Clinical benefit of a precision medicine based approach for guiding treatment of refractory cancers

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ABSTRACT

Patients and methods: Patients with metastatic solid tumors who had progressed on at least one line of standard of care therapy were referred to the Indiana University Health Precision Genomics Program. Tumor samples were submitted for DNA & RNA next-generation sequencing, fluorescence in situ hybridization, and immunohistochemistry for actionable targets. A multi-disciplinary tumor board reviewed all results. For each patient, the ratio of progression-free survival (PFS) of the genomically guided line of therapy divided by the PFS of their prior line was calculated. Patients whose PFS ratio was ≥ 1.3 were deemed to have a meaningful improvement in PFS.

Results: From April 2014–October 2015, 168 patients were evaluated and 101 patients achieved adequate clinical follow-up for analysis. 19 of 44 (43.2%) patients treated with genomically guided therapy attained a PFS ratio ≥ 1.3 vs. 3 of 57 (5.3%) treated with non-genomically guided therapy (p < 0.0001). Similarly, overall PFS ratios (irrespective of cutoff) were higher for patients with genomically guided therapy vs non-genomically guided therapy (p = 0.05). Further, patients treated with genomically guided therapy had a superior median PFS compared to those treated with non-genomically guided therapy (86 days vs. 49 days, p = 0.005, H.R. = 0.55, 95% C.I.:0.37-0.84).

Conclusion: Patients with refractory metastatic cancer who receive genomically guided therapy have improved PFS ratios and longer median PFS compared to patients who do not receive genomically guided therapy.

INTRODUCTION

Precision medicine approaches that seek to match genomic aberrations to potential treatment avenues are rapidly reshaping treatment for cancer patients [1-3]. Advances in genomic technology have enabled clinical application of these approaches, where large scale DNA & RNA profiling of cancer genes from small amounts of archival tissue is now possible [4-6]. In patients with advanced cancer where treatment options are limited, genomic sequencing is frequently being used to identify targets which are potentially clinically actionable with FDA approved drugs or clinical trials. In many cases these targets would have been overlooked under standard clinical practice due to the rarity or novelty of the target, or its presence in a cancer lineage not normally associated
with the target (e.g. HER-2 amplification in colon cancer). These observations and the evolution of targeted therapeutics have spurred a paradigm-shift that recognizes molecular drivers as increasingly powerful additions to the traditionally dominated organ and histology-centric therapeutic standard.

While several publications to date have determined that many cancers have targetable pathways that may benefit from a precision medicine approach [7], formal assessment of clinical benefit of these approaches has been lacking. Published case reports demonstrating exceptional responses to targeted therapy explained by key genomic mutations have brought promise to the field [8-10], but analyses of larger cohorts and randomized controlled clinical trials are still needed to help solidify the utility of cancer genomics in clinical practice. Survival analyses of cohorts with mixed cancer lineages (and mixed genomic aberrations) have been fraught with bias secondary to the widely varying nature of expected survival outcomes for different cancer types. In 2010, Von Hoff et al. [11], reported early experience with identifying molecular targets in patients’ tumors to direct treatment and also performed a formal analysis of clinical benefit by calculating the ratio of the PFS of the patients’ genomically-guided therapy divided by the PFS of their prior line of therapy. This approach took into account both the natural history of the patient’s individual cancer, as well as the fact that subsequent lines of therapy nearly always lead to declining PFS times. In their analysis, each patient was able to serve as his or her own control. Therefore, a PFS ratio in excess of 1.0 (or more conservatively, in excess of 1.3 to minimize false-positive fluctuations) provided a reference point of activity relative to the last line of therapy.

