

REPORT

Alignment-free tools for metagenomics-data analysis

Robert Deibel

Abstract

Metagenomics; as the study and analysis of microorganisms of biotopes, like the human gut, is a field of vast research where researchers have to deal with the giant sets of data gathered through NGS-methods. Since the amount of data puts stress on computation and time resources, the development of fast and light analysis tools is required. In this report I introduce the two main branches of analysis tools, while setting the focus on alignment-free methods.

While the alignment-based approach has its foundation in the alignment of a target sequence against a database – as seen with Smith-Waterman or BLAST – alignment-free methods have different approaches. Here I will showcase a selection of statistical and machine learning approaches and test selected methods on metagenomic data.

$D2z$, Hao and d_2^* are statistical approaches based on k -tupel count frequencies, similarity is measured by applying a metric and through interpretation with current biological knowledge.

Laczny *et al.* used k -mers as vectors in high-dimensional space and the BH-SNE of van der Maaten visualizing related data in two dimensional scatter plots, resulting in a tool with high accuracy for simulated as well as real-world metagenomes.

Especially for analysis of novel data, sampled from microbioms, alignment-free applications of metagenomics are essential for understanding the cooperation of microorganisms and for further research in immunology.

Keywords: alignment-free; machine learning; statistic; metagenome; report

Introduction

Metagenomics

A puddle of mud The metagenome is the whole set of genomes, coding or noncoding, of a population of microorganisms in a microbiome sample. The DNA of organisms is isolated from these samples. Metagenomics is the study and analysis of these metagenomes [1].

A microbiome consists of countless bacteria, archaea and viruses; for which >90% are unculturable, using sequencing and metagenomic analysis as a way to study these.

NGS – Next Generation Sequencing The sheer amount of metagenomic data – Kakirde *et al.*[2] states 10 Tb of DNA in a soil sample – resulted in advances of sequencing.

Nowadays new high throughput methods – also Next Generation Sequencing or NGS for short – are used to compute comparable data from real-world samples.

NGS is a term for methods of rapid parallelized sequencing, producing thousands or millions of reads concurrently.

Data analysis can then be carried out on these reads.

What do we want to achieve? Metagenomics is used in the design of antibiotics and medicine or to analyze the metabolism of microorganisms and its hosts; making it a rapidly developing field of research.

Here, I'm going to summarize two approaches to data analysis and showcase one of those in more detail.

The "classical" approach

Alignment-based methods are proven but don't include all possibilities The best known approach to analyze reads are the various alignment-based methods.

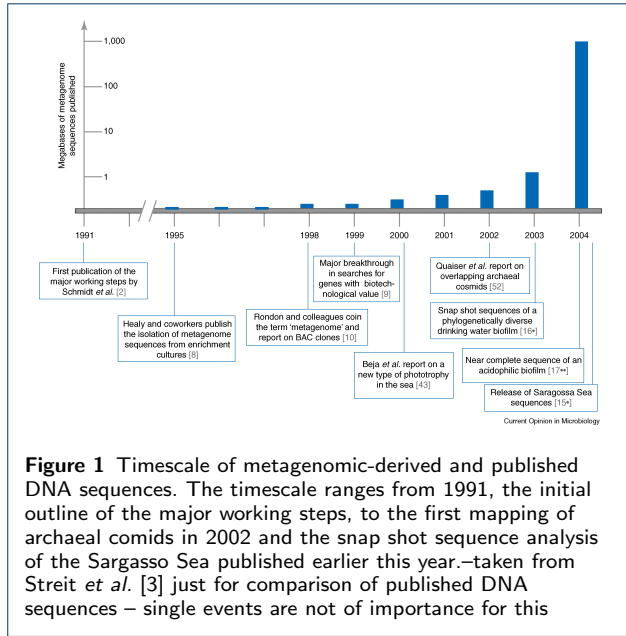
Sequences are aligned against a database of known genomes, the resulting profiles are analyzed based on several factors.

