Genome Research

Whole-genome and multisector exome sequencing of primary and post-treatment glioblastoma reveals patterns of tumor evolution


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Glioblastoma (GBM) is a prototypical heterogeneous brain tumor refractory to conventional therapy. A small residual population of cells escapes surgery and chemoradiation, resulting in a typically fatal tumor recurrence ~7 mo after diagnosis. Understanding the molecular architecture of this residual population is critical for the development of successful therapies. We used whole-genome sequencing and whole-exome sequencing of multiple sectors from primary and paired recurrent GBM tumors to reconstruct the genomic profile of residual, therapy resistant tumor initiating cells. We found that genetic alteration of the p53 pathway is a primary molecular event predictive of a high number of subclonal mutations in glioblastoma. The genomic road leading to recurrence is highly idiosyncratic but can be broadly classified into linear recurrences that share extensive genetic similarity with the primary tumor and can be directly traced to one of its specific sectors, and divergent recurrences that share few genetic alterations with the primary tumor and originate from cells that branched off early during tumorigenesis. Our study provides mechanistic insights into how genetic alterations in primary tumors impact the ensuing evolution of tumor cells and the emergence of subclonal heterogeneity.

[Supplemental material is available for this article.]

The presence of multiple cancer cell clones within a single tumor has been explained as a Darwinian process in which different clones compete for limited resources, and the most phenotypically fit cells eventually prevail (Greaves and Maley 2012; Yates and Campbell 2012; Aparicio and Caldas 2013). It has been suggested that such heterogeneity allows a tumor to respond to local and systemic selective pressures, such as those exerted by therapeutic interventions (Nowak and Sigmund 2004; Greaves and Maley 2012; Bozic et al. 2013). For example, the presence of subclonal driver mutations in cancer cells was indicative of rapid disease progression in chronic lymphocytic leukemia (Landau et al. 2013). Using single-cell sequencing or massively parallel sequencing, clonal architectures ranging from complex polyclonal structures to monoclonal tumors have been described in cancer lineages such as those of the breast, kidney, and blood (Navin et al. 2011; Ding et al. 2012; Shah et al. 2012; Landau et al. 2013; Gerlinger et al. 2014). Distinct subclonal tumor cell populations relating to mosaic amplification of receptor tyrosine kinases were reported in glioblastoma (GBM), suggesting a similarly dynamic architecture for this disease (Snunder et al. 2011; Nickel et al. 2012; Sztupl et al. 2012; Sottoriva et al. 2013).

GBM is the most common malignant brain tumor in adults (Van Meir et al. 2010; Dunning et al. 2012) and is standardly treated with surgical resection followed by concomitant radiotherapy and administration of the alkylating agent temozolomide (TMZ) (Stupp et al. 2005). Despite this aggressive treatment regimen, the median time
to disease recurrence is 6.9 mo, with >90% of GBM tumors recurring at the original site (Wen and Kesari 2008). Therapy targeting the epithelial growth factor receptor variant III (EGFRvIII) led to an improved overall survival time among patients with GBM; however, 82% of these patients lost EGFRvIII expression when the tumor recurred, which suggests a competitive advantage for non-EGFRvIII expressing clones in these tumors (Sampson et al. 2010). Achieving a better understanding of the clonal structure of cancer cells is thus of vital importance and may inform the development of additional targeted therapies for rapidly lethal forms of cancer, such as GBM.

Here, we analyzed genomic profiles of 252 GBM samples from The Cancer Genome Atlas (TCGA) (Brennan et al. 2013), and 60 biopsies taken from 23 pairs of pre- and post-treatment GBMs, to understand (1) the intratumoral clonal compositions of primary GBM; and (2) how GBM responds to therapeutic intervention. Our results provide a molecular portrait of GBM recurrence.

Results

Sample characteristics and mutation calling

In this study, we performed an analysis of genomic data from 252 untreated GBM samples from The Cancer Genome Atlas (cohort I). To study tumor responses to treatment, we obtained a second cohort of tumor samples, for which we collected pairs of primary and first recurrent GBM samples from 21 patients and added pairs of secondary GBM and next disease occurrence samples from two patients (cohort II). Prior to disease recurrence, 21 patients in cohort II had received radiotherapy, and 17 of them had also received adjuvant TMZ. A variety of treatments, including carmustine and anti-inflammatory agents, were administered to the remaining patients in cohort II. An IDH1 R132 mutation was detected in two cases. The clinical data for cohort II is summarized in Supplemental Table 1.

