HiCanu: accurate assembly of segmental duplications, satellites, and allelic variants from high-fidelity long reads

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Complete and accurate genome assemblies form the basis of most downstream genomic analyses and are of critical importance. Recent genome assembly projects have relied on a combination of noisy long-read sequencing and accurate short-read sequencing, with the former offering greater assembly continuity and the latter providing higher consensus accuracy. The recently introduced Pacific Biosciences (PacBio) HiFi sequencing technology bridges this divide by delivering long reads (>10 kbp) with high per-base accuracy (>99.9%). Here we present HiCanu, a modification of the Canu assembler designed to leverage the full potential of HiFi reads via homopolymer compression, overlap-based error correction, and aggressive false overlap filtering. We benchmark HiCanu with a focus on the recovery of haplotype diversity, major histocompatibility complex (MHC) variants, satellite DNAs, and segmental duplications. For diploid human genomes sequenced to 30× HiFi coverage, HiCanu achieved superior accuracy and allele recovery compared to the current state of the art. On the effectively haploid CM13 human cell line, HiCanu achieved an NG50 contig size of 77 Mbp with a per-base consensus accuracy of 99.999% (QV50), surpassing recent assemblies of high-coverage, ultralong Oxford Nanopore Technologies (ONT) reads in terms of both accuracy and continuity. This HiCanu assembly correctly resolves 337 out of 341 validation BACs sampled from known segmental duplications and provides the first preliminary assemblies of nine complete human centromeric regions. Although gaps and errors still remain within the most challenging regions of the genome, these results represent a significant advance toward the complete assembly of human genomes.

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HiFi sequencing libraries are size selected for a target fragment size (currently up to 25 kbp).

Although the resulting read lengths are modest by the modern long-read sequencing standards—PacBio CLR reads frequently exceed 50 kbp, and ultralong Oxford Nanopore Technologies (ONT) reads can exceed even 100 kbp (Jain et al. 2018b)—HiFi is a major leap forward in terms of long-read read accuracy. As the accuracy of other long-read technologies have not exceeded 95%, the median accuracy of current HiFi reads can exceed 99.9% (>Q30), making them a promising data type for separating highly similar repeat instances and alleles.

Early studies adopting HiFi sequencing showed improved variant calling and repeat resolution (Wenger et al. 2019; Vollger et al. 2020). However, these early assemblies were limited to resolving repeats with >1% sequence divergence, owing to limitations of existing tools (Wenger et al. 2019). The recently developed Peregrine assembler (Chin and Khalak 2019) greatly reduced assembly runtime and improved consensus accuracy, removing the need for postprocessing, but did not address the issue of suboptimal repeat resolution or allele separation. Other recent work combined HiFi sequencing with complementary data types, such as parental information (Wenger et al. 2019), Hi-C (Garg et al. 2019), and Strand-seq (Porubsky et al. 2019), to obtain chromosome-scale, haplotype-resolved assemblies.

In the following sections, we present HiCanu, a modification of the Canu assembler (Koren et al. 2017) designed to take full advantage of the high accuracy of HiFi reads. We evaluate HiCanu’s ability to resolve near-identical genomic repeats, with a focus on centromeric repeats and SDs by comparing our results to other HiFi and recent ultralong Oxford Nanopore-based human assemblies (Miga et al. 2020; Shafin et al. 2020). Furthermore, we evaluate HiCanu’s ability to capture both alleles in large phase blocks in a diploid human genome.

Results

HiCanu overview

HiCanu builds on the original Canu pipeline, replacing or significantly modifying its core methods. Here we provide an overview of the new pipeline, highlighting the introduced changes, although a more detailed description of individual steps can be found in the Methods section. Whereas the original Canu pipeline starts with read self-correction, which can homogenize reads from different alleles or near-identical repeat instances, HiCanu begins by compressing all consecutive copies of the same nucleotide to a single base (e.g., “AA...” becomes “A”). In accordance with the earlier observation that misestimation of homopolymer length is the primary error mode of HiFi technology (Wenger et al. 2019), the resulting homopolymer-compressed reads (or “compressed reads” for short) accurately encode the transitions between different bases of the underlying genomic regions. The compressed reads are then trimmed based on their overlaps to other reads to remove any chimeric sequences or sequencing adapters (see overlap based trimming in Koren et al. 2017), and the overlaps are recomputed on the trimmed reads. The overlap error adjustment (OEA) module (Holt et al. 2002; Koren et al. 2017) examines read overlap pileups to identify remaining sequencing errors in the individual reads and recomputes overlap alignment identities. Following our observation that microsatellite repeat arrays are also prone to HiFi read errors, the OEA procedure was modified to ignore any differences within these regions when computing the final alignment identity of two overlapping reads. Compared with the initial reads (Fig. 1A), homopolymer compression (Fig. 1B), pileup-based read correction (Fig. 1C), and ignoring differences in microsatellite repeats (Fig. 1D) result in a drastic reduction of observed sequencing errors.

Draft contigs are then formed from the adjusted overlaps using Canu’s Bogart module (Koren et al. 2017), modified to better handle heterozygous variants and consider only high-identity read overlaps. The final contig sequences are obtained by computing a consensus across the original, uncompressed reads, arranged according to the layout of their compressed versions. Similar to many modern assemblers, when faced with a diploid genome, HiCanu outputs contigs as “pseudohaplotypes” that preserve local allelic phasing but may switch between haplotypes across longer distances. A single set of contigs representing all resolved alleles is output regardless of ploidy, and additional processing with a tool such as Purge_dups (Guan et al. 2020) is required to partition the contigs into primary and alternate allele sets.

Drosophila genome assembly

We first evaluated HiCanu on a 24-kbp HiFi library from a Drosophila melanogaster F1 hybrid (ISO1×A4; see Data access). To match typical coverage, the HiFi data set was down-sampled to

Figure 1. Impact of HiCanu processing on observed read quality. (A) Two hypothetical reads are shown with sequencing errors highlighted in red. (B) The first step of HiCanu is to compress homopolymers, which obscures homopolymer length errors but retains enough information to accurately distinguish reads from different genomic loci. (C) Overlaps are then computed for the compressed reads, and remaining errors are identified by examining the alignment pileups (gray rectangle). (D) Finally, after correcting the identified errors (blue) and ignoring indels in regions of known systematic error (gray), the resulting overlap is 100% identical. (Right) Sequence identity of reads from a 20-kbp HiFi library measured against the CHM13 Chromosome X reference sequence v0.7 (Miga et al. 2020) after each step of HiCanu processing (Supplemental Note 1). Separate boxplots are shown for initial raw HiFi reads (init), homopolymer-compressed reads (compressed), OEA-corrected reads (corrected), and corrected reads after ignoring differences in microsatellite repeats (masked). The median read identity, indicated by solid segments, increases from <99.9% to 100% (note the plot shows y-range of 99.65%-100%). Supplemental Table S1 also shows how HiCanu processing increases the percentage of perfectly aligned (100% identity) HiFi reads from <1% to >97%. 

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Because Purge_dups can split and/or trim the initial contigs but v5.02 (Gurevich et al. 2013) to compare the assemblies against NG50 over Canu).

