

## 1. Project Information

Program	Microbial/CSP 2012
PMO Project	0
Seq Proj ID	1027097
Sequencing Project Name	NAG2 archaeon OSP418 JGI 000156CP-F08
JGI Project ID	0

## 2. Read Statistics

### Illumina Std PE Statistics

File name	7666.6.80850.CGTACG.fastq
Library	TGTA
Number of reads	30,145,848
Sequencing depth <sup>†</sup>	904X
Read type	2x150 bp

<sup>†</sup> 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	30,145,848	100
Contam removed	116	0.0
Artifact removed	342,116	1.1
Total removed	10,145,848	33.7
Total remaining	20,000,000	66.3

### List of Contaminants Removed

Description	Num Reads	Pct Reads
human_chr2	90	0.00
gi 357579577 Canis_lupus_familiaris_chr3	80	0.00
gi 357579535 Canis_lupus_familiaris_chr20	22	0.00
gi 357579571 Canis_lupus_familiaris_chr5	14	0.00
human_chr3	2	0.00
human_chr5	2	0.00

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

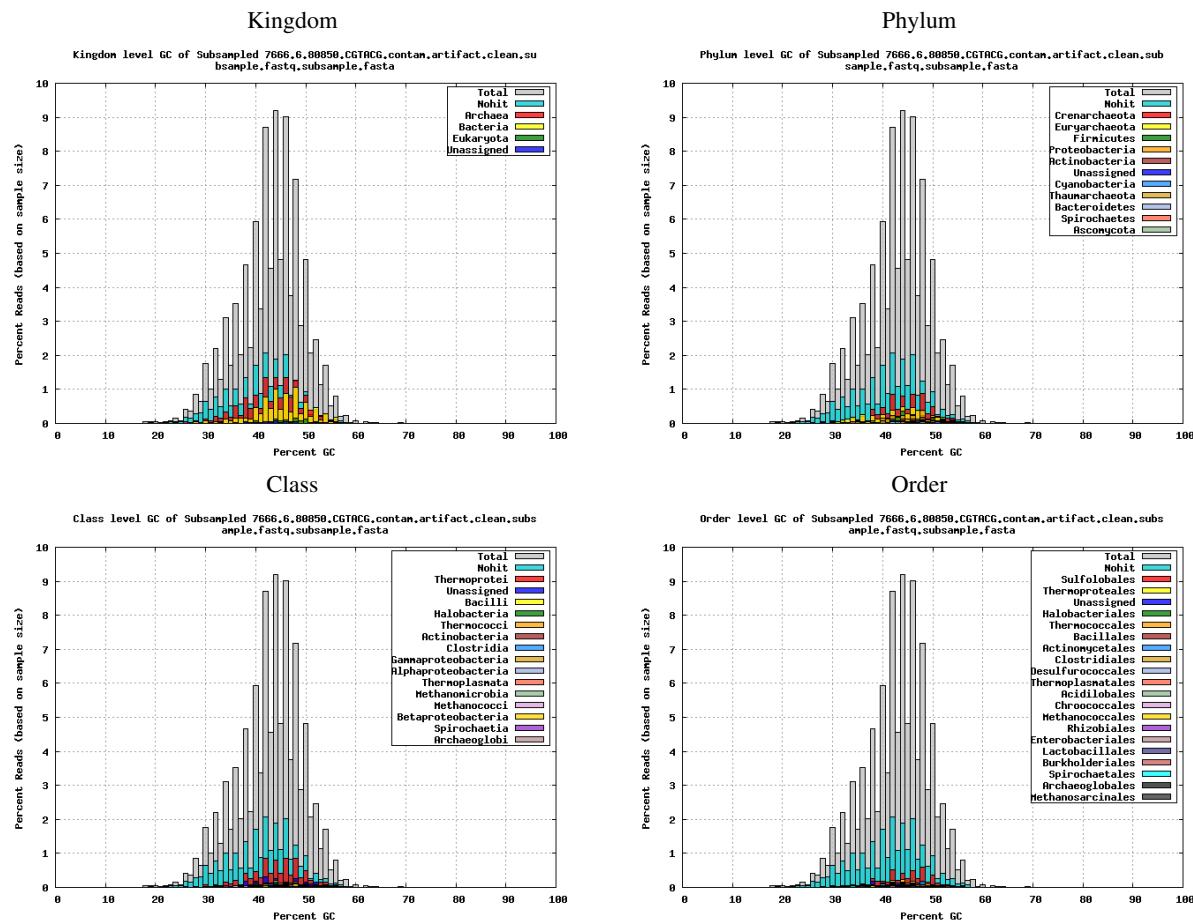
### Illumina Std PE Contamination Identification Statistics

