The haplotype-resolved genome and epigenome of the aneuploid HeLa cancer cell line

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The HeLa cell line was established in 1951 from cervical cancer cells taken from a patient, Henrietta Lacks. This was the first successful attempt to immortalize human-derived cells in vitro. The robust growth and unrestricted distribution of HeLa cells resulted in its broad adoption—both intentionally and through widespread cross-contamination—and for the past 60 years it has served a role analogous to that of a model organism. The cumulative impact of the HeLa cell line on research is demonstrated by its occurrence in more than 74,000 PubMed abstracts (approximately 0.3%). The genomic architecture of HeLa remains largely unexplored beyond its karyotype, partly because like many cancers, its extensive aneuploidy renders such analyses challenging. We carried out haplotype-resolved whole-genome sequencing of the HeLa CCL-2 strain, examined point- and indel-mutation variations, mapped copy-number variations and loss of heterozygosity regions, and phased variants across full chromosome arms. We also investigated variation and copy-number profiles for HeLa S3 and eight additional strains. We find that HeLa is relatively stable in terms of point variation, with few new mutations accumulating after early passaging. Haplotype resolution facilitated reconstruction of an amplified, highly rearranged region of chromosome 8q24.21 at which integration of the human papilloma virus type 18 (HPV-18) genome occurred and that is likely to be the event that initiated tumorigenesis. We combined these maps with RNA-seq and ENCODE Project data sets to phase the HeLa epigenome. This revealed strong, haplotype-specific activation of the proto-oncogene MYC by the integrated HPV-18 genome approximately 500 kilobases downstream, and enabled global analyses of the relationship between gene dosage and expression. These data provide an extensively phased, high-quality reference genome for past and future experiments relying on HeLa, and demonstrate the value of haplotype resolution for characterizing cancer genomes and epigenomes.

We generated a haplotype-resolved genome sequence of HeLa CCL-2 using a multifaceted approach that included shotgun, mate-pair and long-read sequencing, as well as sequencing of pools of fosmid clones using a multifaceted approach that included shotgun, mate-pair and long-read sequencing, as well as sequencing of pools of fosmid clones (Supplementary Table 1). To catalogue variants, we carried out conventional shotgun sequencing to 88 x non-duplicate coverage and reanalysed 11 control germline genomes in parallel (Supplementary Tables 2 and 3). Although normal tissue corresponding to HeLa is unavailable, the total number of single-nucleotide variants (SNVs) identified in HeLa CCL-2 (n = 4.1 x 10⁶) and the proportion overlapping with the 1000 Genomes Project (90.2%) were similar to controls (mean n = 4.2 x 10⁶ and 87.7%, respectively), suggesting that HeLa has not accumulated appreciably large numbers of somatic SNVs relative to inherited variants. Indel variation was unremarkable after accounting for differences in coverage (Supplementary Fig. 1). Short tandem repeat profiles of HeLa also resembled controls, consistent with mismatch repair proficiency (Supplementary Fig. 2).

After removing protein-altering variants that overlapped with the 1000 Genomes Project or the Exome Sequencing Project, similar numbers of private protein-altering (PPA) SNVs were found in HeLa (n = 269) and controls (mean n = 391). Gene ontology analysis found that all terms enriched for PPA variants in HeLa (P ≤ 0.01) were also enriched in at least one control (except for ‘starlike response’ in HeLa), suggesting that known cancer-related pathways are not perturbed extensively by point or indel mutations (Supplementary Fig. 3). Although a previous study of the HeLa transcriptome reported an enrichment of putative mutations in cell-cycle- and E2F-related genes, subsequently generated population-scale data sets contain all variants that we observed in these genes, suggesting that they are inherited and benign rather than somatic and pathogenic.

