The III International Conference "Bioinformatics: from Algorithms to Applications" (BiATA2019) has established itself as one of the few conferences in the field of bioinformatics that brings together both the programmers creating tools for modern studies in multiple areas of life sciences and the researchers conducting those experiments interested in finding reliable and easy to use tools for data analysis. BiATA gives the international community a platform to present the latest achievements in bioinformatics and provides an excellent opportunity for researchers to discuss their pressing needs directly with software developers, while demonstrating the results they have already managed to achieve. This type of interaction is completely unique, because algorithm authors and users habitually tend to attend completely different conferences, which prevents this kind of information exchange.

The conference aims to popularize bioinformatics and promotes active application of bioinformatics in agricultural and biomedical fields of research; identify new trends in the fields of bioinformatics, computational genomics and transcriptomics, as well as in sequencing of biologically active molecules and the application of mathematical methods and algorithms in the life sciences.

Topics covered within the framework of the conference include but are not limited to:

1. Algorithms for the assembly of metagenomic data
2. Big data metagenomics
3. New algorithms for assembling and analyzing long reads obtained via new sequencing technologies
4. Computer biology and agriculture: analysis of soil and air microbiota
5. Human microbiota: nutrition and health
6. Bioinformatics of virome

The event also pays a great deal of attention to the most important task of all genomic research - restoring the primary sequence of genomic DNA from short fragments obtained as a result of using modern DNA sequencing technologies. Despite the fact that the restoration of the primary structure of DNA is not in itself the ultimate goal of research, all subsequent analyses depend on its quality. The quality of genome assembly becomes even more important when dealing with sequencing data generated from the combined genome of natural communities of microorganisms (microbiota) that inhabit a variety of different natural environments (soil, water, air, plants, etc.). Metagenomics - analytical methods and approaches that allow studying total genomes (microbiomes) - deals with large volumes of very complex data and requires specialized methods for solving scientific problems in such important areas as agriculture, medicine, etc.

The timeliness of the subject matter and the high level of quality of the conference can be evidenced by the level of speakers who took part in BiATA (http://biata2019.spbu.ru/). The conference brought together more than 100 participants from Russia, Belgium, Canada, China, Great Britain, France, Israel, Italy, Japan, Latvia, Lebanon, Spain, Singapore and the USA.

O1 Probabilistic model of CDR3 junctions formation in human Ig heavy chain genes and its application
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Immunoglobulins (Igs) play a crucial role in the adaptive immune system. Igs are composed of polypeptide subunits: light and heavy chains. The latter contains a variable domain that is important for an antigen binding. The coding sequences for Ig heavy chain are produced through a complex process, including VDJ recombination and somatic hypermutation (SHM). The latter masks initial segments, which complicates a precise sequence analysis of Ig genes in B-cells. Thus, in this research we focus on such a robust parameter of Ig genes sequence as the length of a V-D/D-J junction, which strongly influences antibodies affinity. As is known, these junctions may be subject to an abnormal recombination, sometimes leading to an autoreactivity and a subsequent lymphomagenesis (e.g. due to VH-replacement). Initially, a junction consists of palindromic (p)-nucleotides (produced by a protein complex of Ku70/Ku80 and Artemis) and non-templated (n)-nucleotides (added by a TdT protein), which further undergoes an impact of exo- and endonucleases.

For all the three stages of V-D junction maturation, we propose simple, yet tractable probabilistic models resulting in a general model describing a distribution of a junction lengths in normal immunoglobulins. The parameters for the developed model may be fitted by means of datasets obtained from healthy individuals, which are available in open databases such as GenBank and ENA. For this purpose, we have developed a pipeline containing the following steps:

1. Ig genes repertoire assembly (PRESTO);
2. clonal families detection and data decorrelation (Partis);
3. sequences demarcation and V/D-D-J junctions extraction (IMGT HighV-QUEST);
4. fitting model parameters via maximum likelihood estimation (custom Python scripts).

An evaluation of the model showed its consistency with the processed samples. The trained model was further applied to datasets describing Ig genes sequences with abnormalities in VDJ process. For
this data a statistically significant divergence with the model was detected. At the same time, no divergence was detected for diseases not related to onco-hematology. This experiment has shown that a V-D/D-J junction length distribution in Ig repertoire may be used as an indicator of the presence of pathological clones in a B-cell population. The possibility of the model application as an early predictor of various diseases presents a significant interest for further research.

O2 Bacteriophage recombination site helps to reveal genes potentially acquired through horizontal gene transfer

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Background The cellulose-synthase gene of ascidians was gained from prokaryote donor and this is the most reliable example of the horizontal gene transfer (HGT). In our previous study a new protein, rusticalin, of ascidian *Styela rustica* was described. Its C-terminal domain coding region was also shown to be inherited from prokaryote ancestor by means of HGT. Both for rusticalin C-terminal domain and for cellulose-synthase catalytic domain it was shown that there coding regions neighbored with bacteriophage recombination site AtpP. Thus we suggested a possible mechanism of HGT by means of bacteriophage insertion. Most of the cases of HGT are described based on sequence similarity alone, but in case of rusticalin we also demonstrated strong evidence of the mechanism of transfer by identifying the recombination site.

