Sequence analysis

Analyzing large scale genomic data on the cloud with Sparkhit

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Abstract

Motivation: The increasing amount of next-generation sequencing data poses a fundamental challenge on large scale genomic analytics. Existing tools use different distributed computational platforms to scale-out bioinformatics workloads. However, the scalability of these tools is not efficient. Moreover, they have heavy run time overheads when pre-processing large amounts of data. To address these limitations, we have developed Sparkhit: a distributed bioinformatics framework built on top of the Apache Spark platform.

Results: Sparkhit integrates a variety of analytical methods. It is implemented in the Spark extended MapReduce model. It runs 92–157 times faster than MetaSpark on metagenomic fragment recruitment and 18–32 times faster than Crossbow on data pre-processing. We analyzed 100 terabytes of data across four genomic projects in the cloud in 21 h, which includes the run times of cluster deployment and data downloading. Furthermore, our application on the entire Human Microbiome Project shotgun sequencing data was completed in 2 h, presenting an approach to easily associate large amounts of public datasets with reference data.

Availability and implementation: Sparkhit is freely available at: https://rhinempi.github.io/sparkhit/.

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Supplementary information: Supplementary data are available at Bioinformatics online.

1 Introduction

Genome and transcriptome projects have provided unprecedented knowledge and insights for life science by interpreting large amounts of next-generation sequencing data (Consortium and Auton, 2015; Peterson et al., 2009; R Genomes Project, 2014; Wyatt et al., 2014). These genomic datasets, sequenced from thousands of samples, take days or weeks to analyze on private clusters (Bao et al., 2014). For researchers with limited computational resources, processing terabytes (TB) of data remains a fundamental bottleneck. Although several existing tools have been developed and utilized on single computers (Bray et al., 2016; Langmead and Salzberg, 2012; Li and Durbin, 2009; Li et al., 2009, 2008; Niu et al., 2011; Wood and Salzberg, 2014), most of the parallelizations on multi-computer networked clusters are done by manually splitting and distributing in smaller batches (Droop, 2016). To be compatible with a cluster, methods involving message passing and graph representation between computers must be re-implemented with higher level programming interfaces (Gropp et al., 1996). To address this challenge, we need both easy-access to large computational infrastructure and bioinformatics tools that are compatible and scalable on such platforms.

Cloud infrastructure and platform services (IaaS and PaaS) have been well established in the informatics discipline (Dean and Ghemawat, 2008; Schadt et al., 2010; Shvachko et al., 2010; Zaharia et al., 2012). To fully exploit distributed cloud computing systems, methods and programs should be scalable, fault tolerant and platform independent. In genomics applications, there are several tools [e.g. ABySS (Simpson et al., 2009) and Ray (Boisvert et al., 2010)] using the message passing interface (MPI) for distributed implementations. However, programming on top of MPI has to tangle with thread synchronization and load balance. Moreover,
there is no complete fault tolerance mechanism built inside of MPI. The conventional Apache Hadoop framework is designed to offer higher scalability and a supervised fault tolerance mechanism by providing a distributed data storage system called HDFS (Shvachko et al., 2010) and a distributed computing engine, Hadoop MapReduce (Dean and Ghemawat, 2008). Cloudbust (Schatz, 2009) is a Hadoop based sequence mapping tool programmed in the MapReduce model. The common seed and extend alignment pipeline is split and implemented in ‘map’ and ‘reduce’ steps, where the ‘map’ step searches K-mer matches (as seeds) and the ‘reduce’ step extends the seeds to apply dynamic alignments. The limitation for such method is that the ‘reduce’ step introduces large data shuffling across cluster nodes that impacts its performance. Crossbow (Langmead et al., 2009), Halvade (Decap et al., 2015) and Myrna (Langmead et al., 2010), on the other hand, directly use Hadoop to invoke existing sequence aligner [Bowtie (Langmead, 2010)] and SNP caller [SOAPsnp (Li et al., 2008) or GATK (McKenna et al., 2010)] for sequence mapping and genotyping on large datasets. The three tools have successfully reduced the run times for mapping, genotyping and gene expression quantification. Yet, their data preprocessing step introduces a heavy overhead and their options for handling distributed data are limited.

