The Use of Metagenomic Approaches to Analyze Changes in Microbial Communities

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Abstract: Microbes are the most abundant biological entities found in the biosphere. Identification and measurement of microorganisms (including viruses, bacteria, archaea, fungi, and protists) in the biosphere cannot be readily achieved due to limitations in culturing methods. A non-culture based approach, called “metagenomics”, was developed that enabled researchers to comprehensively analyse microbial communities in different ecosystems. In this study, we highlight recent advances in the field of metagenomics for analyzing microbial communities in different ecosystems ranging from oceans to the human microbiome. Developments in several bioinformatics approaches are also discussed in context of microbial metagenomics that include taxonomic systems, sequence databases, and sequence-alignment tools. In summary, we provide a snapshot for the recent advances in metagenomics approach for analyzing changes in the microbial communities in different ecosystems.

Keywords: metagenomics, sequencing, microbial diversity, bioinformatics, microbial changes

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Introduction

In the natural environment, constant polymicrobial interaction(s) occur between bacteria, viruses, protozoa, protists, archaea and fungi. These microbes do not exist in isolation and are often found in a dynamic “consortia” of different microbial species populations.\(^1\) Understanding microbial population dynamics in a consortium will benefit the genomic information of all coexisting members. Isolating and sequencing the genome of an individual organism from a consortium might not be adequate as the single isolate cannot be a representative of the full genetic and metabolic potential of its associated members. Moreover, achieving culture conditions for isolating a single member from a consortium would be a daunting task. Traditional microbiologists were always dependent on culture-based techniques for the identification of microbes in environmental samples. The challenge of identifying uncultured organisms was totally ignored. However, an explosion of knowledge in the field of microbial physiology and genetics happened during 1960s to mid-1980s wherein some scientists came to believe that cultured microorganisms did not represent the whole microbial world. This was evidenced by the “great plate count anomaly” showing discrepancy in the microbial numbers between dilution plating and microscopy.\(^2\) From then on, several independent studies supported the rise of this uncultured world of microbes.\(^3\)

New non-culture based approaches have recently been developed that can be extensively used for comprehensive analysis of different communities in a microbial consortia.\(^1,4-6\) Metagenomics or genomic studies of microorganisms refer to an non-culture based approach for collectively studying sets of genomes from a mixed population of microbes.\(^1\) The term “Metagenomics” was first coined by Handelsman and his colleagues in their study of natural products from soil microbes.\(^1\) Community genomics, environmental genomics, and population genomics are often used as synonyms for metagenomics. The field of metagenomics was initially started by an idea from Pace in 1985\(^5\) that subsequently lead to several studies, starting from the first cloning of DNA directly from environmental samples in a phage vector,\(^8\) and culminating in the direct random shotgun sequencing of environmental DNA.\(^9,10\) Since the use of the metagenomics approach in these independent studies, several other studies have come to use this approach to study microbial populations in a wide range of samples ranging from the oceans to humans.\(^8,11-22\)

Metagenomics has also provided significant information on the “changes” in the microbial community. For example, using a metagenomic approach, studies have elucidated changes in the microbial composition in humans fed on different diets.\(^22,23\) Similarly, metagenomics have provided information in the changes of microbial composition in ticks collected from different geographic regions.\(^15\)

Metagenomic studies can be grouped into four categories based on different screening methods: (a) shotgun analysis using mass genome sequencing; (b) genomic activity-driven studies designed to search for specific microbial functions; (c) genomic sequence studies using phylogenetic or functional gene expression analysis; and (d) next generation sequencing technologies for determining whole gene content in environmental samples.\(^4,6,18,24-28\) These four methods can be sub-classified under unselective (shotgun analysis and next generation sequencing) and targeted (activity-driven and sequence-driven studies) metagenomics.\(^4,6,15,24,26,28,29\) Some studies have used an unselective metagenomic approach extensively because of its cost-effectiveness and simplicity in DNA sequencing.\(^30\)

In this review we summarize recent advances in the field of metagenomics in studying changes in bacterial and viral communities from different ecosystems, provide a snapshot of metagenomic analysis and applications of the metagenomic approach, and discuss the development of some of the approaches to answer the challenges faced in accessing metagenomic data.

Experimental Design for Metagenomic Analysis

A common sequence-based metagenomic approach involves steps that are outlined in Figure 1. Due to high experimental costs incurred in metagenomics projects, there is a definite requirement of proper experimental design with appropriate replication and statistical analyses. For example, the approximate cost to produce metagenomic data from one gram of soil requiring 6000 HiSeq2000 runs would cost $ 267 million.\(^31\) A proper experimental design should ideally start with a question rather than technical or
operational restriction. As the ultimate aim of metagenomic projects is to link functional and phylogenetic information of microbial communities to the chemical, physical, and other biological parameters that characterize the environment, suitable reference samples for comparison should be considered and emphasized in the experimental design. The biological or technical variations that may arise during the experiment should not be neglected and should instead be considered carefully in planning the experiment. As microbial systems are dynamic, temporal sampling from an environment can have substantial impact on data and interpretation. Proper replicates need to be included in the experimental design and should also consider the level at which replication takes place. In summary, a well-planned experimental design in metagenomic projects would facilitate integration of data sets into new or existing ecological models.

