Integration of genomics and histology revises diagnosis and enables effective therapy of refractory cancer of unknown primary with PDL1 amplification

The Harvard community has made this article openly available. Please share how this access benefits you. Your story matters

Citation

Published Version
doi:10.1101/mcs.a001180

Citable link
http://nrs.harvard.edu/urn-3:HUL.InstRepos:29625966

Terms of Use
This article was downloaded from Harvard University’s DASH repository, and is made available under the terms and conditions applicable to Other Posted Material, as set forth at http://nrs.harvard.edu/urn-3:HUL.InstRepos:dash.current.terms-of-use#LAA
Integration of genomics and histology revises diagnosis and enables effective therapy of refractory cancer of unknown primary with \textit{PDL1} amplification

Stefan Gröschel,1,2,3,19 Martin Bommer,4,19 Barbara Hutter,3,5 Jan Budczies,5,7 David Bonekamp,8 Christoph Heining,1,2,3 Peter Horak,1,2,3 Martina Fröhlich,3,5 Sebastian Uhrig,3,5 Daniel Hübschmann,9,10,11 Christina Georg,1,3,12 Daniela Richter,1,3 Nicole Pfarr,13 Katrin Pfütze,1,3,12 Stephan Wolf,3,14 Peter Schirmacher,3,15 Dirk Jäger,16 Christof von Kalle,1,2,3,12 Benedikt Brors,3,5 Hanno Glimm,1,2,3 Wilko Weichert,13,17 Albrecht Stenzinger,15,18,19 and Stefan Fröhling1,2,3,19

1Department of Translational Oncology, National Center for Tumor Diseases (NCT) Heidelberg, German Cancer Research Center (DKFZ), Heidelberg 69120, Germany; 2Section for Personalized Oncology, Heidelberg University Hospital, Heidelberg 69120, Germany; 3German Cancer Consortium (DKTK), Heidelberg 69120, Germany; 4Klinikum am Eichert, Department of Hematology, Oncology and Infectious Diseases, Göppingen 73035, Germany; 5Division Applied Bioinformatics, DKFZ and NCT Heidelberg, Heidelberg 69120, Germany; 6Institute of Pathology, Charité University Hospital, Berlin 10117, Germany; 7DKTK, Berlin 10117, Germany; 8Department of Radiology, DKFZ, Heidelberg 69120, Germany; 9Division of Theoretical Bioinformatics, DKFZ, Heidelberg 69120, Germany; 10Department for Bioinformatics and Functional Genomics, Institute for Pharmacy and Molecular Biotechnology and BioQuant, Heidelberg University, Heidelberg 69120, Germany; 11Department of Pediatric Immunology, Hematology and Oncology, Heidelberg University Hospital, Heidelberg 69120, Germany; 12DKFZ, Heidelberg Center for Personalized Oncology (HIPO), Heidelberg 69120, Germany; 13Institute of Pathology, Klinikum rechts der Isar, Technische Universität München, Munich 81675, Germany; 14Genomics and Proteomics Core Facility, High Throughput Sequencing Unit, DKFZ, Heidelberg 69120, Germany; 15Institute of Pathology, Heidelberg University Hospital and NCT Heidelberg, Heidelberg 69120, Germany; 16Department of Medical Oncology, NCT Heidelberg, Heidelberg University Hospital, Heidelberg 69120, Germany; 17DKTK, Munich 80539, Germany; 18Department of Pathology, Center for Integrated Diagnostics, Massachusetts General Hospital, Harvard Medical School, Boston 02114, USA

Abstract Identification of the tissue of origin in cancer of unknown primary (CUP) poses a diagnostic challenge and is critical for directing site-specific therapy. Currently, clinical decision-making in patients with CUP primarily relies on histopathology and clinical features. Comprehensive molecular profiling has the potential to contribute to diagnostic categorization and, most importantly, guide CUP therapy through identification of actionable lesions. We here report the case of an advanced-stage malignancy initially mimicking poorly differentiated soft-tissue sarcoma that did not respond to multiagent chemotherapy. Molecular profiling within a clinical whole-exome and transcriptome sequencing program revealed a heterozygous, highly amplified \textit{KRAS} G12S mutation, compound-heterozygous \textit{TP53} mutation/deletion, high mutational load, and focal high-level amplification of Chromosomes 9p (including \textit{PDL1} [CD274] and \textit{JAK2}) and 10p (including \textit{GATA3}). Integrated analysis of molecular data and histopathology provided a rationale for immune checkpoint inhibitor (ICI) therapy with pembrolizumab, which

