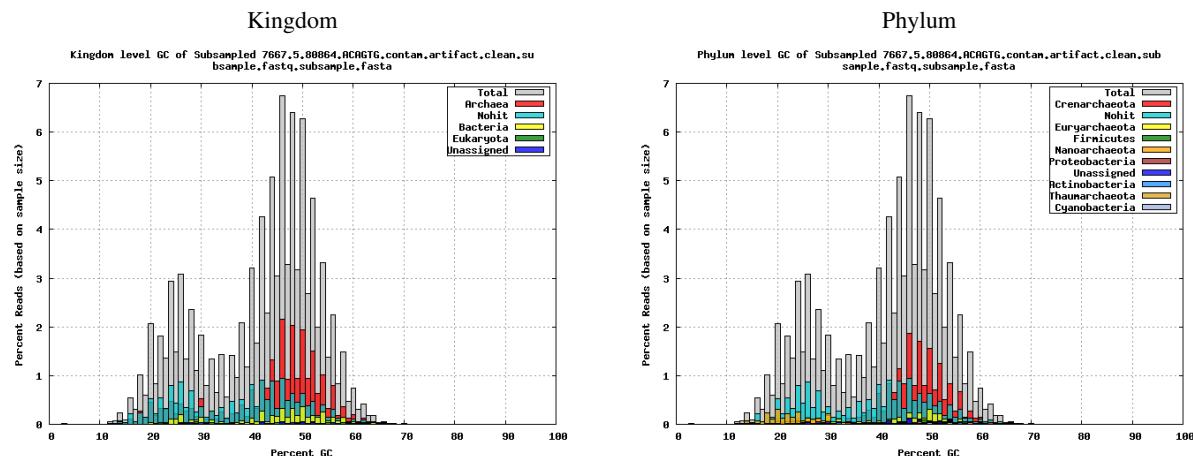
Illumina Std PE Contamination Identification Statistics

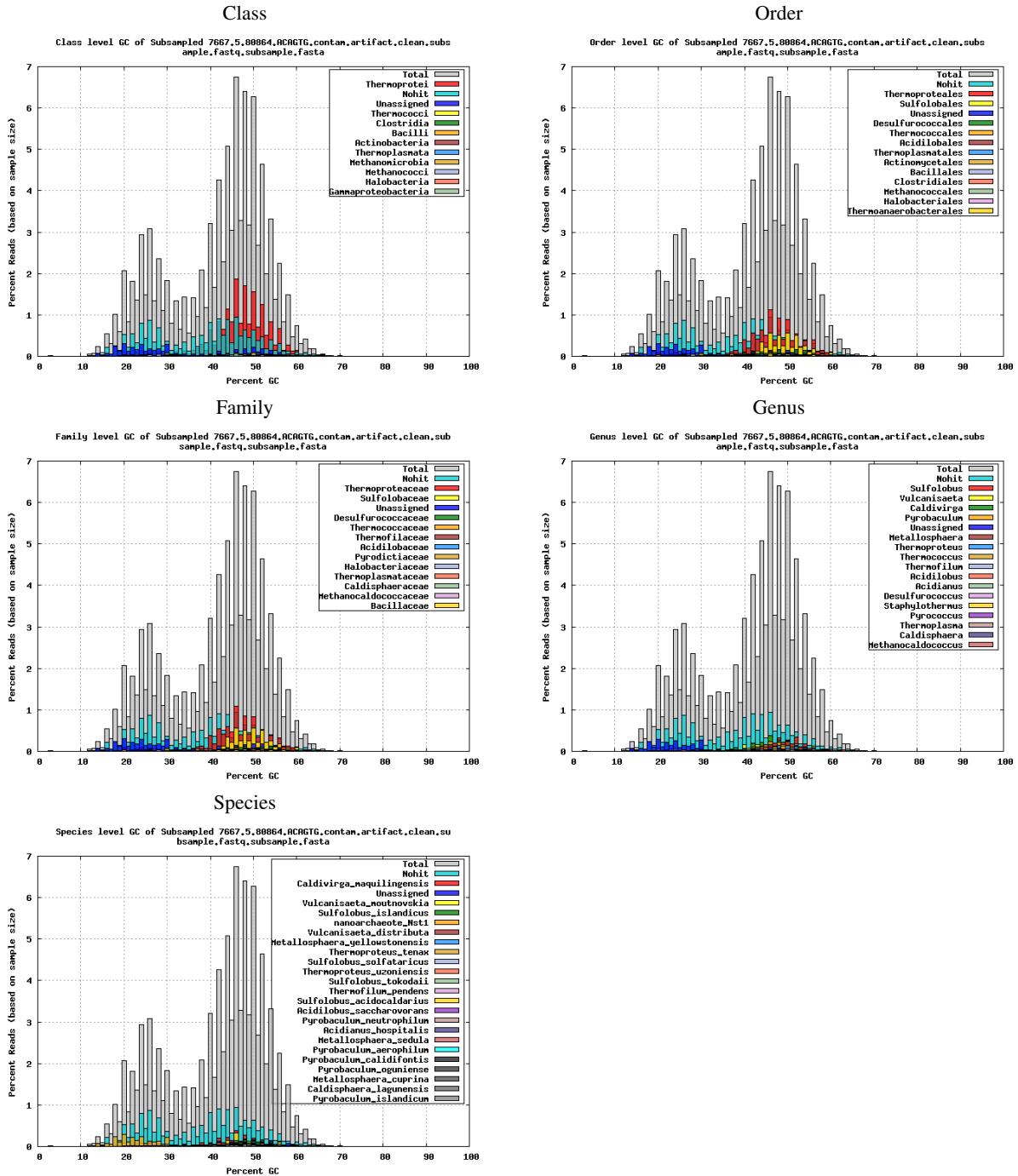
Description	Num Reads	Pct Reads
Input	26,674,140	100
Contam identified	2	0.0