The overlap between PPA variants and the Catalogue of Somatic Mutations in Cancer (COSMIC) was similar for HeLa (n = 1) and control genomes (mean n = 2.6). The gene-level overlap with the Sanger Cancer Gene Census (SCGC) was also similar for HeLa (n = 4) and control genomes (mean n = 8.7). Canonical tumour suppressors and oncogenes were notably absent among the five SCGC genes with PPA variants in HeLa (BCL11B (B-cell CLL/lymphoma 11B (zinc finger protein)), EP300 (E1A binding protein p300), FGFR3 (fibroblast growth factor receptor 3), NOTCH1 and PRDM16 (PR domain containing 16), Supplementary Tables 3–6). However, three are associated with HPV-mediated oncogenesis (FGFR3, EP300, NOTCH1) and may be ancillary to the dominant role of HPV oncoproteins in HeLa and other HPV cervical carcinomas. Mutations in FGFR3 have been noted previously in cervical carcinomas, although infrequently and at different residues than observed here. Both EP300 and NOTCH1 are recurrently mutated in diverse cancers and are involved in Notch signalling, a pathway that is dysregulated in HeLa. EP300, which encodes the transcriptional co-activator p300, interacts directly with viral oncoproteins such as HPV-16 E6 and HPV-18 E7 (ref. 16). Although the in-frame deletion of a highly conserved amino acid in EP300 seems to be somatic (heterozygous within a loss-of-heterozygosity (LOH) region), it is still possible that the others are rare, inherited variants or passenger mutations. Further studies are required to resolve their functional relevance and to assess whether these genes are recurrently altered in HPV cervical carcinomas.

Aneuploidy and LOH, which are hallmarks of cancer genomes, were mapped in HeLa by constructing a digital copy-number profile at kilobase resolution (Fig. 1, Supplementary Fig. 4 and Supplementary Table 7). Read coverage profiles were segmented by a Hidden Markov Model (HMM) and recalibrated to account for widespread aneuploidy (Supplementary Figs 5 and 6). Sixty-one per cent of the genome has a baseline copy number of three, and only a small minority (3%) has a copy number of greater than four or less than two (Supplementary Table 8). LOH encompassed 15.7% of the genome, including several entire chromosome arms (5p, 6q, Xp, Xq) or large distal portions (2q, 3q, 6p, 11q, 13q, 19p, 22q) (Supplementary Fig. 7 and Supplementary Table 9), consistent with previous descriptions of LOH in cervical carcinomas. The overall profile is consistent with published karyotypes of various HeLa strains, suggesting that the hypertriploid state arose either during tumorigenesis or early in the establishment of the HeLa cell line.
Structural variants were identified by clustering discordantly mapped reads from 40-kb and 3-kb mate-pair libraries (Supplementary Fig. 8). Twenty interchromosomal links were identified, including links for marker chromosomes M11 (9q33–11p14) and M14 (13q21–19p13). In addition, 209 HeLa-specific deletions and 8 inversions were found (Supplementary Figs 9 and 11, and Supplementary Table 10). Only two genes that are impacted by HeLa-specific structural rearrangements (Supplementary Table 11) intersected with SCGC (STK11 (ref. 18), FHIT), both of which are recurrently deleted in cervical carcinomas18,19.

Conventional whole-genome sequencing fails to resolve haplotype phase, an essential aspect of the description and interpretation of non-LOH HeLa genome (scaffold N50 of 44.8 megabases (Mb); 550 kb containing 90.6% of heterozygous variants that were probably inherited. Most of the HeLa genome is present at an uneven haplotype ratio (for example, 2:1 in regions in which copy number = 3). We sought to exploit the resulting allelic imbalance to phase consecutive haplotype blocks (Supplementary Fig. 13). We first calculated the cumulative allelic ratio among shotgun reads for the SNVs residing in each haplotype block, which clustered closely with the underlying haplotype. When overlapping clones from distinct pools were merged, this resulted in haplotype blocks with an N50 (the contig size above which 50% of the total length of the haplotype assembly is included) of 550 kb containing 90.6% of heterozygous variants that were probably inherited.