Integrative analysis identifies clonal and subclonal mutations

To investigate the clonal architecture of GBM, we classified somatic mutations into clonal and subclonal categories by integrating variant allele fraction, DNA copy number, genotype, and tumor purity (Methods). We used PyClone, a Bayesian clustering method that simultaneously estimates the distribution of the cellular frequency for each mutation (Roth et al. 2014). After correcting for tumor cell content, the cellular frequency of each mutation was used to infer the mutation clonality. By definition, clonal mutations occur early in or before malignant transformation and thus are present in all tumor cells. In contrast, subclonal mutations occur later in tumor expansion and are present in only a subset of tumor cells. Of 17,636 somatic single nucleotide variations (sSNVs) detected in cohort I (Brennan et al. 2013), we found 67.9% to be clonal ($\mu = 11,973$, median across samples; 68.3% ± 19.3%) and 29.8% ($\mu = 5249$) to be subclonal. A minor fraction of mutations could not be classified (2.3%). To validate our classification approach, we presumed that clonal mutations were present in all tumor compartments and thus should be spatially ubiquitous. We sequenced the exomes of two nonoverlapping sectors from 13 tumors in cohort II, classified detected mutations as clonal or subclonal, and compared both types of sSNVs with spatial distribution. The percentage of ubiquitous mutations (present in both tumor sectors) per sample ranged from 43.1% to 74.7% (59.3.5% ± 10%) (Supplemental Fig. 1A), and the majority of ubiquitous mutations (85.6%) were classified as clonal. In contrast, 41.2% of the private mutations were classified as clonal (Fig. 1A; Supplemental Table 2), which was a significant difference ($P = 2.2 \times 10^{-16}$, $x^2$ test). In a single recurrent tumor sample that had three biopsies available, we found a similar percentage of ubiquitous mutations (45.3%) (Supplemental Fig. 1B). Separating patients into discrete age groups by intervals of 10 yr, we found a significant linear correlation between clonal mutations and age ($P = 3.69 \times 10^{-7}$, multiple group analysis of variance) (Fig. 1B), suggesting that the majority of clonal mutations were acquired before gliomagenesis and accumulated over the life span of the patient at time of diagnosis. This observation was corroborated by the dominant presence of C > T transitions, reminiscent of germline substitutions, in the clonal mutation spectrum (Supplemental Fig. 2A; Nik-Zainal et al. 2012; Alexandrov et al. 2013; Tomasetti et al. 2013). We did not observe a correlation between age and subclonal mutations ($P = 0.62$). We evaluated the clonal status of significantly mutated GBM genes and recurrent cancer genes from the cancer gene census (nonsilent mutation frequency greater than three in the cohort) (Supplemental Fig. 2B). Very few genes were entirely clonal or subclonal (Andor et al. 2014). However, the majority (90.5%) of TP53 and PIK3CA/PIK3R1 mutations were clonal, suggesting a founder role for these events in GBM. In contrast, receptor tyrosine kinases such as EGFR, PDGFRα, and AKT pathway regulator PTEN were more evenly distributed.

![Figure 1](https://www.genome.org)
between the clonal and subclonal mutation groups across cohort I, suggesting heterogeneous deregulation timing.

Subclonal mutations are associated with p53 pathway alterations

To identify genetic aberrations that are associated with subclonal progression, we juxtaposed somatic mutations, DNA copy number alterations, and clonal compositions in cohort I. Interestingly, deregulation of the p53 pathway, in particular TP53 somatic mutations (n = 78) and MDM2 amplifications (n = 21), was strongly associated with an increased fraction of subclonal mutations (P = 6 × 10^-3, Wilcoxon rank-sum test) (Fig. 2A), and this association was independent of age (Methods). G-CIMP positive tumors, a subclass of GBM characterized by a hypermethylator phenotype, mutations in isocitrate dehydrogenase I (IDH1), and mutations in TP53 (Noushmehr et al. 2010), accordingly showed a high proportion of subclonal mutations. Since subclonal mutations were associated with unfavorable outcome in chronic lymphocytic leukemia (Landau et al. 2013), we compared the event-free survival of patients represented in our data set whose primary tumor had a dominance of clonal mutations to patients with tumors in which high fractions of subclonal mutations were detected. Interestingly, the subclonal tumor group showed significantly longer event-free survival than the clonal tumor group (P = 0.025, log-rank test), but only when limiting the analysis to patients younger than 55 yr of age at time of diagnosis (Fig. 2B; Supplemental Fig. 3). TP53 mutation status alone did not correlate with outcome (data not shown). The increased fraction of subclonal mutations in GBM harboring p53 pathway alterations may be indicative of an elevated tolerance to DNA damage or apoptosis suppression (Offer et al. 2002; Kojima et al. 2005; Pant et al. 2013).

