RESEARCH ARTICLE

Robust long-read native DNA sequencing using the ONT CsgG Nanopore system [version 1; referees: 2 approved]

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Abstract

Background: The ability to obtain long read lengths during DNA sequencing has several potentially important practical applications. Especially long read lengths have been reported using the Nanopore sequencing method, currently commercially available from Oxford Nanopore Technologies (ONT). However, early reports have demonstrated only limited levels of combined throughput and sequence accuracy. Recently, ONT released a new CsgG pore sequencing system as well as a 250b/s translocation chemistry with potential for improvements. Methods: We made use of such components on ONTs miniature 'MinION' device and sequenced native genomic DNA obtained from the near haploid cancer cell line HAP1. Analysis of our data was performed utilising recently described computational tools tailored for nanopore/long-read sequencing outputs, and here we present our key findings. Results: From a single sequencing run, we obtained ~240,000 high-quality mapped reads, comprising a total of ~2.3 billion bases. A mean read length of 9.6kb and an N50 of ~17kb was achieved, while sequences mapped to reference with a mean identity of 85%. Notably, we obtained ~68X coverage of the mitochondrial genome and were able to achieve a mean consensus identity of 99.8% for sequenced mtDNA reads. Conclusions: With improved sequencing chemistries already released and higher-throughput instruments in the pipeline, this early study suggests that ONT CsgG-based sequencing may be a useful option for potential practical long-read applications.

Keywords

Nanopore sequencing, long read sequencing, CsgG, R9.4, Oxford Nanopore, complex genomes, HAP1
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Introduction

In 1977, Sanger and colleagues at the Laboratory for Molecular Biology in Cambridge, England, described an efficient sequencing-by-synthesis (SBS) approach to determine the order of nucleobases in DNA molecules (Sanger et al., 1977). This event would revolutionise genetics research, including enabling the first near-complete sequencing of the human genome (Lander et al., 2001; Venter et al., 2001). The practical success of the method is perhaps only equaled by another SBS-based technique, also developed in Cambridge by Balasubramanian, Kleneman and colleagues (Bentley et al., 2008), and which was later commercialised to huge commercial success by the biotech company Illumina. While Illumina sequencing, as well as other second generation sequencing methods, enable high-throughput yields with very good to excellent accuracy, they all suffer from the major drawback of short read lengths, which place some important limitations on their practical applicability. For example, long-read sequencing could enable more complete and efficient de novo assembly of complex genomes (Goodwin et al., 2016), and may conceivably allow for efficient haplotype phasing to characterise clinically-relevant mutations (Goodwin et al., 2016; Hussain, 2015). For RNA sequencing applications, a more reliable method for profiling the alternative isoform composition of complex transcriptomes might also be possible (Steijger et al., 2013).

The development of long read sequencing methods using ‘nanopores’ has in fact been the focus of several academic laboratories for over two decades (Akeson et al., 1999; Cherf et al., 2012; Derrington et al., 2010; Kasiannovicz et al., 1996; Meller et al., 2000; Stoddart et al., 2009). More recently, the technique has been further developed and commercialised by Oxford Nanopore Technologies (ONT) (Brown & Clarke, 2016). ONT sequencing works by placing a nanopore, which in the case of the ONT platform is a protein pore, in a conducting electrolyte solution and applying a small potential difference across the pore. Nucleotide kmer-specific signatures of current fluctuations as a nucleic acid strand passes through a nanopore are then recorded to determine the sequence. A critical consideration is the size and characteristics of the sensing aperture of the pore, which determines how many nucleotides present in the pore contribute to the recorded current (Derrington et al., 2010). With smaller/ more optimal sensing apertures, less nucleotides influence the characteristics of the recorded current, making distinguishing nucleotide sequences subject to much less noise. Thus using an optimally structurally configured protein pore for sequencing is a key determinant of sequencing accuracy. A second critical factor that influences sequencing accuracy, as well as throughput, is the speed and manner of DNA translocation through the pore, and in order to exert control over such parameters motor enzymes that are able to ratchet DNA into the pore at a suitable speed are employed (Cherf et al., 2012).