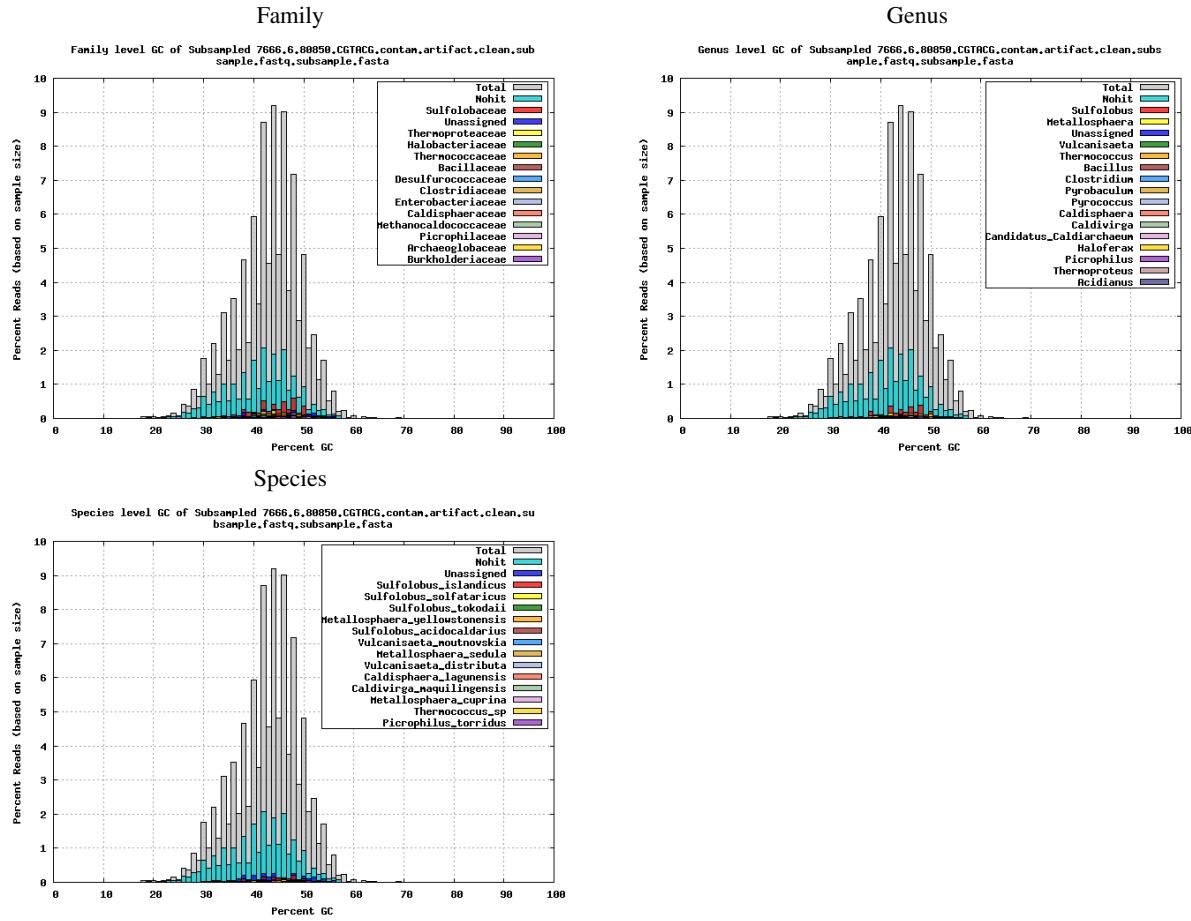
Description	Num Reads	Pct Reads
Input	30,145,848	100
Contam identified	4	0.0