In April 2014, the Indiana University Health Precision Genomics Program was initiated to offer next-generation sequencing for patients with metastatic refractory or rare solid tumors in order to provide genomically guided treatment recommendations. Decisions regarding therapy were derived from a multi-disciplinary advisory board with consideration given to strength of genomic association, co-morbidities, prior drug exposure, and inherited genetic variants that would predict increased risk from a drug. Patient demographics, tumor characteristics, information on duration of prior therapies, genomic information, and clinical follow-up were prospectively collected for all patients since the inception of the program. Herein, we report our first formal efficacy analysis to determine the clinical benefit of a precision medicine approach in patients treated with genomically-guided therapy vs. non-genomically guided therapy.

RESULTS

Patient characteristics

Figure 1 details the Consolidated Standards of Reporting Trials (CONSORT) diagram demonstrating the disposition of the 168 patients who were referred to the Indiana University Simon Cancer Center (IUSCC) Precision Genomics Clinic. Of these, 67 were excluded from the final analysis: the commonest reasons being not receiving any therapy post genomic profiling or lost to follow-up (n = 40). Table 1 lists the characteristics of the 101 patients who were referred to the Precision Genomics clinic and are part of the analyzed cohort. Patient demographics were well balanced between groups. The majority of patients had a diagnosis of soft tissue sarcoma, breast cancer, pancreatic cancer or colorectal cancer. The median number of previous therapeutic regimens was 4 for both groups, and 96% of both groups had an ECOG Performance Status (PS) of 0 or 1.

Molecular characteristics

DNA mutation, copy number variation, Immunohistochemistry (IHC), and mRNA overexpression findings that were considered by the Precision Genomics Tumor Board in determining potential therapies are depicted in Figure 2. The most notable observations that led to clinical interventions included: DNA mutations in IDH1 and EGFR; copy number variation in fibroblast growth factor and Human Epidermal Growth Factor Receptor-2 (HER2); IHC positivity for PD-L1 and MET; and mRNA overexpression for amphiregulin (AREG), epieregulin (EREG), E-Cadherin (CDH1), hENT1, Topoisomerase IIa (TOPOIIa), and Vascular Endothelial Growth Factor-A (VEGFA).

Progression free survival (PFS) analyses

43.2% (19 of 44) of patients treated according to genomic recommendations were found to have a PFS ratio of > 1.3 compared to 5.3% (3 of 57) of patients who did not receive treatment guided by genomic recommendations (p < 0.0001) (Figure 3A). Patients who did not receive treatment guided by genomic recommendation included the following: a lack of actionable genomic target, inaccessibility to treatment recommendation, or physician choice not to pursue genomic based treatment. A secondary endpoint analysis of PFS ratio >1 (equal to at least the previous regimen) occurred in 50% (22 of 44) and 19.3% (11 of 57) of patients who received genomically directed vs. non-genomically directed therapy, respectively (p = 0.0011).
Table 1: Patient Demographics and Clinical Characteristics

