Recurrent patterns of DNA copy number alterations in tumors reflect metabolic selection pressures

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Abstract

Copy number alteration (CNA) profiling of human tumors has revealed recurrent patterns of DNA amplifications and deletions across diverse cancer types. These patterns are suggestive of conserved selection pressures during tumor evolution but cannot be fully explained by known oncogenes and tumor suppressor genes. Using a pan-cancer analysis of CNA data from patient tumors and experimental systems, here we show that principal genes. Using a pan-cancer analysis of CNA data from patient tumors, and experimental systems, here we show that principal component analysis-defined CNA signatures are predictive of glycolytic phenotypes, including component analysis-defined CNA signatures are predictive of glycolytic phenotypes, including 18F-fluorodeoxyglucose (FDG) avidity of patient tumors, and increased proliferation. The primary CNA signature is enriched for p53 mutations and is associated with glycolysis through coordinate amplification of glycolytic genes and other cancer-linked metabolic enzymes. A pan-cancer and cross-species comparison of CNAs highlighted 26 consistently altered DNA regions, containing 11 enzymes in the glycolysis pathway in addition to known cancer-driving genes. Furthermore, exogenous expression of hexokinase and enolase enzymes in an experimental immortalization system altered the subsequent copy number status of the corresponding endogenous loci, supporting the hypothesis that these metabolic genes act as drivers within the conserved CNA amplification regions. Taken together, these results demonstrate that metabolic stress acts as a selective pressure underlying the recurrent CNAs observed in human tumors, and further cast genomic instability as an enabling event in tumorogenesis and metabolic evolution.

Keywords aneuploidy; DNA copy number alterations; genomic instability; glycolysis; metabolism

Subject Categories Cancer; Genome-scale & Integrative Biology; Metabolism

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Introduction

Cancer cells differ from normal cells in that they exhibit aberrant proliferation, resist apoptosis, and invade other tissues (Hanahan & Weinberg, 2011). Modern cancer classification relies on molecular...
PCA revealed a high degree of similarity in CNA profiles between basal breast invasive carcinoma (BRCA basal), lung squamous cell carcinoma (LUSC), ovarian serous cystadenocarcinoma (OV), and serous uterine corpus endometrial carcinoma (UCEC serous) (Fig 1A and Appendix Fig S1). In tumor type-specific PCA, analyzing the four tissue-defined tumor sets of BRCA, LU (lung cancer consisting of LUSC and LUAD [lung adenocarcinoma]), OV, and UCEC revealed two strong PCA-based CNA signatures, termed signatures A and B, in all four cases. Notably, signature A was highly consistent across each tumor type, and reflected the pattern of pan-cancer PCI loadings (Figs 1B and C, and EV2A–C). In BRCA, signature A tumors were enriched for the basal subtype, p53 point mutations, high numbers of genomic breakpoints, and thus subchromosomal alterations (Figs 1B and D, and EV2D; $P < 0.001$, $P < 0.001$, $P = 2 \times 10^{-5}$, respectively). Signature B BRCA tumors, in contrast, were enriched for luminal type tumors ($P < 0.001$) and exhibited amplifications of the oncogenes MYC and MDM2 and deletion of the tumor suppressor CDKN2A (Fig 1B). Amplification of MDM2 and loss of CDKN2A were generally mutually exclusive in signature B tumors (Fig EV2E), reflecting alternate mechanisms for disabling the p53/ARF axis (Sherr & Weber, 2000). In the other tissues, signature A tumors were enriched for lung squamous cell carcinomas, the proliferative subtype of ovarian cancer (The Cancer Genome Atlas Research Network, 2011), and the serous subtype of uterine cancer (Fig EV2A–D). Overall, signature A tumors demonstrated enrichment of p53 mutations, more genomic breakpoints (BRCA, LU, and UCEC), and a higher degree of copy number alterations (LU, OV, and UCEC) than signature B tumors (Appendix Fig S2A and B, $P$-values indicated in figure). In signature A tumors, the per tumor average segment size is on the scale of $1 \times 10^5$ base pairs (approximately one-tenth of a chromosome, containing on the order of 100 genes). Overall, the segment sizes in signature A span from focal to arm-length/whole chromosome scale. The per tumor average segment size for signature B BRCA, LU, and UCEC tumors is statistically larger by $1.3\text{-fold}$ to sevenfold (Appendix Fig S2C–E; $P = 2 \times 10^{-5}$ or less). Unlike signature A tumors, the signature B CNA patterns were quite distinct between tumor types, although some commonalities were observed including point mutations in oncogenes such as KRAS (LU and UCEC) and amplification of MYC (BRCA, LU, and OV). An alternative approach using hierarchical clustering confirmed the existence of the shared pan-cancer CNA signatures across multiple tumor types (signature A), as well as distinct signature subtypes within each of the BRCA, OV, UCEC, and LU tumor types (signature A vs. signature B) (Fig EV3 and Appendix Fig S3, $P$-values of concordance between PCA and clustering approaches indicated in figure). In summary, PCA revealed the CNA signature A as a pattern shared across a subset of tumors in multiple tissue types, as well as several tumor type-specific cases of more distinct signature B patterns (Appendix Fig S1D).

Because altered metabolism is a hallmark of human tumors, we next tested whether the shared CNA signature A was enriched for genes from metabolic pathways. Using CNA-based gene set enrichment analysis over all metabolic pathways defined by KEGG (Kanehisa et al., 2014), we found that the conserved profile of core signature A tumors (i.e., OV, BRCA, UCEC, LU) (Fig 1C) was significantly enriched for DNA amplifications of core glycolysis pathway genes (Fig 1E and F, and Table EV1; $P = 0.024$). For example, BRCA
Figure 1.
A Pan-cancer PCA of copy number data from a balanced, random sampling of tumors of 15 tumor types from The Cancer Genome Atlas (TCGA). The average tumor PC scores for each tumor subtype are shown. Pan-cancer PC1 scores primarily separate diploid from highly aneuploid tumors, while PC2 distinguished GBM from the other tumor types (Appendix Fig S1A and B). Four tumor subtypes with similar PC scores are labeled as “signature positive (+)”. Tumor type abbreviations are as defined by TCGA: ovarian serous cystadenocarcinoma (OV), breast (BR), bladder urothelial (Bl), and thyroid (TH) carcinoma (“CA”), uterine corpus endometrial carcinomas (UCEC), lung (LU), and head and neck (HN) squamous carcinomas (“SC”), skin cutaneous melanoma (Skmcm), lung (LU), rectal (RE), colon (CO), stomach (ST), and prostate (PR) adenocarcinomas (“AD”), glioblastoma (GBM), low-grade glioma (LGG), and kidney renal clear cell carcinoma (KIRC).

B Copy number profiles of 873 breast invasive carcinomas (BRCA). Tumors (rows) sorted by tumor-specific principal component 1 (PC1) score with genomic locations listed across the top. PCA identified and distinguished two signatures with similar degrees of variance in the first component. The triangle marks the transition from signature A through diploid samples to signature B. The membership of each tumor with known molecular subtypes [e.g., basal/luminal] and mutant TP53 status are indicated on the right with red horizontal bars and corresponding permutation-based enrichment P-values. Basal and luminal subtype classifications are from a published gene expression–based determination (The Cancer Genome Atlas Research Network, 2012b). Signature A was moderately enriched for claudin-low tumors, whereas the HER2-enriched subtype was not significantly associated with either signature A or B (Fig EV2B). Similar PCA-based distinctions were observed in the LU, OV, and UCEC datasets (Fig EV2A–C).

C Pan-cancer PC1 loadings and signature A summary profiles for ovarian (OV), breast (BRCA), uterine (UCEC), and lung (LU) tumor types. Summary signatures are normalized gene loci signal-to-noise ratios (SNRs) of the top 10% of PC1-based core signature A tumors compared to normal (non-tumor) samples. Pan-cancer PC1-3 loadings are shown in Appendix Fig S1C. Consistency signatures of conserved amplification and deletions (consistent regions) are indicated by their signed absolute minimum consistency score (SAMCS). The SAMCS is non-zero when all CNA summary signatures have the same sign across all tumor types, and is derived from the absolute value-based minimum summary metric, and then re-signed positive for amplification or negative for deletion (see Materials and Methods). In the bar graph, red and blue denote consistently amplified and deleted regions, respectively. The number or percentage of consistent regions, genome coverage, and gene loci are indicated. The positions of canonical oncogenes, proto-oncogenes, tumor suppressor genes, and telomerase components (TERC, TERT) present in these regions are indicated.

D Breast signature A tumors have more DNA breakpoints per chromosome than signature B tumors, but similar levels of copy number alteration (as measured by an integrated CNA score proportional to the extent and absolute value magnitude of amplifications and deletions). P-values are indicated (top, Mann–Whitney U-test; bottom, Student’s t-test), and data presented in box (median, first and third quartiles) and whisker (extreme value) plots. The number of breakpoints is inversely proportional to the mean DNA segment length.

E KEGG metabolism pathway enrichment analysis based on consistent CNA patterns in the core signature A tumors of (A). The OV, BRCA, UCEC, and LU signatures from (C) were sequentially added and only directionally consistent CNA changes were retained (see Materials and Methods). The combined glycolysis–gluconeogenesis (glycolysis) and pentose phosphate pathway was included based on our prior mRNA work identifying the predictive value of this gene set (Palassak et al, 2011). Core glycolysis is a KEGG–defined gene subset (M00001). Table EV1 lists enrichment results for all KEGG metabolism pathways.

F Schematic showing average signal-to-noise (SNR) metrics of core glycolysis pathway genes plus TIGAR, the tumor suppressor p53, and other enzymes in core signature A tumors from (C) and (E). A more detailed listing of the gene names can be found in Materials and Methods.