This approach well established under various conditions and implemented numerous times. BLAST for example, while not used for metagenomics anymore,

Correspondence: robert.deibel@student.uni-tuebingen.de

Eberhard-Karls Universität, Tübingen, DE

Full list of author information is available at the end of the article



has an accuracy well over 80%[4] and similar values are expected for other alignment-based tools. However, NGS supplies researchers with a lot of data to be analyzed. The analysis of metagenomes is computation heavy. BLAST – and therefore BLAST-like tools – aligns queries with the entries in a database. Applied on metagenomic data this method can be time consuming and has led to the development of faster and more effective methods for data analysis. Research of novel data, not listed in databases, is a main focus of metagenomics. This data stays unanalyzed following an alignment-based approach, resulting in a high demand for lightweight tools independent of databases.

Here, I will showcase methods with differed approaches to the analysis of such data.

An alternative for alignment-based analysis

Apart from alignment of sequences another way is basing the analysis on different factors associated with metagenomic data. In this report I reflect the work of Song *et al.* [5] and Laczny *et al.* [6] both presenting methods for the analysis of metagenomic-data using alignment-free approaches. Their work is based on statistical methods, and visualization and machine learning respectively.

Methods

In this section "power" refers to the statistical term, which is the probability that a test correctly rejects the null hypothesis. With H_0 the sequences are unrelated and H_1 the sequences are related under the underlying model. If not stated differently.

k -tupels as a measure of similarity

In the work of Song *et al.*[5] different methods based on k -tupel occurrences in sequences are presented. A k -tupel is a substring of sequences with length k .

By counting the occurrences of these k -tupels and applying a distance or dissimilarity metric, the tupels are clustered and these clusters analyzed using current biological knowledge. A metric would be based on the resulting counted k -tupel frequencies.

The next section will focus on methods of k -tupel counts as a measure of similarity for sequences.

The D_2 statistic and normalization by $D2z$ Torney *et al.*[7] introduced D_2 using k -tupel matches between sequences to define similarity.

$$D_2 = \sum_{w \in \mathcal{A}^k} X_w Y_w$$

where X_w and Y_w are the number of occurrences of string w in the corresponding sequence and \mathcal{A} is the alphabet.

Kantrovitz *et al.*[8] stated that the D_2 statistic depends on the underlying sequence model and performed a normalization to remove the bias. The resulting statistic is called $D2z$ and is defined as

$$D2z(A, B) = \frac{D_2(A, B) - E(D_2)}{\sqrt{Var(D_2)}}$$

The expected value and variance are calculated by Markov models for the used sequences.

$D2z$ was compared to five other measures of similarity – [5, 8] – through analysis of *cis*-regulatory modules (CRM), outperforming all of them. However $D2z$ requires two parameters; apart from k , r has to be specified, where r is the order of the sequence Markov chain.

Phylogenetic trees through statistics – CVTree Another approach utilizes the expected count of a k -tupel under the $(k - 2)$ -th order Markov chain, estimated by

$$E_w^X = \frac{X_w X_{w_2 \dots w_k}}{X_{w_2 \dots w_{k-1}}}$$

where w is a substring of length k , w_i is the letter at index i in w and X_w is the number of occurrences of w in a sequence A .

The correlation coefficient of the relative difference vectors with the expected count is then used to measure similarity of sequences.

$$Hao = \frac{1}{2} \left(1 - \frac{\sum_w \left(\frac{X_w - E_w^X}{E_w^X} \right) \left(\frac{Y_w - E_w^Y}{E_w^Y} \right)}{\sqrt{\sum_w \left(\frac{X_w - E_w^X}{E_w^X} \right)^2 \sum_w \left(\frac{Y_w - E_w^Y}{E_w^Y} \right)^2}} \right)$$

Notation is taken from Song *et al.*[5].