First, 99.7% of informative read pairs from 3-kb mate-pair sequencing pool. Alleles that were present at distinct heterozygous sites within a given clone were assigned, or 'phased', to the same inherited haplotype, and the unobserved alleles were implicitly phased to the opposite haplotype. When overlapping clones from distinct pools were merged, haplotype blocks (Supplementary Fig. 13). We first calculated the cumulative allelic ratio among shotgun reads for the SNVs residing in each haplotype block, which clustered closely with the underlying haplotype ratio. For example, in non-LOH regions with a copy number of 3 that have ratios of 2:1 or 1:2, allelic ratios calculated for each block had ratios of 2:1 or 1:2, allelic ratios calculated for each block had
gested by high overall breakpoint concordance between strains (81% of
tions were more common. These may correspond to variability in copy
(Supplementary Figs 28 and 29)), but smaller amplifications and dele-
CCL-2 (61%). Gains or losses of entire chromosome arms were
across the eight strains (range 38–60%), similar to its prevalence in
the dominant state consistently, with a median of 52% of the genome
some strain-specific differences (Supplementary Figs 25–27), consis-
HeLa strains that we sequenced lightly (3.5 to 4.3
were also shared, though to a lesser degree: 72.4% of clone-confirmed
Supplementary Fig. 23 and Supplementary Table 15). Somatic mutations
segmental duplications, and requiring
an estimated sensitivity of 61.1%, placing an upper bound on the point-
tary Fig. 22). In summary, 8,165 somatic mutations were validated with
haplotype, with little regional variation in the abundance (Supplemen-
type between the two or more amplified copies of the same germline
haplotype is amplified but both remain present, the majority of observed
were present at copy number four, at a haplotype ratio of 3:1. To char-
acterize the structure and copy number of the insertion, we included
the HPV-18 genome alongside the human reference during align-
ment of clone-pool reads. By analysing patterns of coverage from
breakpoint-spanning fosmid clones, read-depth data and breakpoint
sequencing, we generated a structural model for the viral integration
(Fig. 2a, b, and Supplementary Figs 30 and 31). Two repeat structures
which we designate R1 and R2) consisting of the partial viral genome
are interspersed with regions of human chromosome 8q24.21 genomic
DNA. The viral genome is present with identical breakpoints on each
copy of the amplified haplotype, to the exclusion of the other haplo-
type, which remains at single copy and lacks integration-associated
rearrangements, confirming that integration and rearrangement pre-
ceded aneuploidy. The integrated structure contains only two-thirds of
the complete HPV-18 genome, including full-length copies of the E6
and E7 oncogenes necessary for telomerase activity (amplified to a
copy number of approximately 12), but lacking a functional copy of
E2, an inhibitor of E6 and E7 (ref. 13) (Fig. 2c). In addition, a distinct
portion of the HPV-18 genome, amplified to a copy number of approximately 30 in HeLa, includes an epithelium-specific enhancer

| Figure 2 | HeLa HPV integration locus. a. Chromosome 8 read depth
flanking the HPV integration site (top, blue line), windowed copy-number ratios (purple points, shaded by segment) and integer copy states (red bars, middle), and corresponding segments and breakpoints (circled numbers with genomic coordinates, bottom). b. Proposed HPV integration structure: per-
segment copy number (top left), non-rearranged haplotype B (copy number = 1, top right), rearranged haplotype A with HPV insertion (copy number = 3, bottom) carrying approximately 3 and 6 tandem copies of repeats R1 and R2, respectively. Hap, haplotype. c. The partial HPV-18 genome and corresponding genes (grey and blue, top) with breakpoints highlighted by numbered circles. For reference, the entire HPV-18 genome is shown (bottom).
Figure 3 | Gene expression by copy number and haplotype in HeLa S3.
a. Transcript abundance (reads per kilobase per million (RPKM), for genes with an RPKM ≥ 1) is positively correlated with gene copy. b. Expression per copy (RPKM per gene copy number) does not correlate with copy number. c. Fractional contribution of haplotype A to overall expression (Hap A/total) (RPKM averaged across megabase windows at phased sites) split by haplotype-resolved copy number. Open circles indicate expected fractions. d. Haplotype-A-specific expression in HeLa S3 but not CCL-2 across S3-specific LOH on chr18q. e. Haplotype A fractional contribution to expression across the genome, colour-coded by underlying haplotype-resolved copy number as in c (point size represents the log2 total RPKM, grey boxes indicate HeLa S3 LOH).

that controls E6 and E7 transcription27, possibly contributing to their high expression (Supplementary Fig. 32).