G Gene copy number alteration distributions of selected glycolysis genes, TIGAR, the tumor suppressor p53, and the p53-associated cancer genes MDM2 and CDKN2A in BRCA tumors. Data are presented in box (median, first and third quartiles) and whisker (extreme value) plots.

Data information: in panels (C, E–G), the top 10% of tumors from each indicated signature based on PC1 scores were used in the analysis. See also Fig EV2 and Appendix Figs S1–S3.

Elimination of passenger genes via cross-species synten mapping

While canonical oncogenes and tumor suppressor genes drive some recurrent DNA copy number alterations, many recurrent CNA regions cannot be fully explained by the presence of known cancer genes (Beroukhim et al, 2010; Muller et al, 2012; Davoli et al, 2013). The conservation of CNA signature A across breast, lung, ovarian, and uterine tumor subtypes suggests the existence of one or more selective pressures that are potentially shared by additional tumor types. In that many of the human signature A conserved regions identified in Fig 1C still span large, chromosome-scale regions, we hypothesized that passenger genes could be diluting the enrichment signal. We subsequently reasoned that a previously reported approach of cross-species comparison of CNA data from human tumors and non-human tumorigenesis models would eliminate passenger genes via synten mapping (Maser et al, 2007; Zhang et al, 2013; Tang et al, 2014).

For the cross-species analysis, we first empirically determined the full set of human signature A-like tumor tissue types that could be included based on CNA similarity as defined by the pan-cancer PC1 scores from Fig 1A. Namely, we sequentially added each human tumor type in order of decreasing pan-cancer PC1 score and tested whether inclusion improved the overall pathway enrichment. In a parallel analysis, at each sequential step we also added the corresponding tissue-matched mouse epithelial cancer model signature if it existed. Mouse CNA signatures obtained from the literature and public repositories included signatures for genetically engineered mouse models of mammary (breast) cancer (Brca) (Drost et al, 2011; Herschkowitz et al, 2012), melanoma (Skmcm) (Viros et al, 2014), glioblastoma/high-grade astrocytoma (Gbm) (Chow et al, 2011), and prostate cancer (Prad) (Ding et al, 2012; Wanjala et al, 2015), as well as in vitro mouse epithelial cell models of bladder (Blica), colorectal (Coad), and kidney (Kirc) cancer (Padilla-Nash et al, 2013).
Tumor CNAs reflect metabolic selection

Nicholas A Graham et al

Using human tumors only, the sequential enrichment analysis showed a general improvement through the addition of the COAD tumor type (Fig 2A, dotted orange line; average enrichment signal of the top 10 metabolic pathways). Moreover, sequential inclusion of the corresponding tissue-matched mouse cancer model CNA signatures substantially enhanced the overall pathway enrichment results with the peak occurring at HNSC, a sequentially adjacent tumor type that has similar pan-cancer PCI score with COAD (Fig 2A, solid blue line and arrow indicating the peak). Overall pathway enrichment then decreased as additional, less copy number similar (lower pan-cancer PCI scores) tumor types were added.

The improvement upon including the mouse signatures supports the hypothesis that the cross-species analysis eliminates passenger genes and reveals pathways whose genes are enriched in the resulting cross-species consistent CNA regions. Taken together, this pan-cancer and cross-species analysis demonstrates that a combination of nine human tumor types (OV, BRCA, UCEC, LU, BLCA, READ, SKCM, COAD, and HNSC) and four corresponding mouse tumor models (Brca, Blca, Skcm, and Coad) gives the strongest overall enrichment signal strength across all metabolic pathways (Fig 2A).

Examining the highly ranked individual pathways from this optimized tumor type combination, we found that the carbohydrate metabolism pathway “amino sugar and nucleotide sugar metabolism” (hsa00520) was ranked first and glycolysis-glucoseogenesis (hsa00010, henceforth called glycolysis) was ranked second among KEGG metabolism gene sets (Fig 2B and Table EV1). As in the average of top 10 pathways (Fig 2A), the enrichment score for the glycolysis gene set improved as more human tumor and mouse models were sequentially included up through the additions of HNSC and Coad (Fig 2C). Compared to our previous enrichment analysis of the four core signature A tumors (Fig 1E), the permutation P-value of the glycolysis gene set improved from 0.05 to 0.001, reflecting that our pan-cancer and cross-species analysis has eliminated passenger genes in the non-consistent CNA regions. Thus, we hereafter refer to the consistent signature pattern from OV up through and including HNSC and Coad as “expanded signature A.” Of particular note, upon expanding our analysis to include 1,321 tumors for BRCA and LU (et al 2011), we predicted to be significantly more glycolytic than signature B tumors using RNA-based enrichment analysis (GSEA). To further explore the RNA expression data, we predicted the glycolytic phenotypes of the core signature A tumor types (BRCA, LU, and OV) using RNA-based weighted gene voting (WGV) (Golub et al, 1999) and our previously defined FDG prediction model (Palaksas et al, 2011). UCEC tumors were again excluded because there were not a sufficient number of samples with paired RNA and DNA data. We found that RNA-based predictions of high glycolysis were associated with the signature A end of tumor type-specific PCI for BRCA and LU (Fig EV4B–D). Signature A tumors were predicted to be significantly more glycolytic than signature B tumors for BRCA and LU (P-values of 2.4 × 10^-17 and 5.7 × 10^-18, respectively, Fig EV4E). There was no significant differential trend in OV tumors, potentially because almost all of these tumors are genomically unstable (integrated CNA scores > 0.2) and the signature B of OV is relatively signature A-like (Fig EV2B and Appendix Fig S1D). The OV predictions were consistent with predicted high glycolysis across all tumors (Fig EV4E). To control

CNA signatures are predictive of glycolysis

In that the CNA-consistent region-defined signature A is enriched for core glycolysis genes, we next tested whether signature A patient tumors in vivo were associated with increased tumor glycolysis. To assign a signature A score to a set of FDG-PET-imaged breast cancers (Palaksas et al, 2011), we projected CNA data from these tumors onto a PCA of the four core human tumor tissue types (OV, BRCA, UCEC, and LU, Fig 1C). We found a strong correlation between the strength of CNA signature A and the measured FDG-PET standardized uptake values (SUVs) (Fig 3A and B; Pearson r = 0.94, P = 5 × 10^-5). A similar analysis using all nine expanded signature A tumor types demonstrated equivalent results (Appendix Fig S4A; Pearson r = 0.92, P = 2 × 10^-4). Thus, the CNA-defined signature A is associated with increased FDG uptake in human primary tumors in vivo.

We next asked whether the CNA signature A-defined tumors associated with high FDG uptake also had RNA-based signatures of increased glycolysis. First, we compared signature A tumors to signature B tumors using RNA-based enrichment analysis (GSEA). We analyzed both BRCA and LU tumors because these tumor types show distinct signature A and signature B subtypes. (OV signature B is highly similar to the pan-cancer signature A pattern, and thus does not provide a differential test (Fig EV2B). UCEC tumors were not included due to a lack of sufficient paired RNA and DNA profiling data.) In the enrichment analysis, we included a gene set consisting of genes from the glycolysis and pentose phosphate pathways that were upregulated in FDG-high BRCA tumors, as defined by our previous work (Palaksas et al, 2011). In the GSEA, the empirically defined FDG-high gene set ranked number one overall (NES = 2.5, permutation P-value = 2 × 10^-4), confirming that Sig A tumors have glycolysis RNA expression profiles matching those of FDG-high tumors (Fig EV4A). Furthermore, BRCA and LU signature A tumors were significantly enriched for genes from the full glycolysis and pentose phosphate pathways (overall rank 4th of 76 pathways, NES = 2.0, permutation P-value = 6.8 × 10^-5), as well as other glycolysis-related pathways (Table EV3). This analysis of RNA expression data is consistent with the enrichment of glycolysis-associated pathways at the DNA copy number level (Fig 1E and 2B), and points to glycolysis-related pathways as selection targets for upregulation in signature A tumors.

To further explore the RNA expression data, we predicted the glycolytic phenotypes of the core signature A human tumors and mouse tumors (Fig 2D). Examination of conserved regions revealed that HK2, TPI1, GAPDH, PGAM2, ENO2, and LDHB glycolysis genes contribute to the cross-species consistency signal. Among canonical oncogenes, MYC and KRAS were also present in the amplification regions of the expanded signature A human tumors and mouse models. Due to the well-documented and clinically relevant role of glycolysis and pentose phosphate pathways in tumorigenesis, we chose to further examine the recurrent amplification of the set of these gene loci in subsequent analyses. Equally important for functional validation of these candidate pathways, the activity of glycolysis can be directly and indirectly measured by many assays in both patients (e.g., FDG-PET imaging) and experimental systems.
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Tumor CNAs reflect metabolic selection  Nicholas A Graham et al

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**Figure 2.**

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**KEGG metabolism**

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**Human core signature A**

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**Human expanded signature A**

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**Mouse models**

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**Canonical Pathway (CP)**

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<tr>
<td>9. SIRP family interactions</td>
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**Figure 2.**

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**A** Top 10 metabolic pathways

**B** KEGG metabolism

**C** Glycolysis

**D** Consistent Regions (SAMCS)

**Human Genome**

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<tr>
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**Synteny graph**

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**Gene set** (n=75)

- Amino sugar & nucleotide sugar metabolism
- Glycolysis
- Glycolysis & Pentose Phosphate
- Core Glycolysis
- Other types of O-glycan biosynthesis
- All metabolic genes
- Starch and sucrose metabolism
- Pyruvate metabolism

**Gene set** (n=1,321)

- 2. Glutathione conjugation
- 6. Maturity onset diabetes of the young
- 7. p38 MAPK events
- 8. Botulinum neurotoxicity
- 9. SIRP family interactions
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Abstract

Pathway DNA copy number alterations from signature A are based predictions above, the glycolysis and pentose phosphate pathway were most predictive of glycolytic phenotypes. We thus derived a panel of immortalized mouse embryonic fibroblasts (MEFs) using the classical 3T9 protocol (Todaro & Green, 1963). In this experimental system, under standard culture conditions (e.g., 21% O₂), diploid cells undergo a crisis-associated event that increases tolerance for genomic instability and allows them to escape senescence and evolve into cells with chromosomal instability and genomic aberrations (Fig 4A and Appendix Fig S5A; Sun & Taneja, 2007). This system has been used to study core cancer phenotypes such as proliferation, anti-apoptosis, and chromosomal instability (Lowe et al, 1993; Gupta et al, 2007; He et al, 2007; Sotillo et al, 2007; Sun et al, 2007; Weaver et al, 2007) and is one of the few experimentally tractable cancer models involving spontaneous genomic instability (Sherr & DePinho, 2000). In addition, because this system is not driven by strong oncogenes (e.g., KRAS mutation), it allows for complex CNA signatures to evolve from a combination of individual, presumably weaker, DNA alteration events.