\textcopyright{} 2016 Gröschel et al. This article is distributed under the terms of the Creative Commons Attribution-NonCommercial License, which permits reuse and redistribution, except for commercial purposes, provided that the original author and source are credited.
resulted in rapid clinical improvement and a lasting partial remission. Histopathological analyses ruled out sarcoma and established the diagnosis of a poorly differentiated adenocarcinoma. Although neither histopathology nor molecular data were able to pinpoint the tissue of origin, our analyses established several differential diagnoses including triple-negative breast cancer (TNBC). We analyzed 157 TNBC samples from The Cancer Genome Atlas, revealing PDL1 copy number gains coinciding with excessive PDL1 mRNA expression in 24% of cases. Collectively, these results illustrate the impact of multidimensional tumor profiling in cases with nondescript histology and immunophenotype, show the predictive potential of PDL1 amplification for immune checkpoint inhibitors (ICIs), and suggest a targeted therapeutic strategy in Chromosome 9p24.1/PDL1-amplified cancers.

[Supplemental material is available for this article.]

INTRODUCTION

“Omics” technologies, most prominently next-generation sequencing (NGS), have entered clinical medicine in the last decade and are expected to affect the standard of care in oncology. However, the identification of patients who truly benefit from genomics-driven approaches to clinical management and the appropriate timing of therapy are still challenging (Roychowdhury and Chinnaiyan 2014). Moreover, it is currently unclear how to effectively integrate these new tools with conventional technologies such as standard histopathology, and the algorithms for integrated diagnostics and molecularly guided treatment remain to be exactly defined.

Comprehensive cancer profiling can particularly assist diagnosis and treatment planning in patients with malignancies of unresolved histology and in metastatic disease, in which the tissue of origin of the primary tumor cannot be discerned. Cancers of unknown primary (CUP) constitute 3%-5% of all cancer diagnoses and rank fourth in the causes of cancer-related deaths worldwide (Varadhachary and Raber 2014). NGS-based molecular characterization holds the promise to gain insight into the pathobiology of these difficult-to-classify tumors. Compared to targeted sequencing of selected cancer genes, whole-exome sequencing (WES) and RNA sequencing (RNA-seq) allow for superior detection of coding sequence variants, amplifications, deletions, and structural rearrangements of chromosomes as well as their consequences on mRNA transcription. Integration of these data with mutational and transcriptional cancer profiles deposited in public genome databases, such as those generated by The Cancer Genome Atlas (TCGA) Research Network (http://cancergenome.nih.gov) and the International Cancer Genome Consortium (https://icgc.org), can aid in identifying the tissue of origin of cancers of otherwise indefinable histology and nominate candidate therapeutic targets.

Here we describe a patient with refractory, metastatic anaplastic CUP, where integrated analysis of histopathology, immunohistochemistry (IHC), fluorescence in situ hybridization (FISH), WES, and RNA-seq revised the diagnosis and facilitated genotype-informed treatment with an immune checkpoint inhibitor (ICI) that resulted in a durable, very good partial remission.

RESULTS

A 44-yr-old Caucasian female presented to our center with metastatic CUP. Two years before presentation, she had been diagnosed with multiple metastatic lesions in the mesentery of
the small intestines and the duodenojejunal flexure. At that time, the tumor lesions had been resected completely. Histopathology analyses were inconclusive, showing a high-grade neoplasm with pleomorphic tumor cells that stained positive for vimentin. MIB-1 was positive in ~30% of the tumor cells. Following central pathology review, a preliminary diagnosis of CUP with sarcomatoid features was considered with the most likely differential diagnosis of sarcoma, not otherwise specified. The patient was treated with four cycles of doxorubicin and ifosfamide according to the EORTC (European Organisation for Research and Treatment of Cancer) 62012 protocol (Judson et al. 2014) at an outside institution, and subsequent imaging studies revealed no evidence of disease.