HiCanu improved over all other assemblies both haplotypes together and output few alternate contigs (total length <21 Mbp). Peregrine and CLR assemblies. The Peregrine assembly collapsed noticeably more continuous than any other assembly as measured by BUSCO completeness (>98%). BUSCO duplication values were 4.54% (error); for example, QV40 = 99.99% accuracy. QUAST diffs reports the number of large structural discrepancies observed between the assemblies. Considering that one of the haplotypes (derived from the A4 parental strain) is expected to produce a low number of haplotype switches and mostly preserve long-range phasing. We used Merqury (Rhié et al. 2020) to split the initial contigs into continuous phase blocks, based on haplotype-specific k-mer markers inferred from parental Illumina reads (Supplemental Note 2). As a baseline, we considered a haplotype-resolved assembly produced by TrioCanu (Koren et al. 2018) generated using a combination of CLR reads and parental Illumina data. The HiCanu primary (alternate) contig set has an estimated phase block NG50 of 7.62 Mbp (4.45 Mbp), a maximal block length of 25.4 Mbp (10.1 Mbp), and a low percentage of discordant markers within predicted haplotype blocks (switch rate) of 0.03% (0.02%). For comparison, the TrioCanu assembly has a paternal-ISO (maternal-A4) phase block NG50 of 13.9 Mbp (21.39 Mbp), a max block size of 24.7 Mbp (27.7 Mbp), and an intra-block switch rate of 0.1% (0.04%). In contrast, the phase blocks of all other considered assemblies are much less continuous (at least a 3.5-fold drop in phase block NG50 compared to HiCanu) and, in the case of Peregrine and CLR assemblies, a much higher switch error (Table 1).

**Human genome assemblies**

We next ran HiCanu, Canu, and Peregrine on three different human data sets (see Data access): a 20-kbp library of the completely homozygous cell line CHM13 (Kronenberg et al. 2018; Miga et al. 2020; Vollger et al. 2020), a 15-kbp library of the Ashkenazic cell line HG002 from the Personal Genome Project (Church 2005; Wenger et al. 2019), and a combined library (12% 10 kbp, 62% 15 kbp, 26% 20 kbp) for the Puerto Rican cell line HG00733 from the 1000 Genomes Project (The 1000 Genomes Project Consortium 2012; Purbsky et al. 2019). All data sets consist of

| Assembly          | Size (Mbp) | NG50 (Mbp) | Quality (QV) | BUSCO complete | Phase block NG50 (Mbp) | Intra-block switch error | QUAST (diffs per Mbp) |
|-------------------|------------|------------|--------------|----------------|------------------------|--------------------------|------------------------|
| Canu + Purge_dups | 141.81     | 14.09      | 37.4         | 98.5%          | 0.42                   | 3.86%                    | 0.018                  |
| CLR               | 128.15     | 0.31       | 35.5         | 86.7%          | 0.25                   | 2.97%                    |                        |
| Peregrine         | 141.59     | 12.68      | 32.9         | 98.2%          | 0.07                   | 1.78%                    |                        |
| HiFi              | 20.53      | 0.00       | 33.5         | 1.0%           | 0.00                   | 3.71%                    |                        |
| Canu + Purge_dups | 145.19     | 13.72      | 51.9         | 98.7%          | 2.04                   | 0.03%                    | 0.015                  |
| HiFi              | 130.23     | 1.28       | 46.9         | 93.7%          | 1.26                   | 0.03%                    |                        |
| HiCanu + Purge_dups | 146.27    | 20.16      | 51.0         | 98.8%          | 7.62                   | 0.03%                    | 0.025                  |
| HiFi              | 132.53     | 4.54       | 46.7         | 95.2%          | 4.45                   | 0.02%                    |                        |

A genome size of 143,726,002 was used for NG50 computation. Only contigs ≥50 kbp were used for all analyses except QUAST. Each assembly is reported after running Purge_dups, with the primary reported on top and als on the bottom of each row. Phred-style consensus quality values are QV = −10 log P(error); for example, QV40 = 99.99% accuracy. QUAST diffs reports the number of large structural discrepancies observed between the assemblies before Purge_dups (and therefore have only one entry per assembly) and the D. melanogaster ISO1 reference genome normalized by the assembly size (in Mbp). The best performing result for each genome in each metric is highlighted in bold.
approximately 30× HiFi sequencing coverage. For the HG002 data set, we reused the best assembly from a recent study (Wenger et al. 2019) as it reflects a Canu 1.7.1 assembly before HiCanu’s development and the associated improvements to Canu’s core modules. We additionally included the most continuous published (pseudo-haplotype) assemblies of the same genomes, which relied on ultralong Oxford Nanopore reads to achieve state-of-the-art repeat resolution (Miga et al. 2020; Shafin et al. 2020). As before, contigs <50 kbp were excluded from analysis. As the sizes of HiCanu assemblies for the diploid data sets HG002 and HG00733 were 5.30 Gbp and 5.46 Gbp, respectively, compared with a haploid genome size of 3.1 Gbp, we again postprocessed all diploid assemblies with Purge_dups and computed statistics for both primary and alternate contig sets.

Per-base consensus quality was again estimated by Merqury (Rhie et al. 2020) using Illumina data from the corresponding genome (Supplemental Note 2; Supplemental Table S3). To assess the structural correctness of the assemblies, we followed the methodology of Shafin et al. (2020). Namely, structural differences reported by QUAST v5.0.2 versus the human reference genome GRCh38 (Schneider et al. 2017) were postprocessed to ignore breakpoints in centromeric regions and annotated SDs, in order to reduce the number of false positives (Methods; Supplemental Table S3). As before, because Purge_dups may introduce or correct misassemblies as it modifies the contigs, the structural correctness assessment was performed on the original assemblies.

Primary contig summary statistics for the three human genomes are presented in Table 2. The continuity of HiCanu assemblies, as measured by NG50, exceeded that of all other HiFi-based assemblies and even rivaled the continuity of Nanopore ultralong-read assemblies. Reported rates of structural differences for assemblies and even rivaled the continuity of Nanopore ultralong-read assemblies. To recognize a BAC as resolved within the assembly, we required 99.5% of the BAC length to be aligned to a single contig by minimap2 (Methods; Li 2018). Assembly sequence accuracy was measured as the median alignment identity of resolved breakpoints in centromeric regions and SDs, in order to reduce the number of false positives (Methods; Supplemental Table S3). As before, because Purge_dups may introduce or correct misassemblies, the structural correctness of the assemblies, we followed the methodology of Shafin et al. (2020). Namely, structural differences reported by QUAST v5.0.2 versus the human reference genome GRCh38 (Schneider et al. 2017) were postprocessed to ignore breakpoints in centromeric regions and annotated SDs, in order to reduce the number of false positives (Methods; Supplemental Table S3).

| Genome   | Assembly | Size (Gbp) | NG50 (Mbp) | Quality (QV) | QUAST (diffs per Gbp) |
|----------|----------|------------|------------|--------------|-----------------------|
| CHM13    | ONT      | 2.98       | 74.06      | 28.9         | 35.2                  |
|          | Peregrine| 2.87       | 37.30      | 50.7         | 40.8                  |
|          | Canu     | 3.03       | 45.63      | 61.7         | 40.6                  |
|          | HiCanu   | 3.05       | 77.12      | 58.1         | 45.9                  |
|          | ONT      | 2.83       | 32.34      | 23.2         | 59.0                  |
|          | Peregrine| 0.07       | 0.00       | 23.4         | 35.7                  |
|          | Canu     | 2.82       | 32.09      | 46.0         | 35.7                  |
|          | 0.04     | 0.00       | 33.5       |              |                       |
|          | HiCanu   | 2.99       | 0.18       | 45.1         | 44.6                  |
|          | ONT      | 2.15       | 4.63       | 51.8         | 44.6                  |
|          | Peregrine| 0.12       | 0.00       | 43.2         |                       |
|          | Canu     | 2.97       | 0.18       | 45.1         |                       |
|          | 0.36     | 0.00       | 41.5       |              |                       |
|          | HiCanu   | 2.32       | 38.43      | 50.6         | 57.5                  |