We profiled the genome-wide copy number of 42 independent MEF sublines by CGH microarray (Fig 4A). Most samples were profiled after immortalization (post-senescence), with a few profiled before or during senescence. Analysis of this CNA data by PCA revealed that the MEF system recapitulated a two-signature pattern (signatures A and B). These two signatures were generally orthogonal, with the exception of a few “mixed” samples that had CNA characteristics of both signatures A and B (Appendix Fig S6B). As anticipated from prior MEF studies, immortalized MEF cells exhibited an increased number of genomic breakpoints and a higher degree of copy number alterations than the diploid, pre-senescent MEF cells (Appendix Fig S5C and D). Importantly, the MEF-derived signature A resembled the shared signature A pattern derived from

Figure 2. Pan-cancer and cross-species analysis reveals enrichment of glycolytic genes in conserved amplification regions.

A Average pathway enrichment signal strength for the top 10 enriched KEGG metabolism pathways based on the consistent CNA patterns across multiple tumor types. The four core CNA signature A patterns are defined as in Fig 1C. CNA signatures for other tissue tumor types are defined by the signal-to-noise ratios (SNRs) of all tumors compared to tissue-matched normal (non-tumor) samples. Starting with the four core signature A tumor types from Fig 1 (OV, BRCA, UCEC, and LU), additional human (orange) or human and mouse (blue) signatures were sequentially added based on their decreasing tissue type pan-cancer PC1 score from Fig 1A (see Materials and Methods). The resulting sequentially restricted consistency signatures are illustrated in (D). A maximal peak (indicated by an arrow) in the enrichment signal of this sequential analysis is seen after adding tumors through human HNSC and mouse Coad (no mouse Hnsc signature was available). Enrichment signal strength is based on the negative log frequency of equivalent enrichment by chance (i.e., permutation P-value). Mouse tumor type abbreviations are shown with lower case letters but otherwise match the human TCGA abbreviations.

B Enrichment analysis of metabolism (75 KEGG pathways) and canonical pathways (CP, 1,321 MSigDB canonical pathways). Permutation P-values for the consistent CNA patterns from the four human core signature A tumors (defined in Fig 1C), the human expanded signature A (from A), and the human expanded signature A combined with corresponding mouse cancer models (from A). FDR calculations are described in Materials and Methods. Tables EV1 and EV2 list all sequential enrichment results for KEGG metabolism and canonical pathways, respectively.

C Enrichment score for the glycolysis pathway upon sequential addition of human tumors and mouse models (as in A) showing a maximal peak in the enrichment signal corresponding to the expanded HNSc–Coad signature.

D Genome view of the sequential consistency signatures used in the enrichment analysis of (A–C). The locations of consistently amplified glycolysis genes, canonical oncogenes, and tumor suppressors are indicated. Genes listed in parentheses (e.g., RPA1, TERC, TERT, APC, and TP53) were consistently altered across the human tumors but not consistently altered in the mouse models, due in part to the inclusion of p53 genetic knockout mouse models (TP53) and known differences in human and mouse telomere maintenance (TERC, TERT) (Sherr & DePinho, 2000). The synteny graph at the bottom indicates the syntenic mouse chromosome number and thus the broken syntenic regions between human and mouse genomes. This genome view chronicles how the sequential pan-cancer and cross-species analysis of conserved amplification and deletion loci greatly reduces the percentage of consistent copy number alterations, thereby reducing the percentage of the genome implicated as candidate driver regions.

Experimental recapitulation of tumor CNA signatures

Having demonstrated that the glycolytic pathway is statistically associated with genome-wide DNA copy number patterns, we sought an experimentally tractable system that would allow us to test the hypothesis that recurrent patterns of DNA amplification reflect metabolic selection pressures. We thus derived a panel of immortalized mouse embryonic fibroblasts (MEFs) using the classical 3T9 protocol (Todaro & Green, 1963). In this experimental system, under standard culture conditions (e.g., 21% O₂), diploid cells undergo a crisis-associated event that increases tolerance for genomic instability and allows them to escape senescence and evolve into cells with chromosomal instability and genomic aberrations (Fig 4A and Appendix Fig S5A; Sun & Taneja, 2007). This system has been used to study core cancer phenotypes such as proliferation, anti-apoptosis, and chromosomal instability (Lowe et al, 1993; Gupta et al, 2007; He et al, 2007; Sotillo et al, 2007; Sun et al, 2007; Weaver et al, 2007) and is one of the few experimentally tractable cancer models involving spontaneous genomic instability (Sherr & DePinho, 2000). In addition, because this system is not driven by strong oncogenes (e.g., KRAS mutation), it allows for complex CNA signatures to evolve from a combination of individual, presumably weaker, DNA alteration events.

We profiled the genome-wide copy number of 42 independent MEF sublines by CGH microarray (Fig 4A). Most samples were profiled after immortalization (post-senescence), with a few profiled before or during senescence. Analysis of this CNA data by PCA revealed that the MEF system recapitulated a two-signature pattern (signatures A and B). These two signatures were generally orthogonal, with the exception of a few “mixed” samples that had CNA characteristics of both signatures A and B (Appendix Fig S5B). As anticipated from prior MEF studies, immortalized MEF cells exhibited an increased number of genomic breakpoints and a higher degree of copy number alterations than the diploid, pre-senescent MEF cells (Appendix Fig S5C and D). Importantly, the MEF-derived signature A resembled the shared signature A pattern derived from

for the general increase in glycolysis predictions at increased levels of genomic instability, we also analyzed the correlation between glycolysis predictions and PCA scores for several range windows of integrated CNA scores. This analysis demonstrated that RNA-based predictions of glycolysis were significantly correlated with PC1 scores in BRCA and LU tumors with high integrated CNA scores (permutation P-value for BRCA and LU < 1 × 10⁻⁶ each, Fig EV4F). Taken together, the DNA-defined signature A tumors exhibit RNA expression patterns consistent with increased FDG uptake. These RNA-based results support the more downstream activity-based findings of elevated FDG uptake in signature A tumors (Fig 3A and B).

Returning to the result of the CNA-defined signature A being associated with increased FDG uptake in human primary tumors in vivo, we next asked which sets of metabolic gene loci copy number levels were most predictive of glycolytic phenotypes. We performed a CNA-based weighted gene voting (WGV) analysis to predict the glycolytic phenotypes of breast tumors and breast cancer cell lines (Neve et al, 2006) using individual gene sets from the KEGG metabolic pathways database (Kanehisa et al, 2014). Gene weights were calculated from each of our four tumor type CNA “training” signatures. Specifically, we tested the ability of individual metabolic pathways to predict (i) FDG uptake in patient primary breast tumors and (ii) the lactate secretion of a panel of 32 breast cancer cell lines (Hong et al, 2016). Averaging results across these two test cases revealed that genes from the glycolysis and pentose phosphate pathway were most predictive of these metabolic phenotypes (Fig 3C and Table EV4; P = 0.01). Moreover, signature A-based predictions were predictive of lactate secretion for basal cell lines more so than for luminal lines (Fig 3D and E, and Appendix Fig S4B), consistent with the observed basal-type tumor enrichment in signature A samples (Figs 1B and EV2D). Thus, consistent with the gene expression-based predictions above, the glycolysis and pentose phosphate pathway DNA copy number alterations from signature A are predictive of glycolytic phenotypes of primary human breast tumors and cancer cell lines.

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Molecular Systems Biology 13: 914 | 2017

7
human breast, lung, ovarian, and uterine tumors. In particular, the MEF-derived signature A was characterized by p53 mutation and chromosome loss at the p53 locus, more genomic breakpoints, subchromosomal-sized alterations, and a higher degree of copy number alteration (Fig 4A and B, and Appendix Table S1). Additionally, this experimental system allowed us to profile the same MEF lines at subsequent passages, and a paired statistical analysis of five evolving signature A lines revealed genomic regions changing from mid- to late passage (Fig 4A and Appendix Fig S6A and B). Similar to the shared human tumor signature A, both the MEF signature A and the evolving MEF signature demonstrated DNA amplifications of glycolysis and glycolysis-related genes (Table EV2). In particular, both human and mouse CNA signatures A included amplification of Hk2, Bpgm, Rpia, Tigar (mouse gene 9630033F20Rik), Enol2, Tpi1,
Gapdh, Ldhb, Kras (mouse chr. 6), and Pkg2 (mouse chr. 17) (Figs 1G and 4A, and Appendix Fig S6C).