Sixteen months after initial diagnosis, the patient developed progressive pain in the right shoulder, and magnetic resonance imaging (MRI) showed a mass of 4 × 2 × 2 cm in the right deltoid muscle and a second metastatic lesion of 1.5 × 2 × 2 cm in the right axilla. Incisional biopsy was performed, and subsequent histopathology assessment with repeat central review suggested metastasis of the previously diagnosed CUP rather than a secondary malignancy. After complete surgical excision of the rapidly progressing lesions, adjuvant radiotherapy was administered to the right shoulder and axilla.

Twenty-one months after initial diagnosis, follow-up MRI studies revealed five new tumor formations (left paravertebral region, right M. infraspinatus, right axilla, right M. latissimus dorsi, left chest wall). Again, surgical resection of multiple lesions was performed. However, the tumor progressed rapidly with new lesions of the right shoulder and thigh; treatment with trabectedin was initiated but had to be stopped because of progressive disease and Grade IV thrombocytopenia.

In an attempt to refine the diagnosis and inform potential therapeutic strategies, the patient was enrolled in NCT (Nationales Centrum für Tumorerkrankungen) MASTER (Molecularly Aided Stratification for Tumor Eradication Research), a cross-entity molecular stratification program for younger adults with advanced-stage cancer and patients with rare tumors (Kordes et al. 2016), and a cryopreserved biopsy of a metastatic shoulder lesion with a tumor cell content of >90% was analyzed by WES and RNA-seq. Peripheral blood mononuclear cells served as a germline control.

WES analysis revealed a high mutational load of 341 nonsynonymous single-nucleotide variants, 39 insertion/deletions, and multiple chromosomal gains and losses (Fig. 1). Known driver mutations were detected in TP53 (p.E135fs) and KRAS (p.G12S), the latter also being highly expressed on the RNA level because of a focal amplification on Chromosome 12p (Table 1). Mutations of unknown significance occurred in other cancer-related genes (e.g., PIK3CD, CDKN2A, NCOA1, FAT2, EGFR, MSH3, ARID1A, MDC1, SETD1A, SETD3, and TET1) (Supplemental Table S1). A focal high-level amplification on Chromosome 9p encompassed JAK2 and PDL1 (also known as CD274 or B7-H1) together with other genes implicated in carcinogenesis (Fig. 1; Kietz et al. 2009; Moon et al. 2011; Zhang et al. 2012; Hoffman et al. 2014; Jovanovic et al. 2014; Kim et al. 2015).

In line with these findings, IHC and FISH analyses showed high-level amplification and subsequent overexpression of PDL1 in the tumor cells. Histologic assessment showed substantial peritumoral infiltration of lymphocytes (Fig. 2A). Correspondingly, PDL1 mRNA levels, as determined by RNA-seq, were highest in this tumor specimen compared to all tumor samples included in NCT MASTER (n = 266; Fig. 2B). A query of publicly available databases for matching cancer mutational profiles comprising TP53 inactivation and amplification of Chromosomes 9p (PDL1, JAK2), 10p (GATA3), and 12p (mutant KRAS) as well as review of the literature suggested lung adenocarcinoma, triple-negative breast cancer (TNBC), or gastric adenocarcinoma as the most probable diagnoses (Cimino-Mathews et al. 2013; Gao et al. 2013; Marquard et al. 2015). Principal component analysis using mRNA expression profiles obtained by RNA-seq data from both TCGA and the NCT MASTER cohorts was inconclusive (data not shown). Based on these findings, we decided to challenge the initial
Figure 1. PDL1 and GATA3 amplification in a patient with cancer of unknown primary. (Upper panel) Copy number plot showing chromosomal coordinates computed as a set of regions based on whole-exome sequencing data (x-axis) and the log2 ratio of copy number changes (y-axis). Color codes of alternating green and black regions indicate segmentation between chromosomes. (Lower panel) Amplified region on chromosome 9p.23–24.1.

