Suitability of melanoma FFPE samples for NGS libraries: time and quality thresholds for downstream molecular tests

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The use of NGS in clinical practice for precision diagnosis requires a quality starting material. Despite the broadly established use of formalin-fixed paraffin-embedded (FFPE) samples in molecular testing, these usually have low-quality DNA. We established a method to determine the suitability of melanoma FFPE samples for an amplicon-based NGS custom panel analysis. DNA was extracted from unstained melanoma samples and wide local excision samples. Amplicon-based libraries were constructed and tested using time and quality parameters as variables. Time elapsed from sample retrieval >7 years, a quality control value >5.63 and a DNA integrity value <2.05 indicated samples were not suitable. A decision tree is provided with rate of samples suitable for analysis according to the combination of these parameters.

METHOD SUMMARY

To maximize sample usability and minimize costs, we proposed a procedure to identify samples with enough quality for massive sequencing. Our proposal included the use of quality control and DNA integrity number quality parameters together with the time elapsed from surgery to develop an affordable protocol that can be easily implemented in research and clinical laboratories.
resis [10,11]. Also, the quality control (QC) value from Illumina (CA, USA) enables the researcher to test the quality of the DNA based on a qPCR, giving values closer to zero when the quality is the highest [12,13].

Our aim was to evaluate the suitability of melanoma FFPE samples for an amplicon-based NGS custom panel analysis according to the storage time, type of sample, QC and DIN values.

Materials & methods

Sample collection

FFPE blocks came from the Biobank at the Instituto Valencia de Oncología (Valencia, Spain). A cohort of 59 samples were analyzed including 37 primary melanoma tumors and 22 wide local excision tissue samples retrieved and stored at the Pathological Anatomy Department from January 2000 to April 2017. Therefore, the time elapsed from the surgery (time of storage) was up to 17 years and the distribution was made based on a CART analysis regarding the library functionality (see below): 42.4% of the samples were referred to as ‘old’ (>7 years) and 57.6% of the samples were referred to as ‘recent’ (<7 years). The time of sample fixation in formalin solution was estimated based on the date of the surgery when the sample was taken. Two categories were defined: <1 day (usually overnight fixation), when the day after the surgery was a work day and >1 day when it was a holiday (usually >1–2 days).

This study took place as part of a bigger project that had the approval of the Ethics Committee at the Instituto Valenciano de Oncología, and patients signed a voluntary cession of the samples to the Biobank.

The main outcome variable was the functionality of the sample, which was defined as the ability of a sample to construct an amplicon-based library with a length of 300–350 bp, visible as a single band on an electrophoresis test.

FFPE processing

The Pathological Anatomy Department of our center had a standardized protocol for the processing of FFPE samples. A first formalin fixation of the sample was followed by the block preparation procedure. This was performed in the Excelsior ES automatic processor (ThermoScientific, CA, USA) and included a formalin fixation-step (30 min), an increasing multiple-step dehydration (9 h 45 min), a triple clearing step with xylene (2 h 15 min), and a final three-step embedding in paraffin wax (4 h). Then, blocks were kept at room temperature at the Biobank.

DNA isolation & quantification

From each FFPE block, a 3-μm section was used for hematoxylin and eosin (H&E) stain. Then, a pathologist evaluated and selected the area with tumor-enriched cells for macrodissection. Using the H&E slide as a reference, three 0.6-mm needle biopsies were taken from every primary tumor. For the wide local excision tissue, three 10-μm sections were cut and collected into 1.5-ml tube (Eppendorf).

DNA was extracted using the QiAamp® DNA Investigator kit (QIAGEN, Hilden, Germany), with the following modifications: given the toughness of the skin, we established an overnight incubation at 56°C for the proteinase K to assure a complete digestion. Also, we introduced the optional carrier addition to maximize the extraction yield.

Quantification was obtained using QuantiTm PicoGreen™ dsDNA Assay kit (Invitrogen, MA, USA). All samples had a concentration above 2.5 ng/μl and were accepted for the study.

DNA repair

The NEBNext® FFPE Repair Mix (New England Biolabs, Hertfordshire, UK) kit was used to repair the C-G > T-A changes induced by nucleotide deamination, usually present in FFPE samples.

Quality assessment tests

Real-time PCR was performed using 1X Sybr Green Master Mix (Applied Biosystems) and FFPE QC Kit (Illumina) following the manufacturer’s instructions. Briefly, 2 μl of diluted DNA (1:100) was added to 8 μl of the mix containing SybrGreen and Illumina primers. All runs were processed in an ABI7500 Fast PCR system (Applied Biosystems) using the default run protocol: 50°C/2 min–95°C/10 min–40 cycles of 95°C/30 s, 57°C/30 s, 72°C/30 s). All reactions were performed in triplicates. The resulting QC value was an indicator of the sample quality, with a lower value being the better the quality indicator.

