KAPA HyperPlus / SeqCap EZ workflow: improving data quality and turnaround times for targeted next-generation sequencing of FFPE DNA

Formalin-fixed paraffin-embedded (FFPE) tissue is an important source of DNA for cancer genomic studies and clinical diagnostics. A major challenge in high-throughput, targeted next-generation sequencing of FFPE DNA on the Illumina® platform is the ability to process samples of variable quality with predictable success rates in competitive turnaround times. The KAPA HyperPlus Kit with integrated enzymatic fragmentation streamlines the SeqCap EZ Target Enrichment workflow, and yields libraries of equal or better quality than those produced from Covaris®-sheared DNA.

Introduction

In clinical settings, improvements to success rates and turnaround times can have a major impact on patient care. The Centre for Molecular Pathology at the Royal Marsden Hospital previously adopted the KAPA HyperPrep Kit for library construction in their targeted sequencing workflow, which employs custom SeqCap EZ Choice capture panels. The highly efficient KAPA HyperPrep chemistry and streamlined protocol reduced library preparation time, and expanded the pool of FFPE samples that could be sequenced with consistent success rates.

Mechanical fragmentation of DNA remained a major bottleneck. With the Centre’s M-series Covaris instrument, this is a time-consuming and labor-intensive process that cannot be scaled or automated. To date, enzymatic fragmentation solutions (including tagmentation-based methods) have not proven to be robust with respect to DNA input and quality, and/or suffer from fragmentation and amplification biases that make it difficult to achieve the coverage depth required to reliably detect low-frequency mutations.¹

In this study, the KAPA HyperPlus Kit from Roche Sequencing Solutions, Inc. was evaluated. The kit provides for enzymatic DNA fragmentation and library construction in a single tube. The integrated protocol is not only easy to scale and automate, but eliminates the loss of input DNA associated with the physical transfer of fragmented DNA from Covaris tubes—thereby enabling further improvements to library complexity and overall sequence coverage. Results from validation experiments—performed with the FORMAT assay, which targets regions of 42 genes associated with gastrointestinal (GI) cancer—confirmed that the HyperPlus workflow with enzymatic fragmentation produces libraries that are equivalent or better than those generated with the KAPA HyperPrep Kit from Covaris-sheared DNA. Further improvements to the HyperPlus / SeqCap EZ workflow were achieved through optimization of the capture library used in the GI assay. Turnaround times were reduced significantly by eliminating a second round of hybridization, without sacrificing coverage depth.
Materials and methods

**Experimental design.** Three paired DNA samples from FFPE tissue and peripheral blood were processed in parallel, using a previously validated library construction workflow with Covaris® shearing, vs. the new workflow with enzymatic fragmentation. Sequencing data were compared with respect to the following key metrics: (i) % duplicates; (ii) % unique on-target reads; (iii) level of coverage at 100X, 250X, and median depth; and (iv) detection of variants at 5% frequency and above.

**Library construction.** FFPE DNA (200 ng) or non-tumor blood DNA (50 ng) was sheared to a mode size of 150–200 bp using a Covaris M220 instrument. Libraries were prepared with the KAPA HyperPrep Kit (Roche) and barcoded SeqCap EZ adapters (Roche), according to the previously validated protocol, which includes bead-based size selection after ligation, and pre-capture amplification with KAPA HiFi HotStart ReadyMix (8 cycles for FFPE, 7 cycles for paired normal controls). In parallel, libraries were prepared from the same samples in a similar manner, but using the KAPA HyperPlus Kit with enzymatic fragmentation for 30 min at 37°C. Pre-capture amplification was reduced to 6 cycles for both sample types. Positive control libraries were prepared from 50 ng male control DNA (Promega), and negative control libraries without any input DNA.

**Target capture.** Pre-capture libraries were quantified by Qubit® (ThermoFisher Scientific), after which pools were prepared for multiplexed target enrichment with the FOrMAT panel (~200 kb). Each pool consisted of 1 µg of library DNA, divided more or less equally between five libraries of a single type (FFPE or blood). Positive control (20 ng) and NTC (~0 ng) libraries were included in FFPE pools. Hybridization reactions (one per pool) were set up with SeqCap EZ reagents according to the previously validated protocol. With the original GI panel, two rounds of overnight hybridization (≥16 hours at 47°C) and capture were performed, with 5 cycles of amplification with KAPA HiFi HotStart ReadyMix after the first round, and 11 cycles of post-capture amplification.