Table 1. Somatic variants of potential interest

<table>
<thead>
<tr>
<th>Chr</th>
<th>GRCh37 position</th>
<th>Ref</th>
<th>Var</th>
<th>Type</th>
<th>Gene</th>
<th>HGVS DNA reference</th>
<th>Predicted effect</th>
<th>HGVS protein reference</th>
<th>Genotype</th>
<th>dbSNP ID</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>27099877</td>
<td>CA</td>
<td>C</td>
<td>FS-DEL</td>
<td>ARID1A</td>
<td>g.27099877CA&gt;C</td>
<td>p.N1253fs</td>
<td>p.Asn1253fs</td>
<td>het</td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>9770641</td>
<td>G</td>
<td>A</td>
<td>SNV</td>
<td>PIK3CD</td>
<td>g.9770641G&gt;A</td>
<td>p.S43N</td>
<td>p.Ser43Asn</td>
<td>het</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>24991253</td>
<td>C</td>
<td>A</td>
<td>SNV</td>
<td>NCOA1</td>
<td>g.24991253C&gt;A</td>
<td>p.T1440N</td>
<td>p.Thr1440Asn</td>
<td>het</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>80160762</td>
<td>G</td>
<td>A Splicing</td>
<td>MSH3</td>
<td>g.80160762G&gt;A</td>
<td></td>
<td></td>
<td></td>
<td>het</td>
<td></td>
</tr>
<tr>
<td>6</td>
<td>150922306</td>
<td>C</td>
<td>G</td>
<td>SNV</td>
<td>FAT2</td>
<td>g.150922306C&gt;G</td>
<td>p.R2794S</td>
<td>p.Arg2794Ser</td>
<td>het</td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>55225447</td>
<td>G</td>
<td>T Splicing</td>
<td>EGFR</td>
<td>g.55225447G&gt;T</td>
<td></td>
<td></td>
<td></td>
<td>het</td>
<td></td>
</tr>
<tr>
<td>9</td>
<td>1461199–14398642</td>
<td>AMP PPAPDC2;PTPRD</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>9</td>
<td>21971208</td>
<td>C</td>
<td>A Splicing</td>
<td>CDKN2A</td>
<td>g.21971208C&gt;A</td>
<td></td>
<td></td>
<td></td>
<td>het</td>
<td></td>
</tr>
<tr>
<td>10</td>
<td>70332663</td>
<td>C</td>
<td>G</td>
<td>SNV</td>
<td>TET1</td>
<td>g.70332663C&gt;G</td>
<td>p.Q190E</td>
<td>p.Gln190Glu</td>
<td>het</td>
<td></td>
</tr>
<tr>
<td>10</td>
<td>93076–14013840</td>
<td>AMP TUBB8; PRPF18</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>12</td>
<td>19242142–30289130</td>
<td>AMP</td>
<td>C12orf39; RASSF8</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>12</td>
<td>25398285</td>
<td>C</td>
<td>T</td>
<td>SNV</td>
<td>KRAS</td>
<td>g.25398285C&gt;T</td>
<td>p.G12S</td>
<td>p.Gly12Ser</td>
<td>het rs1219135</td>
<td></td>
</tr>
<tr>
<td>16</td>
<td>30977120</td>
<td>G</td>
<td>C</td>
<td>SNV</td>
<td>SETD1A</td>
<td>g.30977120G&gt;C</td>
<td>p.G640R</td>
<td>p.Gly640Arg</td>
<td>het</td>
<td></td>
</tr>
<tr>
<td>17</td>
<td>7577057</td>
<td>T</td>
<td>C FS-INS</td>
<td>TP53</td>
<td>g.7577057T&gt;TC</td>
<td></td>
<td></td>
<td></td>
<td>p.E135fs</td>
<td>p.Glu135fs</td>
</tr>
</tbody>
</table>

The complete list of somatic mutations can be found in Supplemental Table S1.