Hao calculates the frequencies of appearances of overlapping k -tupels indicated with X_w and subtracts a random background using the $(k-2)$ -th order Markov chain; this is to minimize the influence of random mutation. After computation of correlation C a normalization was defined by subtraction from 1 and multiplication with $\frac{1}{2}$.

$$C = \frac{\sum_w \left(\frac{X_w - E_w^X}{E_w^X} \right) \left(\frac{Y_w - E_w^Y}{E_w^Y} \right)}{\sqrt{\sum_w \left(\frac{X_w - E_w^X}{E_w^X} \right)^2 \sum_w \left(\frac{Y_w - E_w^Y}{E_w^Y} \right)^2}}$$

C describes the cosine of the angle between the sequences, where $C = 1 \Leftrightarrow A = B$ and $C = 0 \Leftrightarrow \forall a_i \in A, b_i \in B : a_i \neq b_i$.

For CVTree a distance matrix is computed by applying *Hao* to each pair of sequences, neighbor joining then constructs a phylogenetic tree to visualize similarity. CVTree was tested by Qi *et al.* on a set of 139 prokaryotic genomes computing a robust result[9].

The application of CVTree on virus data provided on the web service is seen in Figure 2.

Similarity through nucleotide frequency A related approach was presented by Karlin and colleagues, where they observed the relative di-nucleotide frequency defined by

$$\rho_{ab}(A) = \frac{f_{ab}}{f_a f_b}$$

where f_w is the frequency of w in a sequence.

It is stated that ρ_w is stable across a genome and differs in different genomes. The extension to tri- and tetra-nucleotides is achieved by

$$\gamma_{abc} = \frac{f_{abc} f_a f_b f_c}{f_{ab} f_{bc} f_{ac}}$$

and

$$\tau_{abcd} = \frac{f_{abcd} f_{ab} f_{ac} f_{ad} f_{bc} f_{bd} f_{cd}}{f_{abc} f_{abd} f_{acd} f_{bcd} f_{ab} f_{ac} f_{ad} f_{bc} f_{bd} f_{cd}}$$

l_p norm was applied as a dissimilarity measure as

$$\delta(A, B) = \sum_{j \in A} |\theta_j(A) - \theta_j(B)|$$

where A, B are sequences, $\theta \in \{\rho, \gamma, \tau\}$ and

$$j = \begin{cases} \{a, b\} & \text{if } \theta = \rho \\ \{a, b, c\} & \text{if } \theta = \gamma \\ \{a, b, c, d\} & \text{if } \theta = \tau \end{cases}$$

Evolutionary studies on viruses, bacteria, plasmids, prokaryotes and eukaryotes were performed using this measure[5].

D_2^S , D_2^* , d_2^S and d_2^* as powerful statistic measures The D_2^S statistic is defined by

$$D_2^S = \sum_{w \in \mathcal{A}^k} \frac{\tilde{X}_w \tilde{Y}_w}{\sqrt{\tilde{X}_w^2 + \tilde{Y}_w^2}}$$

where \tilde{X}_w and \tilde{Y}_w are the normalization of X_w and Y_w respectively

$$\tilde{X}_w = X_w - \bar{n} p_w^X$$

$\bar{n} = n - k$ and p_w^X is the probability of the k -tupel w under the background model of a sequence A . This is based on Shepp[12]. Where it was observed that for two normal random variables with mean zero, $XY/\sqrt{X^2 + Y^2}$ is also normally distributed.

D_2^* defined by

$$D_2^* = \sum_{w \in \mathcal{A}^k} \frac{\tilde{X}_w \tilde{Y}_w}{\sqrt{\bar{m} \bar{n} p_w^X p_w^Y}}$$

utilizes the idea that the number of occurrences of w is approximately Poisson distributed and mean and variance are the same.

Through simulations and theoretical studies the null hypothesis H_0 was tested against H_1 ; the conclusions were:

- 1 D_2^S and D_2^* have higher power than D_2 increasing with sequence length
- 2 D_2^* has the highest power when the length of k equals the *motif* length
- 3 For short sequences the power of D_2^* is higher while for long sequences D_2^S is generally more powerful.