Results It is possible that bacteriophage recombination site can help finding yet other new cases of HGT in eukaryotic genomes. Unfortunately the length of bacteriophage recombination site AtpP is 43 nucleotides which is too short to find it reliably in big databases. Still we know that in rusticalin related gene AtpP-like site is situated inside the cysteine-rich repeats coding region. Based on that we performed a remote similarity search HMMER using amino acid sequence of cysteine-rich repeats. Cysteine-rich repeats appeared to be part of larger proteins. Therefore conserved domains associated with cysteine-rich repeats were classified.

In spite of the fact that cysteine-rich repeats are found almost exclusively in eukaryotic proteins, they are usually associated with domains typical for prokaryotes or bacteriophages (in 98 proteins out of 124). Among them in 20% (26 proteins) cysteine-rich repeats are associated with phage-lysozyme (PF00095), 14% (17 proteins) with amidase_2 (PF01510). In general nine different domains associated with cysteine-rich repeats can be classified as bacterial cell-wall hydrolyzing enzymes. It is worth to mention that phage-lysozyme domain is found together with cysteine-rich repeats in proteins of different species and even of different taxa as Fungi and Metazoa.

Conclusions Based on that observations we can conclude that cysteine-rich repeat in Eukaryotic proteins is usually accompanied by typical prokaryotic domains. The explanation of that might be the presence of bacteriophage recombination site inside cysteine-rich repeat coding sequence, which can facilitate HGT. The 98 genes potentially acquired through HGT from prokaryotes is found as the result.

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O3 SPAligner: alignment of long diverged molecular sequences to assembly graphs

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Many popular short read assemblers [9,10,11] provide the user not only with a set of contig sequences, but also with *assembly graphs*, encoding the information on the potential adjacencies of the assembled sequences. Naturally arising problem of sequence-to-graph alignment has been a topic of many recent studies [2,3,5,6,7,8,13,14]. Identifying alignments of long error-prone reads (such as Pacbio and ONT reads) to assembly graphs is particularly important and has recently been applied to hybrid genome assembly [1,4], read error correction [12], and haplotype separation [3]. At the same time, the choice of the practical aligners supporting long nucleotide sequences is currently limited to vg [2] and GraphAligner [3], both of which are under active development. Moreover, to the best of our knowledge, no existing graph-based aligner supports alignment of amino acid sequences.

Here we present the SPAligner tool for aligning long diverged molecular (both nucleotide and amino acid) sequences against assembly graphs produced by the popular short-read assemblers. The project stemmed from our previous efforts on the long-read alignment within the hybridSPAdes assembler [1]. Our benchmarks on various Pacbio and Oxford Nanopore datasets show that SPAligner is highly competitive to vg and GraphAligner in aligning long error-prone reads. We also demonstrate SPAligner’s ability to accurately align amino-acid sequences (with up to 80% amino acid identity) onto complex assembly graphs of metagenomic datasets. To further motivate this application we show how SPAligner can be used for identification of biologically important (antibiotic-resistance) genes, which remain under the radar of conventional pipelines due to assembly fragmentation (e.g. genes exhibiting high variability in complex environmental samples). SPAligner is available for download at [http://cab.spbu.ru/software/spaligner/](http://cab.spbu.ru/software/spaligner/).

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O4 HEDGE: GPU-powered protein-protein docking pipeline with accurate energy metrics as a primary scoring function

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Protein-protein interactions play key roles in living systems: cell signaling, immune system reactions, microelements transport and many other processes are based on protein-protein complexes functions. Thus, protein-protein complexes prediction is very important task especially in terms of drug discovery. For instance, in silico optimization stages of antibody-based drug development process requires to solve the problem hundreds times. To perform in silico optimization accurately the docking problem must be solved with high accuracy in short time ranges. But it is one of the hardest, both methodologically and computationally, structural bioinformatics problems.

Recently we developed a pipeline called HEDGE, briefly it can be described as follows: 1) scanning translational solution space using FFT correlation theorem with energy-like correlation function; 2) clustering of solutions by RMSD as a distance metric; 3) refinement of full complex structures with minimization of potential energy; Polak-Ribière-Polyak conjugate gradient method [1] is used to solve optimization problem, optimization target is OPLS [2] force field with additional GB and SA terms; 4) Finally we rank solutions by change of Gibbs free energy (ΔG), which can be considered as the most accurate metric to rank predicted complexes.

Each step of the pipeline above is well-parallelizable, so, the full power of GPUs (graphics processing units) is utilized, thus, overall computation time decreased significantly. Moreover, different rotations of molecules can be processed independently, therefore, multi-GPU mode is supported to scale linearly and achieve maximal performance on multi-GPU supercomputers.

Accuracy was tested on a subset of CAPRI [3] dataset showing about 50% of correct predictions. Time required for prediction of one complex in rigid mode (without minimization) is about 7 minutes on Tesla V100 GPU. Flexible mode requires much more calculations and takes about 1.5 hours on Tesla V100.