The more recent Apache Spark framework addresses these weaknesses with its unique data sharing primitive, called resilient distributed dataset (RDD) (Zaharia et al., 2012). RDD offers a ‘cache’ function to store distributed data in the memory across computers on a cluster, thus, avoiding run time overhead of iterative data I/O (input and output). Moreover, Spark has more build-in functions for RDD to facilitate methods implementation and data handling via its application programming interface (API). These advantages make Spark suitable for large scale genomic data analysis. Nevertheless, existing Spark based bioinformatics tools have their own limitations. For instance, SparkBWA (Abuin et al., 2016) adopts the same idea of Crossbow by using Spark to invoke the BWA (Li and Durbin, 2009) aligner. However, it does not provide data preprocessing functions for large amounts of compressed sequencing data. Thus, manually decompressing the sequencing data introduces a significant run time overhead. Instead of directly invoking existing aligners, MetaSpark (Zhou et al., 2017) re-implemented a fragment recruitment algorithm (Rusch et al., 2007). It has the same ‘seed and extend’ pipeline as Cloudbust, but implemented its algorithm on top of Spark. Thus, it also introduces large data shuffling in the reduce step that impacts its run time performance. Moreover, it requires a self-defined input format rather than the standard Fastq and Fasta format, which introduces an overhead to manually covert large Fastq files (Table 1).

We present Sparkhit, an open source computational framework that is easy to use on a local cluster or on the cloud (Fig. 1). Sparkhit is built on top of the Apache Spark platform, integrates a series of analytical tools and methods for various genomic applications: (i) We have natively implemented a metagenomic fragment recruitment tool and a short-read mapping tool (Sparkhit-recruiter and Sparkhit-mapper). The short-read mapper implements the pigeonhole principle to report the best hit of a sequencing read. Whereas the fragment recruitment tool implements the q-Gram algorithm to allow more mismatches during the alignment, so that extra information is provided for the metagenomic analysis; (ii) For using external software on Sparkhit, we built a general tool wrapper (Sparkhit-piper) to invoke and parallelize existing executables, biocontainers [e.g. Docker containers (Merkel, 2014)] and scripts; (iii) For downstream data mining, we integrated and extended Spark’s machine learning library. All methods and tools are programmed and implemented in a new MapReduce model extended by Spark, where parallelization is optimized (load balanced) and supervised (fault tolerance).

Our benchmarks on Sparkhit demonstrated its high scalability. In comparison, Sparkhit ran 18–32 times faster than Crossbow on data preprocessing and Sparkhit-recruiter ran 92–157 times faster than MetaSpark on fragment recruitment (Figs 2 and 3). Utilizing a powerful compute cloud, Sparkhit can quickly analyze large amounts of genomic data (Fig. 4). Our test case presents a 21 h ‘pay-as-you-go’ cloud application that analyzed 100 TB genomic data from 3 genome projects and a transcriptomics study (Wyatt et al., 2014). The analysis on the Human Microbiome Project (HMP), associated public ‘big data’ with private datasets, demonstrates how Sparkhit can be widely applied in different genomic studies (Supplementary file S2, Figs S1, S2 and S3). Thus, our framework enables the broader community to engage genomic investigations by leveraging cloud computing resources.

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**Table 1. Features of different cloud based bioinformatics tools**

| Tools          | Platform  | Implementation                  | Application                  | Methods and tools       |
|----------------|-----------|---------------------------------|------------------------------|-------------------------|
| Cloudburst     | Hadoop    | Native Java re-implementation   | Mapping                      | Seed and extend         |
| Crossbow       | Hadoop    | Invoking external utilities     | Mapping and genotyping       | Bowtie and SOAPsnp (Li et al., 2008) and GATK (McKenna et al., 2010) |
| Halvade        | Hadoop    | Invoking external utilities     | Mapping                      | BWA and GATK (McKenna et al., 2010) |
| Myrna          | Hadoop    | Invoking external utilities     | Mapping and gene expression profiling | Bowtie (Langmead, 2010) |
| CS-BWAMEM      | Spark     | Native Scala re-implementation  | Mapping                      | Burrows-Wheeler transform Smith-Waterman BWA (Li and Durbin, 2009) |
| SparkBWA       | Spark     | Native Scala re-implementation  | Mapping                      |                        |
| MetaSpark      | Spark     | Native Scala re-implementation  | Fragment recruitment         | Seed and extend         |
| Sparkhit       | Spark     | Both native and external        | Fragment recruitment, mapping, genotyping, taxonomy profiling and gene expression profiling | Seed and extend a collection of tools |
2 Materials and methods