In contrast, the MEF-derived signature B was characterized by amplification of Mdm2 or deletion of Cdkn2a (Ink4a/Arf) and fewer overall copy number alterations. Mdm2 amplification and Cdkn2a loss are alternative mechanisms for inactivating p53 function in human tumors and in MEFs (Sherr & DePinho, 2000), but as found here result in a distinct CNA signature as compared to the p53 mutation-associated signature A. As in human BRCA (Fig EV2E), amplification of Mdm2 and loss of Cdkn2a were generally mutually exclusive in signature B MEFs (Appendix Fig S6D), reflecting alternate mechanisms for disabling the p53/ARF axis (Sherr & Weber, 2000). Additionally, Mdm2 amplification tends to co-occur with Hk1 amplification, both loci being located on mouse chr. 10. Thus, the signature B cases are associated with an alternate HK amplification, similar to our finding in signature B human tumors (HK1 or HK3 rather than HK2) (Fig 1G and Appendix Figs S2G and S6C). Immunoblotting confirmed that signature A MEFs and signature B MEF lines generally had increased expression of Hk2 and HK1 protein, respectively (Appendix Fig S6E and F).

To further characterize the association between p53 loss and CNA signatures, we profiled the CNA patterns of 29 independent p53−/+ MEF sublines derived in standard 3T9 culture conditions. Comparison of the p53−/+ MEF CNA patterns to wild-type MEFs by PCA demonstrated that the CNA patterns of p53−/+ MEFs resemble the wild-type MEF signature A pattern, with no signature B-like sublines observed (Appendix Fig S7). p53−/+ MEFs do not undergo senescence (Olive et al., 2004), and consistent with this, we observe that they tended to have less strong copy number alterations. In summary, strong p53 functional loss (p53 mutation or genetic loss) tends to lead to the CNA signature A pattern, which is associated with a higher degree of copy number alterations (higher integrated CNA score and more breakpoints) and Hk2 amplification, while weaker or less complete p53 functional loss (e.g., mediated by Mdm2 amplification or Cdkn2a loss) is associated with an alternative signature (signature B).

**Numerical and structural chromosomal abnormalities**

Next we sought to understand how CNA signatures revealed by CGH relate to numerical and structural aneuploidy (i.e., whole chromosomal and subchromosomal gains or losses, respectively). Using propidium iodide-based DNA staining, we found that immortalized MEF cells had increased total DNA content compared to pre-senescent MEFs (Appendix Fig S8A–C). Similar results were observed in a comparison between a genome-stable, close to diploid, immortalized human mammary epithelial cell line (MCF10A) and a human breast cancer cell line with a high degree of DNA copy number alterations (Hs578T). In addition, we used spectral karyotyping (SKY) to assess the numerical and structural chromosomal aberrations in immortalized MEF cells. Profiling a representative signature A MEF subline just after immortalization (subline H1; passage 25, P25), SKY revealed increased total DNA content (105 ± 36 chromosomes) with nearly all chromosomes having experienced whole chromosome gains (Appendix Fig SBD, E and H). In addition, there was substantial cell-to-cell heterogeneity in the number and type of chromosomes at passage 25. Profiling the same MEF cell line 23 passages later (P48), SKY revealed that the average number of chromosomes had slightly decreased and stabilized (84 ± 12 chr), as has been reported previously (Hao & Greider, 2004), and clonal markers had begun to emerge (e.g., translocation (3:16)) (Fig 4C and Appendix Fig SBF, G and I). We also observed a substantial number of double minutes in some of the cells examined at P48. As expected, the most strongly amplified chromosomes by SKY were scored as gains in CGH data (e.g., chrs. 3 and 6 in subline H1 at P48/49). Since array CGH analysis is normalized by input DNA quantity, genomic regions scoring as negative on a log2 CNA plot can be greater than diploid if most other chromosomes have amplified to an even greater extent (e.g., trisomy chrs. 1, 2, and 7 in subline H1 at P48/49). Since array CGH analysis is normalized by input DNA quantity, genomic regions scoring as negative on a log2 CNA plot can be greater than diploid if most other chromosomes have amplified to an even greater extent (e.g., trisomy chrs. 1, 2, and 7 in subline H1 at P48/49). Similar results have been reported for karyotyping and CGH results of human cancer cell lines (Kytölä et al., 2000). Taken together, these data reveal that immortalized MEFs experience substantial numerical and structural chromosomal abnormalities, similar to what is observed in human tumors, and further support that selection for optimized rearranged genomes occurs.

**Senescence-associated oxidative stress as a selective force for copy number alterations**

The MEF system allowed us to investigate the selective pressures driving copy number changes during immortalization. Because MEFs cultured under physiological oxygen conditions (3% oxygen) undergo little to no senescence and exhibit less DNA damage...
Tumor CNAs reflect metabolic selection
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Figure 4.

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Molecular Systems Biology 13 5 1 2017

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Figure 4.
Because the MEF signature A is qualitatively similar to the human expanded signature A, we next asked how inclusion of our MEF signatures would affect metabolic and canonical pathway enrichment analysis. Although the MEF immortalization system utilizes fibroblast cells and the mouse tumors are epithelial in origin, both types of models share a similar cross-species consistency with the expanded signature A tumor types (Appendix Fig S9B). When the human tumors signatures were combined with the MEF signatures, consistent genome regions were reduced to 3.6% of the genome spread over 13 conserved regions (Fig 4E). In cross-species pathway enrichment analysis, MEF CNA signatures added a similar amount of enrichment signal to human tumor signatures as do the non-MEF mouse model signatures, and when combined together, the enrichment was even stronger (e.g., glycolysis and core glycolysis pathways, Fig 4F). When all signatures are used, glycolysis was the top-ranked enriched pathway out of 1,321 MSigDB canonical pathways (Table EV2). Thus, including the MEF signatures in the cross-species analysis further implicates the glycolysis pathway in shaping the cancer genome.

Examination of the cross-species genomic regions conserved in human tumors, mouse models of cancer, and the MEF immortalization system revealed consistent amplification of five regions containing the COSMIC database-enumerated oncogenes GATA2 (human chr. 3; Fig 4E, last SAMCS line), TRRAP (chr. 7, region 4), MYC (chr. 8, region 1), CHD4 (chr. 12, region 1), and KRAS (chr. 12, region 2) (Forbes et al., 2015). Notably, four of the cross-species conserved CNA genomic regions included genes from the glycolysis...
pathway: HK2 (chr. 2), GCK and PGAM2 (chr. 7, region 1), ENO2, TPI1, and GAPDH (chr. 12, region 1, which includes TIGAR), and LDHB (chr. 12, region 2, which also includes Kras). In both human tumors and the MEF immortalization system, the genomic region harboring TIGAR–GAPDH–TPI1–ENO2 was separated from the LDHB–Kras region by a deletion-prone region that includes the tumor suppressor CDKN1B (Appendix Fig S9C–E).

To investigate whether amplification of the DNA cross-species conserved CNA regions results in upregulated RNA expression levels, we examined the correlation between DNA copy number and RNA expression levels in BRCA, LU, and OV tumors for glycolytic genes and, as a point of comparison, known oncogenes (Appendix Fig S10). We examined the average correlation across BRCA, LU, and OV signature A tumors for all genes in the 12 cross-species conserved regions. We found that three glycolytic genes (LDHB, TPI1, and GAPDH) and one oncogene (Kras) exhibited strong DNA–RNA correlation ($r > 0.66$), three glycolytic genes (HK2, ENO2, and PGAM2) and three oncogenes (MYC, CHD4, and Trrap) exhibited moderate correlation ($0.2 < r < 0.5$), and only one glycolytic gene (GCK) and one oncogene (Gata2) exhibited weak DNA–RNA correlation ($r < 0.2$) (Appendix Fig S10A and B). This analysis indicates that gene copy number alterations at the DNA level generally lead to increased RNA expression in signature A tumors in BRCA, LU, and OV tumors, with a similar degree of correlation observed for both glycolysis genes and oncogenes. Finally, we compared the upregulation of glycolytic genes and oncogenes on two cross-species conserved regions of human chr. 12 that contain both types of genes (Appendix Fig S10C). Within the centromere-distal region of chromosome 12, the glycolytic genes TPI1 and GAPDH show strong correlation (Spearman rank correlation = 0.71 and 0.66, respectively) while the oncogene CHD4 and the glycolytic gene ENO2 exhibit moderate correlate (Spearman rank correlation = 0.42 and 0.32, respectively). Within the more centromere-proximal region, LDHB and Kras both exhibit strong correlation (Spearman rank correlation = 0.68 and 0.66, respectively). Importantly, the correlation of RNA expression with DNA amplification is not stronger for oncogenes than for glycolytic genes within these cross-species regions. Taken together, these results support a model in which the selection pressures shared during immortalization and tumorigenesis result in cross-species conservation of the glycolysis gene loci copy number alterations (Fig 4E and F, and Table EV2).

Alteration of CNA signatures by exogenous expression of metabolic enzymes

The presence of core glycolysis genes in cross-species conserved amplification regions suggests that these metabolic gene loci drive the amplification of these regions. To test this hypothesis, we transformed pre-senescent MEFs with either wild-type HK2 or HK1, kinase-dead HK2 (D209A/D657A) (McCoy et al., 2014), or wild-type ENO2 and allowed the cells to senesce and immortalize in the presence of these exogenously expressed proteins (Appendix Fig S11). Analyzing the signature A set of sublines, we found that the endogenous HK2 locus (chr. 6) was less amplified in cells expressing exogenous wild-type hexokinase ($P = 0.048$) (Fig 5A and B). As a control, a signature A MEF cell line expressing kinase-dead hexokinase did not show reduced amplification of the HK2 locus ($P = 2 \times 10^{-4}$). In that MEF lines exogenously expressing hexokinase still demonstrated positive selection for the centromere-proximal half of chr. 6 (Fig 5A), we examined the ratio of HK2 gene locus copy number to the maximal amplification on chromosome 6 (HK2:Chr6 max). In this analysis, we found that cells expressing exogenous hexokinase demonstrated significantly reduced HK2:Chr6 max ratios ($P = 3 \times 10^{-5}$) (Fig 5A and C). Additionally, a MEF subline expressing exogenous ENO2 exhibited deletion rather than amplification of the Eno2 locus on chr. 6 ($P = 0.02$) (Fig 5A and D). Analyzing the signature B set of sublines, we found that the endogenous HK1 locus (chr. 10) was copy number neutral, rather than amplified, in a cell line expressing exogenous hexokinase ($P = 0.17$), whereas a signature B MEF cell line expressing kinase-dead hexokinase did not show reduced amplification of the HK1 locus (Fig 5E and F). Fisher’s combined statistical analysis of these results yielded $P$-values of 0.001 or less (Fig 5).