Sample Processing for Metagenomic Analysis
Sample processing is the first step of any metagenomic project. The DNA that will be used for metagenomic analysis should be representative of all cells present in the sample and should be ideal for generation of genomic libraries. High quality DNA extractions that include robust DNA extraction procedures are now readily available. Some of the common DNA extraction procedures, such as use of fractionation or selective lysis for isolating target DNA associated with a host, physical separation, and isolation of cells from the samples (e.g., soil samples) or Direct lysis of cells in the soil matrix, have been reported. Metagenomic analysis requires high nanogram to microgram amounts of DNA. In the case of samples that yield less DNA, amplification methods for the DNA is recommended. Multiple displacement amplification using random hexamers and phage phi29 polymerase has been reported to successfully amplify femtograms of DNA in order to produce micrograms of product.

Metagenomic Sequencing
The metagenomics approach was originally focused on bacterial communities, but since been used to explore a wide range of microorganisms.
Recently, several methods including Shotgun sequencing have been extensively used in metagenomic studies. In Shotgun metagenomics, DNA isolated from an environmental sample is randomly sheared, sequenced in short fragments, and reconstructed into consensus sequences. With this method, detection of several microbes that would otherwise go unnoticed in culturing techniques was successful in environmental samples. With the recent development of next-generation sequencing (NGS) (both 454/Roche and Illumina/Solexa systems), the whole of metagenomic sequencing has shifted from Sanger sequencing technology. However, Sanger sequencing is still considered for sequences with large insert sizes and a read length exceeding 700 base pairs. Emulsion polymerase chain reaction is performed to clonally amplify random DNA fragments that are then attached to microscopic beads when NGS is performed using 454/Roche sequencer. The Beads attached to DNA fragments are deposited into picotitre plate followed by individual and parallel pyrosequencing. In the case of the Illumina/Solexa system, DNA fragments are immobilized on a surface and then solid-surface PCR amplification is performed. The amplified DNA fragments are then sequenced using reversible terminators in a sequencing-by-synthesis process.

A typical bacterial metagenomic analysis of environmental bacteria survey requires the use of the whole 16S ribosomal RNA (rRNA) gene. However, due to the read length restriction in NGS procedures, most surveys are aimed at characterizing selected hyper-variable regions of the 16S rRNA gene. The primary and secondary structures of the 16S rRNA gene show nine hyper-variable regions flanked by relatively conserved regions. This property makes hyper variable regions of 16S rRNA gene an optimal species molecular marker. Recent studies have shown comparable results between sequencing of hyper-variable regions and sequencing of a full-length 16S rRNA gene. Based on these studies, it is recommended to design oligonucleotides for the V1-V3 region or V4-V7 region for Archaea and the V1-V3 region or V1-V4 region for bacteria.

**Metagenomic Sequence Assembly, Binning, and Annotation**

The sequenced DNA fragments are then processed for assembly using one of the two strategies, either reference-based assembly (co-assembly) or de novo assembly. Software packages such as Newbler (Roche), AMOS (http://sourceforge.net/projects/amos/), or MIRA can be employed to perform reference-based assembly. For de novo assembly, tools based on the de Bruijn graphs are created to handle very large amounts of data. In addition, two new assembly programs (Meta Velvet and Meta-IDBA) have been developed to deal with the non-clonality of natural populations. The sequenced information is then processed for binning to sort DNA sequences into taxonomic groups that might represent individual or closely related genomes. Several algorithms employing different methods of grouping sequences have been developed, including but not limited to Phylopythia, S-GSOM, PCAHIER, TACAO, IMG/M, MG-RAST, MOTHUR, MEGAN, TANGO, CARMA, SORT-ITEMS, MetaPhyler, PhymmBL and MetaCluster. These algorithms have been developed depending on the type of input data generated from metagenomic sequencing.