A genome size of 3,098,794,149 was used for computing NG statistics. As in Table 1, only contigs ≥50 kbp were used for all analyses except QUAST. Each assembly is reported after running Purge_dups, with the primary reported on top and alts on the bottom of each row. Structural differences in centromeric regions and SDs were ignored owing to instability in these regions, and diffs were normalized by assembly size (in Gbp). Supplemental Table S4 includes results for a different 10-kbp HiFi library. Superscripts mark previously published assemblies: *Miga et al. (2020); Shafin et al. (2020); Wenger et al. (2019). All ONT assemblies were generated by Canu, as it achieved the highest NG50 and BAC resolution. The best-performing result for each genome in each metric is highlighted in bold. Statistics for assemblies from Garg et al. (2019) and Porubsky et al. (2019) are included in Supplemental Tables S3 and S5.
The criteria for considering a BAC “resolved” is described in the main text and Methods. The alignment identity of each resolved BAC was computed individually and the median of these values reported as a Phred-style quality value. No validation BACs were available for HG002. Superscripts mark previously published assemblies: *Miga et al. (2020); †Shafin et al. (2020). All ONT assemblies were generated by Canu, as it achieved the highest NG50 and BAC resolution. The best-performing result for each genome in each metric is highlighted in bold. Statistics for assemblies from Garg et al. (2019) and Porubsky et al. (2019) are included in Supplemental Table S5.

BACs. Note that HiCanu resolved the highest number of BACs across all considered assemblies and also achieved the highest BAC alignment quality (Table 3; Supplemental Tables S4, S5).

A deeper investigation of the unresolved CHM13 BAC sequences indicated that 11 BACs likely contain assembly errors or cloning artifacts themselves (Supplemental Note 3; Supplemental Figs. S2–S5). Manual inspection of HiFi read alignments did not reveal any standard misassembly signatures in the corresponding regions of the HiCanu assembly, providing evidence that HiCanu was correct in these cases and able to resolve 337 out of 341 (99%) of the CHM13 challenge BACs (Supplemental Table S6).

Although the challenge BACs are useful for validation, they do not represent the full landscape of human repeats. To further assess the ability of HiFi reads and different assemblers to resolve genome repeats, we used the method of Vollger et al. (2019) to identify collapsed repeat instances in the CHM13 assemblies. We identified ∼21.7 Mbp of collapsed repeats corresponding to at least 56 Mbp of unresolved repetitive sequence. The HiCanu assembly had the lowest number of bases in regions identified as collapsed repeats, as well as the smallest amount of repetitive sequence predicted to be missing from the assembly (Supplemental Table S7; Supplemental Fig. S6). A complementary mapping-based analysis confirmed the comparatively high completeness of the HiCanu assembly and classified the majority (80%) of missing sequence as satellite repeats, suggesting good recovery of all other human repeat classes (Supplemental Fig. S6).

Human haplotype phasing

When assembling a diploid genome, an assembler must choose to either collapse allelic sequences into a single sequence or preserve them as two separate sequences. Collapsing heterozygosity results in a mosaic consensus that may not faithfully represent any allele and can introduce frameshifting errors within coding sequence.

HiCanu assemblies of the diploid human genomes included >2 Gbp of alternate contigs, with high BUSCO completeness for both primary and alternate contig sets (>94% and >75%, respectively) (Supplemental Table S8). We again used Merqury (Rhie et al. 2020) to analyze the phase blocks using parental Illumina data (Supplemental Note 2). The phase block NG50s of HiCanu primary (0.6 Mbp) and alternate (0.1 Mbp) contig sets were the highest across all considered assemblies (2.5-fold higher than next best) (Supplemental Table S8). Note that the human phase block NG50s are significantly shorter than for the D. melanogaster F1 hybrid but are longer than a typical human gene. For comparison, Supplemental Table S8 also includes statistics for the recently obtained haplotype-resolved assemblies of HG002 (Garg et al. 2019) and HG00733 (Porubsky et al. 2019). These recent studies have shown that multimegabase NG50 phase blocks can be obtained by integrating HiFi data with long-range linking information derived from Hi-C (Garg et al. 2019) or Strand-seq data (Porubsky et al. 2019).

To assess the phasing accuracy, we used a gold-standard variant set from the Genome in a Bottle (GIAB) consortium (Supplemental Table S9; Zook et al. 2019). HiCanu has an F1 score of 95.37%, which is >50% higher than the next best HiFi assembly and similar to another assembly combining both HiFi and Hi-C data (97.7%) (Garg et al. 2019). Finally, we validated the recovery of complex, clinically relevant alleles; we compared assembly typing results for the six classical human leukocyte antigen (HLA) genes (Dilthey et al. 2019) to the known alleles for HG002 and HG00733, obtained by previous studies (Supplemental Table S10; Chin et al. 2019; Shafin et al. 2020). Only HiCanu and TrioCanu were able to recover all alleles with 100% sequence identity (Supplemental Tables S10, S11). The HiCanu contigs expectedly switch between the haplotypes, but there is only one switch in the MHC region. The Hi-C-phased HG002 assembly from Garg et al. (2019) is phased across the length of the MHC region but contains consensus errors (e.g., both HLA-DRB1 alleles). The Strand-seq–phased HG00733 assembly from Porubsky et al. (2019) is also phased across the length of the MHC region but incorrectly represents HLA-A and HLA-B as homozygous (with both alleles in the assembly matching one ground-truth allele). Both the Garg et al. (2019) and Porubsky et al. (2019) methods rely on initially collapsed assemblies that are then phased using the long-range data. These results suggest that separation of haplotypes early in the assembly process (rather than trying to recover them from collapsed assemblies) may improve the accurate recovery of heterozygous variation.

Complex regions of the CHM13 human genome

The CHM13 HiCanu assembly (Tables 2, 3; Fig. 2) exceeded the predictions of our prior model of human assembly continuity (Supplemental Note 5; Supplemental Fig. S7). To validate this result, we focused on the performance of HiCanu within some of the most difficult-to-assemble regions of the genome, namely, centromeres and SDs. Unlike past assemblies of the human genome, including clone-based assemblies, HiCanu generated several contigs spanning mega-bases of satellite DNA. The CHM13
HiCanu assembly contains nine of 23 (39%) expected centromere regions: Chromosomes 2, 3, 7, 8, 10, 12, 16, 19, and 20 (Supplemental Note 6; Supplemental Table S11). The structure of these regions was consistent with an expectation of one or more higher-order repeat (HOR) array(s) flanked by more divergent tracts of monomeric satellite DNA (Willard and Waye 1987; Schueler et al. 2001; She et al. 2004). Mapped read depth of both HiFi and ultralong Oxford Nanopore data (Miga et al. 2020) across these contigs shows relatively uniform sequence coverage that spans the α-satellite HOR array(s) into the unique sequences on the p- and q-arms (Fig. 3; Supplemental Note 6; Supplemental Fig. S8). The structure and length of the centromeric HOR array(s) in each contig is highly concordant with those reported in the literature (for review, see McNulty and Sullivan 2018).