Taken together, these results demonstrate that exogenous expression of metabolic enzymes can alter the copy number status of the endogenous genomic loci, supporting these metabolic genes as drivers within the conserved amplification regions observed in human tumors and mouse models. In addition, these results support a model in which the net propensity for a chromosomal region to be amplified or deleted is in part related to the sum of the fitness effects of the genes present (Davoli et al., 2013). For example, when HK2 is exogenously expressed, the centromere-distal half of chromosome 6 had a low copy number value in early culture but after additional culture demonstrated increased copy number, supporting that the other gene loci of this region (such as Eno2, Gapdh, and other glycolytic genes) do have a remaining pro-fitness benefit (Fig 5A).

Metabolism and growth phenotypes of CNA signatures

To test whether there exist phenotypic differences between signature A and signature B MEFs, we characterized 11 wild-type MEF lines representative of either signature A or B and one mixed signature line. We found that signature A MEFs generally had higher rates of glucose consumption and lactate production than signature B MEF lines (Figs 6A and EV5F). Plotting the PC1 score versus glucose consumption, we found that CNA-based signature A was highly predictive of glucose consumption in the MEF signature A lines (Fig 6B). In contrast, signature B was only moderately predictive of the glucose consumption of signature B lines, and was not as accurate as signature A in predicting the glucose consumption of signature A lines (Figs 6B and EV5A). In addition, we noted that the signature A cell lines generally exhibited significantly higher rates of proliferation than the signature B cell lines (Fig 6C). Similar to glucose consumption, signature A was predictive of the growth rates of signature A MEFs and signature B was predictive of the growth rates of signature B MEFs, while cross-signature predictions had less power (Figs 6D and EV5B). Furthermore, we observed a general coevolution of higher growth rates and increased CNA signature strength in MEF lines that were profiled at different passage numbers (Fig 6D). As noted above, the evolving MEF CNA signature pattern was enriched for DNA amplifications of genes in the core glycolysis and glycolysis-associated pathways (Table EV2), particularly due to amplification of chromosome 6, which contains multiple metabolic gene loci including HK2 and Eno2 (Fig 4A and E, and Appendix Fig S6B).
Figure 5.

To test whether signature A and signature B MEFs differentially use glucose, we cultured MEF cells with [1,2,13C]-labeled glucose and conducted metabolomic profiling by mass spectrometry (Fig EV5C-G and Table EV5) (Metallo et al., 2009). In all MEF lines tested, we observed a low percentage of single heavy labeled carbon [M1 isopomer compared to M2] in pyruvate, lactate, and alanine, indicating that the contribution of glucose-derived carbon from the oxidative arm of the pentose phosphate pathway to these metabolites was relatively low (Fig EV5D). Nonetheless, the patterns of heavy isotope labeling revealed differences in nutrient utilization between signature A and B MEF cell lines. On average, metabolites of early glycolysis, the pentose phosphate, and nucleotide synthesis pathways showed a higher percentage of glucose-derived heavy carbon labeling in signature A MEFs (Figs 6E and EV5E). In contrast, signature A cells had a lower percentage of glucose-derived heavy carbon labeling in metabolites of the serine synthesis pathway and the TCA cycle. When compared to signature B MEFs, signature A cells also tended to exhibit increased consumption of serine, glutamine, and other amino acids (Fig EV5F and G). The increased glutamine consumption in signature A MEFs reflects similarity with basal breast cancer cell lines (also signature A associated), which exhibit increased glutamine consumption relative to luminal breast cancer cell lines (Timmerman et al., 2013). In general, the percentage of individual metabolites from the early glycolysis, the pentose phosphate, and nucleotide synthesis pathways that incorporated heavy, glucose-derived carbon metabolites was positively correlated with glucose consumption rates (e.g., fructose-1,6-bisphosphate, Fig 6F). Conversely, metabolites from the serine synthesis pathway and the TCA cycle showed a negative correlation between glucose consumption rates and the percentage of each molecule containing a heavy, glucose-derived carbon (e.g., 3-phosphoserine, Fig 6F). While caution must be exercised when interpreting metabolite labeling results, we overall observed a consistent trend of differences between signature A and B lines. Of note, the differences between signature A and B lines were mainly in regard to scale, with the strongest signature B lines demonstrating similar glycolysis and proliferation rates, as well as similar metabolic profiles, as the weakest signature A lines.

Taken together, these results demonstrate that signature A MEFs, which resemble signature A human tumors, exhibit increased glycolysis and to a somewhat lesser extent increased proliferation, and have an increased relative proportion of glucose-derived carbon in metabolites of pentose phosphate-associated biosynthetic pathways such as nucleotide synthesis. These findings are consistent with published mouse model studies in which tumor cells that are channeling glucose toward nucleotide biogenesis achieve faster rates of proliferation (Boros et al., 1998; Ying et al., 2012).

Thus, our cross-species and pan-cancer CNA analysis revealed conserved amplification regions shared by the majority of tumor types studied that are enriched for genes involved in core glycolysis. To aid others in pan-cancer and cross-species CNA signature comparisons, we have created an interactive web-interface resource available at http://systems.crump.ucla.edu/cna_conservations/.

**Discussion**

Chromosomal instability and high glycolysis characterize some of the most aggressive tumors, and the complexity and plasticity of the genomes in aggressive tumors can hinder molecularly targeted therapies (Nakamura et al., 2011; McGranahan et al., 2012; Shi et al., 2012). While the glycolytic changes associated with tumorigenesis were one of the early defining phenotypes of cancer cells (Warburg, 1956), they have not previously been linked to recurrent DNA copy number patterns. Taken together, our experimental and computational data support a model in which glycolysis-linked selective pressures encountered during tumorigenesis (e.g., redox stress and senescence) shape the highly recurrent DNA copy number alterations found in aneuploid human tumors (Fig 6G). We found that CNAs in core glycolysis enzymes (e.g., HK2) and other cancer-linked metabolic enzymes such as TIGAR are coordinately enriched in tumors with distinct CNA signatures. These CNA signatures are predictive of glycolysis, including patient FDG-PET activity and cell line proliferation phenotypes. The strong correlation of CNA-based principle component scores to uptake of the glucose analogue FDG in breast cancer patients and the predictive power of CNA signatures for breast cancer cell metabolism (Fig 3) provide support that the CNAs affecting metabolic gene loci collectively act as a copy number-based driver of metabolic differences. Importantly, in that exogenous expression of hexokinase and enolase enzymes reduced
Figure 6.
Chromosomal instability and aneuploidy—positive and negative impact on tumor cell fitness

Most solid human tumors exhibit both numerical and structural aneuploidy (Holland & Cleveland, 2012). Paradoxically, chromosomal instability can act either as an oncogene or as a tumor suppressor depending on the context (Weaver et al., 2007). Moreover, addition of a single chromosome in MEF cells induces a stress response that impairs proliferation and immortalization (Williams et al., 2008). However, numerical aneuploidy can lead to chromosomal instability (Nicholson et al., 2015) which results in subchromosomal gains and losses as observed in human tumors, in mouse models of cancer, and in immortalized MEF cells. Thus, aneuploidy can cause an initial fitness loss due to the costs of dealing with non-optimized chromosome numbers, gene copy numbers, and resulting proteome imbalances. However, the associated state of chromosomal instability enables further evolution of the genome toward fitness gains through refinement of gene copy number levels. In sum, these fitness gains either outweigh or offset the fitness losses due to aneuploidy. In this context, our data support a model (Fig 6G) in which metabolic selection forces and metabolic loci contribute to the recurrent patterns of DNA copy number alteration observed in human tumors.

Redox stress, biomass accumulation, and associated glycolytic changes in tumorigenesis

Tumorigenesis is a complex, multistage process during which cells must acquire the capability to maintain redox balance while accumulating the macromolecular precursors required for proliferation (DeBerardinis et al., 2008; Hanahan & Weinberg, 2011). Numerous stimuli, including RAS mutations, matrix detachment, altered metabolism, and hypoxia, induce the accumulation of intracellular ROS (Lee et al., 1999; Schafer et al., 2009; Weinberg et al., 2010; Anastasiou et al., 2011). Because increased ROS levels can trigger replicative senescence and subsequent cell cycle arrest (Lee et al., 1999; Takahashi et al., 2006), tumors must maintain pools of reduced glutathione using NADPH in part produced via the pentose phosphate pathway. Additionally, increased levels of ROS can divert glycolytic flux into the pentose phosphate pathway through, for example, oxidation and inhibition of the glycolytic enzyme PKM2, thereby supplying cells with the reducing power and precursors for anaerobic processes (Boros et al., 1998; Anastasiou et al., 2011).
Consistent with this published knowledge on the role of metabolism in tumorigenesis, our study suggests that the metabolic stress associated with senescence (Fig 4A and D) and the metabolic demands of rapid proliferation (Fig 6) are components of the selective pressures underlying recurrent CNA changes.

**Experimentally and computationally deciphering CNA patterns**

Our experimental and bioinformatic approaches complement existing approaches for testing hypotheses for the selection pressures underlying recurrent CNA patterns observed in human tumors. Other CNA-analysis approaches target different resolutions of the genome. Statistical algorithms such as GISTIC (Genomic Identification of Significant Targets in Cancer) have identified many strong individual driver genes and candidate regions (Mermel et al., 2011). Integrating CNA data with RNA knockdown screens and gene expression data has further identified driver genes missed by statistical analysis of CNA data alone (Sanchez-Garcia et al., 2014). RNA knockdown-based analyses have also been used to support a more systems-level model in which the selection for amplification and deletion of a particular DNA region is based on the cumulative effects of many positive and negative fitness gains from multiple genes within that genomic region (Solimini et al., 2012; Cai et al., 2016). Subsequent computational extensions have incorporated somatic mutation patterns to infer the cumulative impact of co-localized genes on fitness, and to successfully predict whole chromosome and chromosome-arm resolution-level CNAs (Davoli et al., 2013).