Thus far ONT have only commercially released their entry-level miniature ‘minION’, currently marketed as a pocket-sized portable sequencing device, and some notable successes have been achieved that have taken advantage of its portability (Quick et al., 2015; Quick et al., 2016). However, given the superiority in producing long reads, it may also be useful to consider the applicability of the minION, and thus the ONT platform in general, for more general laboratory research. Previous benchmarking works have reported limited success in yielding combined sequence accuracy and throughput (Ip et al., 2015; Laver et al., 2015), but these published studies have utilised older versions of protein pores referred to by ONT as ‘R6’ or ‘R7’. While the identity of the R6/R7 pores remain undisclosed, a low raw sequencing accuracy meant that reads needed to be sequenced in ‘2D’, where two complementary DNA strands are joined by a hairpin adapter allowing for their sequential sequencing through the nanopore. More recently, ONT have released an ‘R9 series’, which they have revealed is based on the CsgG bacterial amyloid secretion pore (Brown & Clarke, 2016). Presumably with a more optimally configured sensing aperture, the CsgG pore is reportedly capable of higher sequencing accuracy. A mutant form of this pore, ‘R9.4’, is the current version favoured by ONT, reportedly currently yielding the highest accuracies and pore stability. Such developments in improving raw sequence accuracy could also potentially mean that 1D sequencing, and thus higher throughput, might be possible without accuracy levels falling unacceptably low. In addition, new motor enzymes that are capable of ratcheting the DNA through pores at higher speeds have also been made available, adding further potential for increased throughput. Here we describe the use of the CsgG R9.4 nanopore system, in use with a sequencing chemistry that operates a translocation speed of 250b/s, to robustly produce long sequence reads from native human genomic DNA obtained from human HAP1 cells.

Methods

Cell culture and genomic DNA extraction

Early passage HAP1 cells (Horizon Discovery) were grown in Iscove’s Modified Dulbecco’s Medium (Thermo Fisher Scientific) supplemented with 10% fetal bovine serum (Thermo Fisher Scientific), and maintained at a temperature of 37°C in a humidified incubator with 5% CO₂. Cells were harvested by washing in PBS and then incubating with Trypsin-EDTA followed by further washing of detached cells in PBS. The PureLink Genomic DNA Mini Kit (Invitrogen) was used to isolate purified genomic DNA, as per the manufacturer’s instructions.

Library preparation and sequencing

In general, we followed ONT protocols for library preparation. Wherever significant deviations were made to their recommendations, these are indicated by an asterix and are explained as a note.

500ng of genomic DNA was fragmented by shearing through a 15 gauge needle* 10 times. Damage to DNA was then repaired using the preCR repair mix (NEB)**, according to manufacturer recommendations and DNA subsequently purified using AMPureXP beads (Beckman Coulter). End-repair of DNA fragments was then performed using the Ultra II End Prep module (NEB). Ligation of ‘E7’ motor protein-complexed AMX adapter (ONT, NSK007) to genomic DNA ends was next carried out using the NEBNext Ultra II Ligation Module (NEB)**. Another round of AMPure XP purification was then performed before the DNA library was eluted and loaded onto a running buffer-primed flow cell for sequencing. Sequencing of the native genomic DNA
was performed on a single R9.4/FLO-MIN106 flow cell on a
MinION Mk1B for 30 hours and base-calling performed using the
cloud-based Metrichor/EPI2ME platform. EPI2ME split reads into
a ‘pass’ folder containing high quality reads and a ‘fail’ folder con-
taining low quality reads.

*To enable generation of long fragments, we utilise DNA shearing
through a 15 gauge needle, as opposed to g-tube shearing which is
recommended in the ONT protocol.

**Although we performed sequencing of native DNA without PCR,
we use this DNA repair step primarily for repairing of nicks that
may otherwise disrupt the sequencing of long fragments.

**ONT workflows recommend the use of the Blunt/TA ligation
module (NEB) for ligation of adapters. We instead use the Ultra II
ligation module (NEB), which is more compatible with end-repair
components performed in the previous step, and further allows the
intervening wash step to be omitted.