It is noteworthy that HiCanu generated a draft assembly of the CHM13 Chromosome 19 centromere (Fig. 3). Historically, this region has been considered to be one of the more challenging centromeres to reconstruct because it carries multiple HOR tracts and shares α-satellite sequences with the centromere regions from Chromosomes 1 and 5 (Hulsebos et al. 1988; Baldini et al. 1989; Pironon et al. 2010; Sullivan et al. 2017; McNulty and Sullivan 2018). HiCanu was not only able to assemble a contiguous span of centromere but also accurately distinguished three distinct HOR tracts (D19Z1, D19Z2, and D19Z3) (Supplemental Note 6; Supplemental Fig. S9). This contig revealed a more complete representation of the HOR structure of the D19Z1 HOR unit (13-mer vs. 10-mer) (Supplemental Figs. S9A, S10; Hulsebos et al. 1988; Puechberty et al. 1999), a Chromosome 19-specific dimeric HOR (D19Z1) (Supplemental Figs. S9B, S10; Baldini et al. 1989; Finelli et al. 1996), and two complex HORs (expected to represent D19Z2) (Supplemental Note 6; Supplemental Fig. S10). Alignment of HiFi sequence data to the corresponding HiCanu contig did not reveal any coverage anomalies (e.g., large dips or spikes) that could indicate the presence of structural errors. However, marker-assisted alignment of ultralong Oxford Nanopore data (Miga et al. 2020), an orthogonal data set, showed a coverage drop coinciding with the D19Z1 array. This may indicate a misassembly, mismapping of the noisy sequencing data, or biases in sequencing coverage. Because of the lack of a validated truth set in such regions, this will require extensive wet-laboratory validation and is left for future work.

Beyond the obvious challenge of centromere assembly, SDs represent another significant impediment and have been estimated to account for 68% of misassemblies and contig breaks in recent long-read genome assemblies, irrespective of the platform or assembly algorithm (Porubsky et al. 2019). To estimate the effect of SDs on the continuity of HiCanu assemblies, we aligned contigs from the CHM13 genome assemblies to the human reference genome (GRCh38) and tested if the ends of contigs mapped within SDs. Compared with the Canu, Peregrine, or ONT assemblies, HiCanu had the fewest contig breaks within SDs (n = 95) and the smallest overall fraction of contig ends mapping to SDs (49%) (Table 4). Of these 95 regions, S9 (62%) correspond to the longest (>10 kbp) and most identical (>98%) copy-number polymorphic duplicated regions of the human genome (Supplemental Fig. S11). These results indicate that SDs are better resolved using HiCanu; however, SDs still contribute disproportionately to the overall number of assembly breaks.

The defensin beta gene family, a set of copy-number variable genes (e.g., DEFB136, DEFB135, etc.) mapping to two locations on Chromosome 8p23.1 (which we refer to as the defensin beta cluster), is a case in point. This ∼6-Mbp region plays an important role...
in immune function and disease (Weinberg et al. 2012; Mohajeri et al. 2016) and is known to be highly repetitive and difficult to assemble (Bakar et al. 2009). Previous reconstructions have relied on a BAC-by-BAC assembly approach (Mohajeri et al. 2016), and the first continuous assembly of this region in CHM13 was obtained via an independent approach using ultralong Nanopore data (GA Logsdon, MR Vollger, PH Hsieh, et al., in prep). Figure 4, A and B illustrates self-alignment dot plots at different stringencies of the defensin beta cluster from the T2T Chromosome 8 v3.0 assembly. Figure 4C shows the de novo assembled contig alignment and correctness against this draft. Both the Canu and HiCanu assemblies of the HiFi data consist of four contigs without structural errors. In contrast, the complex inverted repeat structures resulted in misassembled and fragmented contigs in all other evaluated assemblies.

The rightmost contig breaks in the HiFi assemblies are likely owing to the presence of long, nearly identical repeats, which would require either longer reads or a careful examination of repeat copy number to resolve. We also investigated the fragmentation of HiCanu and Canu contigs at position 10.4 Mbp, which is not part of any observed repeat structure. Alignment of the raw HiFi reads onto this region with minimap2 (Li 2018) revealed the presence of a 450-bp region covered by only two HiFi reads (Supplemental Fig. S12), with a coverage drop present in both the 10- and 20-kbp HiFi libraries. This coverage drop is flanked by a >250-bp simple-sequence repeat (AAAGG). Suspecting a possible bias in the HiFi datatype, we further examined Chromosome X, for which we have a complete CHM13 reference sequence available (Miga et al. 2020). HiFi assemblies of the CHM13 Chromosome 19 centromere (D19Z1, D19Z2, and D19Z3; marked with black bars). These HOR arrays are 606 kbp, 289 kbp, and 3.96 Mbp in length, respectively, and are composed of a 13-mer, a complex higher-order HOR, and a dimeric HOR unit, respectively. The HOR repeat underlying D19Z2 shares limited sequence identity with the pC-A16 repeat previously described (Hulsebos et al. 1988; Choo et al. 1991; Finelli et al. 1996) and, therefore, is designated with a question mark. The α-satellite HOR arrays have relatively uniform coverage of HiFi and ultralong Oxford Nanopore data, except for a drop in Oxford Nanopore sequencing coverage over the D19Z1 array, which may be owing to a misassembly, read mismapping, or biases in sequencing. The HiFi coverage plot shows fold coverage of the most common base (black) and the second most common base (red).

Figure 3. HiCanu assembly of the CHM13 Chromosome 19 centromere. RepeatMasker (Smit et al. 2013) of tig00006497 reveals three α-satellite HOR arrays that reside within the Chromosome 19 centromere (D19Z1, D19Z2, and D19Z3; marked with black bars). These HOR arrays are 606 kbp, 289 kbp, and 3.96 Mbp in length, respectively, and are composed of a 13-mer, a complex higher-order HOR, and a dimeric HOR unit, respectively. The HOR repeat underlying D19Z2 shares limited sequence identity with the pC-A16 repeat previously described (Hulsebos et al. 1988; Choo et al. 1991; Finelli et al. 1996) and, therefore, is designated with a question mark. The α-satellite HOR arrays have relatively uniform coverage of HiFi and ultralong Oxford Nanopore data, except for a drop in Oxford Nanopore sequencing coverage over the D19Z1 array, which may be owing to a misassembly, read mismapping, or biases in sequencing. The HiFi coverage plot shows fold coverage of the most common base (black) and the second most common base (red).

Table 4. CHM13 contig ends found within SDs

| Genome   | Assembly | Total no. of aligned contig ends | Total no. of contig ends within SDs | Percentage of contig ends within SDs |
|----------|----------|----------------------------------|------------------------------------|-------------------------------------|
| CHM13    | Canu ONT | 202                              | 137                                | 68%                                 |
|          | Canu     | 322                              | 170                                | 53%                                 |
|          | Peregrine| 468                              | 398                                | 85%                                 |
|          | HiCanu   | 192                              | 95                                 | 49%                                 |
| HG002    | Canu ONT | 608                              | 427                                | 70%                                 |
|          | Canu     | 1290                             | 862                                | 67%                                 |
|          | Peregrine| 862                              | 741                                | 86%                                 |
|          | HiCanu   | 418                              | 234                                | 56%                                 |
| HG00733  | Canu ONT | 606                              | 492                                | 81%                                 |
|          | Canu     | 842                              | 421                                | 50%                                 |
|          | Peregrine| 874                              | 716                                | 82%                                 |
|          | HiCanu   | 376                              | 181                                | 48%                                 |

Previously published assemblies are indicated by superscripts: *Miga et al. (2020), **Shafin et al. (2020). All ONT assemblies were generated by Canu, as it achieved the highest NG50 and BAC resolution. The best-performing result for each genome in each metric is highlighted in bold.
On this chromosome, we identified at least four additional cases of HiFi coverage dropout, with all four instances associated with long, low-complexity (A/G or T/C-rich) sequences. As our HiFi assembly of Chromosome X is split into just 13 large contigs (Supplemental Fig. S13), this coverage bias appears to be a current weakness of the HiFi chemistry.