Generally, metagenomic sequences are annotated in two steps: (a) Feature prediction is performed by identifying characteristics of interest within genes; and (b) functional annotation is performed by assigning putative gene functions and taxonomic neighbors. Several tools such as MG-RAST, IMG/M, FragGeneScan, MetaGeneMark, Metagene, and Orphelia have been developed for classifying sequence stretches as either coding or non-coding. BLAST-based searches are also used for potentially identifying any missing information from these programs. Some of the other tools that are employed for predicting non-protein coding genes are tRNAs, Signal peptides, and CRISPRs. Other primary online sources for obtaining annotated nucleotides sequence information include the International Nucleotide Sequence Database Collaboration (INSDC), the DNA Data Bank of Japan, the European Nucleotide Archive, GenBank, and the Sequence Read Archive (SRA). By mid-September 2010, the SRA had accumulated more than 500 billion reads consisting of 60 trillion base pairs available for download. SRA contained 80% of the sequencing data from the Illumina GA platform, as well as 15% and 5% from the SOLiD TM and Roche/454 platforms, respectively. Functional annotation of the metagenomic data is a major challenge, as only a small percentage of metagenomic
sequences are annotated. The sequences that cannot be annotated, either because they might simply reflect erroneous coding sequences, because they might be real genes but encode for unknown biochemical functions, or because they may not have homology to known genes, are all grouped as ORFans. Additional reference databases such as KEGG, eggNOG, COG/KOG, PFAM, and TIGRFAM are all available online tools that can be used to study functional properties of ORFans.

Statistical Analysis and Data Sources
A typical metagenomic project contains an enormous amount of data that needs careful evaluation using proper statistical methods. Primer-E-Package is a popular tool that can perform a range of multivariate statistical analysis. This package includes generation of multidimensional scaling plots, analysis of similarities (ANOSIM), identification of the species, and identification of gene functions (SIMPER). There is also a web-based tool called Metastats that has been used in recent studies. The Shotgun-FunctionalizerR package also provides several statistical programs to evaluate functional differences between samples. Due to the increasing number of metagenomic studies, it is important to deposit large sets of metagenomic data into databases. Deposition of metagenomic data in centralized services would not only facilitate comparative analysis of different metagenomic data but also facilitate a new level of organization and collaboration among researchers. Services like IMG/M, CAMERA, and MG-RAST are three prominent metagenomic databases that are available for large-scale metagenomic analysis.

Metagenomics to Study Microbial Diversity in Environment
In the last decade, several studies have used the metagenomic approach and provided comprehensive data on microbial communities in different ecosystems. It is estimated that, depending on the sample and methods used, the number of bacteria in soil may vary from 467 species to 500,000 species. Curtis and colleagues have speculated that bacterial content may range up to $4 \times 10^9$/ton of soil and the numbers of bacteria are unlikely to exceed $2 \times 10^8$ in the sea. These comparisons clearly suggest that microbial content is several orders of magnitude less in the sea in comparison to soil environments. The members of the archaeal phylum Crenarchaeota are shown to be predominant microorganisms found in the depth of ocean with estimated numbers $1.3 \times 10^{28}$ in global oceans, and total bacterial cells are estimated to be $3.1 \times 10^{28}$. It has also been noticed that in certain regions and at certain times, 50% of the bacterial community in surface waters consists of members belonging to SAR11 clade. Metagenomic ocean surveys have also led to several surprising discoveries. For example, using anchored chromosome walking, a 130 Kb BAC clone was isolated from uncultivated SAR86 bacterium (a bacterium belonging to alpha proteobacteria that is abundantly found in ocean surface waters). Sequencing of the 130 Kb fragment resulted in the identification of a new class of genes of the rhodopsin family for the first time in bacteria. Further studies on this class of genes in Escherichia coli proved its function as a light-driven proton pump. In summary, this study discovered a new type of light-driven energy generation in oceanic bacteria which subsequently led to the identification of several photoproteins in the Sargasso Sea.

Soil is one of the most challenging environmental sources to analyze microbial diversity. Several parameters of soil, such as particle size, permeability, porosity, water content, mineral composition, and plant cover, can influence microbial composition. In addition, other factors such as collection and storage of soil sample, DNA extraction methods, host-vector systems used for DNA cloning, and representative soil sampling, can also influence the results of microbial content. With the advent of various technical developments, several landmark studies have been performed using the metagenomics approach. By direct cloning into plasmid, cosmid, or BAC vectors, novel genes from soil microbes that encode enzymes and antibiotics have been discovered. These genes share little homology with known genes, thus illustrating the enormous potential of soil metagenomics in isolating novel classes of genes. Some of the genes that were isolated from soil microorganisms include lipases, proteases, oxidoreductases, amylases, antibiotics, antibiotic resistance enzymes, and membrane proteins.

Using a metagenomics approach several studies have provided a wealth of information on microbial diversity in extreme environmental conditions.
Studies from Barns and colleagues have provided information on microbial diversity in hot spring environments.\textsuperscript{94,95} Archaea similar to \textit{Crenarchaeota} phylotype are found to be the abundant species in Yellowstone National Park hot springs.\textsuperscript{94,95} Analysis in the same hot spring revealed more bacterial numbers distributed in twelve new division-level lineages.\textsuperscript{94,95} Furthermore, Blank and colleagues showed differences in microbial content in the samples collected from different Yellowstone National park hot springs at close proximity with similar temperatures and comparable pH values.\textsuperscript{13} Sequencing of polar ice caps has revealed the presence of algal population and several heterotrophic bacteria in ice matrix at low temperatures and low levels of light.\textsuperscript{14,96} Similar findings were noted in the analysis of microbial composition in the cryoconite hole of a glacier.\textsuperscript{16} Recent studies have also found dominance of archaea \textit{Salinibacter ruber} in hyper-saline environments.\textsuperscript{97} Non-thermal environments with extreme acidic conditions have also been shown to contain archaea of \textit{Ferroplasma} and \textit{Thermoplasm} groups. In addition, several bacterial species that include \textit{Acidiphilium}, \textit{Acidithiobacillus}, \textit{Leptospirillum} and \textit{Sulfobacillus} have also been found to be abundant in extreme acidic environments.\textsuperscript{11}