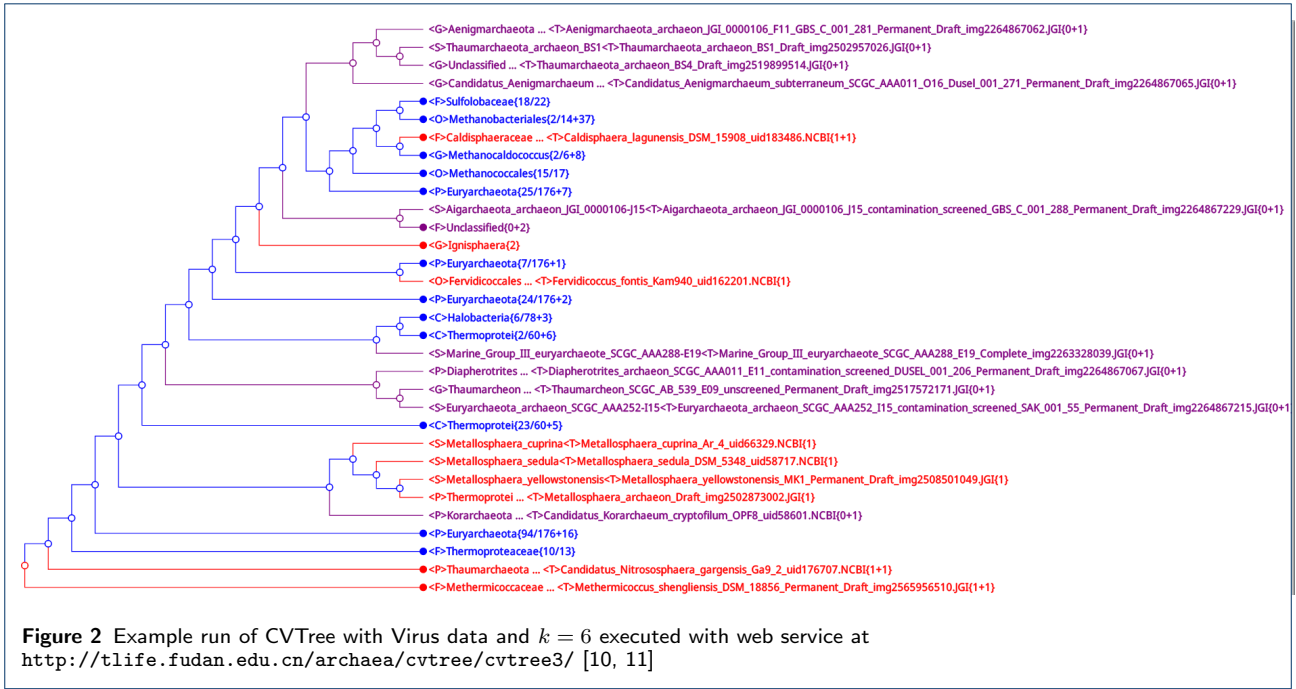
where *motifs* are significantly enriched word patterns[5].

Further normalization of D_2^S and D_2^* removes the property that the magnitude strongly varies depending on different factors. The resulting statistics are d_2^S and d_2^* respectively.

$$d_2^S = \frac{1}{2} \left(1 - \frac{D_2^S}{\sqrt{\frac{\sum_{w \in \mathcal{A}^k} \tilde{X}_w^2}{\sum_{w \in \mathcal{A}^k} \tilde{X}_w^2 + \tilde{Y}_w^2}} \sqrt{\frac{\sum_{w \in \mathcal{A}^k} \tilde{Y}_w^2}{\sum_{w \in \mathcal{A}^k} \tilde{X}_w^2 + \tilde{Y}_w^2}}} \right)$$

and

$$d_2^* = \frac{1}{2} \left(1 - \frac{D_2^*}{\sqrt{\frac{\sum_{w \in \mathcal{A}^k} \tilde{X}_w^2}{\bar{n} p_w^X}} \sqrt{\frac{\sum_{w \in \mathcal{A}^k} \tilde{Y}_w^2}{\bar{m} p_w^Y}}} \right)$$



d_2^S and d_2^* now hold the property that they are 0 when the sequences are the same and close to 1 if they are anti-correlated. Through this d_2^S and d_2^* can be used to cluster sequences of interest.