Computational analysis

The basecalled pass/high quality and fail/low quality reads were
processed separately through poretools (0.6.0; https://poretools.
readthedocs.io/en/latest/) (Loman & Quinlan, 2014) to obtain the
embedded FASTQ data. The respective FASTQ data were mapped
to the reference human genome (GRCh38 Primary Assembly;
GenBank: GCA_000001405.15) using GraphMap (0.3.1;
https://github.com/isovic/graphmap) (Sović et al., 2016) with the
--sensitive parameter enabled on a High Performance Comput-
ing cluster. High quality reads were mapped in 2d:17h using
14 threads. The resulting alignments were processed using sam-
tools (1.4; http://www.htslib.org/) (Li et al., 2009) to obtain
basic mapping statistics and BAM files visualised using Tablet
(1.16.09.06; https://ics.hutton.ac.uk/tablet/) (Milne et al., 2013).

The high quality read alignments were processed through Quali-
Map (2.2.1; http://qualimap.bioinfo.cipf.es/) (Okonechnikov
et al., 2016) to obtain further metrics, such as GC Distribution
(Figure 1C). We employed Quinlan’s python scripts at https://
github.com/arq5x/nanopore-scripts (Quick et al., 2014) to

Figure 1. Overview of throughput and quality of mapped reads. (A) Summary of the sequencing output and mapping performance.
General alignment characteristics are displayed in Figure S1. (B) Read length distribution of high quality reads counted in 1 Kbp bins and
represented on a log10 scale (y-axis). Note that reads shorter than 80 bp are not considered for mapping by GraphMap by default. (C) GC
content distribution as output by Qualimap. (D) Kernel density plot displaying the distribution of mapped read identity with the mean average
(85%), indicated with a vertical line. (E) Pie chart breaking down the different types of errors estimated from AlignQC; further details on the
context of the errors detected is provided in Figure S2.
determine identity to reference (Figure 1D) and complementary metrics, such as alignment profiles (Figure S1). A more detailed estimate of the error types was also obtained using AlignQC (1.2; https://www.healthcare.uiowa.edu/labs/au/AlignQC/) (Weirather et al., 2017) as summarised in Figure 1E (also see Figure S2).

Coverage data were extracted using samtools and bedtools (2.26; http://bedtools.readthedocs.io/en/latest/) (Quinlan & Hall, 2010) and plotted using R with ggplot2 and reshape2 packages (R Core Team, 2016; Wickham, 2007; Wickham, 2009). Gene coordinates were obtained from the GENCODE Release 25 Primary GFF annotation (https://www.gencodegenes.org/releases/25.html). To visualise the influence of coverage on consensus quality with current ONT chemistry and methodology, we performed random downsampling of the mitochondrial genome alignments (which presented the best coverage) using samtools to obtain a range of different coverages. Consensus calling was then performed using Ivan Šović’s majority consensus calling script (https://github.com/isovic/samscripts/src/consensus.py) (Šović et al., 2016) with a minimum coverage of 1, returning the consensus differences with the reference.

Results
Yield and mapping characteristics
The experimental strategy employed in this study was aimed at optimising combined MinION output with regards to throughput, read-length and mapping identity from a limited amount (500ng) of non-reference (a cancer genome) starting DNA material. Though ultra-long reads (>100kb) are theoretically possible on the ONT platform, we opted for needle shearing for fragmentation of genomic DNA with the aim of yielding long sequence reads, the computational analysis tools for which have been previously optimised (Šović et al., 2016). To further improve throughput we employed a 1D sequencing strategy; though this would inevitably lead to a loss in base-calling accuracy compared to 2D sequencing, we reasoned this would likely be compensated by improvements in raw sequencing accuracy offered by the new ONT CsgG R9.4 system employed.

From the single 30 hour minION run of HAP1 native genomic DNA, a total of ~329,000 1D reads comprising ~2.8 billion bases were sequenced (Figure 1A). ~247,000 were high quality reads, and of these ~97% successfully mapped to human reference genome. Thus ~240,000 mapped high quality reads consisting of ~2.3 billion bases were taken forward for analysis in this study. The read length distribution yielded a mean of ~9.6kb and an N50 of ~17kb, and the longest successfully mapped read obtained was ~113kb in length (Figure 1B). No negative bias in terms of sequencing GC-rich sequences was apparent in our dataset (Figure 1C). A mean mapping identity of 85% to reference was achieved from the 1D reads obtained from the native cancer genome (Figure 1D), and most called errors were either mismatches or deletions including homopolymer-associated deletions (Figure 1E). Indeed, ONT protein nanopores have previously generally displayed difficulties in resolving homopolymers that exceed the sensing aperture length of the pore (Jain et al., 2016), and our results further suggest that this problem persists in data generated using the R9.4 nanopore.