List of Contaminants Identified

Description	Num Reads	Pct Reads
<i>Shigella</i>	2	0.00

GC histogram of the reads subsampled to 10k, overlaid with GC of hits based on BLASTX, shown for different taxonomic levels.



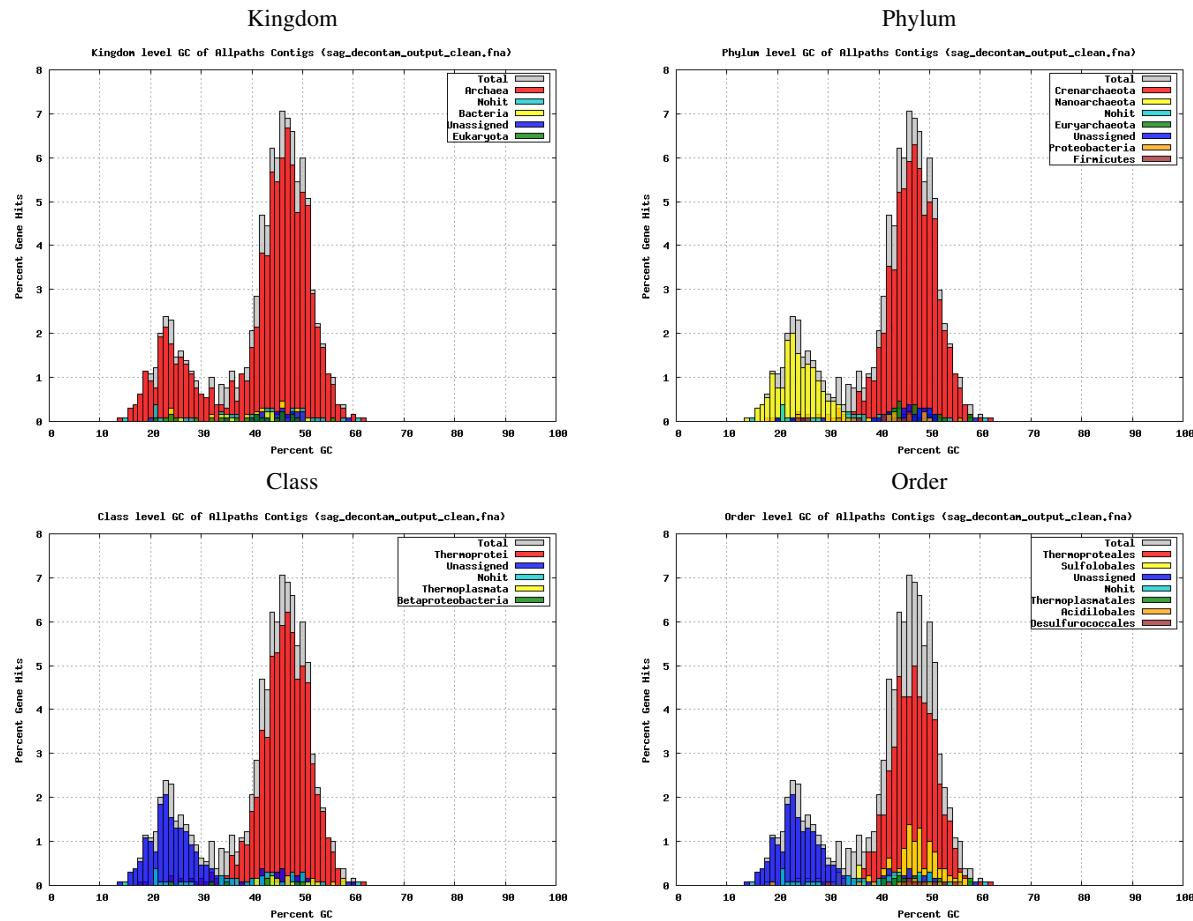


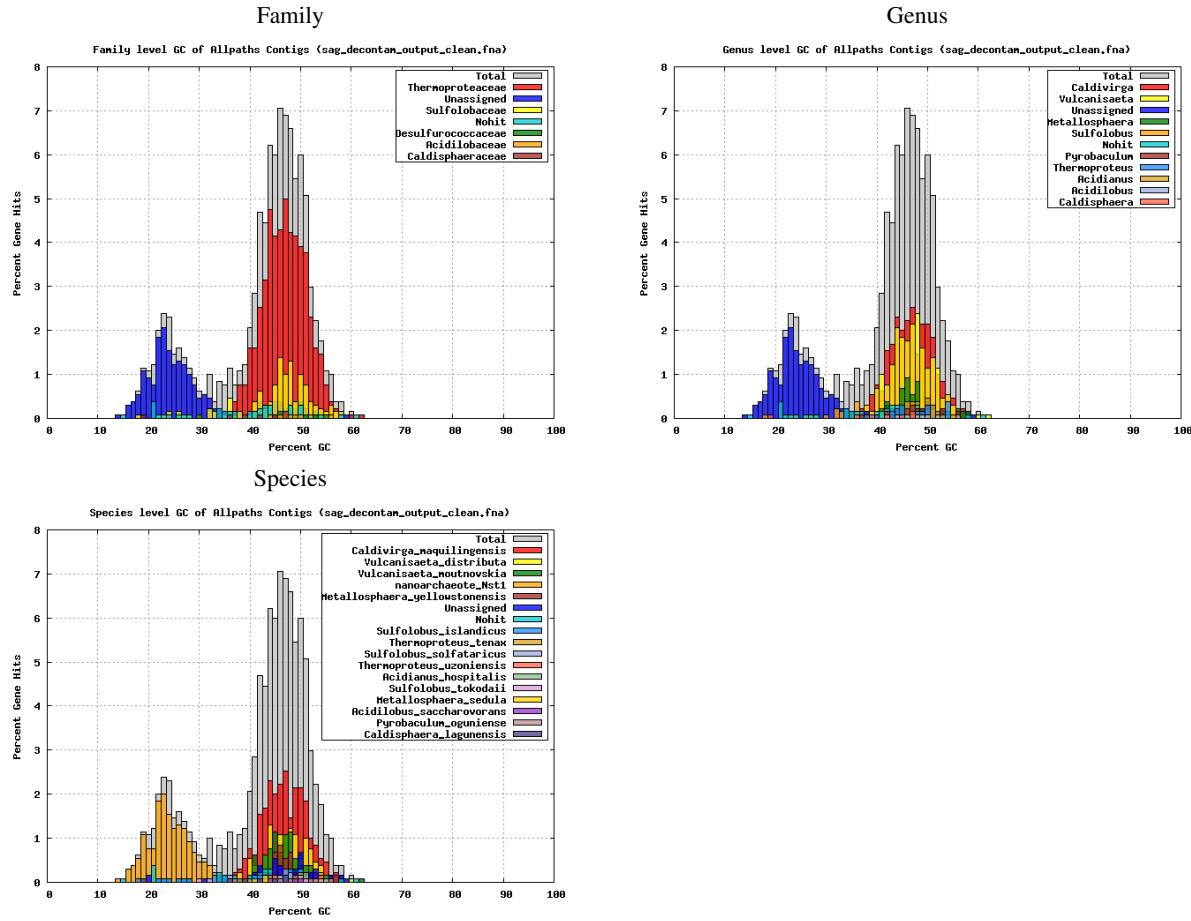
4. Assembly Statistics

Assembly method	SPAdes with auto decontamination
Scaffold total	92
Contig total	92
Scaffold sequence length	1.2 Mb
Contig sequence length	1.2 Mb (0.0% gap)
Scaffold N/L50	25/15.3 kb
Contig N/L50	25/15.3 kb
Largest Contig	57.9 kb
Number of scaffolds >50 kb	2
Pct of genome in scaffolds >50 kb	9.3
Pct of reads assembled (raw)	87.7
Pct of reads assembled (decontam)	77.4

