

The Genome Sequencer FLX System

Metagenomics and Microbial Diversity



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Get the complete picture with long sequence reads

Obtaining an unbiased view of the phylogenetic composition and functional diversity within a microbial community is a central objective of metagenomic analysis. Until now, technical and economic constraints limited the depth of analysis necessary for obtaining a representative picture of microbial and viral communities, their metabolic profiles, and their adaptation dynamics.

The Genome Sequencer FLX System, with its long, highly accurate reads, enables a comprehensive view into the diversity and metabolic profile of your metagenome of interest. Learn how 454 Sequencing systems can help you answer your research questions.

Microbial and metabolic diversity

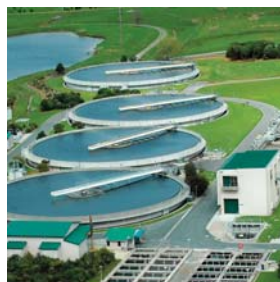
**Metatranscriptomics -
Environmental
expression profiles**

Gene content



**Microbial
abundance**

Viral pathogens



Signature profile of a specific environment

Microbial and metabolic diversity – Obtain an unprecedented picture of diversity with long specific reads

- Gain an in-depth perspective into the diversity and metabolic profile of an environmental habitat with rapid shotgun or bacterial 16S sequencing.
- Use long 400 bp sequence reads to achieve the specificity needed for comparison with nucleotide and protein databases.
- Survey an environment in a single GS FLX Titanium run or less, without traditional cloning steps.
- Assemble long reads into contigs to gain whole or partial genome information.

Microbial abundance – Gain unbiased insights into the relative representation of species within a habitat

- Count environmental gene tags (sequence reads specific for a species or a gene) to analyze the relative abundance of microbial species under varying environmental conditions.

Gene content – Discover new genes in unculturable species

- Discover genes quickly and make proper functional predictions within gene-rich microbiomes through the GS FLX System's sequencing depth and long, accurate reads.
- Screen microbiomes for enzymes of interest such as proteases or polymerases, and rapidly amplify them from the sample.

Metatranscriptomics – Analyze gene expression within microbial communities

- Analyze microbial diversity and gene regulation simultaneously within diverse environments through rapid gene expression profiling.
- Perform functional annotation with long sequence reads that unambiguously distinguish similar transcripts in a complex sample.
- Survey an environmental transcriptome in a single GS FLX Titanium run or less.

Viral pathogen detection – Only long reads can find “the needle in the haystack”

- Analyze viral outbreaks quickly and accurately by sequencing fragments of amplified RNA preparations from infected individuals.
- Identify the viral pathogen by matching the long reads to nucleotide or protein databases.

Signature profiling – Identify the molecular footprint of specific environments

- Take advantage of the GS FLX Titanium's long shotgun reads for straight forward analysis of a wide range of parameters unique to a specific environment.
- In a single instrument run, establish the signature profile of a microbial community by determining its members, predicting their function, and detecting their relative abundance.

Publicly available bioinformatic tools for 454 metagenomic analysis:

1. The SEED database/metagenomics RAST server: <http://metagenomics.nmpdr.org/>
2. NCBI non-redundant Clusters of Orthologs, COG: <http://www.ncbi.nlm.nih.gov/COG/>
3. Kyoto Encyclopedia of Genes and Genomes, KEGG: <http://www.genome.jp/kegg/>
4. MEGAN: <http://www-ab.informatik.uni-tuebingen.de/software/megan/welcome.html>
5. CARMA: <http://www.cebitec.uni-bielefeld.de/brf/carma/carma.html>
6. Integrated Microbial Genomes with Microbiome samples, IMG/M: <http://img.jgi.doe.gov/m>
7. CAMERA: <http://camera.calit2.net>
8. Publicly available 16S databases: *e.g.*, <http://rdp.cme.msu.edu/>

Selected References

Microbial and metabolic diversity

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Microbial abundance

McKenna P *et al.* (2008) The macaque gut microbiome in health, lentiviral infection, and chronic enterocolitis. *PLoS Pathog.* **4**(2):e20.

Gene content

Turnbaugh PJ *et al.* (2006) An obesity-associated gut microbiome with increased capacity for energy harvest. *Nature*. **444**:1027-1031.

Metatranscriptomics

Frias-Lopez J *et al.* (2008) Microbial community gene expression in ocean surface waters. *PNAS*. **105**(10):3805-10.

Viral pathogen detection

Cox-Foster DL *et al.* (2007) A metagenomic survey of microbes in honey bee colony collapse disorder. *Science*. **318** (5848):283-7.

Palacios G *et al.* (2008) A new arenavirus in a cluster of fatal transplant-associated diseases. *N Engl J Med*. **358**(10):991-8.

Signature profiling

Dinsdale EA *et al.* (2008) Functional metagenomic profiling of nine biomes. *Nature*. **452**(7187):629-32.

More references can be found at www.454.com

“The 454 Sequencing platform is rapidly democratizing high-throughput sequencing by allowing individual investigators to produce datasets on the level of a genomic center. My lab can now go into a new ecosystem and determine what the microbes and viruses are doing in a matter of weeks, instead of years. Not only can we figure out the functions of the microbes at a much faster rate, we can do a much better job because of the more detailed information provided by the long, accurate reads.”



— Forest Rohwer,
Associate Professor,
San Diego State University

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