Gel electrophoresis was performed using Genomic DNA ScreenTape in a 4200 TapeStation (Agilent Technologies). Briefly, 1 μl of DNA and 3 μl of sample buffer were added to each well.

DIN value obtained was an indicator of the integrity of the DNA, thus a higher DIN value meant a better quality.

Library construction

Low input DNA libraries of the gene panel containing 21 melanoma-related genes were constructed according to the manufacturer’s instructions using a custom GeneRead DNAseq Panel (QIAGEN). Shortly, DNA fragments were amplified in a multiplex PCR to obtain a total of 633 amplicons of 200 bp in length (GeneRead DNAseq Panel PCR kit V2 Qiagen). At this point, a normalization step was included and 100 ng of each sample continued the process. The ends of the molecules were enzymatically repaired and universal adapters were ligated, then unique combinations of MID adapters were ligated (NEBNext Ultra DNA Library Prep Kit, New England Biolabs).

Library functionality

Final library size was checked with a bioanalyzer using D1000 DNA ScreenTape (Agilent Technologies). The final amplicon size including the MID adapters made an average of 350 bp. Thus, the presence of a single band in the range of 300–350 bp classified the library as functional.

Statistical analysis

The statistics were performed using IBM SPSS Statistics 20.0. Normal distribution of continuous variables was checked using the Kolmogorov–Smirnov test. Pearson test was used to study the correlation among parametric variables and Spearman test was used for non-parametric variables. A 1-factor ANOVA test was used to compare means of continuous variables and qualitative ones. Also, continuous variables were categorized with a CART analysis using library functionality as a filter. Diagnostic parameters, including sensitivity, specificity, predicted positive and negative value, accuracy, and area under the curve from a ROC test were calculated to evaluate the capacity of each parameter or algorithm to predict library functionality.
Results & discussion

The study included a total of 59 unpaired FFPE samples corresponding to 37 melanoma primary tumors and 22 unmatched wide local excision tissues stored for a median of 5 years (range: 1–17 years). The characteristics of the samples are displayed in Table 1.

Influence of date of surgery on library functionality

There was a great variability in library yield independent of the tissue origin or the time of storage, despite the existence of a normalization step to include 100 ng in the end reparation step. A 1-factor ANOVA test showed that the mean storage time was significantly lower in samples with functional library than in samples with non-functional library (4.93 vs 9.49 years, respectively; p < 0.001). The best cut-off that differentiated functional from non-functional samples was established at 7 years by CART analysis, and recent samples (≤7 years) showed a significantly higher percentage of library functionality than old samples (>7 years; 70.6 vs 20%; p < 0.001) (Figure 1A). The fixation time did not influence the functionality (Table 1).

QC & DIN values as quality predictor parameters

QC values were inversely correlated with time of storage (r = -0.616; p < 0.001) (Figure 1B) and library functionality (r = -0.334; p = 0.009). A 1-factor ANOVA test showed that QC means were statistically different between functional and non-functional samples (4.03 vs 5.7; p = 0.002).

According to the CART analysis, cut-off for QC was established at 5.63 (93.1% of samples with a QC ≤ 5.63 produced functional libraries compared with 6.9% of samples with a QC > 5.63; p = 0.004) (Figure 1C).

DIN values showed a direct correlation with the time of storage (r = 0.523; p < 0.001) (Figure 1D), as well as with library functionality (r = 0.319; p = 0.016). We established a cut-off for DIN value at 2.05 based on CART analysis, and results showed that DIN values greater than 2.05 gave functional libraries in a higher proportion than those less than or equal to 2.05 (82.1 vs 17.9%; p = 0.023) (Figure 1E). As expected, QC and DIN were inversely correlated (r = -0.535; p < 0.001) (Figure 1F).

When looking at the possible differences between tumor tissue and wide local excision samples within old and recent groups, it was found that for the old cohort, lower QC values were more frequent in tumor samples (13/24; 54.2%) than in wide local excision samples (1/14; 7.1%; p = 0.009). No difference was found for DIN or functionality in this group. For the recent cohort, functional libraries corresponded in a higher proportion to the wide local excision (15/24; 62.5%) rather than tumor samples (9/24; 37.5%). No difference was found for DIN or QC in this group.

Convergence of parameters in a decision tree

Time of storage, QC and DIN were simultaneously assessed by CART test to analyze their impact on the library functionality, and a decision tree was developed (Figure 2). The storage time was the parameter that better discriminated the library functionality. Samples stored for 7 years or less gave functional libraries in 70.6% of the cases. For samples stored for more than 7 years the value of QC in the first place, and of DIN in the second place, discriminated samples by their functionality. Thus, samples with QC less than or equal to 5.63 and DIN greater than 2.05, allowed identification of a group with 44% of functional libraries.

The diagnostic parameters were evaluated for each variable individually and for the algorithm obtained by CART analysis. The latter was also evaluated only for the oldest samples (Table 2).