### List of Contaminants Identified

Description	Num Reads	Pct Reads
<i>Pseudomonas</i>	4	0.00

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





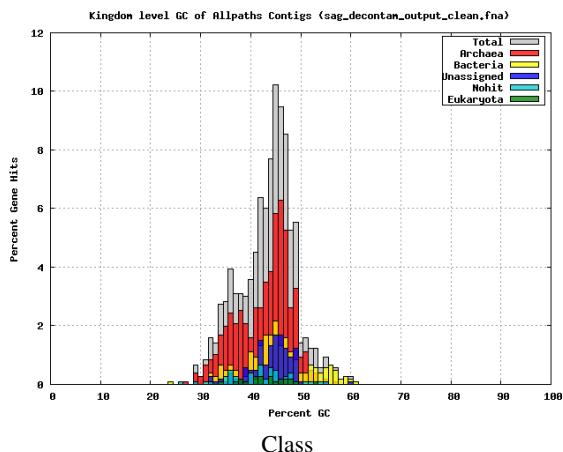
## 4. Assembly Statistics

Assembly method	SPAdes with auto decontamination
Scaffold total	84
Contig total	84
Scaffold sequence length	995.4 kb
Contig sequence length	995.4 kb ( 0.0% gap)
Scaffold N/L50	28/12.8 kb
Contig N/L50	28/12.8 kb
Largest Contig	31.2 kb
Number of scaffolds >50 kb	0
Pct of genome in scaffolds >50 kb	0.0
Pct of reads assembled (raw)	90.2
Pct of reads assembled (decontam)	79.6