Our approach using phenotypic data, functional gene sets, and cross-species syntenic mapping has yielded additional insight into the selective pressures shaping tumor CNA patterns, namely coordinated alteration of genes involved in glycolytic metabolism. Incorporation of copy number data from genetically engineered mouse models allowed synteny constraints to substantially reduce passenger amplifications and define a minimal collection of stringently conserved copy number regions. Additionally, to our knowledge, our experimental approach is the first reported system in which CNA and associated phenotypes have been followed and repeatedly sampled as non-immortalized cells undergo spontaneous genomic instability and proceed from a diploid state to an immortalized aneuploid state. Using this approach, we were able to validate genomic regions that are (i) associated with increased glycolysis and to a lesser but significant extent proliferation, (ii) enriched for genes from the core glycolysis pathway, and (iii) conserved in both human tumors and mouse models of cancer. Metabolic pathways are known to be coordinately regulated by modest changes in mRNA expression of functionally related genes (Mootha et al., 2003; Palaksas et al., 2011). The coordinated alterations of metabolic genes at the DNA level adds an additional mechanism, namely conserved sets of CNA changes, by which glycolysis is dysregulated to promote tumorigenesis.

A strength of our PCA-based approach is the ability to unbiasedly reveal distinct CNA subsignatures within a tumor type. Observed subsignatures, confirmed using an independent clustering analysis, were found to be associated with previously known pathology- or profiling-defined tumor subtypes (e.g., basal/luminal CNA signatures in BRCA) (Bergamaschi et al., 2006) (Figs 1B and EV2A–D). Across multiple tumor types, loss of p53 function through p53 mutation is associated with the high breakpoint signature A pattern. In contrast, loss of the p53/ARF axis via other mechanisms (MDM2 amplification or CDKN2A deletion) in BRCA human tumors is associated with a different CNA signature (signature B) that in general has fewer breakpoints. In our experimental follow-up, the MEF system recapitulated a two-signature pattern. Notably, the two mouse signatures and their associated phenotype strengths were defined by the initiating loss of tumor suppression event, namely Trp53 mutation versus either Mdm2 amplification or Cdkn2a loss. Thus, while the consequences of TP53 mutation and “MDM2 amplification/CDKN2A loss” are considered functionally similar and therefore mutually exclusive (Wade et al., 2013), our findings indicate they are not fully equivalent in terms of genomic instability and subsequent metabolic evolution. The tolerance of more highly disrupted and rearranged genomes upon p53 mutation appears to allow more flexibility in the evolution of aneuploid cancer genomes, thereby resulting in stronger glycolysis and somewhat enhanced proliferation. The specific combinations of CNA changes occurring in enzyme isoforms defining a metabolic pathway may be considered “onco-metabolic isoenzyme configurations” with differential potency, and the sets of combinations possible may be limited in part by the degree or specifics of tolerance to genomic rearrangements.

In summary, our work illustrates the value of cross-species comparisons in the analysis of DNA copy number data, much as recent pan-cancer and integrated genomic approaches have uncovered novel cancer subtypes and driver genes (Gatza et al., 2014; Hoadley et al., 2014). The broken chromosome synteny between human and mouse genomes reduced the size of potential driver regions by fivefold on average, and identified a relatively small number of amplification and deletion regions highly conserved between mouse and human (Figs 2D and 4E). These conserved regions are consistently found across the majority of tumor types examined, have subchromosomal scale with a median size on the order of 10 megabase pairs, and include the glycolysis pathway enzyme CNAs observed in our experimental work.

**Therapeutic and diagnostic implications**

The most copy number aberrant tumors tend to have fewer point mutations in canonical oncogenes (e.g., KRAS, Fig EV2A and C) and less canonical oncogene amplification (e.g., MYC, Fig 1B). Hence, genomic instability and subsequent coordinate alterations in multiple genes within a functional pathway may provide an alternate, more complex, pathway to acquisition of aggressive tumor phenotypes—with tumor evolution and selection guiding the trajectory (Ciriello et al., 2013). In that KRAS mutation and MYC amplification can drive glycolysis (Ying et al., 2012; Dang, 2013), the findings that signature A tumors are de-enriched in these events relative to signature B tumors and enriched for glycolysis gene loci CNA amplifications support that tumor cells can meet their metabolic demands through distinct mechanisms, or combinations thereof. Future models of the most aggressive cases of cancer will need to incorporate aspects of spontaneous genomic instability (mediated by distinct instability mechanisms) and resulting copy number alterations. Translationally, understanding how CNA patterns alter cancer genomes and impact cancer phenotypes will aid in the identification of metabolic or other “hard-wired” vulnerabilities that can
be therapeutically targeted. Furthermore, the relative stability of DNA samples, combined with the growing linkage between highly recurrent copy number changes and phenotypes, supports the potential for molecular classification and diagnostic tests based on DNA copy number patterns (Hieronymus et al., 2014; Cai et al., 2016).

**Materials and Methods**

**Cell culture and mouse strains**

**Tissue culture**

CD1 mouse embryonic fibroblasts, E14.5, were purchased from Stem Cell Technologies. p53<sup>fl/fl</sup> MEFs were obtained at day E14.5 from p53<sup>fl/fl</sup> (FVB.129P2-Trp53<sup>tm1Brd</sup>/Nei) crossed with C57BL/6-129/SV mice. MEF cells were maintained in Dulbecco’s modified Eagle’s medium without pyruvate and supplemented with 10% FBS and 1% SPF. Cells were lifted and re-plated at a density of 9 × 10<sup>5</sup> viable cells/60-mm dish every 3–4 days (i.e., 3T9 protocol; Todaro & Green, 1963). Cells were grown in atmospheric oxygen unless otherwise indicated. Breast cancer cell lines were obtained from the laboratory of Frank McCormick and extensively profiled both genomically and transcriptionally by the laboratory of Joe Gray (Neve et al., 2006; Hong et al., 2016).

**Genetic engineering**

Overexpression of HK1, HK2, kinase-dead HK2 (D209A/D657A), or ENO2 glycolysis enzymes and the MYC oncogene or the control protein RFP in CD1 MEFs was accomplished by transduction of non-immortalized cells with pDS-FB-neo retrovirus, followed by selection in 600 μg/ml G418. Deletion of p53 in p53<sup>fl/fl</sup> MEFs was induced by infection of non-immortalized cells with either retroviral Cre-GFP or Cre-ERT2 plus treatment with 1 μM 4-OHT.

**ROS protection**

To protect cells from reactive oxygen species (ROS), cells were cultured either at physiological oxygen concentrations (3% O<sub>2</sub>), with 250 U/ml catalase from bovine liver (Sigma-Aldrich), or with both 3% O<sub>2</sub> and catalase.

**Genome profiling**

**Array CGH profiling**

Genomic DNA was harvested using the DNeasy kit (Qiagen). DNA from MEF lines and reference genomic DNA from C57BL/6J mouse tissue were hybridized to Agilent SurePrint G3 Mouse CGH 4 × 180 k CGH microarray chips at the UCLA Pathology Clinical Microarray Core. Bioconductor analysis tools were used for data processing: Moving minimum background correction and print-tip loess normalization were performed in snapCGH package (Smith et al., 2009); circular binary segmentation (with a minimum of three markers per segment) was performed on smoothed and log2-transformed copy number profiles using DNAcopy package (Seshan & Oshlens, 2016); segmented data were converted into a matrix by genes for downstream analyses using mus musculus 9 RefSeq reference genome from 2011.08.11 in CNTools (Zhang, 2016). Copy number profiles are presented using the Integrative Genomics Viewer (IGV) (Thorvaldsdottir et al., 2013). CNA public datasets and TCGA tumor type abbreviations are available in the Data availability section.

**Bioinformatic and statistical analysis**

**PCA**

Principal component analysis (PCA) was performed using the mean-centered matrix of CNA values per gene locus. Genes with identical profiles across samples were collapsed to a single representative gene. Techniques such as PCA and the related singular value decomposition (SVD) have been applied to copy number data previously (Sankaranarayanan et al., 2015).

**Mutation and tumor subtype enrichment analysis**

To test for enrichment of mutations or tumor subtypes within CNA-defined PC scores, TCGA tumors were queried for mutations and tumor subtype designations using the cBioPortal for Cancer Genomics (Gao et al., 2013). For each tumor type, we included all genes with significant q-values as calculated by MutSigCV (Lawrence et al., 2013). We also included the gene POLE, a catalytic subunit of DNA polymerase epsilon, in our analysis because of its frequent mutation in UCEC (The Cancer Genome Atlas Research Network, 2013). Molecular subtypes tested were as follows: BRCA: basal, luminal, claudin-low, and HER2-enriched (The Cancer Genome Atlas Research Network, 2012b); LI Jad: bronchioid, magnoïd, and squamoid (The Cancer Genome Atlas Research Network, 2014); LI Jad: basal, classical, primitive, and secretory (The Cancer Genome Atlas Research Network, 2012a); OV: proliferative, immunoreactive, differentiated, and mesenchymal (The Cancer Genome Atlas Research Network, 2011); and UCEC: PO LE ultramutated, microsatellite instability hypermutated, copy number low, and copy number high (The Cancer Genome Atlas Research Network, 2013). Tumors were sorted by their PC1 score, and we calculated a Kolmogorov–Smirnov statistic against the expected distribution of mutations or tumor subtypes. The statistical significance of enrichment was determined by permutation analysis.