p53 pathway status affects mutational burden in recurrent tumor

To further examine the effect of TP53 mutations on tumor evolution, we performed exome sequencing and DNA copy number analysis on 23 pairs of first GBM and matched recurrent tumors in cohort II. In line with a recent report (Johnson et al. 2014), we found that the majority of primary GBM mutations could also be detected in the tumor after disease relapse (median overlap 67%; range 18%-85%) (Supplemental Table 3). Five recurrent tumors showed a very high number of clonal and/or subclonal sSNVs and were considered hypermutated (12–131 mutations per Mb). Despite the high overall population frequency of genetic alterations in TP53 (27.9%), EGFR (57.4%), and CDKN2A (57.8%) (Brennan et al. 2013), copy number alterations and sSNVs in these and other GBM driver genes were frequently present at time of diagnosis but absent in the tumor recurrence, mutated at a difference base, or deleted/amplified with different copy number breakpoints (Fig. 3A). It is therefore unlikely that any of these genes can be unambiguously flagged as an initiating mutation of IDH1 wild-type glioblastoma, and their frequent alterations can be best explained by intratumoral evolutionary pressures resulting in convergent evolutionary events. For example, we observed focal amplification of the Rb pathway regulator CDK4 in three primary GBM tumors but not in two of the three corresponding recurrent tumors (Fig. 3A). Interestingly, both cases had acquired a homozygous deletion of upstream Rb pathway regulator CDKN2A, suggesting a compensatory effect to maintain the deregulation of the Rb pathway. We speculate that one common pathway of tumor initiation starts with loss of 10q and gain of Chromosome 7, which predispose to deactivation of PTEN and activating mutations/amplifications in major oncogenes.

Mutations in TP53 were detected in seven primary GBMs, and in two cases, the DNA binding domain mutation was no longer detected in the recurrent tumor (mutant locus coverage 279× and 46× in the respective recurrent samples). In one case, the original TP53 mutation was replaced with a different variant. One recurrent tumor acquired a TP53 mutation that was not found in the matching primary, an observation confirmed by ultradep sequencing of the diagnostic tumor sample (average mutant locus coverage 1100×) (Supplemental Fig. 4). We compared the mutation frequencies (mutations per megabase, adjusted for coverage) at diagnosis and after disease relapse, and found disparate mutagenesis trends between TP53 mutant tumors and TP53 wild-type tumors (Fig. 3B). When compared to the matching primary tumor, TP53 mutant recurrent GBM showed an increase in subclonal mutation frequency, whereas GBM with wild-type TP53 did not, and the difference was statistically significant (P = 0.0015; Wilcoxon rank-sum test) (Fig. 3C). The clonal mutation frequency was unaffected (P = 0.23; Wilcoxon rank-sum test). Tumor purity, estimated by allele-specific copy number analysis of tumors (ASCAT) (Van Loo et al. 2010), did not confound the results (Supplemental Fig. 5). This increase in subclonal mutation frequency in TP53 mutant but not TP53 wild-type tumors after disease relapse further exposed the association between TP53 mutation and subclonal tumor progression.

Whole-genome sequencing inferred tumor evolution

To comprehensively cover GBM tumor heterogeneity and disease progression, we performed whole-genome sequencing on 10 primary-recurrence pairs from cohort II (Supplemental Table 1). We identified a median number of 4224 mutations (range from 2063 to 142,984) in the genome-wide sequencing data from 20 tumors (Supplemental Table 4). One recurrent tumor had an extremely high number (142,984) of mutations, and this sample was among the five recurrent tumors with hypermutated exomes (Fig. 3A).