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O5 Million sequences indexing

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Most methodological contributions handling sequencing data now acknowledge the need to scale up to the terrific throughput that we face nowadays. Since BLAST, a plethora of tools have been developed to handle the massive amount of available reference sequences. Recently, new structures have been proposed to link a short sequence such as a transcript or a gene to sequencing datasets or reference genomes. The challenge of such structures is to be able to index hundreds of thousands of datasets with a reasonable amount of memory while being able to perform fast queries. A prosperous state of the art of efficient tools quickly emerged, SBT SSBT, HowDeSBT, BIGSI, ... based on different combinations of bloom filters in order to link to k-mer to its associated datasets. Those approaches are incredibly efficient: BIGSI was able to index half a million bacterial genomes with 1.5TB. However, not able to scale to all known genomes or all transcriptomes collections.

We aim to propose a new data structure that could use an order of magnitude less memory than BIGSI while being able to perform similar queries in terms of accuracy and throughput. Instead of indexing all k-mers of a dataset, we choose to rely on local sensitive hashing methods to index a small subset of the input k-mers. This choice allows the scaling of the methods with a satisfying accuracy on medium-sized queries (1kb or larger). Furthermore, we rely on a matrix-like structure similar to BIGSI, that offer fundamental properties for such an index:

- Constant time insertion of a new reference sequence by adding a new row to the matrix
- Queries rely on reading columns that can be compressed column for lighter structure and faster queries
- Easy to balance structure where memory/accuracy trade-off can be precisely chosen

In this presentation, we show the design of such a structure using the Min-Hash scheme. We present preliminary results on a hundred thousand bacterial genomes on a proof of concept implementation.

We compare our performances to BIGSI and Mashscreen, showing that the proposed structure can achieve a comparable accuracy with a better scaling in memory or throughput. We finally discuss the incoming improvements and what can potential pitfalls from this scheme, and its potential applications in mega-scale sequences indexing, clustering, or genome assembly. The open-source implementation is under development and available on Github at https://github.com/Malfoy/Mieikki.

O6 Revealing mechanisms of Bacillus thuringiensis host specificity via molecular modelling of the Cry toxin-receptor interactions

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Proteins possessing cytotoxic properties and commonly referred to simply as toxins comprise a vast group of bacterial virulence factors. For instance, Bacillus thuringiensis, a spore-forming bacterial pathogen of insects and several other invertebrate taxa, produces at least four major classes of proteinaceous toxins. Of these, crystalline pore-forming toxins produced at sporulation stage and known as Cry toxins, pose a particular interest because of the wide range of host species they affect. Cry toxins consist of three domains flanked with unstructured terminal sequences, with the N-terminal domain participating in pore formation and the other two involved in binding to a respective host’s receptor. To date, more than 700 Cry toxins affecting species of four Insecta orders as well as those of Nematoda phylum have been discovered, and several insect membrane proteins, have been proposed to serve as Cry receptors; however, little is known about the molecular mechanisms underlying both mode of action and specificity of these toxins. In this work, we analyze coevolutionary substitution patterns in toxins and alanyl aminopeptidase receptors to reveal molecular mechanisms of the specificity of toxin-receptor interactions. To enlarge the toxin dataset, we developed a novel HMM-based tool for searching Cry toxins and annotating their domain structure, which outperforms its analogs. Launching this tool with all Bacillus-related sequences we discovered 340 new toxins and defined domain layout of all known Cry proteins. Next, amino acid substitutions distinguishing protein sequences in both toxin and receptor epitope subsets were mapped onto minimum distance transition graphs which were then aligned to expose similarities in their
topology. Subsequent in silico docking of toxins and insect N-α-aminopeptidases revealed sites unequivocally involved in toxin-receptor interactions and effects of amino acid substitutions in these sites. Obtained data might be useful for designing novel Cry toxins efficient against particular hosts.

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O7

Indexing De Bruijn graphs with minimizers

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The need to associate information to words is shared among a plethora of applications and methods in high throughput sequence analysis and became fundamental. However, indexing billions of k-mers may lead to scalability issues, as exact associative indexes can be memory expensive. To leverage this issue, recent works take advantage of the properties of the k-mer sets [1]. They exploit the overlaps shared among k-mers by using a De Bruijn graph as a compact k-mer set to provide lightweight structures.

Contribution: we propose a scalable and exact index structure able to associate unique identifiers to indexed k-mers and to reject alien k-mers. The proposed index combines an extremely compact representation along with high throughput. Moreover, it can be efficiently built from the De Bruijn graph sequences. The index implementation we provide achieved to index the k-mers from the human genome with 8GB within 30 minutes, and was able to scale up to the enormous axolotl genome (32 Gbp [2]) with 63GB within 10 hours. Furthermore, while being memory efficient, the index allows above a million queries per second on a single CPU in our experiments, and its throughput can be raised using multiple cores. Finally, we also present the index ability to practically represent metagenomic and transcriptomic sequencing data to highlight its wide applicative range.

Availability: the index is implemented as a header-only library in C++, is open source under AGPL3 license and available at https://github.com/Malfoy/Blight. It is designed as a user-friendly library and comes along with sample code usage.