Although these studies indicate the important role of microorganisms in biogeochemical cycles, many details remain unclear; until we fully understand the nature of microbial diversity in different environments, this will remain as an important area of investigation.

**Viral Metagenomics**

The development of metagenomic approaches has revolutionized evaluation of viral particles in environmental samples. The results from more than 24 independent studies have already been published.\textsuperscript{98} These studies highlight that 50\% of viral sequences are “unknown”. Of the remaining 50\% “known” sequences, many had low amino acid similarities to known viral proteins and thus represent an uncategorized group.\textsuperscript{20} These findings suggest a more complex diversity of viral genomes in comparison to bacteria in environmental samples. This is consistent with the findings that 30\% of the open reading frames in sequenced viral genomes are ORFans, compared to 9\% ORFans from bacteria.\textsuperscript{99} Despite these challenges, viral metagenomics have developed methods to catalogue viruses in environmental samples based on identifiable sequences. Full genome sequences of novel viruses that were identified from different environments have already been reported and assembled.\textsuperscript{100} Based on genomic structure and taxonomic metagenomic analysis, some of the studies have linked viruses with their potential hosts.\textsuperscript{101,102}

Over the past decade, several studies using metagenomics have provided a substantial amount of information in the identification of new viruses from human samples.\textsuperscript{103–106} Most of the infectious diseases caused by viruses were documented before the identification of their causative agent. For example, Egyptian literature from approximately 3700 BC provided information on poliomyelitis. However, the causative agent for this disease was identified as poliomyelitis virus in 1909 AD.\textsuperscript{107} Similar descriptions of clinical conditions likely caused by Smallpox were found in ancient literature from India 1500 BC long before the isolation of the Variola virus.\textsuperscript{108,109} With the steady rise in the development of viral metagenomics, several novel viruses have been isolated within a short amount of time that are associated with disease outcomes in humans.\textsuperscript{98,104–106} Novel viruses including Borna virus, Arena virus, Paralysis virus, LUJO virus, Astroivirus as etiology of mink shaking syndrome, Simian hemorrhagic fever virus, and Klassevirus have been identified by metagenomic approaches as causes of diseases in humans and other mammals.\textsuperscript{110–116} In addition, a recent study has provided important information in the identification of several viruses in a public-health setting.\textsuperscript{104} These studies highlight future perspectives on the use of metagenomic approaches for generating enormous amounts of data in the identification of unknown and potentially infectious agents to humans, all in a short amount of time. Recent metagenomic analysis also addressed changes in the viral communities in Cystic Fibrosis and compared them to those of non-cystic fibrosis individuals.\textsuperscript{105} In addition, interest in tapping the vast novelty of viral genetic information, especially phages, has brought great attention to the use of metagenomics in this field.\textsuperscript{101} Overall, metagenomics has provided substantial insights to virus–host interactions and viral diversity in different environments.

**Tick Metagenomics**

Ticks are medically important arthropod vectors that transmit pathogens causing various human diseases.\textsuperscript{117}
The advancement of metagenomic approaches has facilitated research in studying microbial communities associated with medically important arthropod vectors.\textsuperscript{15,118} Using 454/Roche and Illumina-based metagenomic sequencing, Carpi et al have evaluated pathogen load and microbiome in \textit{Ixodes ricinus} ticks. \textit{MEGAN} comparison of the bacterial taxonomic profiles determined that a total of 108 genera belonging to all bacterial phyla were present in \textit{I. ricinus} ticks.\textsuperscript{15} Their study determined that, in addition to mutualistic bacteria such as \textit{Wolbachia} and \textit{Rickettsiella}, pathogenic bacteria such as \textit{Borrelia}, \textit{Rickettsia} and \textit{Candidatus Neoehrlichia} were also present in ticks. The bacterial content varied in ticks collected from different geographic regions and at different life stages, which might be due to the changes in environmental factors and host-selection behaviors of ticks.\textsuperscript{15} Metagenomic-based studies such as this one would not only facilitate epidemiological surveillance of several zoonotic pathogens, but would also lead to the development of better strategies to control vector-borne human diseases.