2.1 Spark cluster’s architecture and cloud deployment

The architecture of Sparkhit is organized by a Spark cluster, which virtualizes each computer and forms a network consists of one or more master node(s) and worker nodes. To organize all virtual compute nodes, a master node is selected to supervise worker nodes, allocate resources and balance workloads, whereas worker nodes carry out assigned tasks (Fig. 1A). On such architecture, Spark extends and integrates the MapReduce model into RDD’s functions, where the ‘map’ step loads, processes and saves data in parallel on each worker node and the ‘reduce’ step sorts, shuffles and aggregates intermediate data. We used Spark-ec2 and BiBiGrid (https://github.com/BiBiServ/bibigrid), a Java-based tool developed in our research group, to easily setup Spark clusters on the Amazon cloud with one command line (see Supplementary file 1 Method-1.2).

2.2 Parallel data downloading

Downloading and decompressing large genomic files are significant bottlenecks before the actual data analysis begins. Spark’s architecture can greatly benefit from high-performance networks during large data transfers. In particular, when commencing a download task, files can be split into chunks and transferred from distributed storage system, such as Amazon simple storage service (S3), to each worker node. This parallel transfer method fully utilizes high network bandwidth of a distributed cluster (Fig. 1A).

When downloading data from Amazon S3 to HDFS, we have used Hadoop Distcp (distributed copy), a tool designed for large inter-cluster copying. For downloading data from Amazon S3 to a shared network file system (NFS), we used a Java-based tool developed in our research group called BiBiS3 (https://wiki.cebitec.uni-bielefeld.de/bibiserv/index.php/BiBiS3). BiBiS3 not only parallelizes downloading jobs to multiple computer nodes, but also applies multi-thread downloading on each computer node to fully exploit the capacities of every computer’s network connection (see Supplementary file 1 Method-1.3).

2.3 Parallel data decompression

Most genomic datasets are compressed and stored on public repositories. To directly access and analyze the compressed data, we also improved the performance on data preprocessing by introducing parallel decompression after downloading the data. We developed a parallel decompression tool, Sparkhit-spadoop, that is built on top
of the Apache Hadoop (Spark’s parallel decompression has a thread safety issue at 2.0.0 version) and included it in our toolkit. Since Bzip2 files are compressed in blocks (900 KB per block by default), a large Bzip2 compressed file with sequencing data can be decompressed in parallel. In particular, each block is an independent component that is delimited by a 48-bit pattern, which makes it possible to find the block boundaries. When block boundaries are found, parallel decompression can be applied to each block and processed by multiple CPUs, so that more compute cores are utilized. Then, we distributed all processes in a Hadoop MapReduce.
job that creates a ‘mapper’ for each chunk of the input HDFS data. Each ‘mapper’ loads a data chunk and commences a decompression
process on the Bzip2 blocks in the chunk. (see Supplementary file 1
Method-1.4)

2.4 Data loading and saving
The Spark RDD stores data chunks in line-based text format, where identifying entries requires finding line boundaries denoted by newline characters. After decompression, most NGS data is stored in line-based text files, e.g. fastq, SAM and VCF files. For SAM and VCF files, each line is an independent unit that contains its corresponding information (mapping records or genotypes). Thus, when loading these files, each line is read and stored as an element of an RDD. However, a fastq file stores its basic information in a four-line unit, where each line is an essential part of a sequencing read. When loading fastq files into Spark RDD, a filter step is applied to check each four-line unit before sending it out to be processed.

Both loading and saving run in parallel on each partition of the RDD (the sequencing data) with a default size of 256 MB per partition. Sparkhit can also directly load and save data from and to both HDFS and Amazon S3 by using the HDFS and the S3 URL scheme.