Modification of d_2^S and d_2^ accounting for mismatches*
The statistics mentioned above consider exact matches of tuples, since mutations are a fundamental part of evolution and DNA replication in general, mismatches should be considered when applying these methods. For a tuple w its neighborhood can be defined as $\zeta(w)$ with $w' \in \zeta(w)$ when w' has up to a certain number of mismatches with w and a weight a is applied, analogously reverse complements can be included in $\zeta(w)$. The statistics are modified as \tilde{X}_w is replaced by $\tilde{X}_{\zeta(w)}$ where $\tilde{X}_{\zeta(w)} = X_{\zeta(w)} - EX_{\zeta(w)}$ and

$$X_{\zeta(w)} = \sum_{w' \in \zeta(w)} a_{w'} X_{w'}$$

modifications for \tilde{Y}_w are analogous. Song *et al.* performed a series of tests to evaluate the effectiveness of the statistics Hao , d_2^S and d_2^* with consideration of mismatches. The neighborhood was defined as

$$\zeta(w) = \{w', rc(w') | dist_{hamming}(w, w') \leq 1\}$$

Testing was based on sequences taken from mouse embryos. The positive set was taken from the forebrain, midbrain, limb and heart tissues, while the negative set

was chosen from random samples of the same length with a maximum of 30% repetitive sequences[5]. For 500 samples of each set the dissimilarity was calculated for each pair in the respective set. A threshold for dissimilarity was applied on the resulting values, a score lower than the threshold was predicted as positive, one above indicated negatives. Through comparison with the real data false predictions were identified. Different parameters like tuple size k , Markov chain order and mismatch weight were applied.

Performance under the mismatch model Conclusions of these test were:

- 1 Hao performed worse than both d_2^S and d_2^*
- 2 d_2^S and d_2^* performed best with $k = 4$ and mismatch weight of around 0.05. However differences through mismatch weight were negligible. For $k = 5 \vee 6$ a weight close to 1 performed best.

Song *et al.* considered additional statistics for testing, which are not discussed in this report and were therefore not further accounted for.

Additional testing using metagenomes and NGS data was carried out[5], since usually the short reads generated through NGS reduce the power of the discussed statistics. The data consisted of 39 fecal samples of 33 mammalian hosts[13] 56 marine samples[14] and 13 human fecal samples[15] for metagenomes and tree species with unknown complete genomes as NGS data. d_2^S outperformed the other tested measures in terms of consistency and separation as was seen through the

tree samples and human feces metagenome respectively.

Overall d_2^S produced the best results compared to all mentioned statistics with Hao and d_2^* having similar outcomes.

Machine learning for alignment-free data analysis

In his work van der Maaten [16] introduced a machine learning variant – BH-SNE – based on the idea that closely related objects have a larger influence upon each other than unrelated ones. While these objects were originally intended to be points in a picture, Laczny *et al.*[6] used reads of metagenomes.

Barnes-Hut-SNE applies the Barnes-Hut algorithm and metric trees to modify the t-SNE method, commonly used in machine learning

Barnes-Hut and vantage-point trees for faster computation The Barnes-Hut algorithm is often used by astronomers to perform N -body simulations[16]. In this algorithm it is assumed that the force of objects with sufficient distance to each another is infinitesimal and thus can be ignored in further computation. Leading –in the case of BH-SNE – to a cut in objects to include in calculations.

Van der Maaten chose these objects based on vantage-point trees, where similar nodes are saved as the left, dissimilar nodes as the right child. After establishing the data structure one can search the tree and apply the given algorithm to the reduced set of nodes of interest

Sequence signatures as objects Observations suggest the existence of species-specific oligonucleotide signatures in genomic sequences[6, 17]. These consist of k -mers and can be represented as vectors in high-dimensional Euclidean space. For human interpretation these vectors need to be transformed in a two or three dimensional space [6].

vector construction is achieved through assignment of joint probabilities to the k -mers and a similarity function to the corresponding points in high-dimensional space. Utilizing a Kullback-Leibler divergence and the optimizations stated before the points can be optimized and learned.