**Hierarchical clustering**

The TCGA tumors were hierarchically clustered using the heatmap package in R (Kolde, 2015). Prior to clustering, the CNA values were filtered for gene loci that had zero values across all tumor samples. Non-centered and non-scaled tumor samples were then clustered using centered Pearson correlation distance and Ward’s method (with dissimilarities squared before cluster updating). The strength of concordance between hierarchical clustering results and PCA-based signatures was assessed using the hypergeometric P-value.

**Consistency signatures**

Consistency signatures of conserved amplifications and deletions regions were determined using stringent consistency criteria. The signed absolute minimum consistency score (SAMCS) was defined as non-zero when all CNA summary signatures have the same sign across all tumor types, and the score is derived from the absolute value-based minimum summary metric and then re-signed positive for amplification or negative for deletion. Consistent amplifications or deletions were combined into a “consistent region”, when absolute SAMCS values greater than 0.05 spanned at least 1 Mbp. Any
two consistent regions separated by < 1 Mb were combined into a single consistent region.

Enrichment analysis and weighted gene voting (WGV)
Metabolic pathway enrichment analysis (gene set enrichment analysis, GSEA; Subramanian et al, 2005) and pathway-specific weighted gene voting (WGV) prediction analysis (Golub et al, 1999) were performed using 75 metabolic pathways defined by the Kyoto Encyclopedia of Genes and Genomes (KEGG) database (Kanehisa et al, 2014), using pathways with seven or more measured genes. In the RNA-based enrichment analysis, we included a gene set consisting of genes from the glycolysis and pentose phosphate pathways that were upregulated in FDG-high BRCA tumors, as defined by our previous work (Palaskas et al, 2011). For CNA data, an expanded run of enrichment analysis was performed using 1,321 canonical pathways (CP) of seven or more measured genes as defined by the Broad Institute’s Molecular Signatures Database (MSigDB). For KEGG-based metabolic pathway enrichment analysis of CNA data, we collapsed metabolic isoenzyme loci (genes with the same enzyme activity; Enzyme Commission [EC] numbers) that were within 100 kilobases from each other into a single representative locus. For mRNA expression analysis, metabolic isoenzymes were not collapsed. In CNA-based enrichment analysis using gene-based versions of the genome consistency signatures defined above, enrichment scores were calculated through the ranked set of consistently amplified genes since after this point the genes that are not consistent across signatures have a consistency value of zero and accordingly have tied ranks. Consistency regions were stepwise restricted by sequentially adding human, and corresponding tissue-matched mouse models of cancer, signatures based on their decreasing tissue type PC1 value from Fig 1A. To combine BRCA and LU into one enrichment analysis for mRNA data, genes were ordered by their average rank in BRCA and LU tumor types. For WGV predictions of metabolic phenotypes, t-scores were used as gene weights. To calculate permutation P-values, we calculated the fraction of 1,000 randomly chosen gene sets of equal size that gave average gene set rankings (column 2 of Fig 3C) better than the true gene set. False discovery rate (FDR) q-values were calculated using the Benjamini-Hochberg procedure. To increase statistical stringency, FDR values for each individual glycolysis–gluconeogenesis gene set (KEGG Glycolysis-Gluconeogenesis, KEGG Core Glycolysis, KEGG Glycolysis-Gluconeogenesis & Pentose Phosphate Pathway, Reactome Gluconeogenesis, Reactome Glucose Metabolism, Reactome Glycolysis, Biocarta Glycolysis Pathway) were calculated while removing the other glycolysis–gluconeogenesis gene sets. CNA and metabolite changes were visualized in the context of metabolic pathway structure using Cytoscape (Smoot et al, 2011).

Core glycolysis and glycolysis-associated gene list (from Fig 1F)
HK—hexokinase (HK1, HK2, HK3, HK4/glucokinase/GCK); GPI—glucose-6-phosphate isomerase; G6PC—glucose-6-phosphatase catalytic subunit (G6PC, G6PC2); FBP—fructose-biphosphatase (FBP1, FBP2); PFK—phosphofructokinase (PFKL—liver type, PFKM—muscle, PFKP—platelet); PFKFB—6-phosphofructo-2-kinase/fructose-2,6-biphosphatase (PFKFB1, PFKFB2, PFKFB3, PFKFB4); TIGAR—TP53 induced glycolysis regulatory phosphatase; ALDO—aldolase, fructose-biphosphatase (ALDOA, ALDOB, ALDOC); TPI1—triosephosphate isomerase 1; GAPDH—glyceraldehyde-3-phosphate dehydrogenase (GAPDH, GAPDHS—spermatogenic); PGK—phosphoglycerate kinase (PGK1, PGK2); PGAM—phosphoglycerate mutase (PGAM1, PGAM2, PGAM4, BPGM—bisphosphoglycerate mutase); ENO—enolase (ENO1, ENO2, ENO3); PK—pyruvate kinase (PKLR—liver and red blood cell, PKM2—muscle); LDH—lactate dehydrogenase (LDHA, LDHB, LDHC, LDHAL6A–LDH A-like 6A, LDHAL6B–LDH A-like 6B); and PDH—pyruvate dehydrogenase (PDHA1, PDHA2, PDHB, DLD—dihydrolipoamide dehydrogenase, DLAT—dihydrolipoamide S-acetyltransferase).

Metrics
Signal-to-noise ratio (SNR) = (µ1 − µ2) / (√(σ12 + σ22)), t-score = (µ1 − µ2) / sqrt[(σ12/n1 + σ22/n2)], where µ = mean, σ = standard deviation, n = number of samples.

Correlation of mRNA and CNA
Copy number alteration and mRNA expression levels for genes found in the cross-species conserved regions in Fig 4E were compared by calculating the Spearman rank correlation coefficient. BRCA, LU, and OV samples with paired mRNA and CNA data were included. UCEC tumors were excluded because there were not sufficient UCEC samples with paired RNA expression data (26% of all samples with CNA data). Known oncogenes were identified by comparison with the Catalogue Of Somatic Mutations In Cancer database (COSMIC, http://cancer.sanger.ac.uk/cosmic; Forbes et al, 2015).

Senescence score
A summary senescence score for each MEF subline was calculated by subtracting the area under the MEF’s growth curve, Z(x), in log scale, from the area under an ideal growth curve, Y(x) (i.e., consistent growth at the fastest observed rate). The area difference was then averaged by dividing by the passage number, p, and log2-transformed, resulting in a normally distributed score.

\[ \text{Senescence Score} = \log_2 \left( \frac{\sum_{i=2}^{p} \left( \frac{Y_{i-1} + Y_i}{2} - \frac{Z_{i-1} + Z_i}{2} \right) }{p} \right) \]

Integrated CNA
A genomic instability score termed “integrated CNA” was calculated by summation of the Circular Binary Segmentation algorithm-inferred absolute mean copy number of segments multiplied by the length of each segment.

\[ \text{Int.CNA}_{\text{sample}} = \sum_{\text{segments}} |\text{segment end} - \text{segment start}| \times \text{segment mean} \]

# base pairs in sample

Box and whisker plots
In box and whisker plots, the box represents the median, as well as the first and third quartiles, and the whisker indicates the extreme values within 1.5 times the inter-quartile range. In cases where the number of samples permitted, individual values are superimposed as jitter plots.

Statistical tests
Indicated P-values were calculated using (i) Student’s t-test for normally distributed data (with normality confirmed by the P-value
of the Shapiro–Wilk test being > 0.05 for both datasets under comparison); (ii) Mann–Whitney U-test for non-normally distributed data; (iii) hypergeometric distribution P-values; and (iv) permutation-based approaches, as described in the figure legends. For data with a single data point in one comparison group, the z-score was used.

**Propidium iodide staining**

Mouse embryonic fibroblasts cells were washed in cold PBS and then fixed in ice-cold 70% ethanol. Fixed cells were washed in PBS and then incubated for 15 min in PBS with 20 μg/ml propidium iodide and 0.1% (v/v) Triton X-100. Data were acquired using a FACSCalibur (Becton Dickinson) analytic flow cytometer in the UCLA Jonsson Comprehensive Cancer Center and Center for AIDS Research Flow Cytometry Core Facility. Cells were gated using forward scatter and side scatter to remove debris and dead cells, and 10,000 cell events were recorded.

**Spectral karyotyping**

Exponentially growing MEF cells were exposed to colcemid (0.04 μg/ml) for 1 h at 37°C and to hypotonic treatment (0.075 M KCl) for 20 min at room temperature. Cells were fixed in a mixture of methanol and acetic acid (3:1 by volume) for 15 min, and then washed three times in the fixative. Slides were prepared by dropping the cell suspension onto wet slides followed by air-drying. Slides were processed for spectral karyotyping (SKY) according to the manufacturer’s protocol with slight modifications using mouse paint probes (ASI, Vista, CA). Images were captured using Nikon 80i microscope equipped with spectral karyotyping software from ASI, Vista, CA; 12–18 metaphases were karyotyped from each cell line.

**Immunoblot analysis**

Cells were lysed in modified RIPA buffer (50 mM Tris–HCl (pH 7.5), 150 NaCl, 10 mM β-glycerophosphate, 1% NP-40, 0.25% sodium deoxycholate, 10 mM sodium pyrophosphate, 30 mM sodium fluoride, 1 mM EDTA, 1 mM vanadate, 20 μg/ml aprotinin, 20 μg/ml leupeptin, and 1 mM phenylmethylsulfonyl fluoride). Whole-cell lysates were resolved by SDS–PAGE on 4–15% gradient gels and blotted onto nitrocellulose membranes (Bio-Rad). Membranes were blocked overnight and then incubated sequentially with primary and either HRP-conjugated (Pierce) or IRDye-conjugated secondary antibodies (Li-Cor). Blots were imaged using the Odyssey Infrared Imaging System (Li-Cor). Protein levels were quantitated using ImageJ (http://imagej.nih.gov/ij/). Primary antibodies used for Western blot analysis included hexokinase 1 (2024, Cell Signaling Technology), hexokinase 2 (2867, Cell Signaling Technology), p53 (NB200-103, Novus Biologicals), and enolase 2 (8171, Cell Signaling Technology).