Extensive sequencing-based functional genomic data have been generated on HeLa and other cancer cell lines by the ENCODE Project7, but these have the potential to be misinterpreted if their analysis does not account for aneuploidy and phase. As HeLa CCL-2 and S3 are nearly identical in genotype, we used haplotype and copy-number maps of CCL-2 to assign phase to publicly available functional data generated on S3 (ref. 7), including transcription-factor binding, chromatin modification and chromatin-accessibility data sets. We also calculated haplotype-specific gene-expression scores using RNA sequencing (RNA-seq) data generated in this study and by others6,7 (Supplementary Figs 33–35). For each data set, aligned reads were phased by comparison to HeLa CCL-2 haplotype blocks. Corresponding peak scores (chromatin immunoprecipitation followed by high-throughput sequencing (ChIP-seq) and DNase I sequencing (DNase-seq)) or gene-expression values (RNA-Seq) called from the full set of reads were divided proportionally based on the abundance of phase-informative mapping to each haplotype, normalized to each haplotype’s estimated copy number. Mapping to the human reference genome imposed a slight bias, favouring the reference allele by an average of 1.08-fold. We constructed two HeLa-specific reference sequences by introducing all SNVs from each haplotype onto one or the other; mapping to this reference mitigated most of the bias (to 1.02-fold, or a 75% reduction; Supplementary Figs 36–38).

Across the HeLa genome, gene expression is significantly correlated with copy number ($P = 0.075$; Fig. 3a, b), suggesting a minimal role for gene-dosage buffering. Moreover, on average, each haplotype copy makes a comparable contribution to the transcriptome, despite uneven amplification and, in some cases, rearrangement (Fig. 3c, e). This trend is also observed for histone modifications, DNase hypersensitivity and transcription factor binding (Supplementary Figs 39 and 40). Transcript allele balances at sites heterozygous in CCL-2 on chromosome 18q closely followed the genomic balance (mean 66% representation of the A allele (two-thirds was expected)), but S3 nearly exclusively matched the A allele (94% of reads), reflecting the S3-specific LOH event (Fig. 3d). However, a small number of regions showed strong imbalances between each haplotype’s contribution to overall patterns of expression, chromatin modification and transcription factor binding (2.4% of ENCODE peaks, excluding those in LOH regions; Supplementary Figs 41–44). Interestingly, the HPV-18 insertion locus and proto-oncogene MYC (separated by approximately

Figure 4 | Haplotype-specific regulation near the HPV integration site
a. Long-range chromatin interactions between the HPV and MYC loci demonstrated by ChIA-PET28 with the RNA polymerase II signal (top) shown for HeLa S3 and an HPV− cell line (K562). Chromatin interactions (middle) are indicated by a green arrow. Bar graphs (bottom) show read counts at phased, informative sites in MYC (red, haplotype A, blue, haplotype B).

b. Transcript abundance in HeLa S3 across the MYC locus measured by RNA-seq. Overall coverage is shown in grey (top) with phased, informative sites highlighted by green ticks (pink text, non-reference alleles). Haplotype contributions at each variant are shown in bar graphs (bottom), as in a.
METHODS SUMMARY

Cells were maintained at 37 °C in DMEM F-12. Shotgun libraries were prepared by conventional ligation-based methods, sequenced on an Illumina HiSeq 2000 instrument. Point variants were called using shotgun sequence reads. Copy-number variants were identified by a reference-free approach. Functional characterization of HeLa cells by spectral karyotyping. J.S. supervised all analyses and wrote the manuscript. A.A., J.B.H., A.P.L., B.K.M., R.Q. and C.L. maintained cell cultures, constructed libraries and performed DNA sequencing. J.S. supervised all analyses and wrote the manuscript. A.A., J.B.H., A.P.L., B.K.M., R.Q. and C.L. maintained cell cultures, constructed libraries and performed DNA sequencing.