Using center log-ratio (CLR)-transformed – a normalization step – oligonucleotide signatures and the BH-SNE approach of van der Maaten, Laczny *et al.* constructed a tool for application on metagenomic-data with sequence length of 1000 nt and 5-mers as oligonucleotide signatures. While these parameters produced the best results Laczny *et al.* stated that 600 nt might be an appropriate length for some applications, but

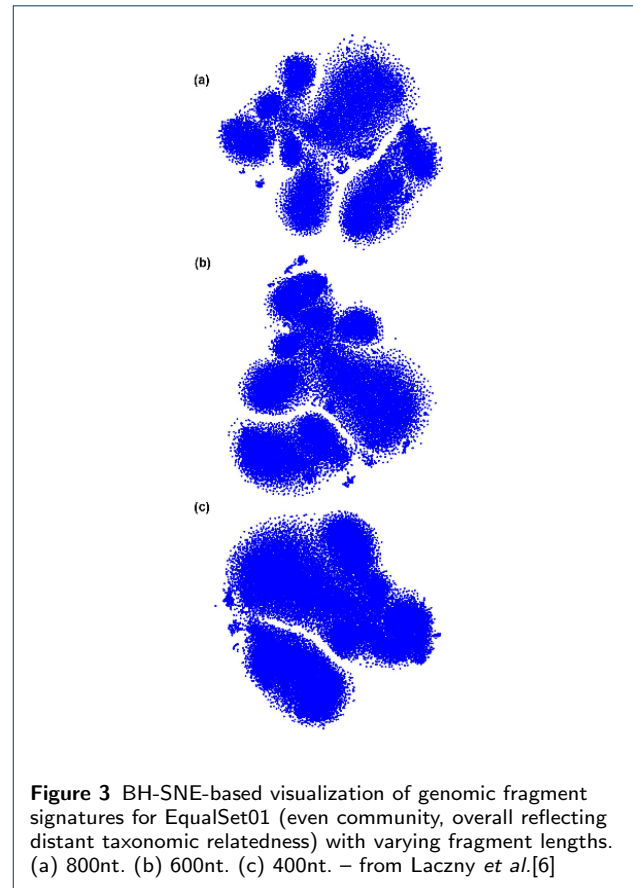


Figure 3 BH-SNE-based visualization of genomic fragment signatures for EqualSet01 (even community, overall reflecting distant taxonomic relatedness) with varying fragment lengths. (a) 800nt. (b) 600nt. (c) 400nt. – from Laczny *et al.*[6]

with lower values the separation would drop remarkably, as can be seen in Figure 3 through lesser separation of the clusters. Implementing their approach using 5-mers produced better congruency compared to transformed and untransformed 4-mers.

For Laczny *et al.* these 5-mers are the before mentioned objects, used for calculation of similarity in BH-SNE.

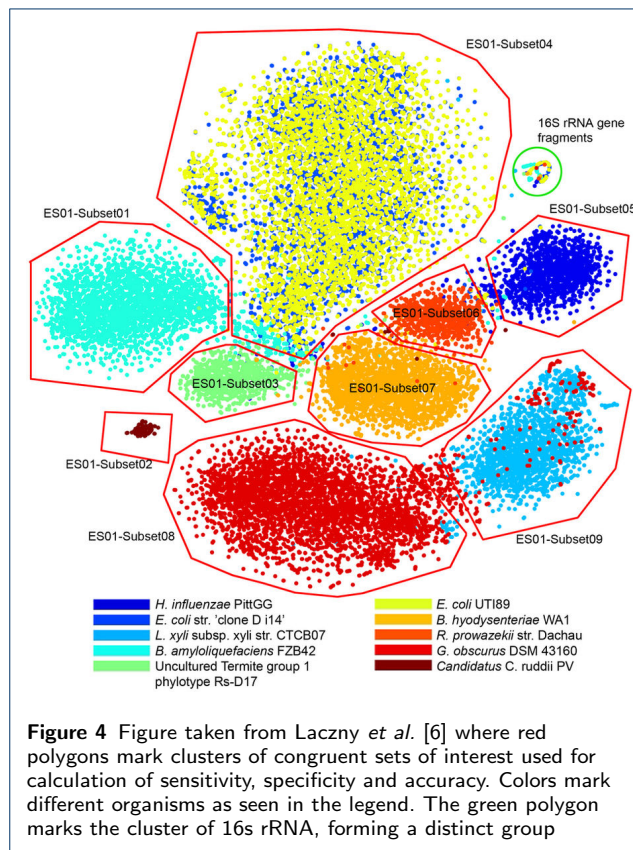
Cluster finding The tool was tested on several simulated data sets; EqualSet01, EqualSet02 and LogSet01. The genomes of organisms in these sets were equally and logarithmically distributed; among the equally distributed sets were genomes with small and high similarity respectively.