**Industrial Metagenomics**

With the advent of the metagenomic approach to discover novel genes that encode various enzymes, antibiotics, photoproteins, and membrane proteins from environmental uncultured bacteria, several industries have shown interest in exploiting these resources for the development of commercially available compounds. Metagenomics has provided access to novel enzymes and biocatalysts that were not initially achievable by conventional cultivable bacteria.\textsuperscript{21,87,119–121} In fact, global sales for enzymes were estimated to be $2.3 million in 2003, a figure that includes sales of enzymes in detergents, food applications, agriculture/feed, textile processing, pulp/paper, leather, and production of fine and bulk chemicals.\textsuperscript{122} In light of increasing energy costs, environmental pollution, public health hazards, and recent global economic crises, the discovery of novel enzymes from metagenomic approaches can be viewed both as an opportunity and as a necessity. For example, Diversa, the largest biotech company focusing on the commercialization of metagenome technologies, has constructed and screened for various nitrilase gene sequences isolated from diverse environmental libraries.\textsuperscript{123} This nitrilase enzyme library was marketed to several fine-chemical and pharmaceutical industries.\textsuperscript{123} In summary, metagenomics has played a significant role in the identification of several bioactive molecules that have attracted interest from both academia and industrial companies.\textsuperscript{1,87,119–121}

**Metagenomic Application to Study Human Gut and Skin Microbiome**

Over the past decade, metagenomics have provided great insights to the human microbiome. Waddington used a metaphor and regarded microbiota as an essential “organ” of the human body capable of performing metabolic functions that human cells might not be able to perform.\textsuperscript{124,125} Several factors such as specific microbial species colonizing the gut, niches they occupy, time, space, factors unique to the environment of each human being such as different dietary needs, and interactions with host cells can all influence taxonomic composition of the human microbiome. Metagenomics have uncovered nearly 1000 human-associated microorganisms’ draft genome sequences, along with 3.3 million unique microbial genes derived from the intestinal tract of over 100 European adults.\textsuperscript{17,126,127} Analysis of intestinal microbial content of humans across various continents revealed that microbes were clustered in 3 groups that are termed as enterotypes.\textsuperscript{17,22,126,127} Metagenomics of the human gut microbiome also revealed interesting functions carried by microorganisms within the gut, ranging from its role in newly discovered signaling mechanisms, vitamin production synthesis, glycan production, amino-acid, and xenobiotic metabolism. Several studies have also reported that microbial composition of the human gut is greatly affected by genetic background, age, diet, and health status of the host.\textsuperscript{17,22,126–128} Differences in microbial content were seen in all age groups of human beings. Babies (breast fed and formula fed), healthy and malnourished infants, youngsters, the elderly, humans that were either lean or obese, and humans with inflammatory bowel diseases (IBD) showed differences in microbial composition.\textsuperscript{23,88,129–131} A metagenomic study from De Filippo et al showed that European children who consumed a diet high in animal protein, sugar, starch, and fat, and low in fiber showed differences in gut microbial content in comparison to children fed on vegetarian diet consisting of carbohydrates, fiber, and non-animal protein.\textsuperscript{127} Interestingly,
the microbiome of European children was enriched with Firmicutes and Proteobacteria, whereas the African microbiome was enriched with Actinobacteria and Bacteroidetes. Members of Xylanibacter and Prevotella were only present in Children from Europe. These results clearly suggest that host dietary habits influence gut microbial content. Metagenomics have also revealed an interesting link with microbial content in the gut to host metabolism and disease development.\textsuperscript{132} The causes of intestinal diseases such as IBD, Crohn’s disease (CD), and ulcerative colitis (UC) have all been linked with both human gene- and microbiome-associated factors.\textsuperscript{132} Several metagenomic and microarray studies have also revealed differences in microbial composition in CD patients in comparison to healthy individuals.\textsuperscript{89,131,133,134}

Recent studies have used metagenomic approaches in looking at the microbial diversity of the human skin.\textsuperscript{135–138} Skin serves as a good host of microbes that include both commensal and pathogenic bacteria. Determination of microbial diversity in skin revealed several interesting findings.\textsuperscript{135–138} Bacteria belonging to Proteobacteria, such as Pseudomonas species and Janthinobacterium species, were found to be abundant in both human and mice skin biopsies.\textsuperscript{136–138} The presence of other bacteria belonging to Alphaproteobacteria, Gammaproteobacteria, Betaproteobacteria, Actinobacteria species, such as Kocuria species, Pripionibacteria species, Firmicutes, and Bacteroidetes were all evident in human skin biopsies.\textsuperscript{136–138} There is substantial evidence that viruses also represent a significant part of the skin microbiome.\textsuperscript{135} The presence of beta and gamma-human papillomaviruses, polyomaviruses, and circoviruses on normal-appearing skin has been reported.\textsuperscript{135} With the increasing interest in understanding the role of the microbiome on human development, aging and disease, the field of metagenomics is rapidly advancing with new techniques and has become a most effective tool in this area of investigation.