**Glucose consumption and lactate secretion measurements**

**BRCA cell lines**

Lactate secretion rates of breast cancer cell lines were measured from the culture media using a colorimetric assay kit (BioVision) (Hong et al, 2016).

**MEFs**

Glucose consumption and lactate secretion rates of MEFs were measured using a BioProfile Basic bioanalyzer (NOVA Biomedical). Data were normalized to the integrated cell number, which was calculated based on cell counts at the start and end of the time course and an exponential growth equation. Because the proliferation rates of MEF sublines vary, each cell line was seeded at the appropriate density so as to give an integrated cell number of approximately 6.5 × 10⁶ cells in a 6-well plate. All samples were run as biological triplicates, and consistent results were seen in multiple independent experiments.

**Mass spectrometry-based metabolomic analyses**

**Sample preparation**

Mouse embryonic fibroblast sublines were seeded onto 6-well plates, and after 24 h, media was replaced with media containing 4.5 g/l [1,2-¹³C]-labeled glucose. Sample collection occurred after 24 h of culture in the labeled glucose media. For intracellular metabolite analysis, cells were washed with ice-cold 150 mM ammonium acetate (NH₄AcO) pH 7.3 and metabolites extracted in 1 ml ice-cold 80% MeOH. The cells were quickly transferred into a microfuge tube, and 10 nmol norvaline was added to the cell suspension for use as an internal standard. The suspension was subsequently vortexed three times over 15 min and then spun down at 4°C for 5 min. The supernatant was transferred into a glass vial, the cell pellet was re-extracted with 200 μl ice-cold 80% MeOH and spun down and the supernatants were combined. Metabolites were dried at 30°C under vacuum and re-suspended in 50 μl of 70% acetonitrile (ACN). For cell culture media metabolite analysis (footprint profiling), 20 μl of cell-free media samples was collected. Metabolites were extracted by adding 300 μl ice-cold 80% methanol, followed by vortexing three times over 15 min, and centrifugation for 10 min at 13,000 rpm at 4°C. The supernatant was transferred to a fresh tube, dried using a vacuum evaporator, and re-suspended in 50 μl of 70% acetonitrile (ACN); 5 μl was used for mass spectrometry-based analysis.

**Mass spectrometry runs**

Samples were run on a Q-Exactive mass spectrometer coupled to an UltiMate 3000RSLC UHPLC system (Thermo Scientific). The mass spectrometer was run in polarity switching mode (+3.00 kV/−2.25 kV) with an m/z window ranging from 65 to 975. Mobile phase A was 5 mM NH₄AcO, pH 9.9, and mobile phase B was ACN. Metabolites were separated on a Luna 3 μm NH₂ 100 Å (150 × 2.0 mm) (Phenomenex) column. The flow was kept at 200 μl/min, and the gradient was from 15% A to 95% A in 18 min, followed by an isocratic step for 9 min and re-equilibration for 7 min.

**Data analysis**

Metabolites were detected and quantified as area under the curve (AUC) based on retention time and accurate mass (≤ 3 ppm) using the TraceFinder 3.1 (Thermo Scientific) software. Relative amounts of metabolites between various conditions, percentage of metabolite isotopomers (relative to all isotopomers of that metabolite), and percentage of labeled metabolite molecules (isotopomer M1 and greater, relative to all isotopomers) were calculated and corrected.
for naturally occurring $^{13}$C abundance (Yuan et al., 2008). Footprinting data were normalized to the integrated cell number as described above, and intracellular metabolite concentrations were normalized to the number of cells present at the time of extraction. All samples were run as biological triplicates, and consistent results were seen in independent experiments. Our analysis focused on metabolite level, percent isotopomer, and percent labeled metabolite measurements with ANOVA $P$-values across the sample panel of < 0.05 in individual experiments, and Pearson correlation coefficients across all samples and between independent experiments of > 0.5.

**Patient tumor samples and quantitative FDG-PET imaging**

Breast cancer patient samples with imaged FDG uptake within 4 weeks prior to surgery, excluding patients with secondary breast cancers and recurrent disease, were collected surgically and processed as previously described (Palaskas et al., 2011). Of eighteen tumors collected in the original study for RNA microarrays (Palaskas et al., 2011), ten samples had sufficient remaining frozen tissue for array CGH profiling. None of the patients received systemic therapy or radiation prior to imaging. $^{18}$FDG tumor uptake was quantified as standardized uptake values (SUVs) and showed the expected wide dynamic range (3.8–18.5). There was no significant difference in patient age, tumor volume, and lymph node involvement between the groups of FDG-high and FDG-low breast cancers. Breast cancers with high $^{18}$FDG-PET SUVs frequently lacked expression of the estrogen receptor (ER) and the progesterone receptor (PR), but hormone receptor-negative tumors were also represented among the tumors with the lowest FDG uptake (Palaskas et al., 2011). We excluded lobular breast carcinomas, because they have been shown to take up less FDG than ductal carcinomas (Avril et al., 2001). We excluded large breast carcinomas (> 5 cm) and breast carcinomas with multifocal FDG uptake because our protocol did not include tissue autoradiography to direct the molecular tissue analysis to areas of distinct radiotracer retention. This study was approved by the Institutional Review Board (IRB) of Memorial Sloan-Kettering Cancer Center, and all participating patients signed the informed consent.

**Data availability**

**CNA dataset**


**CNA conservation web resource**

An interactive website for user-defined pan-cancer and cross-species CNA conservation analysis to perform analysis analogous to that in Figs 1C, 2D, and 4E using any combination of tens of available CNA signatures from human tumors and mouse models (and additional signatures as they become available) and/or the inclusion of uploaded CNA signatures (http://systems.crump.ucla.edu/cna_conservation/). Signatures and genome reference files used in the interactive website are additionally available through the Biostudies repository, accession number S-BSTT7.

**Metabolomics dataset**

Provided in Table EV5.

**TCGA CNA dataset**

The Cancer Genome Atlas (TCGA) CNA profiles were downloaded from the TCGA portal in September 2012 (https://cancergenome.nih.gov/). Copy number profiles obtained were pre-processed level 3 data based on human genome 19, with copy number variations (CNVs) removed. TCGA tumor type abbreviations and number of samples analyzed: bladder urothelial carcinoma (BLCA, 97 samples), brain lower grade glioma (LGG, 181), breast invasive carcinoma (BRCA, 873), colon adenocarcinoma (COAD, 447), glioblastoma multiforme (GBM, 593), head and neck squamous cell carcinoma (HNSC, 308), kidney renal clear cell carcinoma (KIRC, 539), lung adenocarcinoma (LUAD, 368), lung squamous cell carcinoma (LUSC, 359), ovarian serous cystadenocarcinoma (OV, 584), prostate adenocarcinoma (PRAD, 171), rectum adenocarcinoma (READ, 168), skin cutaneous melanoma (SKCM, 256), stomach adenocarcinoma (STAD, 162), thyroid carcinoma (THCA, 333), and uterine corpus endometrial carcinoma (UCEC, 492).

**TCGA mRNA expression dataset**

The Cancer Genome Atlas mRNA expression data were downloaded from the TCGA portal in May 2014. Gene-based mRNA expression levels were pre-processed, normalized Level 3 RNA Seq V2 RSEM values. TCGA tumor type abbreviations and number of samples analyzed: BRCA (865), LIJAD (357), LUSC (358), OV (263).

**Mouse tumor model CNA datasets**

The genetically engineered mouse models with characterized CNA were obtained from public datasets: mammary (breast) tumors (Brca, 57 samples, GSE30710; 62 samples, GSE43997; 44 samples, GSE27101) (Drost et al., 2011; Herschkowitz et al., 2012); melanoma (Skcm, 30 samples, GSE58265) (Viros et al., 2014); glioblastoma/high-grade astrocytoma (Gbm, 72 samples, GSE22927) (Chow et al., 2011); and prostate tumors (Prad, 18 samples GSE35247; 55 samples, GSE61382) (Ding et al., 2012; Wanjala et al., 2015). Additionally, in vitro epithelial murine cell lines modeling human carcinomas were obtained from public datasets: transformed colon cells (Coad, seven samples, GSE70970) and transformed bladder and kidney cells (Blca and Kirc, 6 and 7 samples, GSE45128; Padilla-Nash et al., 2013). Abbreviated mouse tumor names match the corresponding tissue-based human tumor abbreviations from the TCGA datasets. The data were obtained from GEO and segmented using the algorithm described above. Datasets for which no mm9 genome annotation was available on the repository were lifted over to mm9 using UCSC web tools (Rosenbloom et al., 2015).

**Expanded View** for this article is available online.

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Author contributions
NAG, AM, AL, AC, DB, HRC, IKM, and TGG designed the study. NAG, AM, AL, AC, MF, SC, SZ, AD, LB, JTH, DB, and HRC performed experiments. NAG, AM, AL, AC, NGB, JC, NP, NS, DB, and TGG performed bioinformatic analysis. NAG, AM, AL, CH, CN, RQ, HW, MM, HRC, and TGG provided or derived reagents. NAG, AM, AL, DB, HRC, and TGG performed and analyzed metabolomics. AM and ASM performed spectral karyotyping. EP, SML, and IKM generated patient data. NAG, AM, DB, HRC, IKM, and TGG wrote the manuscript. The laboratories of HRC and IKM contributed equally. Experiments were performed at UCLA.

Conflict of interest
The authors declare that they have no conflict of interest.

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Tumor CNAs reflect metabolic selection

Nicholas A Graham et al


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Tumor CNAs reflect metabolic selection


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Molecular Systems Biology 13:914 | 2017 25