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500 kb) were among the regions with the most highly haplotype-imbalanced regulation in the genome (Supplementary Fig. 45). Phased RNA-seq data indicate that MYC is highly expressed, but almost exclusively from the HPV-18-integrated haplotype (mean ratio, 95:1; see Fig. 4b and Supplementary Fig. 46). Phased ENCODE tracks and long-range chromatin interaction data (ChIA-PET (chromatin interaction analysis with paired-end tag sequencing))24, Fig. 4a and Supplementary Fig. 47) across the region indicate that transcription-factor occupancy, active chromatin marks and long-distance physical contacts are also nearly exclusive to the HPV-integrated, transcriptionally active haplotype. Taken together, these data implicate viral integration as a strong activator of MYC expression29, acting in cis rather than in trans and possibly mediated by the epithelium-specific viral enhancer amplified to a copy number of approximately 30 within the R1 repeat structure (Fig. 2b)27. This strong cis interaction — between the amplified, integrated genome of a DNA tumour virus and a canonical proto-oncogene — may underlie the robust growth characteristics of the HeLa cell line, and provides indirect support for the hypothesis that inherited risk loci for cancer at chromosome 8q24 operate through activation of MYC30.

In summary, we present a haplotype-resolved genome and a haplotype-resolved epigenome of a human cancer. Our study not only provides an overuse genomic analysis of the human cell line that is possibly the most commonly used in biomedical research but also represents a unique view into a cancer genome and epigenome enabled by the acquisition of haplotype information.

Supplementary Information

is available in the online version of the paper.

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Author Contributions

A.A., J.B.H., J.K, and J.S. designed the experiment. A.A., J.B.H., and J.K. performed the experiments. A.A. and J.K. analyzed the data. A.A. and J.K. wrote the manuscript. J.S. supervised all analyses and wrote the manuscript.

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**METHODS**

**HeLa cell culture.** HeLa cell cultures (HeLa ATCC, CCL-2 (laboratory stock); HeLa S3 ATCC, CCL-2 (laboratory stock); Chang liver ATCC, CCL-13; L132 ATCC, CCL-5; KB ATCC, CCL-17; HEP-2 ATCC, CCL-23; WISH ATCC, CCL-25; Intestine 407 ATCC, CCL-6; FL ATCC, CCL-62; AV-3 ATCC, CCL-21) were maintained in DMEM F-12, HEPEPS (Gibco) media supplemented with fetal bovine serum (FBS) to 10% and a 1X final concentration of pen-strep antibiotic (Gibco).

**Shotgun sequencing, alignment and variant calling.** All shotgun libraries were constructed using standard ligation chemistry methods and sequenced on an Illumina HiSeq 2000. Reads were aligned to the human reference genome (hg19, b37) using BWA35 followed by duplicate removal, quality score recalibration and local indel realignment using GATK32. SNVs were called using samtools33, indel variants were called using GATK32 and short tandem repeats (STRs) were called using LobsTR33 (Supplementary Note 1). Indel detection as a function of coverage was investigated further as described in Supplementary Note 2. Gene ontology term analysis was carried out using DAVID35. Data sets used for each analysis are depicted as a flow chart in Supplementary Fig. 48.

**Read depth copy number analysis.** Shotgun reads for HeLa and Human Genome Diversity Project (HGDP) control genomes along with a similarly prepared control library with a matched G+C profile were aligned using mrsFAST37, processed as described previously38 to generate read-depth-based copy number predictions within non-overlapping windows of singly unique nucleotide k-mers (SUNK windows; Supplementary Note 3). Copy-number calling in HeLa was carried out at high (approximately 1.5-kb) and low (approximately 77-kb) resolution using an HMM (Supplementary Note 4), and a recalibration process was then used to account for widespread aneuploidy (Supplementary Note 5). Short amplifications and deletions were identified using a sliding-window approach (Supplementary Note 6). Copy-number calling was also carried out on HeLa S3 at both high and low resolutions, as well as on the eight additional HeLa strains at low resolution, and profiles were compared between strains (Supplementary Note 7). Regions of LOH were identified using a two-state HMM that used the fraction of homozygous SNVs in non-repetitive regions across low-resolution copy-number windows described above (Supplementary Note 8).

