

# Lawrence Berkeley National Laboratory

## Recent Work

### Title

The US DOE Joint Genome Institute Microbial Genome Program

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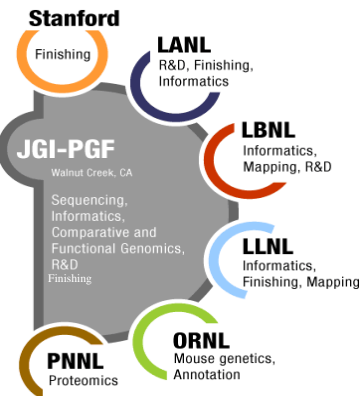
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# The US DOE Joint Genome Institute Microbial Genome Program



Alla Lapidus, Patrick Chain, Cliff Han, Thomas Brettin, Alex Copeland, Chris Detter, Samuel Pitluck, Tijana Glavina del Rio, Susan Lucas, Kerrie Barry, Miriam Land, Frank Larimer, Nikos Kyrpides, Natalia Ivanova, Jeremy Schmutz, Victor M Markowitz, David Bruce, Paul Gilna, Jim Bristow, Eddy Rubin, Paul Richardson.  
DOE Joint Genome Institute



**Programs:**  
**DOE Microbial Program**  
**DOE GTL Program (GTL)**  
**Community Sequencing Program (CSP)**  
**JGI Internal Program**  
**LSP- Lab Science Program**

**Goal: to provide the scientific community access to high throughput sequencing and to operate as a Genomic Infrastructure for American Science**

## The Community Sequencing Program

<http://www.jgi.doe.gov/CSP/index.html>

### Types of projects:

A wide range of projects. The most important factor for acceptance is a project's scientific merit. The deliverables can range from raw sequence traces to well-annotated assembled genomes

JGI is a leader in performing sequences to support the number of U.S. Department of Energy (DOE) Microbial Programs. The Microbial Genome and GTL programs were established to determine the complete genome sequence of a number of microbes selected for their relevance to DOE missions. Community Sequencing Program recently started by JGI (<http://www.jgi.doe.gov/CSP/index.html>) also includes sequence and detailed analysis of the genomes of the different representatives of the microbial world.

A workflow procedure for all microbial programs has been formalized to process samples from DNA prep through sequencing, assembly, finishing, quality assurance, annotation and analysis. To date, the JGI has sequenced over 300 microbes to draft quality, finished over 60 and is currently working on more than 100 additional microbial projects. Most projects are now targeted for complete finishing.

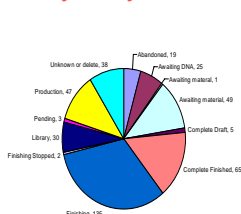
A completely sequenced, high quality genome is a perfect starting point for the genome annotation (<http://img.jgi.doe.gov/v1.1/main.cgi>), microarrays, knockout experiments, etc. Despite the fact that draft genomes are useful on their own, a completed genome is, overall, a better product, especially if it will be used to analyze previously unknown and difficult-to-cultivate microbes; for the comparative analysis of clinical isolates, or for the creation of microbial strains overproducing different proteins and amino acids. Knowledge of the completely finished genome will allow scientists to modify specific regions of the genome and, therefore, to affect the expression of the gene being studied. Thus, in order to be able to realize these and many other studies, it is necessary to close most (if not all) of the genomes being sequenced at JGI.

The Integrated Microbial Genomes (IMG) system was created to provide a framework for comparative analysis of the genomes sequenced by the Joint Genome Institute. Its goal is to facilitate the visualization and exploration of genomes from a functional and evolutionary perspective. Currently IMG includes 674 genomes from archaea to eukarya sequenced by JGI (134) and other centers.

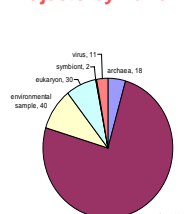
This work was performed under the auspices of the US Department of Energy's Office of Science, Biological and Environmental Research Program, and by the University of California, Lawrence Livermore National Laboratory under Contract No. W-7405-Eng-48, Lawrence Berkeley National Laboratory under Contract No. DE-AC02-05CH11231 and Los Alamos National Laboratory under Contract No. W-7405-ENG-36. LBNL-57373 Poster II