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O8

Synteny paths for assembly graphs comparison

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Despite the recent developments of long-read sequencing technologies, it is still difficult to produce complete assemblies of eukaryotic genomes in an automated fashion [1]. Genome assembly software typically output assembled fragments (contigs) along with assembly graphs, that encode all possible layouts of these contigs. Graph representation of the assembled genome can be useful for gene discovery, haplotyping, structural variations analysis and other applications. To facilitate the development of new graph-based approaches, it is important to develop algorithms for comparison and evaluation of assembly graphs produced by different software.

In this work, we introduce synteny paths: maximal paths of homologous sequence between the compared assembly graphs. We describe Asgan – an algorithm for efficient synteny paths decomposition, and use it to evaluate assembly graphs of various bacterial assemblies produced by different approaches. Similarly to synteny blocks [2] (that are commonly used in comparative genomics studies), synteny paths reveal structural similarities between the compared genomes, but are robust to assembly fragmentation. We define the problem of finding the minimum number of synteny paths between two assembly graphs and prove that the exact solution is NP-hard. Instead, we propose an efficient approximate algorithm for synteny path decomposition via transforming two assembly graphs into a colored breakpoint graph.

First, we use synteny paths to compare between Flye [3] and Canu [4] assemblies of 21 bacterial genomes from the NCTC collection. The final graphs produced by two assemblers from the same genome might not be identical, due to the different approaches to repeat resolution. However, if graphs are free from errors, they should encode the original genome sequence as path. Indeed, in 14 out of 21 NCTC datasets each connected component in assembly graphs was covered by a single synteny path, revealing the putative genome walks.

Secondly, we apply synteny paths to compare between the assemblies of 15 Drosophila genomes with extensive structural variations. We show that synteny paths reveal longer homologous segments, comparing to synteny blocks reconstructed using the fragmented contigs. Interestingly, the length distribution of synteny paths was highly correlated with the evolutionary distances between the compared genomes. This allowed to reconstruct the phylogenetic tree of 15 Drosophila genomes using pairwise synteny paths similarities as a distance metric.

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O9

Extending rnaSPAdes functionality for hybrid transcriptome assembly

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De novo RNA-Seq assembly is a powerful method for analysing transcriptomes when the reference genome is not available or poorly annotated. However, due to the short length of Illumina reads it is often impossible to reconstruct complete sequences of complex genes and alternative isoforms. Recently emerged possibility to generate long RNA reads, such as PacBio and Oxford Nanopores, may dramatically improve the assembly quality, and thus the consecutive analysis. While reference-based pipelines were already developed and applied to long RNA reads [1, 2], there are not many possibilities for de novo assembly of such data. Among available methods, Trinity
Another member of SPHMM family is PathRacer-Seq that produces frameshift-tolerant alignments of amino acid pHMM to nucleotide sequences significantly improving the accuracy of gene recovery out of assemblies obtained from long noisy reads.

O11
Local sequence alignment using intra-processor parallelism

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Local alignment of DNA sequences is a fundamental problem of bioinformatics. Standard solutions include fast heuristics, as well as the more time-consuming exact methods. An efficient exact local alignment technique, based on a “sliding window” approach, was previously developed at Warwick [1], resulting in biologically significant results [2-4]. The efficiency of that implementation was achieved, in particular, by low-level intra-processor parallelism.

In recent years, microprocessor architecture has been developing rapidly, culminating with Intel’s AVX-512 [5], an instruction set taking intra-processor parallelism to a new level of efficiency and sophistication, while also being surprisingly well-suited for speeding up the “braid combing” sequence alignment technique developed by the second author [6-7]. We present a prototype software tool [8] that is, to our knowledge, the first sequence alignment software taking advantage of AVX-512 parallelism. Our approach allows one to produce sliding window alignments between a short fragment (“pattern”) and a long sequence (“text”), using braid combing and intra-processor parallelism.

In the simplest case of unweighted alignment, the braid combing algorithm can be described as growing an object called a braid, embedded in the grid defined by the input sequences (Figure 1). The combing logic is as follows: we iterate over cells of the grid left-to-right and top-to-bottom, extending the braid to the current cell. Two strands enter the current cell, one horizontally, the other vertically. In a match cell, the two strands pass through the cell and exit it without crossing (the cell that entered horizontally exits vertically, and vice versa). In a mismatch cell, the strands cross and keep their directions; otherwise, they behave as in a match cell. In AVX-512, this logic can be implemented efficiently by processing the cells in parallel, iterating through the grid in an antidiagonal frontier of independent cells. The frontier is represented by two integer vectors: one storing the indices of the horizontal, the other of the vertical strands. 32-bit integers suffice for all realistic local alignment scenarios. The crossing rules correspond to pairwise sorting of strand indices via vector intrinsic _mm512_mask_min_epu16 / _mm512_mask_max_epu16, using a mask indicating whether individual frontier cells are match or mismatch ones (Table 1). For rational-weighted alignments, the blow-up technique [9] can be used to reduce the problem to the unweighted case.

In future, we plan to extend our implementation to a fast exact local sequence aligner.

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inactivation of \textit{mgrB}, a gene encoding a negative regulator of the PhoPQ two-component system (TCS), and \textit{crrAB} (a sensory TCS) have recently gained attention as mediators of colistin resistance.