The equally distributed data was used for reasons of simplicity, in real-world metagenomes, DNA is never evenly distributed, the logarithmic set should simulate this real-world data with varying quantities of different genomes. High and low similarity sets test the discrimination capabilities of the tool.

After applying their tool on the simulated metagenomes their results showed distinct clustering for different species as depicted in Figure 4 for EqualSet01 and

LogSet01, respectively. Clustering of EqualSet02 resulted in overlapping of closely related organisms and separation of more distant relatives.

Overall the runs on simulated data resulted in high sensitivity, specificity and accuracy. A selection of values is outlined in Table 1. The calculation was performed by enclosing clusters with polygons as seen in Figure 4. Points inside represent the positives, points outside the negatives. Similar outputs were achieved by fitting (semi-)automated Gaussian Mixture models to calculate these values.



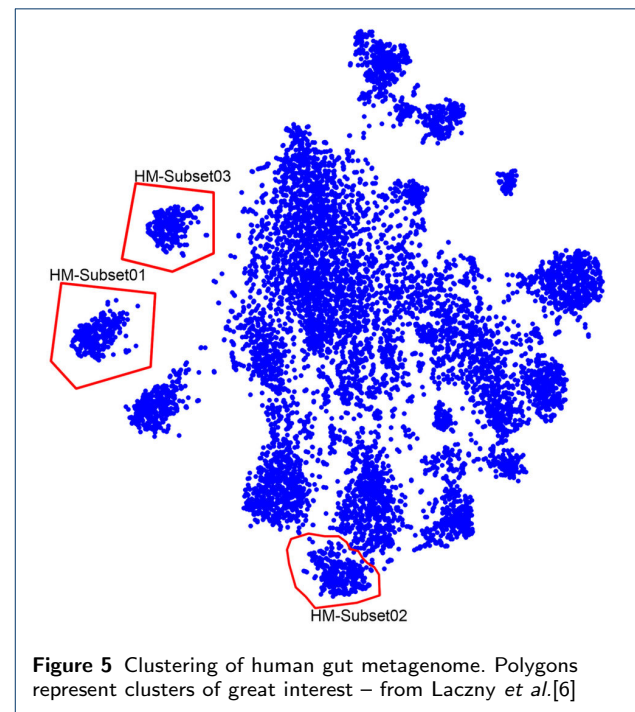
Application on real-world metagenomes With great results on simulated data, Laczny *et al.* also performed testing on real-world metagenomes of ground water[18], the human gut [19] and the deep sea[20]. They reported similar clustering (Figure 5) compared to simulated data with sensitivity, specificity and accuracy well above 90% for all subsets of the human gut metagenome except for one, where accuracy was slightly below 80%.

The values were calculated using polygons to mark clusters and verifying these by comparison with the NCBI non-redundant nucleotide database.

The ground water metagenome also produced distinct

clusters, as seen in Figure 6. Calculation of sensitivity, specificity and accuracy could not be carried out since they reported a lack of characterized reference genomes. Instead they used what they called "essential genes" which can indicate the completeness of a genome. They reported four out of eight of these essential genes as over 80% complete, indicating a positive result for their tool.

As for the marine sample, the clusters, as seen in Figure 7, identified by the tool were linked to yet uncharacterized data.



Compared to an ESOM-based approach, Laczny *et al.* reported better clustering of metagenomic-data while also significantly reducing runtime from around 3.8-fold to 50.4-fold[6] for their used metagenomes and good visualization capabilities as tested on simulated and real-world data.