UCRL-POST-218486

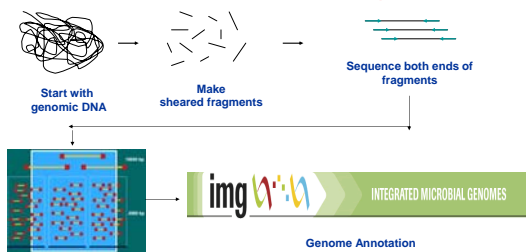
### Projects by Status



### Projects by Taxonomy



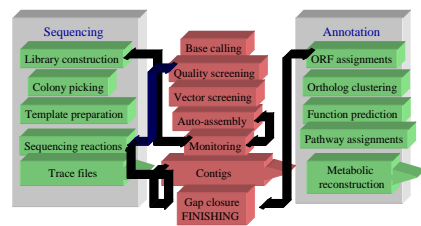
## Genome Sequencing



## Finished Genomes

- 2005 DOE Microbe Projects**
  - Rhodopseudomonas palustris* HaA2
  - Acidobacterium* Ellen 345
  - Rhodopseudomonas palustris* BisB18
- 2005 CSP Microbe Projects**
  - Ignicoccus* sp. KIN4/1
- 2005 WFO Microbe Projects**
  - Chlamydia trachomatis* Ds2923
  - Chlamydia trachomatis* G9301
- 2004 DOE Microbe Projects**
  - Pelodictyon luteolum* DSM 273
  - Methanospirillum hungatei*, JF1
  - Nitrosospora multiformis* Surinam
  - Nitrosococcus oceani* ATCC 19707(ex *Nitrosomonas* oceani)
  - Nitrobacter winogradskyi*, Nb-255
  - Frankia* sp., Cc13
  - Anaeromyxobacter dehalogenans*, 2CP-C
  - Thiomicrospira crunogena* XCL-2
- 2001 Microbes**
  - Cytophaga hutchinsonii* ATCC 33406
  - Methanosarcina barkeri* fusaro
  - Pseudomonas fluorescens* PFO-1
- 2002 Microbes**
  - Ehrlichia canis*
  - Saccharophagus* (ex *Microbulbifer*) *degradans* 2-40
  - Ralstonia eutropha* JMP134
  - Dechloromonas aromatica* RCB
  - Pseudomonas syringae* B728a
  - Rhodospirillum rubrum* ATCC 11170
  - Desulfovibrio desulfuricans* G20
  - Geobacter metallireducens* GS-15
  - Psychrobacter* sp. 273-4
  - Methanococcoides burtonii*
  - Haemophilus somnus* 129PT
- 2004 GTL**
  - Rhodoferrax ferrireducens* DSM 15236
  - Pelobacter carbinolicus* DSM 2380
- Organism: 2003 Microbes**
  - Prochlorococcus* sp. MIT9312
  - Methylobacillus flagellatus*
  - Chlorobium chlorochromatii* Cd3 (ex *Chlorochromatium aggregatum*)
  - Moorella thermoacetica* ATCC 39073
  - Anabaena variabilis*
  - Synechococcus elongatus*
  - Methylobium petroleophilum* PM1(ex *Rubrivivax gelatinosus* PM1)
  - Burkholderia strain* 383(ex *R-18194*)
- 2001 Microbes**
  - Rhodobacter sphaeroides* 2.4.1
  - Thermobifida* *Fusca* YX
  - Burkholderia xenovorans* LB400 (ex *Burkholderia fungorum*)
  - Nostoc punctiforme* ATCC 29133
  - Novosphingobium* (ex *Sphingomonas*) *aromaticivorans*
  - Synechococcus* *WH8102*
  - Rhodopseudomonas palustris* CGA009
- 2003 Microbes**
  - Nitrosomonas europaea*
  - Prochlorococcus marinus* MED4
  - Prochlorococcus marinus* MIT9313
  - Bacillus anthracis* Sterne
  - Bacillus thuringiensis* 9727
  - Bacillus thuringiensis* ALH
  - Bacillus thuringiensis* Zebra Killer
  - Brucella abortus* 2308
  - Francisella philomiragia* 2773039
  - Francisella tularensis* OR-960463

## Steps in the Sequencing Process



## JGI-wide Finishing Standards

- All low quality areas (<Q30) should be reviewed and re sequenced.
- Final error rate should be < 0.2 per 10 Kb.
- No single clone coverage, i.e. minimum of 2X depth everywhere.
- Manually inspect and quantify single stranded regions.
- Check all high quality discrepancies.
- Verify all repeats (paired ends and PCR if necessary).
- Make sure to check ends of final contigs (chromosomes, plasmids)
- Using Assembly Viewer and phrapViewer tools check correctness of final assembly. Confirm questionable areas with PCR.

135 Microbial projects in finishing  
65 – Microbial genomes finished

## Genome Analysis

finished/draft	JGI	Total
Bacteria	43/67	256/117
Archaea	1/3	23/3
Eukarya	0/0	12/3
Viruses	0/0	260/0
All Organisms	44/90	551/123
Grand Total	134	674

Next IMG release: March 1, 2006