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Genomic Directed Group (n = 44), no. (%)</th>
<th>Non-Genomic Directed Group (n = 57), no. (%)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male Sex</td>
<td>47.7%</td>
<td>45.6%</td>
<td>0.83</td>
</tr>
<tr>
<td>Female Sex</td>
<td>52.3%</td>
<td>54.4%</td>
<td></td>
</tr>
<tr>
<td>Mean (+/- SD) Age, years</td>
<td>55.5 (12.5)</td>
<td>58.4 (11)</td>
<td>0.22</td>
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<tr>
<td>ECOG PS</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>12 (27.3)</td>
<td>16 (28.1)</td>
<td>0.93</td>
</tr>
<tr>
<td>1</td>
<td>31 (70.5)</td>
<td>39 (68.4)</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>1 (2.3)</td>
<td>2 (3.5)</td>
<td></td>
</tr>
<tr>
<td>Mean (+/- SD) time until progression on previous regimen, days</td>
<td>122 (103)</td>
<td>130 (97)</td>
<td>0.75</td>
</tr>
<tr>
<td>No. of Prior Regimens</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1 to 2</td>
<td>15</td>
<td>25</td>
<td>0.32</td>
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<tr>
<td>3 to 4</td>
<td>21</td>
<td>24</td>
<td></td>
</tr>
<tr>
<td>5 or greater</td>
<td>8</td>
<td>8</td>
<td></td>
</tr>
<tr>
<td>Median (range) number of regimens</td>
<td>4 (2-6)</td>
<td>4 (2-6)</td>
<td></td>
</tr>
<tr>
<td>Tumor type</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Soft tissue sarcoma</td>
<td>10 (22.7)</td>
<td>11 (19.3)</td>
<td>0.67</td>
</tr>
<tr>
<td>Breast</td>
<td>8 (18.2)</td>
<td>8 (14.0)</td>
<td>0.57</td>
</tr>
<tr>
<td>Pancreas</td>
<td>1 (2.3)</td>
<td>7 (12.3)</td>
<td>0.13</td>
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<tr>
<td>Colorectal</td>
<td>7 (15.9)</td>
<td>5 (8.8)</td>
<td>0.36</td>
</tr>
<tr>
<td>Bladder/Urothelial</td>
<td>2 (4.5)</td>
<td>4 (7.0)</td>
<td></td>
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<tr>
<td>Prostate</td>
<td>0</td>
<td>1 (1.8)</td>
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<tr>
<td>Renal cell</td>
<td>1 (2.3)</td>
<td>0</td>
<td></td>
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<tr>
<td>Cervical</td>
<td>1 (2.3)</td>
<td>0</td>
<td></td>
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<tr>
<td>Cholangiocarcinoma</td>
<td>1 (2.3)</td>
<td>1 (1.8)</td>
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<tr>
<td>Esophageal</td>
<td>2 (4.5)</td>
<td>0</td>
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<tr>
<td>Head and Neck</td>
<td>1 (2.3)</td>
<td>1 (1.8)</td>
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<tr>
<td>Hepatocellular</td>
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<td>1 (1.8)</td>
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<tr>
<td>Melanoma</td>
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<td>3 (5.3)</td>
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<tr>
<td>Neuroendocrine</td>
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<td>2 (3.5)</td>
<td></td>
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<tr>
<td>Non-small cell lung cancer, adeno</td>
<td>0</td>
<td>4 (7)</td>
<td></td>
</tr>
<tr>
<td>Non-small cell lung cancer, squamous cell</td>
<td>0</td>
<td>2 (3.5)</td>
<td></td>
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<tr>
<td>Small cell lung cancer</td>
<td>1 (2.3)</td>
<td>1 (1.8)</td>
<td></td>
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<tr>
<td>Squamous cell carcinoma, NOS</td>
<td>0</td>
<td>1 (1.8)</td>
<td></td>
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<tr>
<td>Unknown Primary</td>
<td>1 (2.3)</td>
<td>0</td>
<td></td>
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<tr>
<td>Thyroid, anaplastic</td>
<td>1 (2.3)</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>Adrenal Cortical Carcinoma</td>
<td>0</td>
<td>1 (1.8)</td>
<td></td>
</tr>
<tr>
<td>Small Bowel</td>
<td>1 (2.3)</td>
<td>0</td>
<td></td>
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<tr>
<td>Ampulla of Vater</td>
<td>0</td>
<td>1 (1.8)</td>
<td></td>
</tr>
<tr>
<td>Chondrosarcoma</td>
<td>1 (2.3)</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>Endometrial</td>
<td>1 (2.3)</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>Ewing</td>
<td>1 (2.3)</td>
<td>0</td>
<td></td>
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<tr>
<td>Glioblastoma multiforme</td>
<td>0</td>
<td>1 (1.8)</td>
<td></td>
</tr>
<tr>
<td>Ovarian</td>
<td>1 (2.3)</td>
<td>2 (3.5)</td>
<td></td>
</tr>
<tr>
<td>Myoepithelial</td>
<td>1 (2.3)</td>
<td>0</td>
<td></td>
</tr>
</tbody>
</table>
Similarly, a PFS ratio >1.5 occurred in 29.5% (13 of 44) of patients with genomic directed therapy versus 5.3% (3 of 57) without \((p = 0.0017)\). When statistically comparing all PFS ratios, patients who received genomically directed therapy had higher PFS ratios compared to non-genomic guided therapy (mean PFS ratio: 1.34 vs 0.8, \(p = 0.05\)). Lastly, Kaplan-Meier PFS analysis demonstrated a median PFS of 86 days in the genomic therapy group vs. 49 days in the non-genomic therapy group \((p = 0.005, \text{H.R.} = 0.55, 95\% \text{C.I.:} 0.37-0.84)\) (Figure 3B).