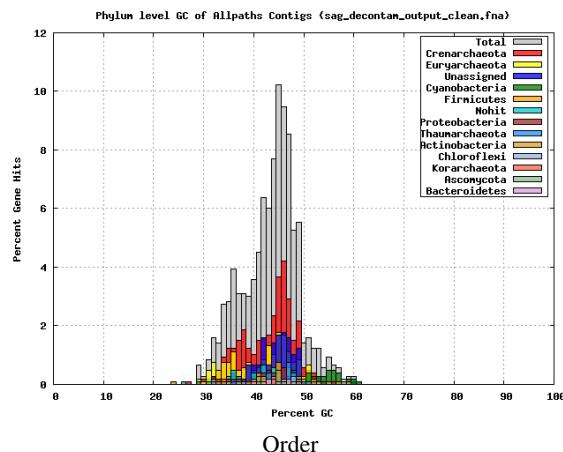
## 5. Assembly QC Results

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

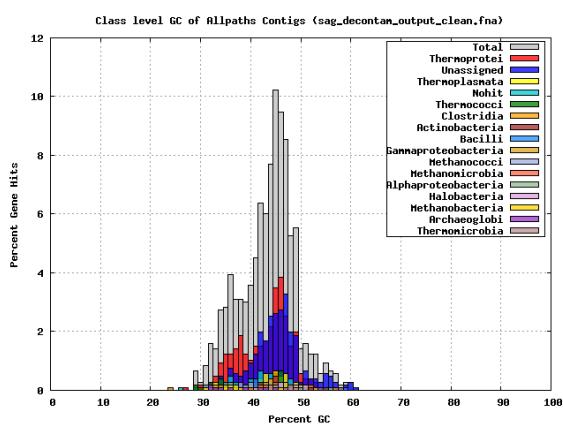
### Kingdom



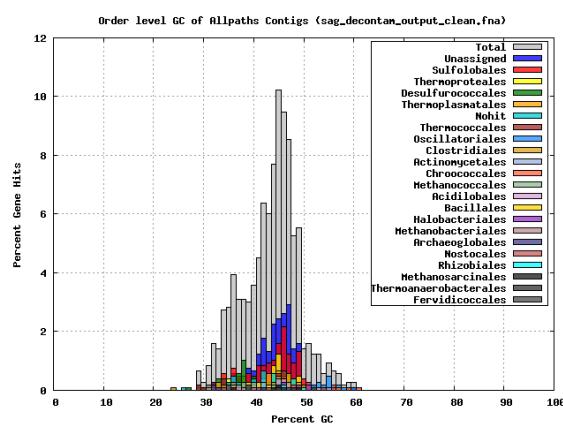
### Phylum



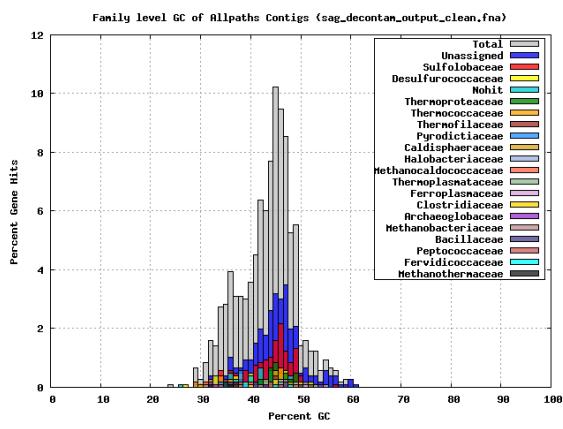
### Class



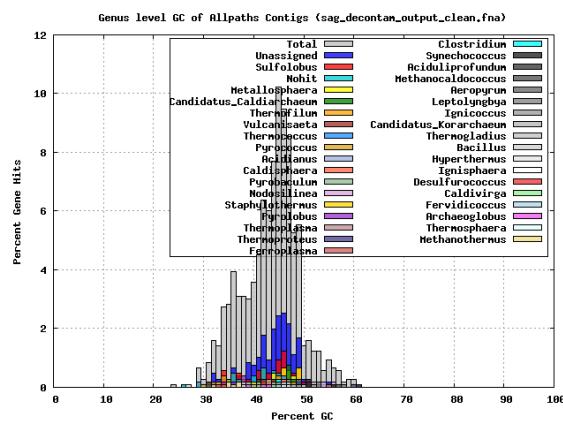
### Order



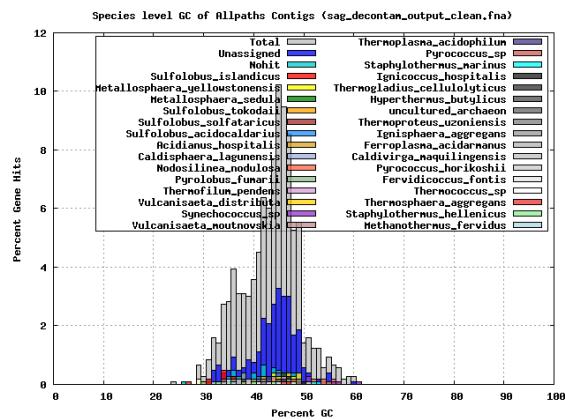
### Family



### Genus

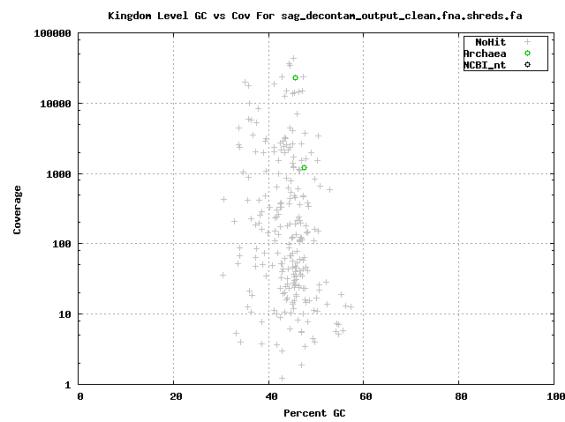


## Species

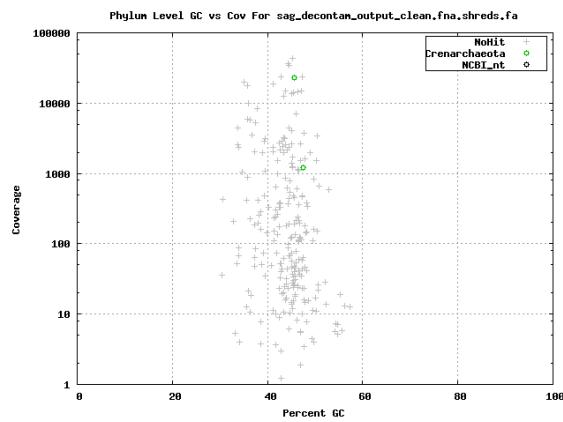


GC vs coverage based on GC of NCBI nt and Greengenes 16S rRNA gene hits to the assembly using megablast, shown for different taxonomic levels.