Barring a 30Mb diploid region spanning a portion of chromosome 15, human HAP1 cells are a fully haploid cell line (Carette et al., 2011; Essletzbicher et al., 2014), and thus generally represent a particularly amenable tool for CRISPR-Cas9 mediated genome editing in potential studies of genetic function. Accordingly, coverage obtained from our dataset along the haploid genome appeared fairly uniform, except for a portion of chromosome 15 which likely corresponds to the disomic region of the genome (Figure 2A and B). With an N50 of ~17Kb in our dataset, as expected, a significant proportion of reads covered the full length of at least a single annotated gene (Figure 2C).

Mitochondrial genome consensus calling
Notably, a particularly high density of reads mapped to the ~17kb mitochondrial genome, for which we achieved 68x coverage. Most of the mtDNA reads covered a significant portion of the mitochondrial genome, with a few reads indeed covering the entire genome (Figure 3A). The mtDNA reads mapped with 84% identity to reference, and as with our observations for the nuclear genome, most called errors were either deletions or mismatches (Figure 3B). Since we had obtained a significant degree of coverage for the mitochondrial genome, we proceeded to inspect how errors can affect final consensus calling at increasing coverage levels, which at full coverage achieves 99.8% mean identity to reference (Figure 3C). Our analysis revealed that ~10X coverage was required to prevent most random mismatches from being called, beyond which mismatches at specific positions remained (Figure 3C). Insertions made up a smaller proportion of the total called errors (Figure 3B), and required a lower coverage for complete correction (Figure 3C). Deletions, which included homopolymer deletions, required a much greater degree of coverage, and some deletions still remained with the highest level of coverage achieved (Figure 3C). However, following consensus calling, the majority of the total called errors remaining were in fact mismatches (Figure 3D). Since we observed that mismatch error trend halted at ~10X coverage, it is quite likely that the remaining mismatches in fact represent genuine non-reference bases, i.e. they are a true reflection of the native DNA sequenced.

Such non-reference nucleotides are likely to include unrepaired DNA damages and single nucleotide variants, and possibly also naturally modified bases.

Discussion
Here we describe the sequencing of native DNA obtained from the near-haploid human cancer cell line HAP1. Utilising recently made available sequencing pores and chemistries, we report significant improvements in combined throughput and raw sequencing accuracy compared to that previously published using older pore-type and sequencing chemistry versions (Ip et al., 2015; Laver et al., 2015). While we achieved only an 85% mapping identity of sequenced bases from 1D reads, there are a couple of relevant considerations. Firstly, we sequenced native DNA, and thus DNA modifications and unrepaired DNA
Figure 2. Overview of coverage achieved over the genome. (A) Coverage deviation per million base pairs for each chromosome. Horizontal gridlines represent the average coverage (0.8 x) with values ranging between a minimum and maximum range of -2 and +2 standard deviations from the mean. (B) Average coverage per million base pairs, the largest increase in chromosome 15 corresponds with the remaining diploid region of HAP1 cells. (C) Counts of GENCODE 25 genes with all positions covered by at least 1 base pair and varying levels of depth for each chromosome.
damages were unlikely to be base-called correctly. Secondly, we used a cancer cell line with unknown identity to reference. According to ONT data (https://nanoporetech.com), ~90% accuracy should be achievable with R9.4 1D sequencing, and our mapping data possibly fits well with such reports. It may further be noted that higher accuracies of ~95%, are potentially achievable with R9.4 2D sequencing, though at the cost of significant drops in throughput. However, in this regard, it is relevant that ONT have more recently released a sequencing chemistry that operates a 450b/s translocation speed, thus reportedly further improving throughput significantly (ONT data; https://nanoporetech.com). Using our fragmentation protocol, a mean read length of ~9.6kb with an N50 of ~17kb was achieved. We opted for needle shearing as a demonstration of a quick and inexpensive method for random DNA fragmentation; however, it may be possible that even longer mappable reads can also robustly be obtained by performing library preparation without an intentional fragmentation step. Indeed, thousands of high quality reads of more than 50kb in length were mapped in our study, further suggesting that robust ‘ultra-long’ read generation should also be readily possible on the platform.