2.5 Fragment recruitment and short-read mapping implementations of Sparkhit
In metagenomic studies, fragment recruitment is a key step to understand the genome structure, evolution, phylogenetic diversity and gene functions of biological samples (Rusch et al., 2007). As a special case of read mapping, fragment recruitment supports more mismatches during the alignment. Thus, the computational complexity for fragment recruitment can be higher than for standard short read mapping.

We implemented a fast and sensitive fragment recruitment tool, called Sparkhit-recruiter. Sparkhit-recruiter extends the Fr-hit (Niu et al., 2011) pipeline and is implemented natively on top of the Apache Spark. The pipeline consists of four steps: (i) building reference index, (ii) searching candidate blocks, (iii) block filtering and (iv) banded alignment (Fig. 1C). When building the reference index, we use a K-mer hash table to store each K-mer’s location on the reference genome (Fig. 1C in blue color). Once the index is built, we extract the K-mers from sequencing reads (Fig. 1C in red color) and search against the reference hash table for exact matches. The matched K-mers will be placed on the genome as seeds to extend candidate blocks. The block filtering step incorporates a q-Gram threshold to remove badly matched blocks, therefore, improves run time performance. After filtering, banded alignment is applied to give a final mapping result.

We also implemented a short-read aligner, called Sparkhit-mapper (Fig. 1B and D). It adopts the same pipeline of Sparkhit-recruiter, but uses a more strict pigeonhole principle for the filtering step. Compared to the q-gram filter implementation in Sparkhit-recruiter, the pigeonhole principle allows fewer mismatches on the sequence so that less identical blocks are filtered and high similarity candidate blocks are preserved (see Supplementary file 1 Method-1.5.5). In this case, less candidate blocks are sent to the next step for banded alignment, resulting in much faster runtime than Sparkhit-recruiter.

To implement the pipelines in a distributed fashion, we split the mapping processes into two parts: building the reference index and
querying sequencing reads (Fig. 1C, D, blue and red dashed boxes). When starting a Sparkhit-recruiter job, the reference index is, firstly, built on the driver node (usually the master node), where the main Spark program runs. Once the reference index is built, the driver program executes a ‘broadcast’ command to ship one copy of the index to each worker node shared by all local querying tasks. On the worker nodes, sequencing data chunks are loaded from HDFS to a Spark RDD. Each partition of the RDD is, then, independently queried to the broadcasted reference index as a ‘map’ step of the MapReduce pipeline. In the end, a ‘reduce’ step summarizes the mapping result (Fig. 1B, see also Supplementary file S1 Method-1.5).

### 2.6 Parallelizing external tools using Sparkhit

To be flexible for different kinds of analyses, we built a general tool wrapper called Sparkhit-piper. It extends Spark’s ‘pipe’ function to invoke existing tools for analyses like sequence mapping, taxonomic profiling, gene expression quantification and genotyping (Fig. 1E). We use Spark RDD to split and distribute NGS data across cluster nodes. Then, distributed datasets are sent to the invoked tool via a standard input (stdin) stream. The tool processes input data in a batch and sends back the result to another RDD via a standard output (stdout) stream. In this case, the Spark RDD splits and distributes NGS data, while external tools carry out correspond computations. Sparkhit-piper is intuitive and flexible for users to parallelize their own scripts or tools directly without modifying their codes.

The implementation is based on Spark RDD’s ‘pipe’ function, where the RDD is able to send its data out of the JVM for processing in the operating system, like a Linux pipe operator (‘|’). The Java code can be expressed as:

```java
JavaRDD<String> MapRDD = FastqRDD.pipe(param.tool + param.toolOptions);
```

where ‘FastqRDD’ is the input RDD that stores sequence data and ‘MapRDD’ is the output RDD that stores mapping result. The ‘param.tool’ represents the full path of the tool executable while ‘param.toolOptions’ represents the corresponding tool parameters. Together, they assemble an external command that runs as an independent process on each partition of the input RDD.

### 2.7 Machine learning library

Downstream genomic data mining usually applies statistical tests and machine learning methods to analyze upstream results (gene abundance, taxonomic profile of different samples and genotypes of a certain cohort). We extended Spark’s machine learning library (Mllib) and integrated a variety of algorithms: (i) clustering, (ii) regression, (iii) chi-square test, (iv) correlation test and (v) dimension reduction (Fig. 1F). These algorithms are implemented with more RDD functions and iterative processes (see Supplementary file 1 Method-1.6).