Discussion

We have shown that HiCanu is capable of generating the most accurate and complete human genome assemblies to date and is able to achieve the resolution of repeats that are up to 99.99% identical. As a result, HiCanu surpasses prior HiFi and Nanopore ultra-long-read assemblies in terms of both repeat resolution and per-base consensus accuracy. HiFi data excels in resolving large highly-similar (but nonidentical) repeat instances. The remaining unresolved sequences seem to primarily represent satellite repeats (Supplemental Fig. S6). In particular, Figure 2 illustrates that HiCanu’s reconstruction of human Chromosomes 1, 7, 9, and 16 notably improves continuity over the previous assembly of ultra-long Nanopore reads (Koren et al. 2017; Kolmogorov et al. 2019; Shafin et al. 2020). These chromosomes contain several SDs exceeding 200 kbp in length, requiring high-fidelity reads to identify variants and separate the individual copies. HiFi data further enabled draft assemblies of nine centromeric regions, which are one of the final challenges of automated telomere-to-telomere assembly. Assembly of other centromeric regions is likely limited by a low frequency of unique markers compared with current HiFi read lengths. In contrast, Chromosome X has the highest count of large (>20 kbp) near-identical (>99.9%) repeats (Bailey et al. 2002) that were better resolved by long, spanning Nanopore reads. Thus, the two technologies are complementary at present, and the best technology depends on the specific characteristics of the repeats and haplotypes being assembled.

HiCanu’s approach to read correction permits considering only the highest identity overlaps during contig construction and is general enough to be applied to other applications such as metagenomic assembly. Although HiCanu is not as fast as some of the competing methods, we note that the number of CPU hours required for assembly of a human genome is under 4000, which could be completed on any modern cloud platform in less than a day for a few hundred dollars. This is 30-fold less than recent Oxford Nanopore assemblies that required more than 100,000 CPU hours (Jain et al. 2018b; Shafin et al. 2020). At the time of writing, the most computationally expensive step of HiFi analysis is generating the data itself, because each individual HiFi read represents a consensus of multiple, aligned sequences of the same DNA molecule. Coupled with the instrument runtime and sequencing cost, HiCanu is a small fraction of the total project cost and duration (Supplemental Note 7). Most prior long-read assemblers have also required a final “polishing” step to improve consensus accuracy, which requires additional computation but can also introduce errors in repeat instances owing to ambiguous read mappings (Miga et al. 2020). Because of the initial accuracy of HiFi reads and because of the precise resolution of allelic variants and repeats, HiCanu assemblies do not require polishing.

When choosing HiFi, the library size should also be considered when beginning a sequencing project. Because HiFi read accuracy...
depends on the size of the sequenced fragments (shorter equals more passes and higher accuracy), one should consider the relative importance of read length versus accuracy. A metagenomic project may aim for shorter, higher accuracy reads to confidently identify low-abundance strains, whereas a vertebrate genome project may benefit from longer reads to span midsized identical repeats. We also identified an apparent bias in the current HiFi chemistry at low-complexity A/G (T/C) repeats, leading to coverage drops and assembly fragmentation. This issue warrants further investigation and may limit the applicability of HiFi sequencing to genomes with large stretches of such repeats. Thus, identifying optimal sequencing strategies and developing methods that can combine multiple technologies remains an area for future research.

HiCanu’s diploid assemblies accurately capture both alleles in long haplotype blocks of very high quality (QV50). In particular, HiCanu consistently recovered both haplotypes for the six canonical MHC typing genes in the human genome, improving upon recently developed HiFi-based methods for haplotype-resolved assembly (Garg et al. 2019; Porubsky et al. 2019). However, because HiFi data alone do not provide long-range linking evidence, HiCanu’s contigs represent pseudohaplotypes that typically require additional information and processing to achieve chromosome-scale phasing. Canu also does not assign contigs to haplotypes and requires postprocessing with a tool such as Purge_dups (Guan et al. 2020) to split the diploid assembly into primary and alternate alleles. Although recent studies have successfully integrated HiFi data with additional long-range linkage information (Garg et al. 2019; Porubsky et al. 2019), we do not expect that significant improvements in phasing can be achieved by HiFi-only assemblies without an increase of HiFi read lengths. One option is postprocessing of HiCanu assemblies by a haplotype-aware scaffold, such as FALCON-Phase (Korenberg et al. 2019), which could potentially correct haplotype switch events and deliver further improvements to phasing accuracy and assembly continuity. In general, we feel that HiFi contigs combined with Hi-C phasing and scaffolding is a promising recipe for phased telomere-to-telomere vertebrate genome assembly, and we plan to integrate these data types in future versions of Canu.

**Methods**

**Mitigating errors in HiFi data**

Although HiFi reads are highly accurate compared with other long-read sequencing technologies, they are not error free, which complicates the identification of reads originating from the same genomic loci during assembly. To identify and remove false read overlaps, we sought to increase the accuracy of HiFi data via read correction.

Wenger et al. (2019) observed that the majority of HiFi errors are in homopolymers, where the number of individually repeating nucleotides is incorrectly counted. To lessen the impact of such errors, HiCanu modifies the input reads by compressing every homopolymer to a single nucleotide. Our approach is similar to run-length encoding (RLE), which has been previously applied to 454 (Miller et al. 2008), PacBio CLR (Li 2016, 2018; Ruan and Li 2020), and Oxford Nanopore (Shafin et al. 2020) reads. However, HiCanu does not explicitly store the lengths of the compressed homopolymer stretches and instead reverts back to the uncompressed reads when needed.

Although the transition to homopolymer-compressed sequence space can reduce the specificity of the read alignment search, the corresponding reduction in the number of observed errors in the reads allow for a more restrictive alignment identity threshold (based on empirical analyses, we require a minimum overlap identity of 99%). Subsequent steps are performed on the homopolymer-compressed sequences, whereas the detailed correspondence between positions of original and compressed versions is generated on the fly when necessary. Compressed reads are first subjected to overlap-based trimming (Koren et al. 2017). Although this step only affected one human assembly during development (HG002 sequel II system with pre-2.0 early access chemistry 15-kbp library available from https://github.com/human-panogenomics/HG002_Data_Freeze_v1.0) and does not appear to be necessary for newer HiFi data sets, we chose to enable it by default on all assemblies in this paper for consistency. This improvement suggests that a significant fraction of reads was structurally incorrect in the poorly performing library owing to a low-quality sequencing library. Because other libraries did not show this problem, it is likely future versions of the HiCanu pipeline can skip this step and reduce runtime by >60%.