HGVS, Human Genome Variation Society; dbSNP, Database for Short Genomic Variations; AMP, amplification; FS-DEL, frameshift-deletion; FS-INS, frameshift-insertion; het, heterozygous; SNV, single-nucleotide variant.
diagnosis of a metastatic sarcoma and reevaluated tissue parameters of the current metastatic lesion. Comprehensive immunohistochemical profiling of 25 different antigens revealed strong expression of cytokeratins (as measured by AE1/3) and CK7. These data are in line with decisive histological features (e.g., focal formation of small tubular structures) and focal positivity with periodic acid–Schiff (PAS) staining leading to the diagnosis of an adenocarcinoma. We also noticed focal positivity of vimentin, which is consistent with poor differentiation and epithelial–mesenchymal transition from a histological and biological point of view, respectively. Tumor cells showed strong GATA3 expression (Supplemental Fig. 1) consistent with the genetic amplification and were negative for estrogen receptor (ER), progesterone receptor (PR), and ERBB2 (also known as HER2). In the context of poor differentiation, negativity for TTF1 and napsin does not rule out the differential diagnosis of lung adenocarcinoma but is not supportive either. Although neither IHC profiling nor molecular analyses allowed us to unequivocally pinpoint the tissue of origin, reevaluation of all parameters suggested metastatic TNBC among the top differential diagnoses, although a definitive categorization based on conventional diagnostic criteria of this CUP case still remained tentative, given the clinical presentation without a primary tumor detectable in either breast, lung, or stomach.

Figure 2. PDL1 protein expression, PDL1 amplification, peritumoral lymphocyte infiltration, and PDL1 mRNA expression in a patient with anaplastic cancer of unknown primary. (A) (Left) Photomicrograph showing PDL1 protein expression in a metastasis (scale bar, 100 µm); (middle) representative fluorescence in situ hybridization signal pattern showing amplification of the PDL1 locus (green signals) relative to the centromere of Chromosome 9 (red signals); (right) CD4 lymphocyte staining (scale bar, 200 µm). (B) Ranking of 266 patient samples analyzed in the NCT MASTER study according to PDL1 (CD274) mRNA levels, as determined by RNA-seq. The red bar indicates the described index patient. RPKM, reads per kilobase of exon model per million mapped reads.
In similarity to our case, immune cell infiltrates and Chromosome 9p amplification have recently been described in a small cohort of TNBC (Barrett et al. 2015). We thus aimed to systematically address the question whether genomic PDL1 gains are recurrent in TNBC and associated with elevated PDL1 expression as observed in our case. To this end, we analyzed PDL1 copy number variation in a TCGA cohort of 937 breast carcinomas (Fig. 3A). Copy number gains were more frequent in TNBC (43.3%) compared to hormone receptor (HR)+/HER2−, HR+/HER2+, and HR−/HER2+ breast cancer (10.9%, 10.3%, and 26.3%; p = 0.00045). Focal amplifications occurred in 23.6% of TNBC samples compared to 1.3%, 2.8%, and 10.5% in HR+/HER2−, HR+/HER2+, and HR−/HER2+ breast cancer specimens. PDL1 mRNA expression analysis was performed in 934 tumors (Fig. 3B) and revealed significantly higher PDL1 levels in tumors with PDL1 copy number gains compared to tumors that were diploid for the PDL1 locus (fold change, 1.25; p = 0.019). Furthermore, triple-negative tumors had significantly increased PDL1 expression compared to HR+/HER2− and HR+/HER2+ tumors (fold change, 1.49; p = 0.00030 and fold change, 1.42; p = 0.0068).

Both amplification and overexpression of PDL1 and the hypermutated tumor genome provided a strong rationale for ICI treatment with the anti-PD1 monoclonal antibody pembrolizumab. The patient received pembrolizumab at a dose of 2 mg/kg body weight for repeated cycles of 3 wk starting from month 24 after initial diagnosis. At the start of treatment, baseline positron emission tomography/computed tomography (PET/CT) imaging showed widely disseminated tumor manifestations (Fig. 4A). Treatment was well tolerated and could be administered without serious side effects. After 2 mo of therapy, PET imaging showed evidence of tumor regression in most lesions while a left gluteal and a left axillary lesion increased in size and metabolic activity, likely representing an immune-mediated phenomenon. Clinically, performance status was improving rapidly, and tumor-related symptoms such as pain and motoric impairment of the left arm decreased. Six months after initiation of pembrolizumab, PET imaging results suggested a near-complete remission, formally qualifying as a very good partial remission according to Response Evaluation