**Sequencing and data analysis.** Final, enriched library pools were quantified by qPCR using the KAPA Library Quantification Kit (Roche). Each pool was diluted to 4 nM, after which one tumor and one normal pool was combined at a ratio of 4:1, denatured and loaded onto an Illumina® MiSeq® instrument. Paired-end sequencing (2 x 150 bp) was performed using a MiSeq v3 chemistry kit (Illumina). Data were processed using a standardized analysis pipeline, which employs MiSeq Reporter for initial data processing and output, and Picard for de-duplication and generation of sequencing metrics. Reads were aligned against the hg19 human reference genome. Variant analysis was performed with Illumina VariantStudio v2.2, and translocations called with Manta.

**Results**

Comparative sequencing (Figure 1) and coverage (Figure 2) metrics for libraries prepared from Covaris-sheared DNA—using the previously validated KAPA HyperPrep / SeqCap EZ workflow vs. the KAPA HyperPlus / SeqCap EZ workflow with integrated enzymatic DNA fragmentation—are given on the next page. These metrics were chosen as they are reflective of library quality and the utilization of sequencing capacity. The HyperPlus workflow outperformed the existing workflow in all five metrics, with notable improvements for the FFPE samples. In addition to eliminating a process bottleneck, the KAPA HyperPlus Kit enabled a significant increase in coverage depth for FFPE samples (from the same amount of sequencing), which is critical for the accurate detection of rare somatic variations.

The final requirement for validation of the KAPA HyperPlus / SeqCap EZ workflow was to compare the detection of sequence variants at a frequency above 5% in libraries prepared with the new workflow vs. the existing protocol with Covaris shearing. The results of this comparison are given in Figure 3. For each of the six libraries, a random number of chromosomal positions were interrogated, and the percentage variant frequency vs. the hg19 human reference genome calculated. Correlations between the two methods ranged between 96.7% and 99.6% for the three FFPE libraries, and between 98.1% and 99.8% for the three libraries prepared from blood DNA. This confirmed that the HyperPlus workflow produced libraries that were functionally equivalent to those produced with the previously validated protocol.
High-quality capture libraries from FFPE samples | 3

Figure 2. Comparative coverage metrics. (A) The level of coverage at 100X coverage depth was high and similar for both protocols and sample types. (B) At higher coverage depth (250X), the HyperPlus workflow outperformed the original protocol. The average HyperPlus coverage at 250X was 89.2% and 86.2% for FFPE and paired controls, respectively—as opposed to 73.7% and 65.7% for the original protocol with Covaris® shearing. (C) Since coverage is impacted by the number of reads obtained for each library in the sequencing pool, the calculated value for each sample was adjusted to reflect the difference in median coverage between the two protocols, should the number of reads have been equal for all libraries. The average median coverage achieved with the HyperPlus workflow for the FFPE samples (1,154X) represents a 3.5-fold improvement over the existing protocol (average median coverage = 328X). The difference in average median coverage for the paired blood samples was much smaller (333X for HyperPlus vs. 302X for the Covaris protocol, or a 1.1-fold improvement).

Figure 3. Variant detection correlations. For each of the three FFPE (tumor) libraries (top row) and its normal paired library (prepared from peripheral blood DNA; bottom row), the variant frequency (vs. the hg19 human reference genome) was calculated for a select number of chromosomal positions with expected variant frequencies between 5% and 100%. Each data point represents the calculated variant frequency (%) obtained with the existing workflow (x-axis) vs. the HyperPlus workflow (y-axis). Regression analysis was performed in Microsoft® Excel.
Elimination of Covaris® shearing reduced library preparation time (fragmentation to pre-capture library) by ~30%. However, the majority (48 hours) of the total turnaround time from tissue to sequence data (~76 hours, excluding data analysis) was still dedicated to the “double capture” protocol, previously shown to be advantageous for small and challenging capture panels.\(^5\)

Optimization of the gastrointestinal cancer target library, through the addition of probes in areas that were historically poorly covered, and removal of probes from non-unique regions, enabled elimination of the second round of hybridization and capture, and the 5 cycles of amplification between the two rounds. This translated into a significant time saving (1 day) with a further decrease in duplication rates (Figure 4A). As expected, a single round of capture data with a small panel resulted in lower on-target rates (not shown), but mean coverage depth remained similar as a result of more uniform coverage achieved with the optimized panel (Figure 4B).

**Conclusions**

Implementation of the KAPA HyperPlus Kit in a targeted sequencing assay employing a small, custom SeqCap EZ Choice target enrichment panel enabled significant improvements to both turnaround time and FFPE library quality. The robust, single-tube, low-bias HyperPlus chemistry eliminates mechanical shearing, and enables streamlining of the capture workflow through improved library construction efficiency. Overall, sample preparation time (tissue to raw sequence data; analysis excluded) was reduced by ~30%, while improving library complexity and coverage depth.

**References**