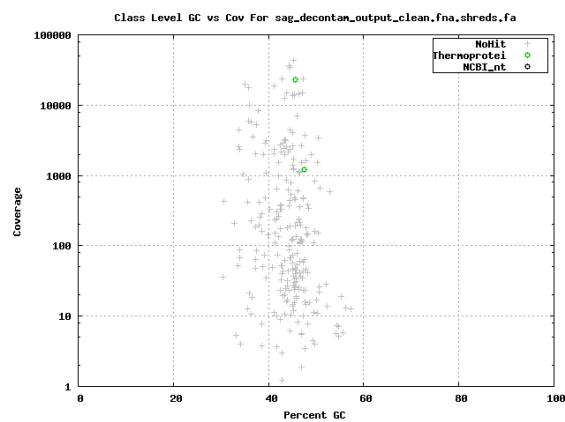
## Kingdom



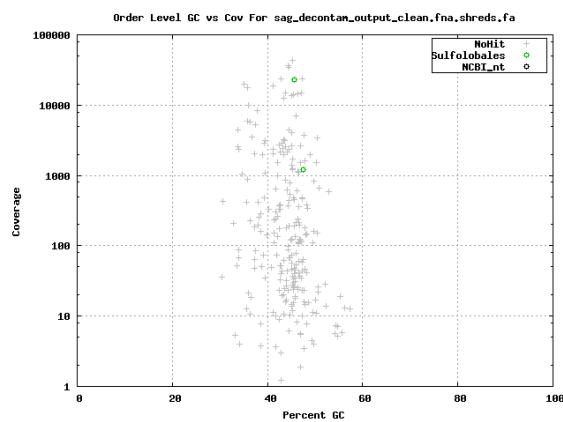
## Phylum



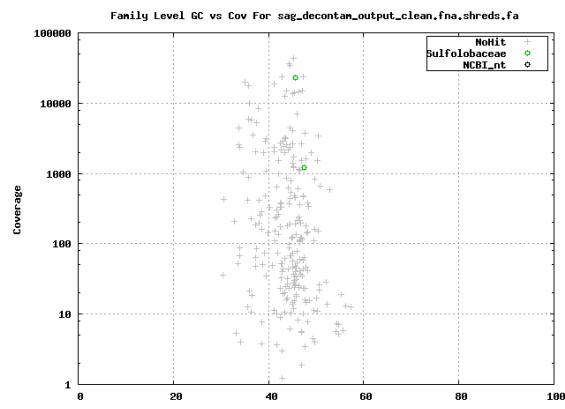
## Class



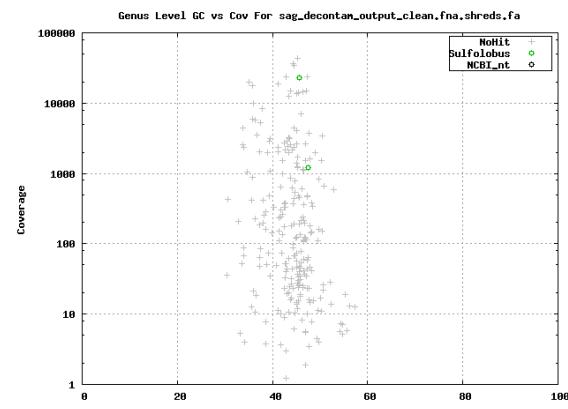