![Figure 3](image-url). PDL1 DNA copy number and PDL1 mRNA expression in the molecular subtypes of breast cancer (TCGA data sets). (A) PDL1 copy number gains are more frequent in triple-negative breast cancer (TNBC) (43.3%) compared to HR+/HER2−, HR+/HER2+, and HR+/HER2− breast cancer (26.3%, 10.9%, and 10.3%, respectively). (B) PDL1 mRNA expression is higher in TNBC compared to HR+/HER2− and HR+/HER2+ breast cancer (fold change, 1.49; p = 0.00030 and fold change, 1.42; p = 0.0068). In the beeswarm plot, each colored dot represents a tumor. The horizontal red lines indicate the first quartile, the median, and the third quartile.
The patient has remained in continuous near-complete remission and free of disease-specific symptoms while being on immunotherapy at the time this report was written (14 mo after commencement of pembrolizumab).

**DISCUSSION**

“Multi-omic” approaches and integrative analyses allow tissue-of-origin molecular profiling and can uncover new diagnostic and therapeutic opportunities in patients with high-risk and
refractory cancers (Roychowdhury and Chinnaiyan 2014). Matching the genetic data obtained in individual patients to the rapidly expanding public repositories of genome, exome, and transcriptome data from large cohorts of well-annotated cancers can inform histopathologic assessment, including characteristic tissue-of-origin IHC and FISH markers, and facilitate the identification of the primary cancer site in difficult cases. In our patient, molecular analysis and histopathology suggested several possible tissues of origin, but a definitive diagnosis could not be reached. This caveat may in part be explained by the limitations associated with comparison of molecular profiles of metastatic tissue with data sets obtained in treatment-naïve tumors, such as those available from TCGA.

Besides its diagnostic implications, WES can identify driver genomic alterations that are missed by interrogating single genes or a panel of cancer-related genes but may represent targets for individualized treatment. Importantly, tissue context may modulate the efficacy of therapies directed against specific molecular lesions, as shown, for example, by the variable responses to vemurafenib seen among different tumor entities with BRAF V600E mutations (Samuel et al. 2014; Corcoran et al. 2015; Hyman et al. 2015; Kopetz et al. 2015). Hence, responses can be difficult to predict, particularly to drugs not developed for a given indication (Le Tourneau et al. 2015). Despite this purported predicament, targeted therapies have become the mainstay of treatment in a range of malignancies (e.g., gastrointestinal stromal tumor, chronic myeloid leukemia, or non-small-cell lung cancer), underscoring the utility of personalized genomic medicine and the incorporation of “omics” data in the clinical decision algorithm on an individual-case basis in patients otherwise refractory to conventional treatment modalities (Tothill et al. 2013; Knoechel et al. 2015; Schwaederle et al. 2015; Kordes et al. 2016). Our diagnostic approach ruled out sarcoma and rendered the diagnosis of poorly differentiated adenocarcinoma. We were, however, unable to trace the definite tissue of origin even when employing both molecular and conventional pathology. This result may in part be explained by the fact that our case is a patient with metastatic CUP who underwent several lines of treatment before molecular profiling, a setting that hampers direct comparison with molecular profiles of treatment-naïve tumors, such as those deposited in the TCGA database. Furthermore, novel data-sharing initiatives, such as AACR (American Association for Cancer Research) Project GENIE (Genomics, Evidence, Neoplasia, Information Exchange; http://www.aacr.org/Research/Research/Pages/aacr-project-genie.aspx), as well as innovative interventional clinical trials, such as NCI-MATCH (National Cancer Institute-Molecular Analysis for Therapy Choice; http://www.cancer.gov/about-cancer/treatment/clinical-trials/nci-supported/nci-match), are expected to broaden the basis for greater individualization of therapy, particularly in the case of rare cancers and rare mutations in common cancers.