We utilized the large number of mutations detected in each tumor to investigate the architecture of disease evolution by analyzing the distribution of purity-scaled variant allele frequencies (PS-VAFs) determined using the SciClone method (Miller et al. 2014). SciClone is a variational Bayesian algorithm to infer genetic composition of subclones from the variant allele frequencies of somatic mutations. To avoid potential confounding effects from DNA dosage, loss of heterozygosity (LOH), and sequencing depth, we limited the analysis to mutations with at least 60× coverage and located in copy-neutral and LOH-free regions. Theoretically, heterozygous mutations with 0.5 PS-VAF are clonal mutations present in all tumor cells. Analysis of PS-VAFs from paired primary and recurrent tumors revealed at least two evident patterns of GBM relapse. The ancestral cell of origin pattern was demonstrated by cases TCGA-06-0152 (Fig. 4A) and TCGA-14-1034 (Supplemental Fig. 6A). In this pattern, primary and recurrent tumors shared a cluster of clonal mutations close to PS-VAF 0.5; whereas at the same time, a large cluster of clonal mutations in the primary tumor disappeared in the matching recurrent tumor. The limited overlap of clonal mutations implicated a common ancestral cell that gave rise to both primary and recurrent tumors. The lack in overlapping subclonal mutations and the wide range of PS-VAFs from 0 to 0.5 in primary and recurrent specific mutations suggested that the two subsequent disease instances evolved independently (Fig. 5A).

A different pattern was observed in cases TCGA-06-0125 (Fig. 4B), TCGA-06-0190, TCGA-06-0221, and TCGA-14-1402 (Supplemental...
Figure 2. Association of subclonal fraction with TP53 mutation and event-free survival. (A) Clonal (yellow) and subclonal (blue) mutation frequencies and their association with G-CIMP methylation phenotype (blue) and p53 pathway-related alterations. The total number of sSNVs per patient is indicated in the top panel. (B) Patients with non-G-CIMP GBM harboring a higher subclonal mutation frequency (top 30% of non-G-CIMPs; Group II; blue) exhibited a longer event-free survival time compared to non-G-CIMP patients, whose tumor cell mutations had higher clonal frequency (bottom 30% of non-G-CIMPs; Group III; red). G-CIMP cases are grouped separately into Group I (dark green) and show event-free survival similar to that of Group II. Only patients younger than 55 yr of age were included in Groups II and III, and all G-CIMP cases were included in Group I.
Figure 3. Spectrum of somatic mutations in primary and recurrent GBM. (A) The number of mutations detected in exome sequencing (top panel), clinical characteristics (second panel), EGFRvIII aberrations (third panel), copy number alterations (fourth panel), and somatic exonic mutations (bottom panel). Only frequently amplified, deleted, and mutated GBM genes are included in the copy number and mutation panels. \( \omega \) (mutation of a different nucleotide in the same gene); \( v \) (DNA copy number alteration defined by different breakpoints). Clonal status for one sample with an extreme number of mutations could not be reliably classified and is shown in gray. (B) Comparison of subclonal (left panel) and clonal (right panel) mutation frequencies between primary or secondary GBM and matching recurrent tumors. Patients treated with TMZ are indicated with a brown triangle. (C) Association of TP53 mutation with subclonal mutational burden. Changes in clonal (left panel) and subclonal (right panel) mutation frequency are indicated as yellow dots for GBM harboring wild-type TP53 and purple triangles for tumors with mutated TP53 in both primary and recurrent tumors. The green diamond represents a patient who acquired a TP53 mutation at GBM recurrence; the blue diamond represents a case where a TP53 mutation was observed in the primary but not the matching tumor recurrence. A P-value was obtained by performing a Wilcoxon rank-sum test comparing subclonal mutation frequency change between TP53 wild-type and mutant groups.
Figure 4. Comprehensive comparison of mutations in four pairs of primary and recurrent GBM. Each row in the upper panels of A–D represents exonic mutations detected in sectors of the primary tumor (P.1, P.2) and exonic mutations detected in sectors of the matching recurrent GBM (R.1, R.2, R.3). Mutations are color-coded for cancer cell fractions (tumor purity-corrected cellular frequencies), in which a cancer cell fraction of 1.0 indicates that a mutation was present in all tumor cells. The columns represent the same chromosomal nucleotide position in all samples displayed. The bottom panels of A–D show scatter plots of the copy-neutral and non-LOH localizing mutations detected in whole-genome sequencing data of the primary and recurrent tumor. The x-axis and y-axis indicate purity-scaled variant allele fractions. Variant allele frequency distribution plots for individual tumors are also displayed at the bottom for primary tumor and at the right side for its matching recurrent tumor.
To address the possibility that the lack of overlap between subsequent tumors and subclones was due to intratumoral heterogeneity, we analyzed whole-exome sequencing data from a second, and in one case a third, sector of seven primary and six recurrent cohort II tumors. On average, 20 additional exonic mutations were detected in the second primary tumor biopsy, which represented a 25% increase on the total number of mutations found in the primary tumor. However, the additionally recovered mutations were in majority unique to the second primary sample (median: 17 of 20) and not detected in the tumor recurrence (Supplemental Table 5). Analysis of a second recurrent tumor biopsy did not alter this result but confirmed that the majority of “second biopsy” mutations were not conserved between time points. We therefore suggest that intratumoral heterogeneity did not explain the large number of mutations that were uniquely detected in primary and recurrence.