Clustering seems robust on the applied metagenomes, while similar data tends to be near to each other in the visualization, 16S rRNA sequences form a distinct cluster, due to the high conservation of these regions in the genome.

As a downside sequences of 1000 nt were required to achieve good clustering, which are yet hard to gather through raw reads. Advancements in sequencing technologies are needed to fully utilize the capabilities of this tool.

Table 1 Sensitivity, specificity and accuracy of EqualSet01 – excerpt from Laczny *et al.*[6]

Subset	Sensitivity (%)	Specificity(%)	Accuracy(%)	Organism
01	90.06	99.99	99.94	<i>B. amyloliquefaciens</i>
02	91.25	100	100	<i>Candidatus C. ruddii</i>
03	95.42	99.90	97.57	Uncultured Termite group1 bacterium
04	98.60	98.23	96.67	<i>E. coli</i>

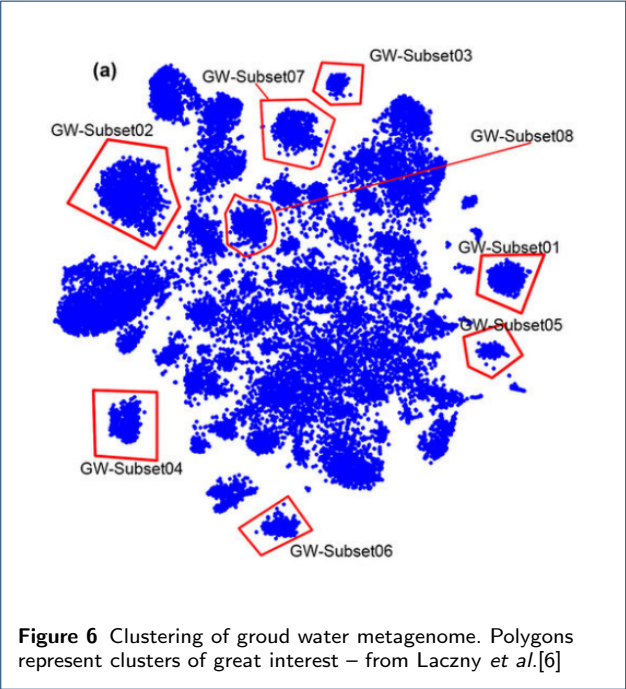


Figure 6 Clustering of ground water metagenome. Polygons represent clusters of great interest – from Laczny *et al.*[6]

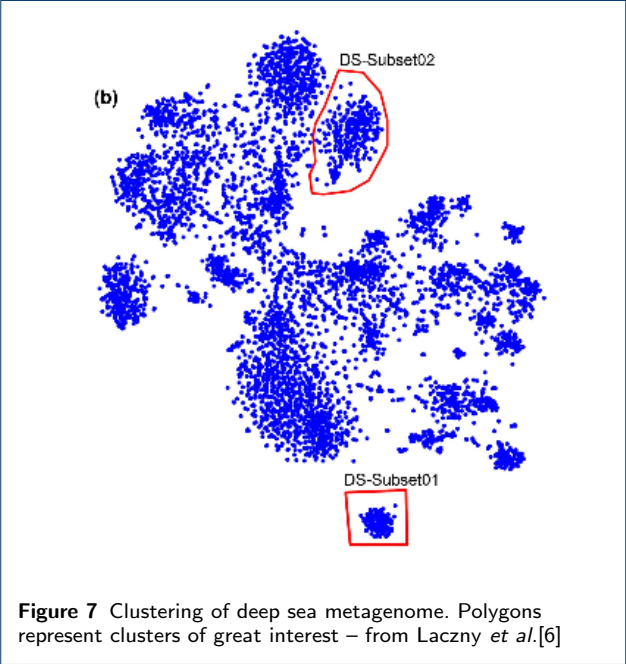


Figure 7 Clustering of deep sea metagenome. Polygons represent clusters of great interest – from Laczny *et al.*[6]

Conclusions

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<http://aem.asm.org/content/75/16/5345.full.pdf+html>