To further reduce the influence of the errors in compressed HiFi reads, we have updated the OEA module of Canu (Holt et al. 2002; Koren et al. 2017). This module identifies errors in individual reads by jointly considering all of their overlapping reads. Every such read votes for the nucleotides at the positions that it covers based on the pairwise alignment of the overlapping regions. A read’s position is considered erroneous if no other reads support the original sequence and the majority of votes agree on a particular change (by default >50%) and at least seven if there is a read supporting the original sequence. After the corrections are introduced, the alignment scores of the overlaps are recomputed, but the actual read sequences stored within the assembler are not modified as doing so would invalidate the previously computed overlap coordinates. Although our naive approach may not always be able to correct errors within highly-similar genomic repeats, such events are rare owing to the low number of errors in compressed HiFi reads and the high identity threshold used for gathering candidate overlaps.

Manual investigation of read alignments during HiCanu development revealed a previously unreported error mode in HiFi reads: incorrect repeat unit counts within microsatellite repeat arrays. Because the incorrect repeat counts are systematic and often supported by multiple reads, the conservative strategy described above is not able to correct them. Recognizing this, we modified the OEA procedure for recomputing overlap alignment scores to ignore sequence differences flanked by a microsatellite repeat in either read. Namely, the difference is ignored if five out of six nonoverlapping flanking k-mers are the same for any k ranging between two and six on either side (starting at zero to k − 1 bp from the difference). We note that this phenomenon deserves a deeper investigation, and our strategy can be improved to capture additional genomic differences, which are ignored by the current approach.

We evaluated the contribution of each of the above corrections using the recently completed CHM13 Chromosome X (Miga et al. 2020) as a reference. Raw, compressed, corrected, and masked 20-kbp HiFi reads were mapped and the mappings filtered to retain high-confidence alignments (Supplemental Note 1). Figure 1 shows the resulting alignment identity values, with each correction step boosting the identity of the aligned sequence. Each step (compression, correction, masking) contributes to this improvement (Supplemental Table S1; Supplemental Fig. S14). Although almost no (<1%) raw HiFi reads map error free, 97.23% of the compressed, corrected, and masked reads map without a single difference. Without correction (compression + masking only), reads have similar median error to just compression, and less than half have perfect alignment. As we did not control for reads
mapping from other chromosomes and as the Chromosome X sequence itself is not error free, this likely represents a lower bound on the percentage of error-free reads. To extend beyond Chr X, we also estimated read accuracy using k-mers from short-read data for all human genomes and found correction improved read accuracy across all data sets (Supplemental Note 1).

**Bogart modifications**

The Bogart module constructs a set of draft contigs from read overlap information. A detailed description is given by Koren et al. (2017). We describe here the modifications made for HiFi data.

**Overlap identity threshold**

Canu’s initial overlap search uses a relaxed identity threshold to account for varying error rates between samples. Because overlap identities are changed by OEA and because we wished to avoid considering false-positive overlaps, Bogart first attempts to select a higher overlap identity threshold. Previously, Canu computed the identity of the best-scoring overlap on each side of every read (where score is defined as the number of matching bases in the overlap alignment) and set a threshold based on the median and MAD of the computed values (Koren et al. 2017). However, during the development, we realized that this way of computing the threshold was not informative for highly accurate reads because both the median and MAD were 100% across all tested data sets. Additionally, with the number of matching bases as a score, the read delivering the highest scoring overlap could come from a different haplotype in genomes with low heterozygosity. As a result, the selected threshold could inadvertently reflect the heterozygosity level of the organism rather than the accuracy of the reads. Based on empirical testing, we opted for an alternative two-step procedure. First, all overlaps with identity below the fixed value T (default 99.97% or three differences in 10,000 bp) are dropped. This step is aimed at removing from consideration the majority of the cross-haplotype overlaps even for low-heterozygosity organisms, for example, human heterozygosity rate of 0.1% (The 1000 Genomes Project Consortium 2012). Next, the identities of the highest scoring overlaps are collected as before, and the final threshold is set as the 90th percentile of this sample. It is possible that 99.97% is too stringent given higher error reads. We could detect this condition when the 90th percentile is too close to 99.97% and rerun the overlap filtering. However, on all data sets evaluated to date, the chosen identity threshold was 100%. To support the desired overlap filtering stringency, the Canu codebase had to be modified to increase the precision with which the overlap identity values are stored.

**Handling heterozygous differences**

Bogart uses the filtered overlaps to identify and eliminate the reads likely representing sequencing artifacts and then constructs the best overlap graph (Miller et al. 2008), using the same overlap scoring function as before. This graph consists of the best-scoring overlap off both the 5’ and 3’ ends of each read, and the nonbranching paths within this graph form the preliminary layouts (arrangements of reads) that we refer to as greedy contigs. Bogart then inspects each greedy contig for long repeat instances that could have been incorrectly traversed. Repeats are detected by considering overlaps between the reads within and outside of the contig. If a suspected repeat has no reads spanning it or if there is a similar-length alternate read overlap, it is broken at the repeat boundary to avoid potential assembly errors as the method of Koren et al. (2017) to form the final draft contigs.

HiCanu aims to reconstruct long pseudohaplotype contigs (Vinson et al. 2005; Chin et al. 2016)—potentially switching between paternal and maternal alleles—and capture the alternative regions as shorter contigs. Unfortunately, the original Bogart approach described above led to the classification of extended homozygous regions within greedy contigs as unscaled repeats, leading to fragmentation of the pseudohaplotype (Supplemental Fig. S15). In Canu, this behavior had been affecting only genomes with >1% heterozygosity, because below this threshold most heterozygous differences were implicitly hidden by the relatively permissive threshold on overlap identity. With the high-accuracy HiFi data, and a correspondingly high overlap identity threshold, this overfragmentation became an issue even for human levels of heterozygosity.

In HiCanu, Bogart has an additional step to identify contigs representing alternative alleles within the set of greedy contigs, which we refer to as bubble contigs. As suggested by the name, the bubble contigs are related to the bubble subgraphs, typically considered by most assemblers. Candidate bubbles are found by identifying reads in each contig that have overlaps to some other, larger, contig. A read within a smaller contig can be placed in the larger contig if the overlaps between it and the reads in the larger contig are below a specified threshold of similar quality to the previously incorporated overlaps (0.1% by default). If the placements for both the first and last reads of a candidate contig are correctly oriented and placed at approximately the correct distance in the larger contig (75%–125% of the candidate contig size), the candidate contig is flagged as a bubble and its reads are excluded from later repeat detection. This avoids fragmentation of otherwise structurally correct pseudohaplotype contigs. Similar strategies have previously been used in short-read assembly (Pevzner et al. 2001; Zerbino and Birney 2008; Li et al. 2010; Gnerre et al. 2011), scaffolding metagenomes (Koren et al. 2011; Ghurye et al. 2019), and long-read assembly (Chin et al. 2016). Bubble contigs are also explicitly marked in the final output; however, because placements are not always found, especially for longer, more heterozygous alleles, we recommend using a postprocessing tool such as Purge_dups (Guan et al. 2020) to classify alternate alleles and remove any false duplications.

**Consensus calculation**

A consensus sequence is computed for all contigs using the uncompressed reads (trimmed to their good regions identified in compressed space). Canu originally used the layout produced by Bogart to estimate the position of each read within the contig and align it only to that location. Because the read layouts are now in homopolymer-compressed space, this strategy is unable to locate the read in uncompressed space. Instead, we compute the correspondence of each position in the compressed read to the original. This is used to update the read positions within the contig and expand the layout to uncompressed space. A modified version of the PBdagon algorithm (Chin et al. 2013), with improved support for long-contig sequences, is used to compute the final consensus sequence.