Evolutionary phylogenetic trees constructed from exome mutation profiles showed that primary and recurrent tumors sectors grouped in separate branches, with varying separation structures (Fig. 6). In one sample, TCGA-06-0125, the tree suggested that the tumor cells seeding the recurrent tumor were derived from primary tumor sector 2, which is consistent with the observations made in our analysis of the whole-genome sequencing data. Similarly, in TCGA-06-0211, the structure suggested that the tumor cells responsible for disease relapse were predominantly found in sector 1, where sectors 2 and 3 branched off linearly at a later time point.

Therapy-induced hypermutation is a subclonal process

We observed temozolomide treatment induced hypermutation, previously described in secondary GBM (Hunter et al. 2006; Johnson et al. 2014), in the recurrent tumor of five patients. The majority of mutations were classified as subclonal, and this was confirmed by the analysis of two independently sequenced biopsies from one hypermutated recurrent tumor, in which we observed 2429 and 5980 somatic point mutations, respectively, confirmed by the analysis of two independently sequenced biopsies, suggesting that mechanisms not related to these genes and resulting in hypermutation could exist. Although the number of samples was too small to identify genes associated with hypermutator phenotype at statistical significance, we noted that the receptor tyrosine kinases EGFR and ALK were among the eight genes mutated in all six
hypermutator samples (Supplemental Table 6). MGMT was methylated in all five primary tumors and in three of the five recurrent tumors, whereas MGMT status was not available for the remaining three recurrent tumors.

Following their second surgery to remove a GBM tumor, the five patients had respective lengths of survival of 35, 64, 107, 191, and 245 d, relative to a median survival of approximately 8 mo after surgery upon disease progression (Wen and Kesari 2008). The clinical outcome of the hypermutators illustrated the lethality of GBM, but the limited number of cases is unable to confirm that a high level of mutation is associated with relatively aggressive disease progression at statistically significant levels.

Discussion

As indicated by the adjective "multiforme," the histopathological features of glioblastoma show a high degree of intratumor heterogeneity. Genomic studies have further illustrated this disease characteristic by demonstrating local variation in the amplification patterns of receptor tyrosine kinases (Snuderl et al. 2011; Szerlip et al. 2012; Sottoriva et al. 2013; Francis et al. 2014) as well as a wide landscape of somatic alterations, expression subtypes, and epigenetic differences across GBM (Noushmehr et al. 2010; Verhaak et al. 2010; Brennan et al. 2013; Zheng et al. 2013). Here, we expand our knowledge of GBM by evaluating heterogeneity in a large number of primary tumor samples from TCGA as well as through a comparison of pre- and post-treatment GBM tumor pairs.