**Future Directions**

Over the past decade metagenomics has undoubtedly benefited the scientific world in rapidly analyzing changes in microbial communities in different environments. Despite exhaustive research efforts, both in financial and intellectual terms, the underlying mechanisms of the relationship between the microbial communities to the environment or to human gut metabolism, aging, and disease remains unclear. Therefore, improvements in metagenomic techniques that involve functional microbiomic approaches need to be developed. In addition, development of novel metagenomic approaches that consider several geochemical parameters is highly warranted to evaluate the complexity of microbial population in extreme environments. Metagenomics has provided identification of several new microbial genes from different environmental samples.\textsuperscript{11,12,14,15,17,19,91,93,135} Heterologous gene expression is an important and challenging approach that is required to identify the function of new genes identified by metagenomic studies.\textsuperscript{139–142} Studies have successfully used a heterologous gene expression system to identify several antibiotic resistance genes.\textsuperscript{140,142–145} E. coli or other domesticated bacteria are commonly used to express genes identified from metagenomic approaches.\textsuperscript{1,140,142,144,145} However, the important limitation in this approach is that many genes, and indeed perhaps most genes, are not expressed in these bacteria.\textsuperscript{1,140,142,144,145} Therefore, improvement in heterologous gene expression systems and production of functional recombinant proteins would speed up the discovery of important biomolecules from different environments. Due to the sheer volume of metagenomic data that is continuously being generated, development of novel methods for analysis, data storage, and sharing is warranted. This would not only facilitate best use of metagenomic data for researchers, but would also lead researchers to the answers to some of the fundamental questions about microbes—namely “who they are” and “what they are doing”—in this complex world of microorganisms.

**Author Contributions**

Conceived and designed the experiments: GN, HS. Analyzed the data: GN, HS. Wrote the first draft of the manuscript: GN, HS. Contributed to the writing of the manuscript: GN, HS. Agree with manuscript results and conclusions: GN, HS. Jointly developed the structure and arguments for the paper: GN, HS. Made critical revisions and approved final version: HS. All authors reviewed and approved of the final manuscript.
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Author(s) disclose no potential conflicts of interest.