\textbf{Materials and methods}

In this study, broth microdilution colistin susceptibility testing and whole-genome sequencing were used to resolve phenotypic and genotypic resistance profiles in 11 clinical carbapenem- and colistin-resistant \textit{K. pneumoniae} (KP). Whole-genome sequencing (WGS) was performed using short-paired end reads technology on an Illumina Miseq. Core genome single nucleotide polymorphisms (cg-SNP) were called by the Snippy pipeline, and recombination events were highlighted using Gubbins. The pan-genome was generated using Roary. Chromosomally encoded genes were also screened for synonymous and non-synonymous mutations, in particular, \textit{pmrAB}, \textit{pmrD}, \textit{pmrc}, and \textit{phoPQ}. The genetic environment of \textit{mgrB} was manually validated by Sanger sequencing. Lipid A was extracted using mild acetic acid hydrolysis and profiled using MALDI-TOF MS to examine noteworthy modifications linked to decreased susceptibility to colistin.

\textbf{Results}

The lipid A major mass ion was observed at (m/z 1840) in all KP isolates. PCR amplification of \textit{mgrB} revealed insertionally inactivated \textit{ΔmgrB} in three of the studied isolates (designated as KP5, KP6, and KP16) showing MICs $\geq$16 mg/L. \textit{Kpn14} was associated with KP5 and KP6, while \textit{IS902} was detected in KP16. Wildtype \textit{mgrB} gene in the remaining 8 isolates might suggest the involvement of other mechanisms underlying their nonsusceptibility to colistin. Recombination analysis highlighted genomic loci involved in both toxin-antitoxin and MFS efflux systems as favored hotspots for recombination. All 11 isolates were negative for the \textit{crrAB} genes. Further biochemical and molecular analysis is in progress to characterize genetic determinants that play key roles in colistin resistance.

\textbf{Conclusion}

Along with the escalating prevalence of CRKP and the lack of novel antibiotics, colistin resistance has imposed a worldwide concern. With the power of WGS and lipidomic approaches, genetic alterations in pathways responsible for lipid A modification can be detected with high precision, enabling us to better understand the molecular mechanisms involved in resistance.

\section*{O12}

\textbf{On the verge of colistin resistance: genetic determinants mediating intermediate colistin resistance in \textit{Klebsiella pneumoniae}}

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\section*{Background}

Colistin is one of the last resort antibiotics used to treat infections by carbapenemase-producing \textit{Klebsiella pneumoniae} (CPKP). Insertional inactivation of \textit{mgpB}, a gene encoding a negative regulator of the PhoPQ two-component system (TCS), and \textit{crrAB} (a sensory TCS) have recently gained attention as mediators of colistin resistance.
specific example: metastasis-related genes for neuroblastoma. Neuroblastoma is a pediatric cancer with a heavy metastasis burden for high-risk patients. However, the type of metastasis is very specific for neuroblastoma and cannot be directly compared to adult metastasized cancers. We show the workflow of mining for the neuroblastoma-related gene set of metastasis-relevant genes and analyze its enrichment in neuroblastoma experimental data. As a comparison, we then run a similar analysis on metastatic samples from breast cancer to illustrate the added value of research-specific gene set enrichment analysis. The gene set analysis tool is part of a broader text mining tool “sina” (search indexed nomenclature associations) that we are developing and is available at https://github.com/dicaso/sina.

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O14
DASE-AG: conditional-specific differential alternative splicing events estimation method for around-gap regions
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Alternative splicing is a mechanism to generate more than one mRNA isoforms from a single locus, and it increases the genetic diversity during post-transcriptional gene regulation. Furthermore, alternative splicing is often differentially regulated across tissues and during development. It suggests that each splicing isoform may have specific spatial and temporal roles in life system.

We have developed the differential alternative splicing variants estimation method, DASE and DASE2. DASE2 uses FPKMs or TPMs as expression quantities. FPKMs and TPMs are read counts normalized with the lengths of transcripts. However, DASE2 had three problems in finding splicing quantities. FPKMs and TPMs are read counts normalized with the lengths of transcripts, DASE and DASE2. DASE2 uses FPKMs or TPMs as expression quantities, but DASE2 treated expression quantities of AG regions tend to up to the 100th rank and DASE2 could not find any of those events. One of the factors is that expression quantities of AG regions tend to be higher than those of the whole sequences of transcripts.

DASE-AG is available at https://github.com/koukiyonezawa/DASE-AG.

P1
Target selection protocol for DNA-machines development
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Deoxyribozymes based DNA-machines are universal approach to cleave mRNA of target gene and can be applied to prokaryotic and eukaryotic organisms. This technique was successfully used against cancer cells and Influenza A Virus [1,2]. Choosing the right target gene is still a fundamental stage for using DNA-machines. In case of eukaryotic organism number of housekeeping genes can reach of thousands, what makes cumbersome subsequent gene analysis by hand.