**Details of responders**

The PFS durations and therapies of those who achieved a PFS ratio ≥ 1.3 \((n = 19)\) are detailed in Figure 4. A number of patients with significant benefit had a diagnosis of sarcoma (6 cases with soft tissue sarcoma, 1 case with chondrosarcoma, and 1 case with Ewing’s sarcoma). Of interest, five of our patients were continuing a positive response at the time of analysis. Details of the three patients who achieved a PFS ratio of ≥ 1.3 who did not receive genomically-directed therapy are as follows: one patient was diagnosed with NSCLC and received previous therapy with vinorelbine followed by therapy with nivolumab on a clinical trial, the second patient was diagnosed with squamous cancer NOS and received previous therapy with cetuximab followed by therapy with nivolumab on a clinical trial. Of note, both of these patients were seen in our program prior to incorporation of PD-L1 IHC testing as part of our panel. The third patient was diagnosed with metastatic colorectal cancer and was previously treated with regorafenib followed by the non-genomically directed therapy of BB1503 as part of a Phase I clinical trial.

**DISCUSSION**

Solid tumors have historically been treated based on the organ site of the cancer’s origin. More recently, it has been appreciated that some disease types have important molecular biomarkers that allow for therapies to be tested and used only in a subset of that histological disease subtype [12, 13]. These have been called “actionable targets”. For some disease types, actionable targets are common; for others, they are rare. In the heavily pretreated metastatic setting, conventional cytotoxic therapies have limited success [14, 15]. Many patients in this setting are guided to early phase trials, where novel compounds are being tested with the hope that a unique mechanism of action might provide benefit. The reality of non-molecularly guided early phase trials is that the average response rate and progression free survival is limited [16]. Targeted therapy, conversely, has yielded some examples of substantial benefit when a driver target is appropriately disrupted. Further, these therapeutic targets have occasionally been found to be successful in some tumor types (HER2+ for both breast and gastric cancer).

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**Figure 1:** CONSORT diagram of evaluable population.
Figure 2: Oncoprint analysis detailing the genomic findings considered by the molecular tumor board for the study population \((n = 101)\). Each column represents a single patient, and each row a gene that harbors a mutation, copy number gain, IHC (immunohistochemistry) positivity, or mRNA overexpression.
but not others (HER2 in lung cancer) [18]. The incomplete dominance of a therapeutic target implies a complex interaction between the target itself and the disease type, perhaps due to convergent phenotypes or because of the degree of driver dependence and is a major hurdle that must be considered [19].

Affordable and fast high-throughput genomic testing has made it possible to evaluate hundreds or even thousands of driver aberrancies simultaneously. This approach allows for the exploitation of a cadre of drugs including both FDA approved drugs and targeted agents in development in clinical trials. Several trials have been conducted and multiple others are underway to more formally explore the relative success for matching drugs with high throughput genomic biomarkers [20-22]. Although these trials will represent critical steps toward

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**Figure 3: Progression free survival (PFS) analysis of patient population.**

**A.** Bar graph comparing the percentage of patients who achieved a PFS ratio equal to or greater than 1.3. 43.2% (19 of 44) of patients on genomically directed therapy achieved a PFS ratio of 1.3 compared to only 5.3% (3 of 57) who received non-genomically directed therapy ($p < 0.0001$).