## Order



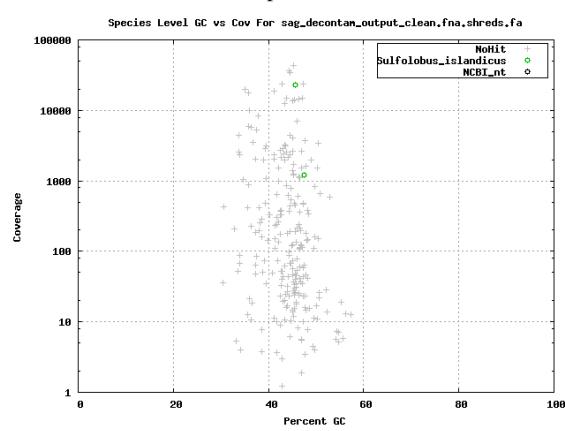
Family



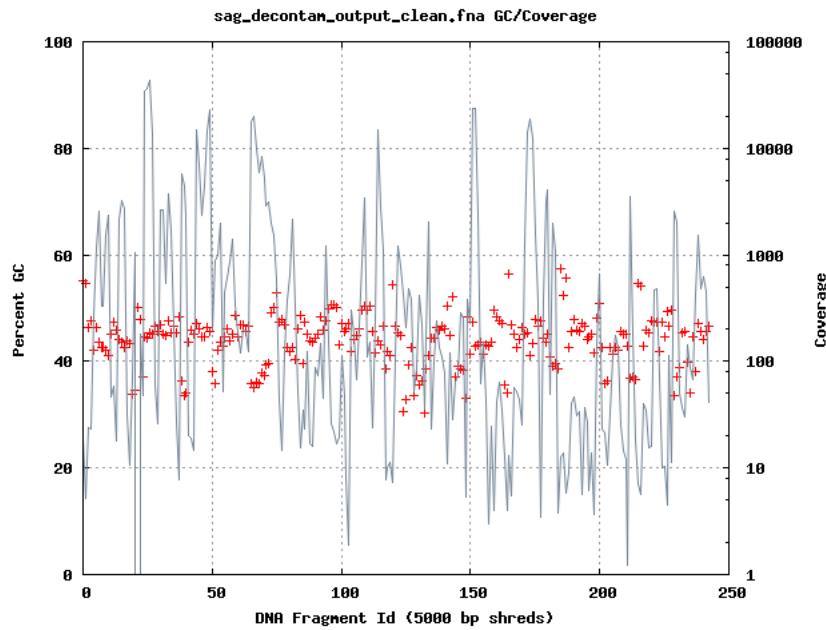
Genus



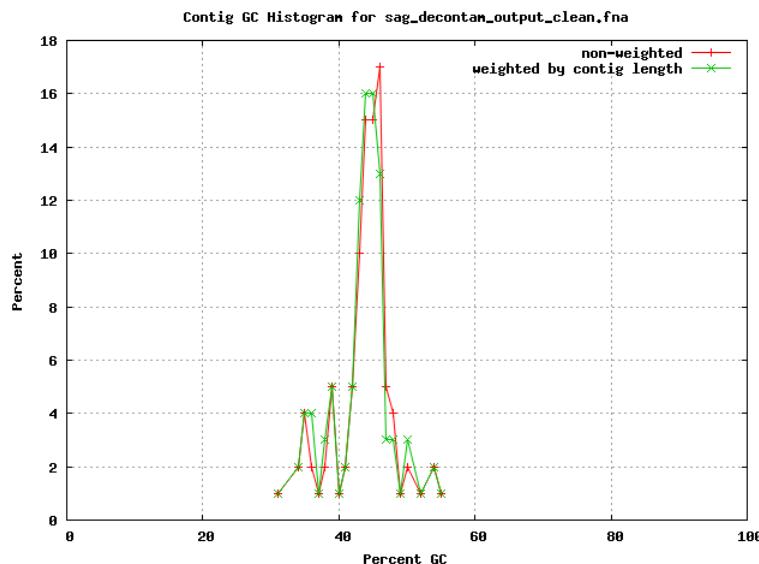
Species



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.