Currently, HiCanu will exclude erroneous reads from large contigs, but these erroneous reads may form their own short, low-coverage contigs. This can slightly reduce average assembly accuracy for homozygous genomes versus a more permissive strategy like that in Canu. However, Canu’s more permissive approach will incorrectly mix haplotypes and similar repeat copies. Further HiCanu consensus gains are possible with better handling of erroneous reads and a more sophisticated approach for predicting homopolymer run length, similar to MarginPolish (Shafin et al. 2020).
Previously generated data and assemblies

When available, previously published assemblies were downloaded and used. This included Oxford Nanopore UL Canu assemblies presented by Shaﬁn et al. (2020) for HG0002 (80× Guppy HAC 2.3.5) and HG00733 (50× Guppy HAC 2.3.5), Canu + Racon assembly presented by Vollger et al. (2020), HG0002 assembly of HiFi reads presented by Wenger et al. (2019), Oxford Nanopore Canu assembly for CHM13 (40× +80× UL Guppy HAC 3.1.5) presented by Miga et al. (2020), HiFi + Hi-C assemblies for HG002 presented by Garg et al. (2019), and HiFi + Strand-seq assemblies for HG0733 presented by Porubsky et al. (2019). In the remaining cases, assemblies were run locally on the NIH Biowulf cluster.

The D. melanogaster HiFi data are available from NCBI Bioproject database (https://www.ncbi.nlm.nih.gov/bioproject/) at PRJNA573706 (SRR10238667; median: 24.4 kbp; mean: 24.4 kbp) and CLR (SRR9969843; median 13.3 kbp; mean: 17.2 kbp). Because of the high coverage, this data set was down-sampled to 40× HiFi data and 200× CLR data. These coverages represent ~25% of the full run output. Because the exact parents of the F1 were not available, we used the previously generated short-read sequencing for binning and analysis (A4: SAMN00849823; ISO1: SRR8670564). The CHM13 Nanopore data are available at https://s3.amazonaws.com/nanopore-human-wgs/chm13/nanopore/release/13.fastq.gz and Illumina at GitHub (https://github.com/nanopore-wgs-consortium/CHM13#genomics-data). The HG002 Nanopore data are available at https://s3-us-west-2.amazonaws.com/human-pangenomics/index.html, HiFi at SRX5327410. HG002 and parent Illumina data are available from GIAB (Zook et al. 2016) at GitHub (https://github.com/genome-in-a-bottle/giab_data_indexes), we only used the 2×250 data sets. The HG00733 Nanopore data are available at https://s3.amazonaws.com/nanopore-human-wgs/chm13/nanopore/release/13.fastq.gz and Illumina at GitHub (https://github.com/nanopore-wgs-consortium/CHM13#downloads).

Software commands

HiCanu was run using Canu branch hicanus_rc with the following command:

```bash
canu -p asm -d asm genomeSize=150m corOut Coverage=100 batOptions=-dg 6 -db 6 -dr 1 -ca 500 -cp 50 -pacbio-raw-reads.fastq.gz
```

For CLR data Canu branch hicanu_rc was run with the following command:

```bash
canu -p asm -d asm genomeSize=G corOut Coverage=100 batOptions=-dg 6 -db 6 -dr 1 -ca 500 -cp 5 -pacbio-raw-reads.fastq.gz
```

All HiFi assemblies required less than 12 wall-clock hours on the NIH Biowulf cluster quick partition with all jobs using <120 GB RAM. We reran HG002 on our cluster limiting the maximum concurrent CPUs to 288, which required 30 h. We estimated the cost of an AWS run using the c5d.18xlarge instance, which costs $3.456/h. Assuming four reserved nodes (for a total of 72 × 4 = 288 CPUs) and an average runtime of 4200 CPU hours with perfect parallelization, the run would complete in 14.5 h. We increase this by a factor of 2.0 to account for any nonparallelized steps based on the experiments above for a cost of $3.456×4×29=$401. We note these estimates limited by differences in CPU and I/O between our cluster and AWS, as well as the overhead of waiting for a job to be scheduled on our cluster. The cost could also be reduced if additional nodes were spun up on-demand for the parallel portions of compute and spin down when not needed (as performed in Canu’s DNAExeXtended implementation). We omit this from the estimate for simplicity. We also note that the assemblies could be completed faster if more nodes were allocated in parallel.

Peregrine assembler and SHIMMER ASMKit (0.1.5.3) was run with the command:

```bash
yes yes | python3 /data/korens/devel/Peregrine/bin/pg_run.py asm
  chm13.1ist 24 24 24 24 24
  -with-consensus -shimmer-r 3 -best_n_ova 8
  --output /
```

This required 7 CPU hours and 29 GB of memory for D. melanogaster, 32 CPU hours and 347 GB of memory for the CHM13 10-kbp library, 58 CPU hours and 449 GB of memory for the CHM13 20-kbp library, 55 CPU hours and 407 GB for HG0002, and 32 CPU hours and 347 GB for HG00733.

Commands for defensin beta cluster and Chr X validation

HiCanu contigs flagged as bubbles were excluded from the analysis. MUMmer (Kurtz et al. 2004) 3.23 was used to identify repeats with the command:

```bash
nucmer -maxmatch -nosimplify delta-filter -i 98 -l 10000
```

and high-stringency repeats

```bash
nucmer -maxmatch -noextend -nosimplify -l 500 -c 1000
delta-filter -i 99.9 -l 10000
```

QUAST alignments were generated as

```bash
quast.py -t 20 --large-skpunaligned-mis-contigs --min-alignment 10000 --min-identity 98.0 --extensive-mis-size 5000 --min-contig 50000
```

Icarus was patched not to show breaks at “small indels” and “stretches of mismatches,” and used to visualize the resulting alignments.

Commands for RepeatMasker

RepeatMasker version 4.1.0 was run with the commands
on each contig ≥50 kbp in the assembly. Centromeric arrays were identified by taking all hits marked as Satellite/cent and merging any hits within 100 bp of each other using BEDTools (Quinlan and Hall 2010). Resulting arrays >800 kbp were reported. There were nine internal arrays whose start and end coordinates were at least 500 kbp away from a contig end. These initial coordinates were manually curated based on reference alignments and are reported in Supplemental Table S12.

**Commands for MHC typing**

HLA’LA version commit 24930dadbd0d2b6bcd69a271401df-c88a5d09f4d was run with the commands

```
HLA-ASM.pl –use_minimap2 1 –assembly fasta $asm –sampleID $prefix –workingDir $pwd/$prefix –truth reference_HLA_ASM/$truth
```

where $asm was the assembly, $prefix was a unique identifier, and $truth was either truth_HG002.txt or truth_HG00733.txt.

**Commands for Purge_dups**

Purge_dups version commit 8f580b41e6a20c99383d66ff19-b8689e93d7490e was run with the commands

```
python pd_config.py asm.fasta ‘pwd’ <pb folder> <10x folder left blank> asm minimap2 -16g -xasm5 -DP asm.split.asm.split > asm.split.self.paf minimap2 -16g -xmap-pb asm.fasta $line > pb.$jobid.paf (for each HiFi cell) pbcstat pb.*.paf calculcts PB.stat > cutoffs 2>calculcts.log purge_dups -2 -T cutoffs -c PB.base.cov asm.split.-self.paf > dups.bed 2> purge_dups.log get_seqs dups.bed asm.fasta > purged.fa 2> hap.fa
```

For *D. melanogaster*, an incorrect threshold was computed for the cutoffs owing to the entire genome being separated and so the cutoffs were manually adjusted to be

50 1 115 2 200.