**Mate-pair library construction, sequencing and analysis.** Library construction for 40-kb mate-pair libraries was carried out starting with fosmid clone DNA pooled within each original fosmid preparation, using a protocol similar to one described previously39 (Supplementary Note 9). Libraries of approximately 3-kb inserts were constructed following protocols described previously39 (Supplementary Note 9). After read trimming and alignment, reads were split into classes based on aligned orientation and insert size, and processed using sliding windows to identify regions of probable structural rearrangement (Supplementary Note 10).

**Fosmid pool construction, sequencing and haplotype phasing.** Three replicate fosmid libraries were prepared as described previously40, and then partitioned by limited dilution into 96 sub-libraries. This was followed by outgrowth, barcoded transposase-based library preparation40, sequencing and alignment (Supplementary Note 11). Clone boundaries were inferred as described previously40, and base calls were made at all heterozygous variant positions as ascertained from whole-genome shotgun sequencing. Overlapping clones were merged to consensus haplotype blocks using an implementation of the ReFHap algorithm40 (Supplementary Note 12). Within the majority of the HeLa genome in which haplotypes are unequally amplified, adjacent blocks were merged to create scaffolds, using an HMM that finds the most likely phase of neighbouring blocks given their shotgun allele frequencies of inherited variants (those found within the 1000 Genomes Project, Supplementary Note 13). This produced a final set of haplotype scaffolds with an N50 size of 44.8 Mb, which was then used in conjunction with copy-number calls to estimate haplotype-resolved copy number for HeLa (Supplementary Note 13). Haplotype scaffolds were analysed for variant population frequencies to investigate the ancestral origin of phased blocks (Supplementary Note 14). Finally, overall copy numbers were compared among all HeLa strains sequenced in this study (Supplementary Note 15).

**Long-read phase validation.** Genomic DNA from HeLa CCL-2 was mechanically sheared using a Covaris G-tube column and standard microcentrifuge following the manufacturer’s instructions, and this produced a mean fragment size of approximately 10 kb. Single-molecule real-time sequencing libraries for the Pacific Biosciences RS sequencer were prepared using the Pacific Biosciences DNA Template Prep Kit (3–10 kb), and the resulting library was sequenced across eight cells using a 90-min movie. Resulting base calls were aligned to the genome with bwa (using parameters ‘-b5 -q2 -r1 -x1’). Reads that overlapped at least two phased SNVs were considered, excluding those within ±10bp of an insertion or deletion in the alignment.

**Identification of putative post-aneuploidy mutations.** We searched for candidate somatic post-aneuploidy mutations by taking the initial set of SNVs called from the shotgun sequencing data and filtering to remove probable germline variants. SNVs that were phased on a duplicated haplotype but that were polymorphic between the two duplicated copies were identified. Common polymorphisms and sequencing artefacts were removed by filtering against repeat annotations and control genomes (Supplementary Note 16).

**HPV-18 insertion characterization.** The HPV-18 integration locus was characterized by aligning all fosmid libraries to a modified genome that included the HPV-18 integration locus. The HPV-18 reference genome as an additional chromosome. Interchrosmosomal read pairs, fosmid-pool coverage profiles, and copy-number calls were used to determine the re-arrangement pattern of different HPV-18 integration loci. A large region of rearrangement was identified by performing an identity-adjusted alignment of data sets for all phased SNVs and assigning haplotype contributions to either peaks (epigenomics data sets) or RPKM (RNA-seq data sets), and then carrying out copy-number normalization (Supplementary Note 20). Reference bias was investigated in all tracks and removed in a subset to identify its impact on outlier calling (Supplementary Note 21). Haplotype-specific peaks were then identified in all data tracks (Supplementary Note 22). Finally, a meta-analysis of all data tracks was used to identify large regions of haplotype imbalance (Supplementary Note 23).

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