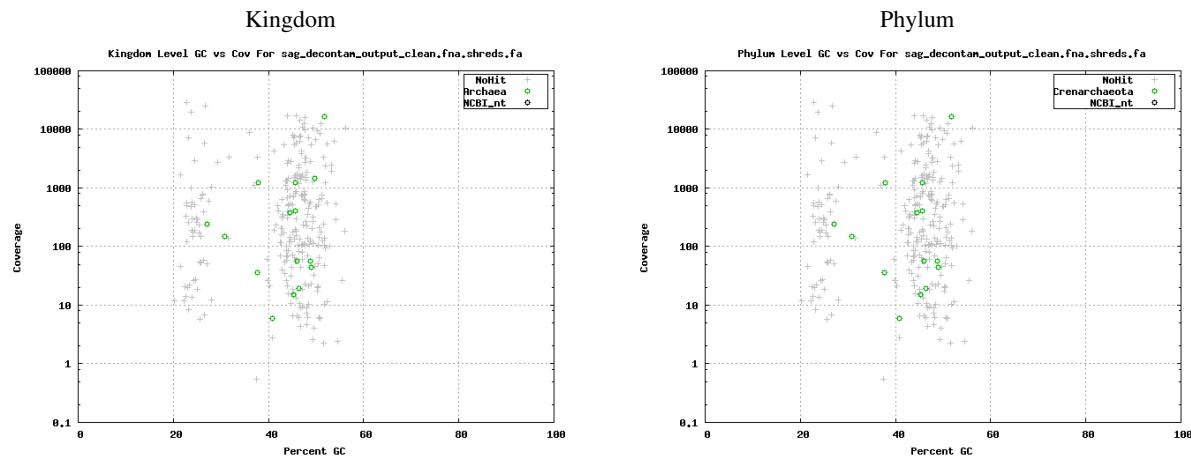
5. Assembly QC Results

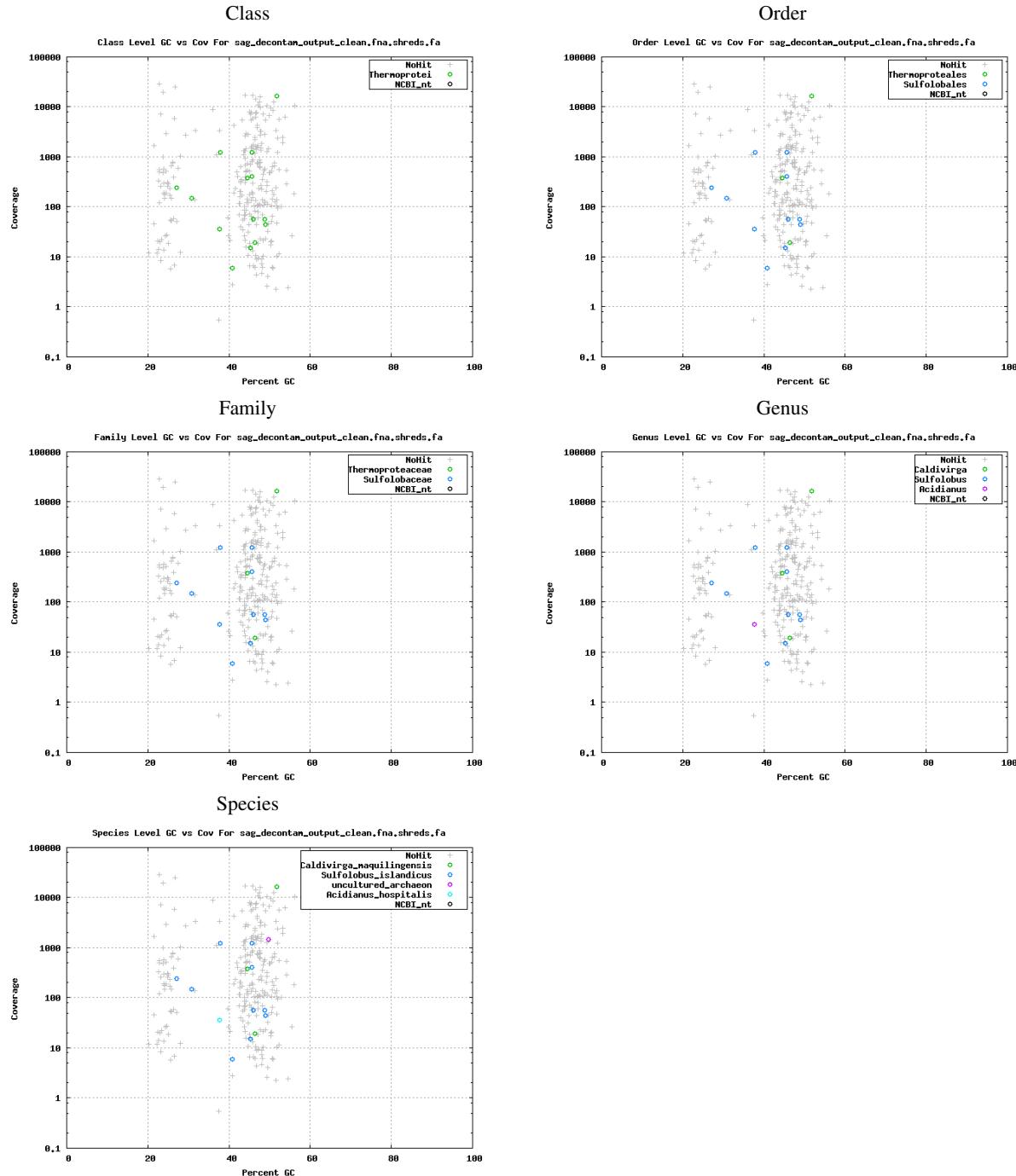
GC histogram of the predicted genes on each contig, overlaid with GC of hits based on BLASTP, shown for different taxonomic levels.