In our patient, the finding of PDL1 amplification and overexpression in conjunction with the high mutational load, and the failure of previous conventional chemotherapeutic approaches, prompted us to administer off-label immunotherapy. Based on our patient’s favorable clinical course, our exploration of potential differential diagnoses, and an analysis of the TCGA data set on TNBC, we speculate that ICIs might be a new treatment modality in this aggressive and clinically challenging tumor type (Buisseret et al. 2015; Le et al. 2015). PD1-inhibitory agents are currently being investigated in TNBC in phase II/III trials that are prospectively collecting data on tumoral PDL1 expression, allowing correlative evaluation of PDL1 copy number changes as a predictive marker (Sharma and Allison 2015; Topalian et al. 2015). The mechanistic rationale underlying PD1 blockade is to reverse tumor immune evasion mediated by PDL1-PD1 engagement of the tumor with antitumor CD8+ T cells, which consequently become anergic. The choice of immunotherapy in our patient was supported by the high density of tumor-infiltrating lymphocytes, PDL1 amplification resulting in strong intratumoral PDL1 expression both on the RNA and protein level, and a high mutational load, which was reported as being predictive for
response to ICI in a range of cancers (Garon et al. 2015; Le et al. 2015; Robert et al. 2015). Although future studies are needed to further establish robust and reliable determinants of response to ICI (Carbognin et al. 2015; Patel and Kurzrock 2015), PDL1 amplification might serve as a rapidly available and inexpensive surrogate marker across histologic entities, as supported by the striking therapeutic activity of PD1 blockade in TNBC (this study) and refractory classical Hodgkin lymphoma (Ansell et al. 2015; Roemer et al. 2016) characterized by Chromosome 9p24.1 gain. In a recent pan-cancer analysis, PDL1 gains were shown to occur in many cancer types and to result in PDL1 mRNA overexpression in some tumors (Budczies et al. 2016). Moreover, a PDL1 core amplification region of 7.8 Mb on Chromosome 9p that extended to the telomere was identified, consistent with the finding in this report. JAK2 is coamplified in this region (Balko et al. 2016), which is of potential interest, because JAK2 has been reported previously as a positive regulator of PDL1 and PDL2 genes (Green et al. 2010). Inactivating JAK2 mutations have recently been implicated in acquired resistance to PD1 inhibitor treatment and immune evasion of cancer cells, possibly related to down-regulation of PDL1 and antigen presentation as a consequence of abolished interferon signaling due to loss-of-function alterations in JAK2 (Zaretsky et al. 2016).

In summary, our observations show that clinical NGS can uncover new therapeutic targets and may eventually help to change the management of a considerable number of patients with CUP. The steadily decreasing turnaround time of WES and even whole-genome analysis, declining sequencing costs, and continuous workflow standardizations create a strong imperative to implement this methodology for the diagnostic workup of CUP patients. Furthermore, our experience illustrates how integration of different methods can optimize diagnostics and identify targets for intervention especially in cases in which it is difficult to arrive at a consensus diagnosis by conventional means.

**METHODS**

**WES, RNA-seq, and Bioinformatic Analysis**

Exome capture was performed using SureSelect Human All Exon V5+UTRs in-solution capture reagents (Agilent). One and one-half micrograms of genomic DNA was sheared sonically to 150–200 bp (paired-end) insert size with a Covaris S2 device. Two hundred and fifty nanograms of Illumina adapter-containing library was hybridized with exome baits at 65°C for 16 h. Paired-end sequencing (101 bp) was performed with a HiSeq 2500 instrument (Illumina) in a rapid mode. RNA-seq libraries were prepared using the TruSeq RNA Sample Preparation Kit v2 (Illumina). Briefly, mRNA was purified from 1 µg total RNA using oligo(dT) beads, poly(A)⁺ RNA was fragmented to 150 bp and converted into cDNA, and cDNA fragments were end-repaired, adenylated on the 3’ end, adapter-ligated, and amplified with 12 cycles of polymerase chain reaction. The final libraries were validated using a Qubit 2.0 Fluorometer (Life Technologies) and a Bioanalyzer 2100 system (Agilent). Paired-end sequencing (2 × 101 bp) was performed with a HiSeq 2500 instrument (Illumina) in a rapid mode. Sequencing data were analyzed using a previously reported bioinformatics workflow (Kordes et al. 2016; see Table 2 for coverage metrics).