The purged.fa output was then used as the primary set reported in the tables. To obtain the alternate set, we ran a second round of Purge_dups using hap.fa as the input assembly instead. This required an average of 20 CPU hours and 7 MB of memory for *D. melanogaster*, 59 CPU hours and 2 MB of memory for HG0002, and 74 CPU hours and 24 MB of memory for HG00733.

**Commands for Mermqury**

Mermqury version commit 154610d19ee64fexd77d077af1e-d7abdbe8866 was used. For each assembly and read set, canonical k-mers were built using meryl available as a binary within Canu:

```
meryl count k=<k-size> <reads.fasq.gz> output <genome>.<k-size>.meryl
meryl count k=<k-size> <asm.fasta> output <asm>.<k-size>.meryl
```

using k=21 for humans and k=18 for *D. melanogaster* based on (Fofanov et al. 2004). QV and k-mer completeness were obtained with

```
eval/spectra_cn.sh
eval/spectra-cn.sh
```

which converts k-mer Jaccard to distance as previously described (Ondov et al. 2019) and to a Phred score (Ewing and Green 1998). Haplotype blocks were estimated by first building parent-specific k-mer databases. K-mers in each parental data set were counted as above, then subtracted to obtain parent-specific k-mers, and finally intersected with the child (in the case of human data sets in which child illumina data was available) with

```
trio/hapmers.sh
trio/phased_block.sh
```

For further information see Supplemental Note 2 and https://github.com/marbl/merqury/wiki.

**Commands used for QUAST**

QUAST 5.0.2 ran with the command

```
quast.py <asm> -o quast_results/<asm> -r <reference> -t 16 -s -large
```

Variants were filtered using the pipeline from (Shafin et al. 2020) to filter errors in varying sites, including known SVs (HG002 only available from GIAB) (Zook et al. 2020) at ftp://ftp-trace.ncbi.nlm.nih.gov/assembly/NIST_SVs_Integration_v0.6/HG002_SVs_Tier1plusTier2_v0.6.1.bed):

```
python3 reference/quast_sv_extractor.py –q quast_results/<asm> -o <asm>/contigs_reports/all_alignments.tsv -c reference/centromere.bed -d reference/GRC38_marked_regions.bed -s reference/empty
```

and

```
python3 reference/quast_sv_extractor.py –q quast_results/<asm> -o <asm>/contigs_reports/all_alignments.tsv -c reference/centromere.bed -d reference/GRC38_marked_regions.bed -s reference/HG002_SVs_Tier1plusTier2_v0.6.1.bed
```

for HG002. We used https://www.ncbi.nlm.nih.gov/assembly/GCF_000001215.4 filtered to remove any unassigned sequences for *D. melanogaster* (Chr 2L, Chr 2R, Chr 3L, Chr 3R, Chr 4, Chr M, Chr X, Chr Y only) and https://hgdownload.soe.ucsc.edu/goldenPath/hg38/bigZips/hg38.fa.gz filtered to exclude alts and unaligned sequences (Chromosomes 1–22, X, Y, and MT only).

Because no filtering file was available for *D. melanogaster*, we modified QUAST parameters to try to avoid false-positive misassembly counts with the command

```
quast.py <asm> -o quast_results/<asm> -r <reference> -large -min-alignment 20000 -extensive-mis-size 50000 -min-identity 90
```

**Commands for BAC validation**

We used the BAC validation pipeline available at GitHub (https://github.com/skoren/bacValidation) run with default parameters. This pipeline aligns reads using minimap2 (Li 2018) and parses the SAM (Li et al. 2009) format to generate summary statistics. Output BAC identity was computed as the median across all BACs marked as correctly resolved. BAC libraries were downloaded from NCBI (CHM13: https://www.ncbi.nlm.nih.gov/nuccore/?term=VMRC59+and+complete, HG0002: https://www.ncbi.nlm.nih.gov/nuccore/?term=VMRC62+and+complete). HiFi read alignments to the assembly and BAC sequences were visualized with the Integrative Genomics Viewer (IGV) (Robinson et al. 2011).
**Commands for variant analysis**

We downloaded trio-phased GIAB (Zook et al. 2019) variant calls for HG002 from ftp://ftp-trace.ncbi.nlm.nih.gov/Reference/Sample/gh/release/AshkenazimTrihHG002_NA24385_sn_NI_STv3.2/GRCh38/HG002_GRCh38_GIAB_highconf_CG-illlb-llse
nietionHC-Ion-10XsentieonHC-SOLIDgatkHC_CHROM1-22_v.3.3.2_highconf_triophased.vcf.gz. We ran dipcall followed by vcfalign to estimate SNP sensitivity and precision (Supplemental Table S8) with the commands:

```
run-dipcall hg002_purge GRCh38_full_analysis_set plus decoy_hla.fa primary.fasta alts.fasta > hg002.mak
make -j1 -f hg002.mak
# exclude chrX/Y since there are no GIAB variants on them
gunzip -c hg002_purge.dip.vcf.gz | grep -v chrX | grep -v chrY |
bgzip -c > hg002_purge.dip.nohom.vcf.gz
# measure statistics
rtg vceval -b HG002_GRCh38_GIAB_highconf_CG-Illlb-Illse
nietionHC-Ion-10XsentieonHC-SOLIDgatkHC_CHROM1-22_v.3.3.2_highconf_triophased.vcf.gz -o hg002_purge.dip.vcf.gz -e HG002_GRCh38_highconf_CG-Illlb-Illse
nietionHC-Ion-10XsentieonHC-SOLIDgatkHC_CHROM1-22_v.3.3.2_highconf_noinconsistent.bed -t GRCh38_h381.sdf -m annotate -o h381
```

To evaluate phasing, we evaluated the number of maternal and paternal variant calls out of the true positive calls in each contig and reported the total fraction of misphased variants (Supplemental Note 8).

**Commands for identifying contig ends**

Alignments were made between assemblies and GRCh38 using the following minimap2 command:

```
minimap2 --secondary=no -a -t eqx -Y -r asm20 -s 200000 -z 10000,50 -r 50000 -end-bonus=100 -O 5,56 -E 4,1 -B 5
```

Contig ends that intersected SDs were identified by parsing the CIGAR string to find the location of contig ends and then intersecting these locations with annotated SDs plus 10 kbp on either side from the UCSC Genome Browser using the following commands:

```
bedtools slop -i {segdups.bed} -b 10000 | bedtools merge -i - > {expanded.segdups.bed} & bedtools intersect -a {contig.ends.bed} -b {expanded.segdups.bed}
```

**Data access**

All raw and processed sequencing data generated in this study have been submitted to the NCBI BioProject database (https://www.ncbi.nlm.nih.gov/bioproject/) under accession number PRJNA530776 (10 kbp: SRR9087597–SRR9087600; 20 kbp: SRR11292120–SRR11292123).

We have posted the down-sampled data sets, generated assemblies, and corrected CHM13 BAC sequences at https://obj.
umiacs.umd.edu/marbl_publications/hicanu/index.html. HiCanu is implemented within the Canu assembly framework and is available as Supplementary Code and from GitHub (https://github.com/marbl/canu).
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