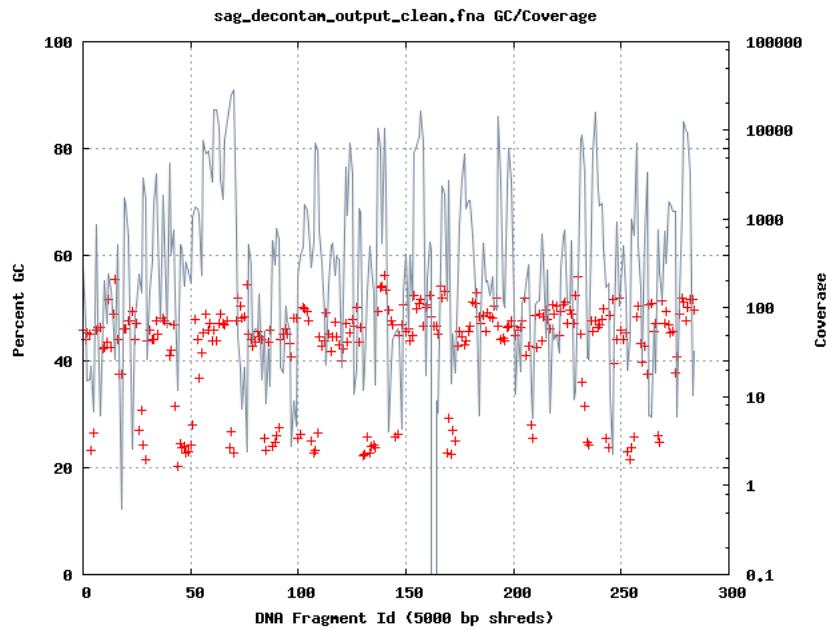


GC vs coverage based on GC of NCBI nt and Greengenes 16S rRNA gene hits to the assembly using megablast, shown for different taxonomic levels.