Following consensus calling, we achieved 99.8% mean consensus identity in reads mapped to the mtDNA genome, for which we had obtained 68X sequencing coverage. Our analysis suggested that the majority of remaining non-reference bases in our mtDNA reads were likely genuine features of the native DNA, indicating that remaining true errors may in fact have been potentially reduced to near-zero levels. However, it should be noted that this observed high efficiency in consensus calling might well have been in part owing to the simple nature of the mtDNA genome. Homopolymer associated error-correction in more complex genomes, for example, may prove less efficient, and efforts focussed on optimising sequencing chemistries and base-calling parameters to help resolve such issues would be most welcome. Nonetheless, very high levels of sequence coverage obtained during ONT
sequencing could conceivably compensate for its lower raw sequencing accuracy for applications associated with genetic variant calling. In this regard, it is relevant that ONT have begun the beta-testing phase for their benchtop sequencer, the PromethION, reportedly capable of yielding 6TB of sequence from a 24-hour run (Jain et al., 2016). The ultra-high-throughput instrument will operate on the same pore type and chemistries to that used for the MinION, i.e. likely similar iterations to those used in the current study. It is likely that high-throughput, long-read sequencing on the ONT platform may enable significant advances in genetics applications relevant to complex genomes to be made in the near future.

Data availability
Sequence data used for analysis in this study is publicly archived at the European Nucleotide Archive (ENA) under accession code ERR189853. Files contain high quality sequence data, as well as associated alignment data.

Supplementary material
Figure S1: Alignment profile of mapped reads. Dotplot generated using Quinlan’s nanopore scripts (https://github.com/arq5x/nanopore-scripts; Quick et al., 2014), which incorporates two factors for each read, the fraction of the read aligned and the identity (or number of matches versus alignment length + insertions). Note that a wide spread of read fractions aligned may be expected when aligning a cancer genome to reference.

Click here to access the data.

Figure S2: Detailed error contexts from AlignQC. AlignQC output displaying if there was a bias towards mismatches/deletions/insertions in specific contexts. The error contexts seem to differ somewhat from those published in the AlignQC paper (Weirather et al., 2017); such differences have likely arisen largely owing to the nature of different material sequenced, i.e. cDNA in the Weirather et al. study and native cancer genome DNA in ours.

Click here to access the data.

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Carter and Hussain have presented MinION nanopore sequencing and analysis results using R9 chemistry. Their choice of HAP1 as DNA substrate is an interesting one, given its (mostly) haploid nature. The error rate measurements in this study agree with measurements reported by other MinION users for R9 chemistry. Given the rapid evolution of MinION sequencing, data from the newer R9.4 MinION chemistry (and soon to arrive R9.5) are superior than R9 data.

However, this study will be useful to the nanopore community in form of library protocols as well as computational analysis tools.

Is the work clearly and accurately presented and does it cite the current literature?
Yes

Is the study design appropriate and is the work technically sound?
Yes

Are sufficient details of methods and analysis provided to allow replication by others?
Yes

If applicable, is the statistical analysis and its interpretation appropriate?
Yes

Are all the source data underlying the results available to ensure full reproducibility?
Yes

Are the conclusions drawn adequately supported by the results?
Yes

Competing Interests: No competing interests were disclosed.

I have read this submission. I believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.
Carter and Hussain present details of the use of Oxford Nanopore's MinION, specifically R9.4 chemistry using the CsgG pore, to sequence HAP1. Though this particular sequencing kit has been surpassed by a faster kit, nevertheless the description of the sequencing serves as an adequate guide for MinION data quality and potential methods for analysing MinION data

Is the work clearly and accurately presented and does it cite the current literature?
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Yes

Are the conclusions drawn adequately supported by the results?
Yes

**Competing Interests:** No competing interests were disclosed.

**Referee Expertise:** Genomics, bioinformatics and sequencing

I have read this submission. I believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.