### 3 Results

#### 3.1 General performance benchmarking for Sparkhit

In the following section we present a series of benchmarks for our framework (Fig. 2).

##### 3.1.1 Run time comparison between different mappers

We first measured the run times of all Sparkhit based mappers and compared them to Crossbow (Fig. 2A-D). Our toolkit ran faster than Crossbow across different numbers of worker nodes (30 and 50), different sizes of input data (1.3 TB and 545 GB) and different sizes of reference genomes (36 MB, 72 MB and 142 MB, see Supplementary file 1 Materials and Methods). Although Sparkhit-recruiter was slower than other Sparkhit based mappers, it recruited many more reads than standard short-read mappers such as BWA (Fig. 2H).

##### 3.1.2 Scaling performance of Sparkhit-recruiter

Sparkhit-recruiter scaled linearly with the increasing amount of input data on a 30 worker nodes Spark cluster (Fig. 2E). When scaled-out to more compute nodes, Sparkhit experienced slight slowdown after we increased the number of worker nodes to 60 (Fig. 2F). The slowdown is introduced by the overhead of building the reference index. However, since metagenomic fragment recruitment applications actively change reference genomes between different studies, the index building overhead is quite small and it runs much faster compared to other Burrows-Wheeler transform (BWT) based methods (Supplementary file S2, Fig. S5 and Discussion section).

##### 3.1.3 Preprocessing comparison with Crossbow

Data preprocessing is a critical step for interpreting cloud stored public datasets. Manually decompressed and distributed large amounts of genomic data on a cluster introduce significant overheads before data analysis. Although, several existing cloud tools have provided preprocessing functions (Decap et al., 2015; Langmead et al., 2009; Schatz, 2009), their preprocessing speeds are limited by their non-parallel implementations. We have compared the run time performances on data preprocessing between Sparkhit and Crossbow. For 338 TB Bzip2 compressed data (Data-1), Sparkhit ran 18 to 32 times faster than Crossbow on 50 and 100 worker nodes (Fig. 2G). Since Sparkhit utilizes all vCPUs for parallel decompression, its run time performance almost doubled from 50 nodes to 100 nodes, whereas Crossbow had similar run times on both clusters (Supplementary file S2, Table S12).

##### 3.1.4 Machine learning library benchmarking and run time performances on different clusters

For modules of the machine learning library, we have compared their run time performances on both a private cluster and the Amazon EC2 cloud (Fig. 21 and J). For the private cluster, data was stored on magnetic disks of a shared network file system (NFS). Whereas on Amazon EC2, data was stored on an HDFS deployed on solid state disks (SSD). We ran every module on both 20 and 40 nodes Spark clusters using a 200 GB VCF file from the 1000 Genomes Project and compared their run times between a private cluster and Amazon EC2. Since each module opened 640 and 1280 I/O tasks (20 nodes and 40 nodes, each node has 32 cores) to read input data and write output result, the run time performance on the private cluster was significantly slower than on the Amazon EC2 cloud (Supplementary file S2, Table S14). We also observed a significant improvement on run time for cached iterative computations (K-means clustering), compared to non-cached ones (Fig. 2K).

##### 3.1.5 Fragment recruitment comparison with MetaSpark

We have compared Sparkhit-recruiter to another Spark based distributed fragment recruitment tool (Fig. 3), MetaSpark (Zhou et al., 2017). Sparkhit-recruiter ran 92–157 times faster than MetaSpark across different numbers of worker nodes (10, 20 and 30), different numbers of input reads (1 million and 6 millions) and different sizes of reference genomes (72 and 142 MB). Although our tool recruited
10-12% reads less than MetaSpark using the same K-mer size, we have adjusted to a smaller K-mer size that recruits more reads than MetaSpark, while still ran 47-124 times faster (see Discussion section).