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References
1. Handelsman J. Metagenomics: application of genomics to uncultured microorganisms. Microbiol Mol Biol Rev. 2004;68:669–85.
2. Staley JT, Konopka A. Measurement of in situ activities of nonphoto-synthetic microorganisms in aquatic and terrestrial habitats. Annu Rev Microbiol. 1985;39:321–46.
3. Whang K, Hattori T. Oligotrophic bacteria from rendzina forest soil. Antonie Van Leeuwenhoek. 1998;54:19–36.
4. Riesenberg CS, Schloss PD, Handelsman J. Metagenomics: genomic analysis of microbial communities. Annu Rev Genet. 2005;38:525–52.
5. Schleper C, Jurgens G, Jousmies Chabotier M. Genomic studies of uncultivated archaea. Nat Rev Microbiol. 2005;3:479–88.
6. Shedure J, Ji H. Next-generation DNA sequencing. Nat Biotechnol. 2008;26:1135–45.
7. Pace NR, Stahl DA, Lane DJ, Olsen GJ. Analyzing natural microbial populations by rRNA sequences. ASM News. 1985;51:4–12.
8. Schmidt TM, DeLong EF, Pace NR. Analysis of a marine picoplankton community by 16S rRNA gene cloning and sequencing. J Bacteriol. 1991;173:4371–8.
9. Tyson GW, Chapman J, Hugenholz P, et al. Community structure and metabolism through reconstruction of microbial genomes from the environment. Nature. 2004;428:37–43.
10. Venter JC, Remington K, Heidelberg JF, et al. Environmental genome shotgun sequencing of the Sargasso Sea. Science. 2004;304:66–74.
11. Baker BJ, Banfield JF. Microbial communities in acid mine drainage. FEMS Microbiol Ecol. 2003;44:139–52.
12. Boja O, Aravind L, Konopka A. Evolution of gene expression by RNAs in marine and terrestrial environments. J Bacteriol. 2008;190:2169–73.
13. Blank CE, Cady SL, Pace NR. Microbial composition of near-boiling silica-depositing thermal springs throughout Yellowstone National Park. Appl Environ Microbiol. 2002;68:5123–35.
14. Bowman JP, McCannonn SA, Brown MV, Nichols DS, McMeekein TA. Diversity and association of psychrophilic bacteria in Antarctic sea ice. Appl Environ Microbiol. 1997;63:3068–78.
15. Carpi G, Cagnacci F, Wittekindt NE, et al. Metagenomic profile of the bacterial communities associated with Ixodes ricinus ticks. PLoS One. 2011;6:e25604.
16. Christner BC, Kvitko BH 2nd, Reeve JN. Molecular identification of bacteria and Eukarya inhabiting an Antarctic cryoconite hole. Extremophiles. 2003;7:177–83.
17. Claesson MJ, Cusack S, O’Sullivan O, et al. Composition, variability, and temporal stability of the intestinal microbiota of the elderly. Proc Natl Acad Sci U S A. 2011;108 Suppl 1:4586–91.
18. Cox-Foster DL, Conlan S, Holmes EC, et al. A metagenomic survey of microbes in honey bee colony collapse disorder. Science. 2007;318:283–7.
19. Curtis TP, Sloan WT, Scannell JW. Estimating prokaryotic diversity and its limits. Proc Natl Acad Sci U S A. 2002;99:10494–9.
20. Edwards RA, Rohwer F. Viral metagenomics. Nat Rev Microbiol. 2005;3:504–10.
21. Knietsch A, Waschkowitz T, Bowien S, Henne A, Daniel R. Metagenomes of complex microbial consortia derived from different soils as sources for novel genes conferring formation of carboxyls from short-chain polycarboxylates on Escherichia coli. J Mol Microbiol Biotechnol. 2003;5:46–56.
22. Maccaferri S, Biagi E, Brüggi P. Metagenomics: key to human gut microbiota. Dig Dis. 2001;29:525–30.
23. Monira S, Nakamura S, Gotoh K, et al. Gut microbiota of healthy and malnourished children in Bangladesh. Front Microbiol. 2011;2:228.
24. Harisemody O, Ng PC, Strausberg RL, et al. Evaluation of next generation sequencing platforms for population targeted sequencing studies. Genome Biol. 2009;10:R32.
25. Turnbaugh PJ, Ley RE, Mahowald MA, et al. An obesity-associated gut microbiome with increased capacity for energy harvest. Nature. 2006;444:1027–31.
26. Palenik B, Ren Q, Tai V, Paulsen IT. Coastal Synechococcus metagenome reveals major roles for horizontal gene transfer and plasmids in population diversity. Environ Microbiol. 2009;11:349–59.
27. Biddle JF, Fitz-Gibbon S, Schuster SC, Brenchley JE, House CH. Metagenomic signatures of the Peru Margin subseafloor biosphere show a genetically distinct environment. Proc Natl Acad Sci U S A. 2008;105:10583–8.
28. Edwards RA, Rodriguez-Brito B, Weyegy L, Haynes M, Breitbart M, et al. Using pyrosequencing to shed light on deep mine microbial ecology. BMC Genomics. 2006;7:57.
29. Turnbaugh PJ, Gordon JL. The core gut microbiome, energy balance and obesity. J Physiol. 2009;587:4153–8.
30. Chen K, Pachtler L. Bioinformatics for whole-genome shotgun sequencing of microbial communities. PLoS Comput Biol. 2005;1:106–12.
31. Desai N, Antonopoulos D, Gilbert JA, Glass EM, Meyer F. From genomics to metagenomics.Curr Opin Biotechnol. 2012;23:72–6.
32. Burke C, Kjelleberg S, Thomas T. Selective extraction of bacterial DNA from the surfaces of macroalgae. Appl Environ Microbiol. 2009;75:252–6.
33. Delmont TO, Rohe P, Clark L, Simonet P, Vogel TM. Metagenomic comparison of direct and indirect soil DNA extraction approaches. J Microbiol Methods. 2011;86:397–400.
34. Thomas T, Rutsh D, DeMaere MZ, et al. Functional genomic signatures of sponge bacteria reveal unique and shared features of symbiosis. ISME J. 2010;4:1557–67. Sesittisch A, Weilharter A, Gerzabek HH, Kirchmann H, Kandelers E. Microbial population structures in soil particle size fractions of a long-term fertilizer field experiment. Appl Environ Microbiol. 2001;67:4215–24.
35. Ishoey T, Woyke T, Stepanauskas R, Novotny M, Lasken RS. Genomic sequencing of single microbial cells from environmental samples. FEMS Microbiol Lett. 2010;30:228–34.
36. Ishoey T, Woyke T, Stepanauskas R, Novotny M, Lasken RS. Genomic sequencing of single microbial cells from environmental samples. FEMS Microbiol Lett. 2010;30:228–34.
37. Eisen JA. Environmental shotgun sequencing: its potential and challenges for studying the hidden world of microbes. PLoS Biol. 2007;5:e82.
38. Metzker ML. Sequencing technologies—the next generation. Nat Rev Genet. 2010;11:31–46.
39. Mardis ER. The impact of next-generation sequencing technology on genetics. Trends Genet. 2008;24:133–41.
40. Goltsman DS, Denef VJ, Singer SW, et al. Community genomic and proteomic analyses of chemoheterotrophic iron—oxidizing “Leptospirillum rubarum” (Group II) and “Leptospirillum ferrooxidantium” (Group III) bacteria in acid mine drainage biofilms. Appl Environ Microbiol. 2009;75:4599–615.