We developed a computational approach to infer the cellular frequency of sSNVs and to classify mutations as clonal or subclonal, which we then validated using multisector sequencing (Gerlinger et al. 2012). Corroborating the findings from previous reports (Nik-Zainal et al. 2012; Alexandrov et al. 2013; Tomasetti et al. 2013), we note that the number of clonal but not subclonal mutations found at the time of diagnosis increased with the age of the patient, and that clonal mutations reflected the signature of germline substitutions. Over the lifetime of the patient, the cancer cell of origin may thus have been subjected to mutational processes before acquiring the necessary alterations to become tumorigenic, and this observation could be used to increase the specificity of algorithms aimed at identifying cancer-contributory genes (Lawrence et al. 2013). Mutation of TP53 has been related to an increased frequency of double-strand breaks and chromothripsis in medulloblastoma (Rausch et al. 2012) and tumor progression from low-grade to high-grade glioma (Ishii et al. 1999; Fulci et al. 2002). We extended our knowledge of the damaging effects of TP53 deactivation by identi-
In conclusion, this study has provided a better understanding of intratumor heterogeneity and disease progression in GBM. Our dissection of the temporal sequence of mutations revealed mutational forces acting on the cancer genome and resulted in new insights into the patterns and dynamics of tumor evolution. Although the therapeutic opportunities for GBM remain limited, continuing efforts to detail the mechanisms of disease relapse will contribute to our ability to provide curative treatment for this lethal disease.

Methods

Whole exome DNA sequencing and targeted resequencing

DNA of multisector and additional non-TCGA primary and recurrent paired tumor samples in cohort II was extracted by the University of Texas MD Anderson Cancer Center DNA Analysis Facility following standard protocols. All specimens were obtained from patients with appropriate consent from the relevant institutional review board. Samples were then multiplexed and sequenced on Illumina HiSeq 2000 by the Sequencing and Microarray Facility at an average target exome coverage of 100× using 76-bp paired-end reads. Targeted resequencing of selected mutations for validation was performed by PCR using a microfluiddic device (Fluidigm Corp.). PCR primers with Fluidigm-compatible tails were designed to flank the sites of interest and produce amplicons of 200 bp ± 20 bp. Oligonucleotides containing Illumina adaptor sequences were mixed with 20–50 ng of each DNA sample along with a sample-specific molecular barcode and a sequence complementary to the primer tails. This mixture was used as the PCR template for each sample amplified on the Fluidigm access array. PCR was performed on the Fluidigm access array according to the manufacturer’s instructions. Barcoded libraries were recovered for each sample in a single collection well on the Fluidigm access array, quantified using PicoGreen dsDNA Quantiitation Reagent (Invitrogen), and concentrations were normalized for use across libraries. Libraries were loaded on the Illumina MiSeq instrument and sequenced using paired-end 150-bp sequencing reads.

Whole genome DNA sequencing

Library construction and sequencing alignment (using the Human Genome Reference Consortium build 37) were performed by the Broad Institute as previously described (Brennan et al. 2013). All whole-genome sequencing BAM files are available through The Cancer Genomics Hub (https://cghub.ucsc.edu/).

Exome sequence data processing and mutation calling

Exome sequencing reads were aligned to the human reference genome (hg19) using Burrows-Wheeler Aligner 0.7.2 (Li and Durbin 2009). BAM files were subjected to duplication marking, indel realignment, and recalibration using Picard version 1.7 (http://broadinstitute.github.io/picard/) and GATK version 2.4.9 (McKenna et al. 2010). On average, 70% of the bases on exonic regions were covered with at least 20 reads of sequence. We then applied MuTect algorithm (version 1.14) (Cibulskis et al. 2013) to identify somatic single nucleotide variations (sSNVs) from tumor and patient-matched normal blood samples. In this manuscript, we limited our analysis to the mutations whose variant allele fractions were at least 0.05. All sSNVs were annotated by ANNOVAR (Wang et al. 2010) and Oncotator (http://www.broadinstitute.org/oncotator). Only mutations in exon regions were included in subsequent analyses. For primary samples from cohort I, sSNVs were publically available and downloaded from the TCGA data portal.
Validation of somatic single nucleotide variants

To validate our somatic single nucleotide mutation calls, we performed targeted resequencing at high coverage (>1100×). We randomly selected 239 unique bases, which had been found to be mutated in primary, recurrence, or both, and sequenced these in both primary and recurrences. These sites corresponded to 367 SNVs and 101 wild-type nucleotides. In total, 343 of 367 mutations called from the exome sequencing data were detected in the high coverage data, resulting in a true positive validation rate of 93.5%. Evidence for a low allelic fraction somatic mutation was observed in nine of 101 wild-type nucleotides. The variant allelic fractions (VAFs), i.e., the number of reads harboring the variant allele divided by all reads covering to that base, of exome and validation sequencing were highly correlated (Pearson correlation = 0.93) (Supplemental Fig. 4).