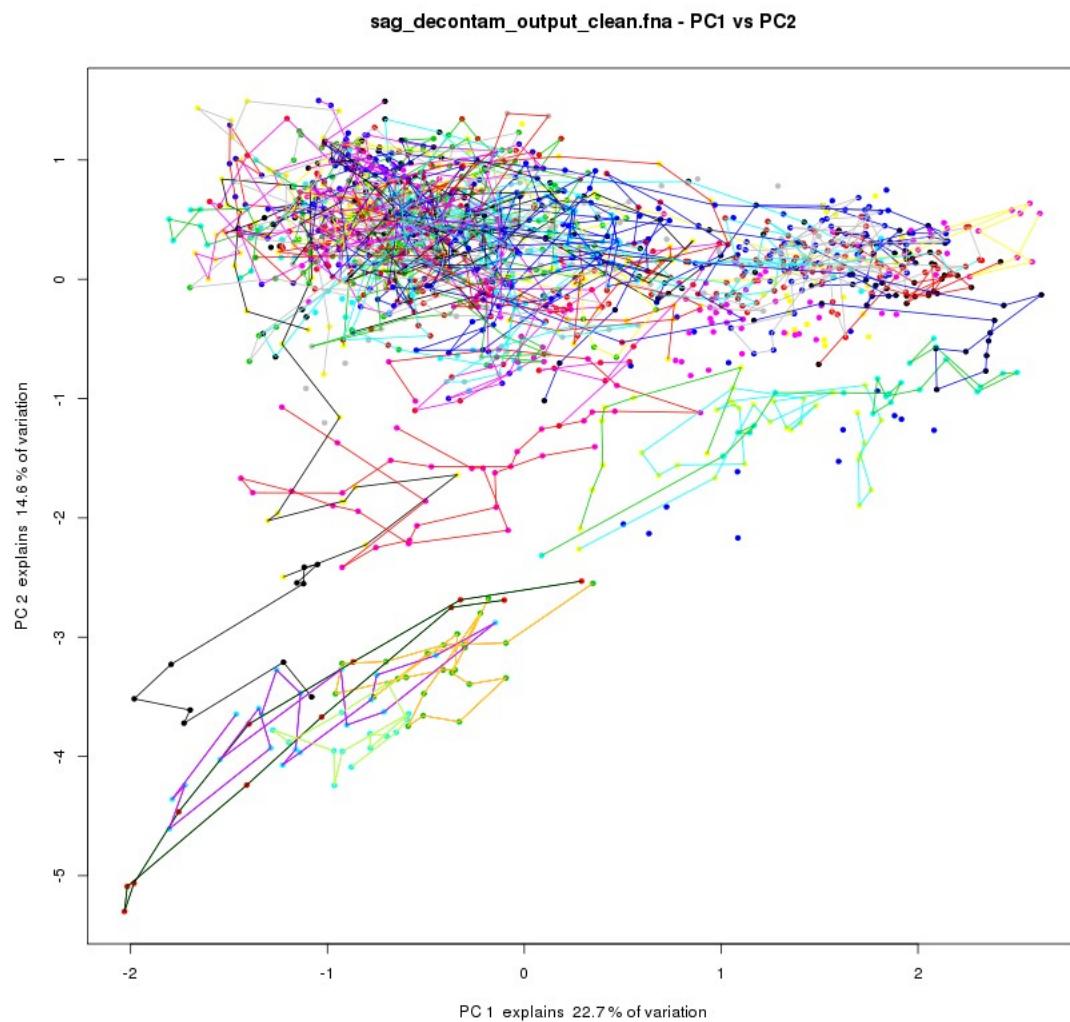


List of contigs and average percent GC, grouped in bins of 5:

Pct GC Bin	Contig Name
30	NODE_34.length_12552 cov_3062.84.ID_71, NODE_49.length_10012 cov_423.118.ID_101 NODE_61.length_.8343 cov_.257.706.ID_133
35	NODE_2.length_28596 cov_.7675.09.ID_3, NODE_7.length_20644 cov_131.751.ID_13, NODE_8.length_20501 cov_111.418.ID_15, NODE_10.length_18497 cov_.638.532.ID_19, NODE_15.length_16058 cov_.671.825.ID_29, NODE_30.length_12799 cov_13.3839.ID_63, NODE_32.length_12758 cov_961.698.ID_67, NODE_39.length_11155 cov_1396.68.ID_81, NODE_41.length_11084 cov_74.9589.ID_85, NODE_51.length_9569 cov_.690.312.ID_115,