To achieve goals of our research we developed a Python script utilizing Entrez library, BLAST software [3] and The NCBI SRA Toolkit to access mRNA sequences and estimate the level of gene expression. Firstly, it downloads and creates local indexed databases via ’sra-toolkit’ [4] and ’makeblastdb’ applications respectively. Secondly, it queries genes sequences in prepared databases to retrieve summary statistics for their occurrences using’blastn’ software. Our software speed up the detection of over-expressed genes, moreover it could deal with both eukaryotic and prokaryotic organisms.

DNA-machine is very sensitive to mismatches in sequences therefore rapid evolution of genes can disrupt the process of mRNA cleaving. MEGA program was used to identify the most conservative genes. Furthermore, to extend time of work DNA-machines we should choose genes with stable mRNA. This characteristic is determined by the half-life of the mRNA. We exclude genes connected with replication process, which may take a long time, to reduce time in vivo experiments.

We verified essentiality of target genes in BioCyc Database Collection demonstrating result of genes knockout. The absence of vulnerable housekeeping gene will lead to cell death.

With the help of the script, we estimated expression level of 3800 housekeeping genes in 2 human transcriptomes. Additionally, we analyzed 206 housekeeping genes in 5 Escherichia coli transcriptomes. On the basis of target selection protocol, we determined the most relevant genes for deoxyribozyme development which are tested in vitro (Figure 3). Further, deoxyribozymes based DNA-machines are tested in vitro and in vivo.

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Improved Architecture of Artificial Neural Network for Secondary Structure Analysis

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The idea about using secondary structure while solving different sequences analysis problems is considered in many works [1-4]. One of the classical ways of describing secondary structure is formal grammars.

An approach for biological sequences processing by combination of formal grammars and neural networks is proposed in the work [5]. While classical way is to model secondary structure of the full sequence by using grammar, the proposed approach utilizes it only for primitive secondary structure features description. These features can be extracted by parsing algorithm and processed by neural network. It is shown that this approach is applicable for real-world data processing and several questions are formulated for future research. In this work we provide answers to some of them.

The first question is whether it is possible to use convolutional neural networks for parsing result processing. The result of matrix-based parsing algorithm for some string and fixed nonterminal is an upper-triangular boolean matrix. The first way to represent this matrix is to drop out the bottom left triangle and vectorize the rest of matrix row by row. The vector can be handled by using dense neural network. It requires the equal length of the input sequences, therefore, we propose to cut sequences or add some special symbol till the definite length. The second way is to represent the matrix as an image: the false bits as white pixels and the true bits as black ones, so, we can process sequences with different length and resize images afterwards. To handle these images we use network with a small number of convolutional layers. After linearization we use the same network as for vectors handling.

The second question is whether it is possible to move parsing to network training step, because parsing is the most time-consuming operation of our solution. We solve this problem by using two-staged learning. Firstly, we prepare a network which takes parsed data as an input. After that we extend trained network with a number of input layers that convert original nucleotide sequence into parsing result. This network handles sequences, so, we require parsing only for training the first network.

We use these improvements for tRNA sequences analysis problems: classification into two classes (eukaryotes, prokaryotes) and four classes (archaea, bacteria, plants, fungi). Accuracy for image- and vector-based classifiers is about 90% and accuracy for extended networks is about 95% on the test dataset.

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P3
Analysis of microbiomes of the ecogenetic series of podzolic soils using an artificial neural network
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Background
The application of molecular-genetic methods is currently one of the necessary steps in the natural microbiome analysis. The use of these approaches in the study of microbial complexes of soil chronosequences is promising in identifying of taxonomic markers and microbiological drivers of pedogenesis. Otherwise, the search for optimal ways of high-throughput sequencing data processing remains relevant today. The aim of the study was the analysis of the microbial composition of the genetic horizons occurring end developing during the long-term soil evolution in the middle taiga zone.

Materials and Methods
Sample set included the podzol soils chronosequences: 1) the initial soil formation (samples of 1-2 years, with no signs of pedogenesis; stages of self-overgrowing (15–20 years with sod-podzol occurrence, 30-35 years old with embryopodzol) and a 2) the ongoing podzol pedogenesis (soils of 70-, 145-, 455, 1590 years). Soil samples were analyzed by classical methods (physico-chemical), DNA isolation was performed using the PowerSoil® DNA Isolation Kit (MO BIO, USA). Amplification of a V4 variable region of the 16S rRNA was carried out with universal primers (F515 and R806) (Bates et al. 2010). Amplicon libraries sequencing was performed by ILLUMINA Miseq. Sequence data processing was carried out using “Trimmomatic” (Bolger et al. 2014) and “QIIME” (Caporaso et al. 2010) software. Samples clustering and concurrent microbial taxonomic composition analysis was carried out using an artificial neural network that is trained using unsupervised learning - Kohonen’s self-organizing map (SOM) (R package ‘kohonen’; Wehrens and Krusselbrink, 2018).

Results
Characteristic patterns of the microbial taxa composition were revealed for the early and late stages of podzolic soils succession. The SOM application allowed to determine the indicative role of microbiome taxonomic structure in marking of solum differentiation during the time of podzol ontogenesis.