41. Bentley DR, Balasubramanian S, Swedlow HP, et al. Accurate whole human genome sequencing using reverse terminator chemistry. Nature. 2008;456:53–9.

42. Al Masalma M, Lonjon M, Richet H, et al. Metagenomic analysis of brain abscesses identifies specific bacterial associations. Clin Infect Dis. 2012;54: 202–10.

43. Lazarevic V, Whiteson K, Gaia N, et al. Analysis of the salivary microbiome using culture-independent techniques. J Clin Bioinformatics. 2012;2:4.

44. Shah N, Tang H, Doak T, et al. Comparing bacterial communities inferred from 16S rRNA gene sequencing and shotgun metagenomes. Pac Symp Biocomput. 2011:165–76.

45. Jeraldo P, Chia N, Goldenfeld N. On the suitability of short reads of 16S rRNA for phylogeny-based analyses in environmental surveys. Environ Microbiol. 2011;13:3000–9.

46. Kim M, Morrison M, Yu Z. Evaluation of different partial 16S rRNA gene sequence regions for phylogenetic analysis of microbiomes. J Microbiol Methods. 2011;84:81–7.

47. Chevreux B, Wetter T, Sahai S. Genome sequence assembly using trace signals and additional sequence information computer science and biology. Proc German Conf Bioinform. 1999:95–45–56.

48. Miller JR, Koren S, Sutton G. Assembly algorithms for next-generation sequencing data. Genomics. 2010;95(6):315–27.

49. Pevzner PA, Tang H, Waterman MS. An Eulerian path approach to DNA fragment assembly. Proc Natl Acad Sci U S A. 2001;98(17):9748–53.

50. Peng Y, Leung HC, Yiu SM, Chin FY. Meta-IDBA: a de Novo assembler for metagenomic data. Bioinformatics. 2011;27(13):94–101.

51. Chan CK, Hsu AL, Halgamuge SK, Tang SL. Binning sequences using very sparse labels within a metagenome. BMC Bioinformatics. 2008;9:215.

52. Diaz NN, Krause L, Goessmann A, Niehaus K, Nattkemper TW. TACOA: taxonomic classification of environmental genomic fragments using a kernelized nearest neighbor approach. BMC Bioinformatics. 2009;10:56.

53. Glass EM, Wilkening J, Wilke A, Antonopoulus D, Meyer F. Using the metagenomes RAST server (MG-RAST) for analyzing shotgun metagenomes. Cold Spring Harb Protoc. 2010;2010(1)pdb.prot5568.

54. Huson DH, Auch AF, Qi J, Schuster SC. MEGAN analysis of metagenomic data. Genome Res. 2007;17(3):377–86.

55. Krause L, Diaz NN, Goessmann A, et al. Phylogenetic classification of short environmental DNA fragments. Nucleic Acids Res. 2008;36(7):2230–9.

56. Markowitz VM, Chen IM, Chu K, et al. IMG/M: the integrated metagenome platform for visualizing and exploring metagenomic data. Nat Methods. 2007;4:95–102.

57. Krause L, Goesmann A, Niehaus K, Nattkemper TW. TACOA: taxonomic classification of environmental genomic fragments using a kernelized nearest neighbor approach. BMC Bioinformatics. 2009;10:56.

58. Zheng H, Wu H. Short prokaryotic DNA fragment binning using a hierarchical classifier based on linear discriminant analysis and principal component analysis. J Bioinform Comput Biol. 2010;8(6):995–1011.

59. Qin J, Li R, Raes J, et al. A human gut microbial gene catalogue established by metagenomic sequencing. Nature. 2010;464(7285):59–65.
139. Handelsman J, Rondon MR, Brady SF, Clardy J, Goodman RM. Molecular biological access to the chemistry of unknown soil microbes: a new frontier for natural products. Chem Biol. 1998;5:R245–9.

140. MacNeil IA, Tiong CL, Minor C, et al. Expression and isolation of antimicrobial small molecules from soil DNA libraries. J Mol Microbiol Biotechnol. 2001;3:301–8.

141. Rondon MR, August PR, Bettermann AD, et al. Cloning the soil metagenome: a strategy for accessing the genetic and functional diversity of uncultured microorganisms. Appl Environ Microbiol. 2000;66:2541–7.

142. Gillespie DE, Brady SF, Bettermann AD, et al. Isolation of antibiotics turbomycin a and B from a metagenomic library of soil microbial DNA. Appl Environ Microbiol. 2002;68:4301–6.

143. Rondon MR, Goodman RM, Handelsman J. The Earth’s bounty: assessing and accessing soil microbial diversity. Trends Biotechnol. 1999;17:403–9.

144. Rondon MR, Raffel SJ, Goodman RM, Handelsman J. Toward functional genomics in bacteria: analysis of gene expression in Escherichia coli from a bacterial artificial chromosome library of Bacillus cereus. Proc Natl Acad Sci U S A. 1999;96:6451–5.