	NODE..67.length..7603cov..2003.94.ID..143, NODE..83.length..5477cov..7.77628.ID..181
40	NODE..1.length..31190cov..5176.4.ID..1, NODE..5.length..23838cov..2686.12.ID..9, NODE..6.length..22615cov..524.849.ID..11, NODE..9.length..18778cov..22217.7.ID..17, NODE..13.length..16975cov..37.8271.ID..25, NODE..14.length..16963cov..75.5506.ID..27, NODE..16.length..15879cov..6.37588.ID..31, NODE..17.length..15656cov..56.3436.ID..33, NODE..19.length..14918cov..250.743.ID..37, NODE..20.length..14903cov..51.2083.ID..41, NODE..22.length..14510cov..12.0939.ID..45, NODE..26.length..13835cov..762.57.ID..55, NODE..28.length..13562cov..1813.88.ID..59, NODE..33.length..12733cov..1527.77.ID..69, NODE..35.length..12467cov..595.704.ID..73, NODE..36.length..12240cov..13301.1.ID..75, NODE..37.length..11988cov..144.538.ID..77, NODE..50.length..9975cov..40.5739.ID..103, NODE..53.length..9260cov..483.477.ID..119, NODE..55.length..9174cov..299.056.ID..107, NODE..56.length..9053cov..83.0051.ID..123, NODE..62.length..8332cov..1273.51.ID..135, NODE..63.length..8148cov..37.1111.ID..137, NODE..73.length..6553cov..18.0114.ID..157, NODE..77.length..5870cov..28.0884.ID..169, NODE..78.length..5824cov..8.25498.ID..171, NODE..86.length..5390cov..15.483.ID..187, NODE..90.length..4814cov..5.88065.ID..195
45	NODE..3.length..28068cov..6926.3.ID..5, NODE..4.length..24749cov..434.024.ID..7, NODE..11.length..18293cov..159.877.ID..21, NODE..12.length..17483cov..1178.95.ID..23, NODE..21.length..14693cov..475.346.ID..47, NODE..23.length..14147cov..922.32.ID..49, NODE..27.length..13595cov..175.849.ID..57, NODE..31.length..12794cov..289.576.ID..65, NODE..40.length..11111cov..293.158.ID..83, NODE..42.length..11022cov..19.9457.ID..87, NODE..43.length..10966cov..220.677.ID..89, NODE..44.length..10963cov..33.4745.ID..91, NODE..45.length..10947cov..19.8674.ID..93, NODE..46.length..10936cov..6.52743.ID..95, NODE..47.length..10909cov..40.5483.ID..97, NODE..48.length..10304cov..61.5664.ID..99, NODE..52.length..9402cov..86.6375.ID..117, NODE..54.length..9234cov..607.897.ID..121, NODE..58.length..8862cov..13.9508.ID..127, NODE..59.length..8798cov..133.195.ID..129, NODE..64.length..8069cov..73.0304.ID..139, NODE..66.length..7643cov..26.4512.ID..111, NODE..68.length..7256cov..29.5284.ID..149, NODE..69.length..7033cov..445.636.ID..151, NODE..70.length..6976cov..35.406.ID..153, NODE..72.length..6570cov..240.465.ID..147, NODE..74.length..6521cov..16.038.ID..163, NODE..75.length..6096cov..25.9396.ID..165, NODE..76.length..6070cov..63.6075.ID..167, NODE..79.length..5740cov..72.5186.ID..173, NODE..81.length..5572cov..1847.51.ID..177, NODE..82.length..5525cov..17.6276.ID..179, NODE..84.length..5308cov..25.7595.ID..183, NODE..87.length..5315cov..5.36958.ID..189 NODE..89.length..5217cov..7.63367.ID..193
50	NODE..25.length..13849cov..958.413.ID..53, NODE..29.length..13499cov..14.6773.ID..61, NODE..38.length..11789cov..8.19397.ID..79, NODE..60.length..8522cov..4.212.ID..131 NODE..65.length..7829cov..6.76962.ID..141
55	NODE..71.length..6697cov..10.1736.ID..155

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



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

HMM	Pct Recovered
bacteria	43.96 %
archaea	67.22 %

## 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 30,145,848 reads totaling 4,521.9 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 84 contigs in 84 scaffolds, totalling 995.4 Kb 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].

1. Bennett S. Solexa Ltd. *Pharmacogenomics*. 2004;5(4):433–8.
2. Mingkun L, Copeland A, Han J. DUK, unpublished, 2011.
3. Bankevich A, et.al. SPAdes: a new genome assembly algorithm and its applications to single-cell sequencing. *J Comput Biol* 2012; 19:455–77.
4. Hyatt D, Chen GL, Lascasio PF, Land ML, Larimer FW, Hauser LJ. Prodigal: prokaryotic gene recognition and translation initiation site identification. *BMC Bioinformatics* 2010; 11:119.
5. Pati A, Ivanova NN, Mikhailova N, Ovchinnikova G, Hooper SD, Lykidis A, Kyrpides NC. GenePRIMP: a gene prediction improvement pipeline for prokaryotic genomes. *Nat Methods* 2010; 7:455–457.
6. Lowe TM, Eddy SR. tRNAscan-SE: a program for improved detection of transfer RNA genes in genomic sequence. *Nucleic Acids Res* 1997; 25:955–964.
7. Pruesse E, Quast C, Knittel, Fuchs B, Ludwig W, Peplies J, Glckner FO. SILVA: a comprehensive online resource for quality checked and aligned ribosomal RNA sequence data compatible with ARB. *Nuc Acids Res* 2007; 35: 2188–7196.
8. INFERNAL. Inference of RNA alignments. <http://infernal.janelia.org>.
9. The Integrated Microbial Genomes (IMG) platform. <http://www.ncbi.nlm.nih.gov/pubmed/24165883>
10. Markowitz VM, Mavromatis K, Ivanova NN, Chen IMA, Chu K, Kyrpides NC. IMG ER: a system for microbial genome annotation expert review and curation. *Bioinformatics* 2009; 25:2271–2278.