Conclusions
The neural networks with training is a promising approach to the analysis of soil microbiomes, which in turn helps getting the biologically relevant information (the adaptive and evolutionary strategies of microorganisms during soil-f ormation) and the practically valuable results.

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analysis. The known genotypes are hidden randomly. The data sets are imputed and the genotypes thus obtained are compared to the original ones. The discordance between the genotypes is counted. We demonstrated several applications of this toolkit. Firstly, it can be applied for benchmarking of imputation software. We applied the tool to the data sets from HapMap and 1000 Genomes Project and compared the quality of imputation made with fastPHASE and BEAGLE softwares. Both programs showed the discordance of about 3%. Secondly, imputeqc can be applied for choosing the model parameters for imputation with fastPHASE. Two parameters were studied: the number of haplotype clusters and the expectation-maximization cycles. The data set represented merged genotypes of CEU, TSI, CHB, and JPT populations from 1000 Genomes Project. The optimal number of haplotype clusters was estimated to be 20 and the number of expectation-maximization cycles to be 25. Thirdly, we demonstrated that the tool can be used in conjunction with hapFLK program. The estimated number of haplotype clusters fits well hapFLK model. Applying it to the pool of CEU, TSI, CHB, and JPT population we observed a strong signal of selection at the region of LCT gene. Finally, imputeqc can be applied to estimate the quality of imputation in GWAS by identifying the single nucleotide polymorphism that can be taken for the studies. The toolkit is implemented as an R package imputeqc and command line scripts. The code is freely available at https://github.com/inzilico/imputeqc under the MIT license. The reported study was funded by RFBR according to the research project No 19-29-01151.

P6
CDSnake: Snakemake pipeline for retrieval of annotated OTUs from paired-end reads using CD-HIT utilities
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Sequencing of 16S rRNA is a commonly used method for cost-efficient research of microbial communities. Illumina paired-end reads are often used as sequencing method. Since even short variable regions of 16S provide sufficient information for microbe identification, sequenced fragment is often shorter than sum of lengths of paired reads. Thus reads of pairs can be merged for downstream analysis. In spite of development of several tools for merging of paired-end reads, poor quality at the 3’ ends in the overlapping region prevents the correct assembly of significant portion of read pairs. Recently CD-HIT-OTU-Miseq was presented as a new approach, avoiding reads merging due to separate clustering of paired reads and discarding of reads voting for non-matching clusters as chimeric. CD-HIT-OTU-Miseq utilities are command line tools written in C++ and Perl. Here we assembled CD-HIT-OTU-Miseq utilities into pipeline using Snakemake workflow. We benchmarked our pipeline with two commonly used pipelines for OTU retrieval, incorporated into popular workflow for microbiome analysis, QIIME2 - DADA2 and deblur. Benchmarking was made on 3 mock datasets, Balanced, HMP, and Extreme, each having highly overlapping paired-end 2 × 250 reads. The Balanced community contained 57 bacteria and archaea at nominally equal frequencies, the HMP community contained 21 bacteria at nominally equal frequencies, and the Extreme community contained 27 bacterial strains at frequencies spanning five orders of magnitude and differing over the sequenced region by as little as 1 nucleotide (nt). CDSnake outputted less OTUs than DADA2 and deblur, since last two tools aim to output sub-OTUs by error processing, and OTU-Miseq doesn’t process errors and tries to output most correct OTUs using clustering. However, on Balanced and HMP datasets number of OTUs outputted by CDSnake was closer to real number of strains which were used for mock community generation, than those outputted by DADA2 and deblur. On Extreme dataset CDSnake, as expected, performed worse than DADA2 and deblur, since clustering algorithm cannot separate sequencing errors from actual 1-nt differences, present between strains in this community.

CD-HIT-OTU-Miseq provides one more approach for amplicon analysis capable to outperform popular tools in certain conditions. We developed Snakemake pipeline for OTU-Miseq utilities, which can be useful for easier automated runs. This work has been supported by the Russian Science Foundation (grant 19-16-00049).

P7
Genome heterogeneity affecting binning of complex fungal communities
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The vast majority of fungi are yet to be described and cultured. Since in nature these species mostly occur mixed with other organisms, accessing genomic information from these fungi is a serious challenge. Shotgun sequencing techniques do not offer a reliable way to extract the genome of a target fungus from a mixed dataset, which might include other eukaryotic genomes. Previously, some standard database-independent binning approaches were applied to metagenomes of complex eukaryotic communities. These methods are based on oligonucleotide frequency distribution and rely on the assumption of homogeneity of sequence composition across any given genome. This assumption, however, might not hold true for some fungi. Genomes of these species show strong intragenomic difference in base composition, a phenomenon thought to be caused by repeat-induced point mutation (RIP). RIP is a mechanism used by fungi against transposable elements, silencing multicopy DNA elements by directed mutational processes. Lichens are complex symbiotic communities including multiple species of fungi, algae, and bacteria, and represent a case where two phenomena, unculturable fungi and heterogeneous fungal genomes, overlap. In our study, we aim to assess the extent to which genome heterogeneity might affect metagenomic binning and propose a strategy to improve the binning of complex fungal communities.