**Triple-Negative Breast Cancer Data Set**

Breast cancer data sets (study brca_tcga_pub2015) including data on DNA copy number status (GISTIC) and mRNA expression (RNA Seq V2 RSEM) of PDL1 were retrieved from the cBioPortal for Cancer Genomics (http://www.cbioportal.org/public-portal) and analyzed as described previously (Budczies et al. 2016). Immunohistochemistry data
for ER, PR, and HER2 as well as FISH data for HER2 were retrieved from the TCGA data portal (https://tcga-data.nci.nih.gov). The study cohort comprised 937 breast carcinomas with available copy number data. Cases were classified into molecular subtypes: 597 (63.7%) HR+/HER2− tumors, 145 (15.5%) HR+/HER2+ tumors, 38 HR−/HER2+ tumors (4.1%), and 157 (16.8%) HR−/HER2− tumors. RNA-seq data were available for 934 of these tumors.

Immunohistochemistry and Fluorescence In Situ Hybridization

Immunohistochemical staining was performed on formalin-fixed and paraffin-embedded whole-tissue slides using a BenchMark XT device (Ventana Medical Systems) according to quality-controlled standard procedures. After antigen retrieval, the following primary antibodies were used: AE1/3: 1:100 dilution, DAKO (#M3515); GATA3: ready to use, Ventana (#760-4897); CK7: 1:50 dilution, DAKO (#M7018); PDL1: 1:100 dilution, Spring (#07309457001); CD4 mouse monoclonal antibody: 1:20 dilution, Novocastra/Leica (clone 1F6); napsin: 1:400 dilution, Novocastra/Leica (clone IP64, #NCL-L-Napsin A); TTF1: 1:100 dilution, Novocastra/Leica (clone SPT24, #NCL-L-TTF1). Either 3,3′-diaminobenzidine peroxide substrate or 3-amino-9-ethylcarbazole served as chromogens.

FISH using a commercially available dual color probe for the PDL1 (CD274) gene (Zytovision; # Z-2179–200) was performed on whole slides according to the manufacturer’s instructions. Four-micrometer-thick sections were cut, mounted on SuperFrost slides, and deparaffinized. The PDL1 probe was labeled with a green fluorochrome and the classical satellite III region of Chromosome 9 (CEN9, D9Z3) was labeled with an orange fluorochrome.

ADDITIONAL INFORMATION

Data Deposition and Access

WES and RNA-seq data derived from the patient specimen have been deposited at the European Genome-phenome Archive (EGA; https://www.ebi.ac.uk/ega/), which is hosted by the European Bioinformatics Institute (EBI), under accession number EGAS00001001846. Point substitution variants were submitted to the Catalogue of Somatic Mutations in Cancer (COSMIC; http://cancer.sanger.ac.uk/cosmic, identifier COSP41747).

Ethics Statement

Patient tissue samples were collected after informed consent was obtained and provided by the NCT Heidelberg Tissue Bank under protocol NCT MASTER, S-206/2011 in accordance with its regulations and after approval by the Ethics Committee of Heidelberg University. Specifically, patients provided written informed consent including permission of tissue banking of tumor and germline specimens and analysis by next-generation sequencing.
technology (i.e., whole-exome sequencing and RNA sequencing and ancillary molecular and histopathological methodologies for validation of results).

Acknowledgments

The authors thank the DKFZ-HIPO and NCT Precision Oncology Program (POP) Sample Processing Laboratory, the DKFZ Genomics and Proteomics Core Facility, and the DKFZ-HIPO Data Management Group for technical support and expertise. We also thank Katja Beck, Karolin Willmund, Roland Eils, and Peter Lichter for infrastructure and program development within DKFZ-HIPO and NCT POP.

Author Contributions


Funding

This work was supported by grants H021 and H063 from DKFZ-HIPO.

REFERENCES