Copy number and LOH analysis

For cohort II, Affymetrix SNP 6.0 arrays were preprocessed by either aroma.affymetrix (Bengtsson et al. 2009) or Copy-NumberInferencePipeline in GenePattern (http://genepattern. broadinstitute.org) and segmented by the circular binary segmentation algorithm (Olshen et al. 2004). Tumor and matched normal samples were used in pairs to obtain somatic alterations. The GISTIC2 algorithm (Mermel et al. 2011) was applied to the segmented copy number profiles for significant aberrations and sample-specific events. For cohort I, GISTIC2 outputs were downloaded from the Broad GDAC Firehose website (https://confluence.broadinstitute.org/display/GDAC/Home). In both cohort I and II, the loss of heterozygosity (LOH) levels were determined by the paired parent-specific circular binary segmentation method (PSCBS, v0.28.1) (Olshen et al. 2011) to raw Affymetrix SNP 6.0 CEL files from paired tumor and normal samples.

Inference of cellular frequency distributions of sSNVs

We defined cellular frequency of a mutation as the fraction of cells harboring the mutation. To estimate cellular frequency in individual tumors, we integrated reference and variant allele counts, LOH, and copy number alterations using the PyClone algorithm (version 0.2) (Shah et al. 2012). In brief, PyClone implements a Dirichlet process clustering model that simultaneously estimates the distribution of the cellular frequency for each mutation. Copy number levels at somatic mutation sites were inferred by thresholded copy number profiles determined by GIStIC 2.0 on a single sample basis. We considered “−2” as homozygous deletion, “−1” as hemizygous deletion, “0” as copy number neutral, “1” as low level gain, and “2” as high level gain. LOH status was inferred from the PSCBS LOH file. PyClone was performed using default settings. The outputs were a pairwise mutation co-occurrence matrix and cellular frequency value distribution per sSNV estimated from Markov-chain Monte Carlo (MCMC) sampling. The median value of the MCMC sampling-derived distribution was used as a representative cellular frequency for each mutation.

Inferring cancer cell fraction and clonal status of sSNVs

When corrected for tumor cell content, the cellular frequency of each mutation can be used to infer the cancer cell fraction of the mutation. We assumed a group of mutations were shared by all cancer cells in a biopsy, and their cellular frequencies reflected the relative abundance of cancer cells (cellularity) in the admixture of infiltrating normal cells. Then, the cancer cell abundance in each sample can be used as a scale factor for rescaling cellular frequencies to the cancer cell fractions of the mutations in the sample. To infer cancer cell abundance, we performed hierarchical clustering of the pairwise mutation co-occurrence matrix and then selected a group of at least 10 mutations with the highest median cellular frequency at a given tree cutoff threshold. Multiple grouping thresholds were applied, and we manually inspected the results to select the most reasonable one. Euclidean distances were used in hierarchical clustering. The distribution of rescaled cancer cell fraction per mutation in a tumor was used to compute the clonal probability of the mutation. We defined a mutation to be clonal if the probability of the mutation being present in >95% of the cancer cells was more than 0.5, or subclonal otherwise. The cutoff of 0.5 was selected based on the empirical distribution of clonal probabilities for our data set.

Tumor purity estimation

In cohort II, tumor purities and ploidy were estimated by the ASCAT algorithm (v 2.1) using Affymetrix SNP 6.0 array data (Van Loo et al. 2010).

Survival analysis

We carried out a survival analysis using the “Survival” package in R version 3.1.0 (R Core Team 2014). Clinical information for cohort I was downloaded from TCGA data portal (Brennan et al. 2013). An event was considered to be either disease progression or death. Survival curves were estimated by the Kaplan-Meier method. The statistical significance of survival differences was calculated using the log-rank test in the “Survival” package.

Statistical analysis

All statistical analysis was performed with R version 3.0.1 (R Core Team 2014).

Data access

SNP6 array and sequencing data from this study have been submitted to the European Genome-phenome Archive (EGA; http://www.ebi.ac.uk/ega/) under accession number EGAS00001001033.

Competing interest statement

M.M. is a founder of and equity holder in Foundation Medicine.

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astrocytic tumors evolve clonally to malignancy and are an unfavorable prognostic factor. *Oncoogene* 18: 5870–5878.


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