P8
Genome-wide analysis of multidrug-resistant Shigella spp. isolated from patients in Lebanon
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Background
Shigella spp. are Gram-negative rod-shaped bacteria belonging to the family Enterobacteriaceae and are a major cause of bacillary dysentery worldwide. In this study, whole-genome sequencing was used for the molecular characterization of ESBL producing Shigella spp. isolates collected from hospitals in Lebanon.

Materials and methods
Polymerase chain reactions (PCRs) were performed to detect β-lactam resistance gene reservoirs and to identify the ones mediating virulence and host adaptation. PCR-based replicon typing (PBRT) was performed to identify patterns of plasmid distribution and multi-locus sequence typing (MLST), whole-genome based single nucleotide polymorphism (SNP) analysis, pan-genome analysis and pulse field gel electrophoresis (PFGE) were performed to determine the phylogenetic relatedness of the isolates and to trace evolutionary lineages.

Results
S. sonnei was the dominant serogroup (8/10 S. sonnei, 1/10 S. boydii, 1/10 S. flexneri). A total of 13 genes conferring resistance to
aminoglycosides, β-lactams, sulfonamides, trimethoprim, tetracycline and chloramphenicol were identified, while all the isolates were susceptible to ciprofloxacin and norfloxacin. Five types of β-lactamase genes were detected \( bla_{CTX-M-15} \), \( bla_{TEM-1B} \), \( bla_{OXA-1} \) and \( bla_{CTX-M-3} \) in cephalosporin-resistant isolates. \( bla_{OXA-1} \) was associated with \( S.\ flexneri \), while \( bla_{CTX-M-15} \), \( bla_{TEM-1B} \), \( bla_{OXA-1} \) and \( bla_{CTX-M-3} \) with \( S.\ sonnei \). \( bla_{OXA-1} \) was linked to class 1 integron integrated on IncFII type plasmid, while \( bla_{CTX-M-3} \) was detected on an IncI1 plasmid. The genetic environments of \( bla_{CTX-M-3} \), \( bla_{CTX-M-15} \) and \( bla_{TEM-1B} \) were also determined. All isolates harbored virulence genes and tested positive for the invasion plasmid antigen H (ipaH). Serine Protease A (SepA), responsible for critically disrupting the intestinal epithelial barrier, was associated with \( S.\ flexneri \), whereas the invasion associated locus (ial) with \( S.\ boydii \). \( S.\ sonnei \) had a larger core genome (by approximately 78kb) compared to \( S.\ flexneri \) and \( S.\ boydii \), both having a smaller core genome but a wider variety of accessory genes.

**Conclusion**

To the best of our knowledge this is the first detailed molecular characterization of \( Shigella \) spp. isolates recovered from patients in Lebanon. Our results revealed the association between antimicrobial resistance and increased virulence-related genes, and the emergence of strains with high levels of resistance to third generation cephalosporins. Although there are still some active antimicrobial agents that can be used to treat shigellosis, further emergence of antibacterial resistance by inappropriate use should be carefully followed and prevented.

**P9**

**Identification of small RNAs derived from commensal microbiota or infections**

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**BMC Bioinformatics 2019, 20(Suppl 17):P9**

The intricate mixture of small RNAs of human and non-human origin obtained from a wide range of biofluids is one of the most complex problems to be resolved in RNAseq data analysis. In order to identify accurately those sRNA reads of human origin, the other species sources (bacteria, fungi and viruses) should be separated. For this purpose, we have developed a new algorithm, which allows reducing false positive matching of reads to improper species by two-pass analysis based on the BLAST output on “nr” database using a representative random subset of reads. The second pass will assign the hit to the species, which were most frequently encountered in the first pass, in case of a similar score. At the same time, valuable research information on accompanying species will be also obtained. Only the genomes of the most represented species in successful BLAST hits will be used for the following alignment step. Contrary to full-length mRNA, sRNAs reads usually align in multiple sites of the genome. Our algorithm aligns the reads allowing multiple alignments per read and then reassigns them taking into account the local coverage using ShortStack algorithm.

Results show that, in general, our approach is more suitable for small RNAs analysis than Kraken2/Sourmash/MetaPhlAn2 due to the fact that the K-mers used to generate their databases are longer than most of the small RNA derived reads. Our approach also demonstrates more sensitive results than Kraken2 for highly damaged DNA, as for example, those obtained from archaeological microbiome samples. Nevertheless, the specificity of the method should be improved. Presented algorithms will be included in the future release of sRNAflow - a software tool for the analysis of small RNAs in biofluids. Besides existing packages for adapter removing, quality control, mapping and counting of reads, differential expression analysis, and miRNA target prediction, this pipeline currently includes the creation of a catalogue of expressed RNA types using human genome annotations and differential expression analysis tools such as DESeq2 for all of the classifiable RNA types. Human genome annotation has been expanded and includes Ensembl database, as well as miRBase, IncRNApedia, piRBase, piRNAdb, piRNAbank, GtRNAdb and GtRNAdb derived tRFs databases. The prioritization algorithm for building a catalogue of expressed RNA types allows solving the problem that exists when employing different annotations database due to annotations overlap. In addition, our pipeline will include identification of non-templated miRNA isoforms.

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