**B.** Kaplan-Meier plot of the PFS of the genomic-directed group (blue line) vs. non-genomic directed group (yellow line). Patients treated with genomic guided therapy had a superior median PFS compared to those treated with non-genomic guided therapy (86 days vs. 49 days, $p = 0.005$, H.R. = 0.55, 95% C.I.:0.37-0.84).
answering the question of the relative efficacy of this approach, to date, these trials have been difficult to execute for a variety of reasons. First, the panel of potential markers and drugs is ever changing and expanding. By the time a trial is complete, the biomarkers and targeted drugs are often outdated. In fact, none to date will have comprehensively included all FDA-approved drugs and all drugs in development. Second, many of these trials have conventional exclusion criteria that don’t mirror the real world scenario where mild organ dysfunction and declining performance status are commonplace in advanced cancer patients. There has also been the creation of clinical programs, similar to that described here, with the goal of maximizing informed decisions. These clinical programs often represent a more generalizable population and may have access to broader selection of therapeutic options.

Despite the advantages highlighted in this type of clinical program, there are several limitations to our study. First, as this was not a randomized-controlled trial, therapeutic selections were recommended (not mandated) and rigorous follow-up with predefined time-points of evaluation were not uniform. Second, targeted genomic sequencing might serve as a prognostic marker rather than a predictive marker. It may be that patients who didn’t receive directed therapy may have also been those patients who were unable to travel to a clinical trial or tolerate therapy because of declining performance status. Lastly, a proportion of those patients not included in the analysis were due to the inability to completely quantify the duration of prior response. This is a unique challenge that traditional clinical trials don’t encounter as the primary measures of response happens during (ORR, PFS) or after enrollment (OS) as opposed to the need to accurately quantify prior duration of response retrospectively. Strengths of this study include the use of a multidisciplinary tumor board to optimize consistency of drug-target selection as well as having a body of experts to routinely review the biomarker literature and evolve as biomarker data and drugs/trials become available in real time. Another strength is the meticulously curated data from the previous treatment experience for each patient and the ability to prospectively follow outcomes.

In our cohort of patients, we found a significant improvement in PFS ratio regardless of cut-point and an improvement in median PFS for the genomically directed group. These findings are congruent with those reported to date and support the evolving body of literature that demonstrates superior outcomes for patients who receive genomically guided therapy [23-25]. Moving forward, programs like ASCO’s Targeted Agent and Profiling Utilization Registry (TAPUR) study (http://www.asco.org/practice-research/targeted-agent-and-profiling-utilization-registry-study) will likely facilitate the speed with which the cancer genomics community is able to determine those target/drug combinations with the most utility and simultaneously those that will be futile. Further, the complex interaction between successful target/drug combination and site of disease can be further elucidated with combined data from multiple collaborating groups. Given the emerging body of consistent data demonstrating similar relative benefits to those patients who have access

![Figure 4: Details of responders.](image)
to targeted sequencing, this should be discussed with appropriate patients as a therapeutic avenue.

**MATERIALS AND METHODS**

**Study objectives**

The primary objective for the study was to compare the frequency of a PFS ratio ≥ 1.3 between patients who received the precision genomics recommended treatment and those who did not. Utilizing the most recent therapy selected by molecular profiling (MP) or clinicians choice of a patient’s tumor (Period B) with the PFS for the most recent therapy on which the patient had just experienced progression (Period A). If the (PFS of period B/PFS of period A) ratio was ≥ 1.3, then the tumor board MP-selected therapy was defined as beneficial for the patient. Secondary objectives included mean PFS ratio, PFS ratio ≥ 1, PFS ratio ≥ 1.5, and median PFS between groups.