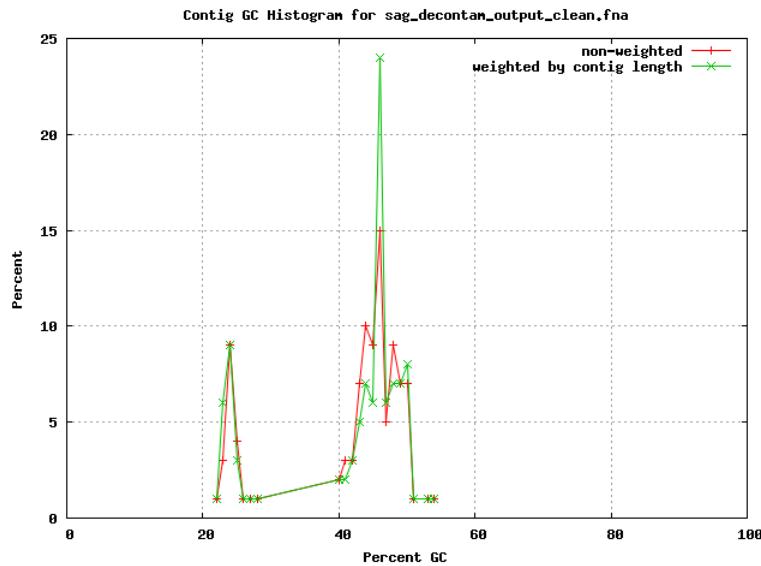




Coverage vs GC. Contigs were shredded into non-overlapping 5kbp and the GC of each shred was plotted as a point, colored by scaffold id. Coverage was calculated by mapping the fragment library to the final assembly and plotted as connected points.



GC histogram of the contigs, including contig length weighted distribution.



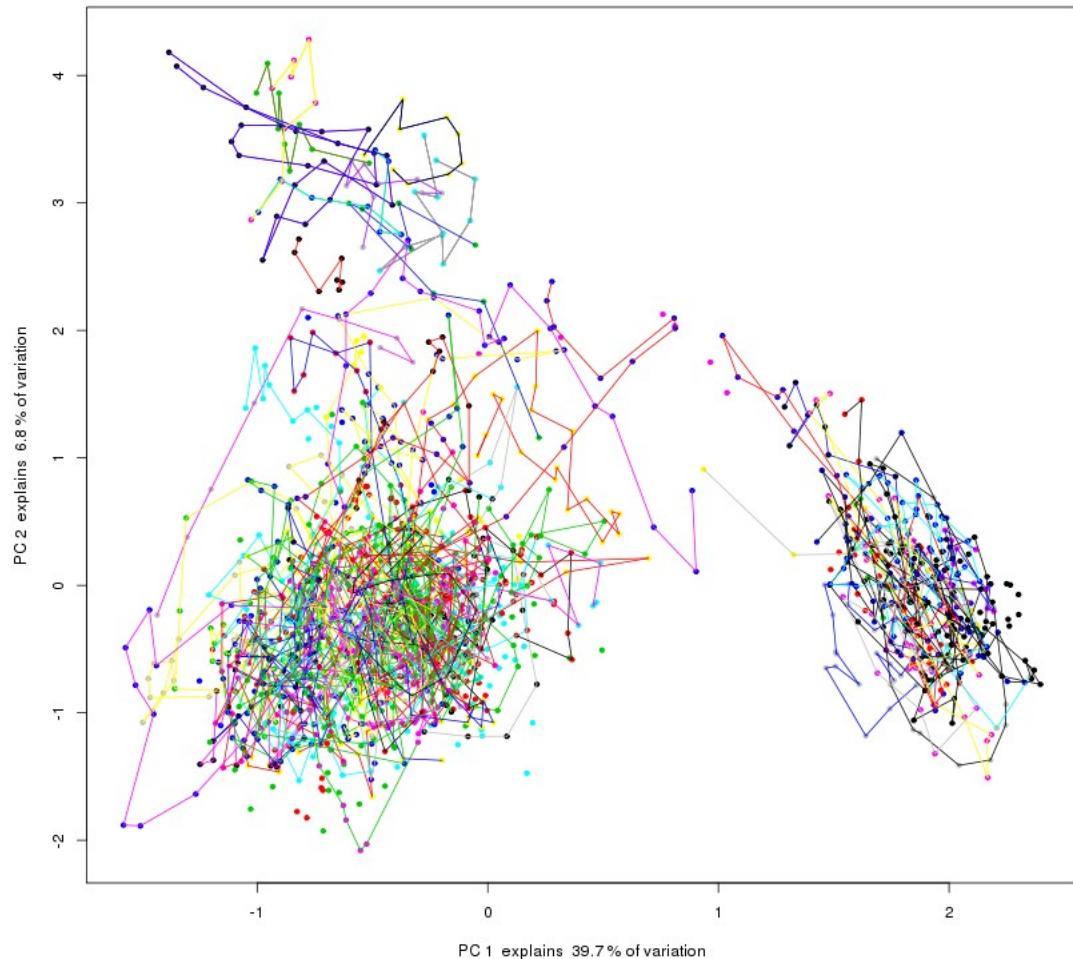
List of contigs and average percent GC, grouped in bins of 5:

Pct GC Bin	Contig Name
20	NODE_3.length_38555_cov_238.463.ID_5, NODE_7.length_22238_cov_169.114.ID_13, NODE_15.length_19889_cov_2312.17.ID_29, NODE_18.length_18333_cov_274.879.ID_35, NODE_35.length_12318_cov_20.0213.ID_69, NODE_39.length_11860_cov_15740.3.ID_81, NODE_48.length_9008_cov_17.9319.ID_97, NODE_49.length_8976_cov_22.6169.ID_99, NODE_51.length_8723_cov_10.6673.ID_103, NODE_53.length_8410_cov_1643.24.ID_107, NODE_57.length_8071_cov_121.125.ID_115, NODE_60.length_7749_cov_10.9062.ID_121

25	NODE.20.length_.17707_cov_382.6.ID.39, NODE.45.length_.9887_cov_941.446.ID.91, NODE.55.length_.8373_cov_267.224.ID.111, NODE.59.length_.7869_cov_393.81.ID.119, NODE.61.length_.7650_cov_142.776.ID.123, NODE.78.length_.6256_cov_7.39445.ID.169 NODE.84.length_.6147_cov_37.1822.ID.181
40	NODE.16.length_.18994_cov_707.422.ID.31, NODE.19.length_.17938_cov_186.006.ID.37, NODE.25.length_.15292_cov_260.769.ID.49, NODE.31.length_.13578_cov_105.933.ID.61, NODE.33.length_.12721_cov_278.611.ID.65, NODE.34.length_.12380_cov_295.146.ID.67, NODE.36.length_.12234_cov_84.5075.ID.71, NODE.42.length_.10518_cov_789.41.ID.87, NODE.44.length_.10040_cov_191.929.ID.75, NODE.50.length_.8808_cov_1269.81.ID.101, NODE.52.length_.8577_cov_77.1368.ID.105, NODE.56.length_.8345_cov_42.6204.ID.113, NODE.58.length_.7940_cov_293.961.ID.117, NODE.62.length_.7498_cov_193.273.ID.129, NODE.64.length_.7235_cov_19.1077.ID.141, NODE.67.length_.7056_cov_31.4845.ID.145, NODE.69.length_.6701_cov_877.414.ID.149, NODE.73.length_.6356_cov_347.857.ID.135, NODE.77.length_.6267_cov_94.2423.ID.167, NODE.79.length_.6246_cov_2339.06.ID.171, NODE.80.length_.6225_cov_648.101.ID.173, NODE.87.length_.5804_cov_32.3242.ID.157 NODE.98.length_.5319_cov_66.2475.ID.209
45	NODE.1.length_.57872_cov_5263.83.ID.1, NODE.2.length_.50783_cov_986.172.ID.3, NODE.5.length_.26827_cov_187.662.ID.9, NODE.6.length_.24626_cov_244.447.ID.11, NODE.8.length_.21666_cov_1831.82.ID.15, NODE.9.length_.21587_cov_45.084.ID.17, NODE.11.length_.21107_cov_124.042.ID.21, NODE.12.length_.20908_cov_639.345.ID.23, NODE.13.length_.20814_cov_2856.37.ID.25, NODE.17.length_.18584_cov_4356.9.ID.33, NODE.21.length_.17109_cov_1078.35.ID.41, NODE.22.length_.16949_cov_2302.73.ID.43, NODE.23.length_.15758_cov_717.659.ID.45, NODE.26.length_.14999_cov_568.39.ID.51, NODE.27.length_.14421_cov_266.389.ID.53, NODE.28.length_.14172_cov_788.282.ID.55, NODE.29.length_.13943_cov_229.87.ID.57, NODE.30.length_.13810_cov_161.98.ID.59, NODE.32.length_.13265_cov_57.5776.ID.63, NODE.37.length_.12222_cov_191.222.ID.73, NODE.38.length_.11892_cov_218.542.ID.79, NODE.40.length_.11594_cov_208.364.ID.83, NODE.43.length_.10312_cov_67.1227.ID.89, NODE.46.length_.9603_cov_1055.57.ID.93, NODE.54.length_.8408_cov_78.9294.ID.109, NODE.65.length_.7205_cov_10.3551.ID.125, NODE.66.length_.7108_cov_34.9321.ID.143, NODE.68.length_.6778_cov_165.66.ID.147, NODE.71.length_.6482_cov_150.409.ID.153, NODE.74.length_.6348_cov_7950.76.ID.161, NODE.75.length_.6303_cov_23.9262.ID.163, NODE.76.length_.6287_cov_11.9801.ID.165, NODE.81.length_.6223_cov_68.8713.ID.175, NODE.82.length_.6199_cov_422.415.ID.177, NODE.83.length_.6153_cov_5.72155.ID.179, NODE.86.length_.5889_cov_18.1046.ID.183, NODE.89.length_.5651_cov_5.87974.ID.187, NODE.94.length_.5424_cov_13.5897.ID.201, NODE.95.length_.5414_cov_4415.96.ID.203, NODE.96.length_.5374_cov_6.35909.ID.205 NODE.99.length_.5272_cov_495.668.ID.211
50	NODE.4.length_.27276_cov_4280.09.ID.7, NODE.10.length_.21500_cov_6272.83.ID.19, NODE.14.length_.20118_cov_208.837.ID.27, NODE.24.length_.15573_cov_2933.75.ID.47, NODE.41.length_.11592_cov_972.217.ID.85, NODE.47.length_.9455_cov_5754.77.ID.95, NODE.63.length_.7462_cov_74.221.ID.131, NODE.70.length_.6494_cov_33.0064.ID.151 NODE.91.length_.5597_cov_4.02237.ID.191