3.1.6 Accuracy and sensitivity of natively implemented tools
To assess the accuracy and sensitivity of Sparkhit-recruiter and Sparkhit-mapper, we used public datasets from the Genome in a Bottle Consortium (GIAB) (Zook et al., 2014) and simulated 6 datasets (doi: 10.4119/unibi/2914921) from three microbial reference genome templates (Ref-1, Ref-2 and Ref-3; Supplementary file S2, Table S2). In general, Sparkhit-recruiter has slightly higher accuracy than Fr-hit, Bowtie2 and SOAP, while having slightly lower accuracy than BWA (Fig. 2M–L and Supplementary file S2, Table S1). For sensitivity, Sparkhit-recruiter, Fr-hit, Bowtie2 and BWA are higher than SOAP on 100 nt simulated reads. For 150 nt simulated reads, all mappers have similar sensitivities. Sparkhit-mapper has slightly higher accuracy and sensitivity than Sparkhit-recruiter on the GIAB data.

3.2.1.3 Mapping 106 samples of the 1000 Genomes Project
The 1000 Genomes Project hosts all WGS data on Amazon S3 (Virginia region). It provides a detailed catalogue of human genetic variants in the studied populations. Before variant detection, a mapping step is required to present all matches and mismatches on the reference genome. To test the run time performance of Sparkhit on whole genome sequence mapping, we downloaded 5.6 TB compressed sequencing data to the Spark cluster. Using Sparkhit invoking BWA (Li and Durbin, 2009), we mapped 106 samples to a human reference genome. After sequence mapping, all results (SAM format) were uploaded to Amazon S3 for persistent storage. The entire process was completed in 5:31:17 h (Fig. 4D and Supplementary file S2, Table S7).

3.2.3 Gene expression profiling on prostate cancer RNA-seq data
Profiling tumor specific gene expression is a critical step for cancer research. To enable fast transcriptome quantification on the cloud, we tested the run time performance of Sparkhit on profiling 3.2 TB compressed RNA-seq data of a prostate cancer transcriptomics study (Wyatt et al., 2014). Since all datasets are archived on the ENA ftp server (see Supplementary file 1 Materials), the downloading process consumed 1:29:28 h, whereas gene expression profiling with Sparkhit invoked Kallisto (Bray et al., 2016) was completed in 21 min and 9 s (Fig. 4E and Supplementary file S2, Table S7). After uploading the final result to Amazon S3, we shut down all EC2 instances.

3.2.4 Recruiting HMP metagenomic data to microbial reference genomes
Metagenomics can capture and obtain genome fragments of uncultivated microbes (the microbial dark matter) by using shotgun sequencing to aggregated microorganisms sampled directly from the environment (Rinke et al., 2013). The Human Microbiome Project applied metagenomic shotgun sequencing to characterize microbial communities at different human body sites. As all HMP data is hosted on Amazon S3, we can directly access and analyze these metagenomic datasets on the cloud.

To demonstrate that associating HMP metagenomic data with private datasets (e.g. cultured microbial genomes or uncultured single cell genomes) can provide more biological insights, we downloaded and recruited all HMP whole genome shotgun data (2.3 TB compressed) to 12 selected microbial genomes (Ref-1, see Supplementary file S2, Table S2). Based on the metadata of the HMP samples, we present the fragment recruitment profile of 7 different microbial genomes across 6 different body sites (15 sub-body sites) in Supplementary file S2, Figure S1. The abundance of each microbial genome was normalized and illustrated across different mapping identities (from 75 percent to 100 percent). In general, sub body sites of the same main body site have similar profiles, but with few exceptions. For instance, the abundances of Neisseria meningitides are different between saliva and gingival plaque from the oral body site. Streptococcus aureus has higher abundance in buccal mucosa, attached gingivae and saliva compared to other sub body sites in the oral body site. By changing the input reference genomes,
users can easily produce the abundance profile of other microbes (see discussion).

3.4 Functional study based on HMP fragment recruitment result

Mapping metagenomic sequences to a new strain could reveal potential structural variances. A common method for functional studies is based on the hypothesis that such structural variances can have functional impact on the gene level. When recruiting HMP data to the reference genomes, the recruitment result was sensitive enough to pick up structural differences between genome sequences of two E.coli strains (Supplementary file S2, Fig. S1A and B). We, then, followed up a functional analysis, gene prediction and pathway enrichment, based on the sequences (see Supplementary file 1 Method-1.10). We extracted all gaps on the E.coli O104: H4 strain and predicted 158 new genes. These genes are enriched in two pathogenic pathways: Shigellosis and Pathogenic E.coli infection (Supplementary file S2, Fig. S2E-G). We annotated all genes and identified two toxic genes (Shiga toxin 2 subunit A and B) present in the E.coli strain that caused the 2011 German E.coli outbreak (Rasko et al., 2011).