**Study design**

The hypothesis for this study was that using a multidisciplinary tumor board to provide treatment recommendations based on each patient’s tumor MP data would favorably change the clinical course for an individual patient. This single institution, prospective, cohort study was conducted in patients with refractory metastatic cancer to compare the PFS of patients receiving a genomic-guided therapy to a cohort receiving a non-genomic guided therapy. The PFS ratio was determined by dividing the PFS of the new therapy (either the genomically guided or the non-genomically guided) by the PFS for the patient during their most recent regimen on which the patient had experienced progression. Patients were evaluated by the Indiana University Health Precision Genomics Program on a referral basis. The study was conducted in accordance with the Declaration of Helsinki and was approved by the Indiana University Institutional Review Board.

**Patient eligibility**

Eligibility criteria included the following: any histologic type of metastatic cancer; progression by RECIST criteria on at least one prior regimen for advanced disease; ability to undergo a biopsy or surgical procedure to obtain tumor or having recent tissue available; age ≥ 18 years; referred to the IUSCC Precision Genomics clinic between April 2014-October 2015; Eastern Cooperative Oncology Group performance status of 0 to 2; measurable or evaluable disease; resistance to last line of therapy (documented disease progression under last treatment; discontinued last treatment for progression); and received treatment following Precision Genomic assessment. Patients were excluded if the immediate previous therapy was received on a Phase I or II clinical trial as this could positively bias the PFS ratios if the experimental agent was not active. All patients who did not go on to receive additional therapy (i.e. died or were referred for hospice with best supportive care) were removed from the analysis as to not bias against the non-genomically guided arm and to be able to calculate PFS ratios that can be compared to the genomically guided arm. Minimum follow-up was defined as either disease progression, or the minimum amount of time for a patient to achieve a PFS ratio of 1.3.

**Molecular profiling of patient samples**

Formalin-Fixed Paraffin Embedded (FFPE) blocks from tumor biopsies or surgical resections were sent to Paradigm Diagnostics (Ann Arbor, MI) for the PCDx assay. PCDx is a comprehensive next-generation sequencing test that has been designed to analyze most known clinically actionable genomic variations in cancer, including: mutations, gene-fusions/rearrangements, copy number variations, mRNA expression as well as protein expression by IHC. Differences in RNA expression were measured by comparison of tumor gene expression compared to a panel of lineage-matched normal tissues. Next-generation sequencing was performed on an Ion Torrent Personal Genome Machine, with an average coverage > 5,000X. All analyses were performed in a CLIA-certified laboratory. Of note, one patient in our study received testing using the FoundationOne test, whose methods are described elsewhere [26].

**Deliberation of genomic results**

A multidisciplinary tumor board of medical oncologists, genomic and pharmacogenomic scientists, oncology clinical pharmacists, an oncologist-ethicist, pathologists, and nurses reviewed the results of the genome analyses. The results were considered in the context of the patient’s prior treatment history, concomitant medications, comorbidities, and germline variants that might predict increased toxicity to a given therapy. Identified targets were chosen based on: first priority, DNA mutation or copy number variation or DNA fusions; second priority, IHC or FISH; and last priority, mRNA overexpression alone. Board recommendations were provided to the referring clinician and the patient was treated according to the treating physician’s standard-of-care usage of that agent or referred to an available clinical trial.
Statistical considerations and methods

Using estimates from prior published data [11], a sample size of 74 patients was required to test the hypothesis that the PFS ratio > 1.3 in 28% of patients receiving recommended MP treatment, with an α risk of 5% and a power of 80%, compared to the assumption that the rate of obtaining a PFS of >1.3 without MP guided treatment was 5%. Time-to-event data were summarized using the Kaplan-Meier method and compared using the log-rank test; continuous data were compared using student’s t-test; categorical data were compared using the chi-squared or Fisher’s exact test, as appropriate.

CONFLICTS OF INTEREST

The authors do not have any conflicts of interest.

GRANT SUPPORT

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Editorial note

This paper has been accepted based in part on peer-review conducted by another journal and the authors’ response and revisions as well as expedited peer-review in Oncotarget.

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