Principal component analysis of tetramer frequencies of contigs. Detectable variations are highlighted in color.

sag_decontam_output_clean.fna - PC1 vs PC2



Estimated genome recovery derived from analysis of universal single-copy genes detected in final assembly.

HMM	Pct Recovered
bacteria	29.58 %
archaea	60.36 %

6. Sequence Data Availability

The following sequence fasta files can be downloaded from our JGI portal website.
<http://www.jgi.doe.gov/genome-projects>

Filename	Description
sag_decontam_output_clean.fna	SPAdes with auto decontamination

7. Annotation Data Availability

The annotation of the assembled contigs can be found within IMG.
<http://img.jgi.doe.gov>

8. Methods

Single Cell Minimal Draft

Genome sequencing and assembly

The draft genome of was generated at the DOE Joint genome Institute (JGI) using the Illumina technology [1]. An Illumina std shotgun library was constructed and sequenced using the Illumina HiSeq 2000 platform which generated 26,674,140 reads totaling 4,001.1 Mb. All general aspects of library construction and sequencing performed at the JGI can be found at <http://www.jgi.doe.gov>. All raw Illumina sequence data was passed through DUK, a filtering program developed at JGI, which removes known Illumina sequencing and library preparation artifacts [2]. Following steps were then performed for assembly: (1) artifact filtered Illumina reads were assembled using SPAdes [3] (version 3.0.0), (3) Parameters for assembly steps were `-t 16 -m 120 --sc --careful --12`. The final draft assembly contained 92 contigs in 92 scaffolds, totalling 1.2 Mb in size. The final assembly was based on 3,000.0 Mb of Illumina data. Based on a presumed genome size of 5.0 Mb, the average input read coverage used for the assembly was 600.0X.

Genome annotation

Genes were identified using Prodigal [4], followed by a round of manual curation using GenePRIMP [5] for finished genomes and Draft genomes in fewer than 20 scaffolds. The predicted CDSs were translated and used to search the National Center for Biotechnology Information (NCBI) nonredundant database, UniProt, TIGRFam, Pfam, KEGG, COG, and InterPro databases. The tRNAscanSE tool [6] was used to find tRNA genes, whereas ribosomal RNA genes were found by searches against models of the ribosomal RNA genes built from SILVA [7]. Other non-coding RNAs such as the RNA components of the protein secretion complex and the RNase P were identified by searching the genome for the corresponding Rfam profiles using INFERNAL [8]. Additional gene prediction analysis and manual functional annotation was performed within the Integrated Microbial Genomes (IMG) platform [9] developed by the Joint Genome Institute, Walnut Creek, CA, USA [10].

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