The decision tree provided better values in the diagnostic parameters (sensitivity = 91.3%; specificity = 72.9%; predicted positive values = 65.1%; predicted negative values = 93.8%; accuracy = 79.5%; ROC area under the curve = 0.733) than each parameter individually. In addition, the decision tree restricted to the old samples also showed acceptable figures (ROC = 0.78).

There are several studies in the literature that have evaluated QC and DIN/RIN values prior to NGS [11]. Yakovleva et al. established in 2017 a cut-off of RIN of > 2.00 for using FFPE samples in downstream processes [10]. Similarly, Bonfiglio et al. proposed in 2016 the use of a DIN cutoff at 3.0 [13]. All these previous studies agree with our findings, but all of them used FFPE samples that had been stored for a maximum of 6 years. Hence, we contribute

Table 1. Sample distribution according to their final library functionality.

<table>
<thead>
<tr>
<th></th>
<th>Total</th>
<th>Functional</th>
<th>Non-functional</th>
<th>p-value</th>
</tr>
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<tbody>
<tr>
<td>QC</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>QC ≤ 5.63</td>
<td>45</td>
<td>76.2</td>
<td>27</td>
<td>93.1</td>
</tr>
<tr>
<td>QC &gt; 5.63</td>
<td>14</td>
<td>23.7</td>
<td>2</td>
<td>6.9</td>
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<tr>
<td>DIN</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>DIN ≤ 2.05</td>
<td>19</td>
<td>33.9</td>
<td>5</td>
<td>17.9</td>
</tr>
<tr>
<td>DIN &gt; 2.05</td>
<td>37</td>
<td>66.1</td>
<td>23</td>
<td>82.1</td>
</tr>
<tr>
<td>Time of storage</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Old</td>
<td>25</td>
<td>51</td>
<td>5</td>
<td>17.2</td>
</tr>
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<td>Recent</td>
<td>34</td>
<td>57.6</td>
<td>24</td>
<td>82.8</td>
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<tr>
<td>Fixation time</td>
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<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;1 day</td>
<td>25</td>
<td>44.6</td>
<td>15</td>
<td>51.7</td>
</tr>
<tr>
<td>≥1 day</td>
<td>11</td>
<td>19.6</td>
<td>5</td>
<td>17.2</td>
</tr>
<tr>
<td>Not available</td>
<td>20</td>
<td>35.7</td>
<td>8</td>
<td>27.6</td>
</tr>
</tbody>
</table>

p-value by Chi-squared test. DIN: DNA integrity value; N: Number; QC: Quality control.
Figure 1. Statistical analysis. Here are presented the graphical distribution of failed (white) and functional (gray) libraries for the different variables (Date of surgery, QC threshold and DIN threshold) (A, C, E). Also, the correlations studied between QC and date of surgery (B), DIN and date of surgery (B), and QC and DIN (F). DIN: DNA integrity value; QC: Quality control.

Table 2. Diagnostic test parameters for each variable and the algorithm obtained by CART analysis.

<table>
<thead>
<tr>
<th>Variable</th>
<th>S (%)</th>
<th>SP (%)</th>
<th>PPV (%)</th>
<th>PNV (%)</th>
<th>A (%)</th>
<th>ROC (AUC)</th>
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<tbody>
<tr>
<td>QC</td>
<td>80.8</td>
<td>68.2</td>
<td>60.0</td>
<td>85.7</td>
<td>72.9</td>
<td>0.33</td>
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<td>DIN</td>
<td>70.3</td>
<td>66.1</td>
<td>62.2</td>
<td>73.7</td>
<td>68.0</td>
<td>0.66</td>
</tr>
<tr>
<td>Time of storage</td>
<td>77.9</td>
<td>73.1</td>
<td>70.6</td>
<td>80.0</td>
<td>75.3</td>
<td>0.75</td>
</tr>
<tr>
<td>CART†</td>
<td>91.3</td>
<td>72.9</td>
<td>65.1</td>
<td>93.8</td>
<td>79.5</td>
<td>0.73</td>
</tr>
<tr>
<td>CART (for old samples)‡</td>
<td>87.7</td>
<td>62.8</td>
<td>44.4</td>
<td>93.8</td>
<td>69.1</td>
<td>0.78</td>
</tr>
</tbody>
</table>

*Cart† includes the results for the diagnostic parameters of the proposed flowchart obtained by CART analysis.
‡Only for old samples, with a date of surgery > 7 years. Positive if QC ≤ 5.63 and DIN > 2.05.
*Positive for the test if date of surgery < 7 years or if QC ≤ 5.63 and DIN > 2.05.
Author contributions
DM, ZG and EN conceived and designed the experiments. JAL provided the technology. JB, CR and VT assessed sample identification. DR and AR managed sample collection and preparation. DM and DR performed the experiments. DM and EN analyzed the data. ZG and EN supervised the study. DM wrote the paper. EN corrected the manuscript.

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