4 Discussion

In this study, we presented a Spark based distributed computational framework for large scale genomic analytics, called Sparkhit. Sparkhit incorporates a variety of tools and methods that are programmed in the Spark extended MapReduce model. We have described (i) the implementation of a fragment recruitment tool and a short-read mapping tool using Spark’s RDD API, (ii) the construction of a general tool wrapper to invoke and parallelize external tools and (iii) the integration of Spark’s machine learning library for downstream data mining. We also presented the architecture of Sparkhit and the utilities that we used for deploying Spark clusters and downloading public datasets. Sparkhit outperforms most Hadoop and Spark based bioinformatics tools in computational run time. Using our framework, we analyzed large amount of public genomic data on the cloud within a short time.

The performance benchmarks demonstrated the scalability of Sparkhit. Sparkhit-recruiter scaled linearly (Fig. 2E) with the increasing amount of input data, as we utilized Spark RDD to balance data distribution and optimized the computational parallelization. In addition, the distributed data I/O via HDFS further reduces latency. On HDFS, data is distributed and loaded locally or from the closest node (depending on the redundancy setting of HDFS), avoiding massive data transfer across the network. We also observed the advantage of using HDFS when comparing the run times of the machine learning library between the Amazon EC2 cloud and the private cluster, which stored input data on an NFS shared by all worker nodes. On NFS, all data was read and written through the network connection to a mounted volume that can saturate the bandwidth. When scaling out to more worker nodes, a slight slowdown was observed (Fig. 2F). The slowdown was caused by the overhead of building the reference index, which runs solely on the driver node. This can be improved by pre-building the reference index using our locally implemented recruiter (a Java-based tool included in our framework). Moreover, the overhead for constructing reference index is small compared to the run time of the fragment recruitment process.

Our tool had excellent run time performance on data preprocessing compared to Crossbow (18–32 times faster) and significant run time improvement on fragment recruitment compared to MetaSpark (92–157 times faster). Although we recruited 10–12% less reads than MetaSpark, we can adjust to a smaller K-mer size that recruits slightly more reads than MetaSpark, while still ran 47–124 times faster (Fig. 3). In addition, our tool has a reasonable accuracy and sensitivity on sequence mapping (Supplementary file S2, Table S1). Sparkhit-recruiter also offers more options for fragment recruitment, such as an option for reporting the best match for each read and an option to choose between global or local alignment, whereas MetaSpark can only apply local alignment.

Spark and Hadoop based bioinformatics tools are not widely used in genomic studies as they require users to comprehend certain amount of knowledge on cloud computing. Therefore, we particularly focus on providing a simple and comprehensive cloud application to directly access and analyze public genomic datasets. We described a simple way to setup a Spark cluster on the Amazon cloud with one line of command. We also facilitated downloading and preprocessing large amounts of data by leveraging distributed Amazon S3 storage and optimizing parallel data decompression. In addition, Sparkhit enables users to parallelize their own tools or public bio-containers. Our large scale data analyses on 100 TB data presented how we completed the entire cloud utilization cycle in only 21 h. Moreover, the fragment recruitment application on all WGS data of HMP was completed in less than 2 h on Amazon EC2.

The fragment recruitment application presented a study model that users can easily query the entire HMP data to a personalized reference dataset on the cloud. In microbial studies, combining metagenomic data with microbial reference genomes has been commonly used (Eloe-Fadrosh et al., 2016). In our use case, we recruited all WGS data of the HMP to two different strains of E.coli: a toxic strain and a non-toxic strain. The intention is to find genome sequence segments that are not presented in the metagenomic samples. These sequence segments, which are unique for the toxic strain, might have functional impact that differs from the non-toxic one. We applied a functional analysis using the sequence segments and reproduced two toxic genes that caused the 2011 German E.coli outbreak. The same method can be applied to other isolates or single cell genomes.

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