A Network Model of a Cooperative Genetic Landscape in Brain Tumors

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MALIGNANT GLIOMAS, with disproportionately high morbidity and mortality, are among the most devastating of human tumors. Particular genomic alterations are fundamental to both their formation and their malignant progression. Although genomic instability (BOX) lends a dynamic character to the human glioma genome, gliomas harbor recurrent chromosomal alterations (Box). Chromosomal alterations presumably exert their tumor-promoting effect on glioma cells by modifying the expression or function of distinct genes, which map to those alterations, so as to deregulate growth factor signaling and survival pathways. For many chromosomal alterations, the biologically relevant target genes remain to be discovered.

Oncogenic research on gliomas has focused on the tumor-promoting or tumor-suppressive function of target genes within individual chromosomal alterations. However, these alterations do not exist in isolation, nor do single genes account for gliomagenesis.2-4,5 Chromosomal alterations precurrent chromosomal alterations in glioma genome, gliomas harbor recurrent chromosomal alterations—that involves altered regions (“territories”) on chromosomes 1p, 7, 8q, 9p, 10, 12q, 13q, 19q, 20, and 22q (false-discovery rate–corrected P<.05). A network model shows that these territories harbor genes with putative synergistic, tumor-promoting relationships. The coalteration of the most interactive of these genes in glioblastoma is associated with unfavorable patient survival. A multigene risk scoring model based on 7 landscape genes (POLD2, CYCS, MYC, AKR1C3, YME1L1, ANXA7, and PDCD4) is associated with the duration of overall survival in 189 glioblastoma samples from TCGA (global log-rank P=.02 comparing 3 survival curves for patients with 0-2, 3-4, and 5-7 dosage-altered genes). Groups of patients with 0 to 2 (low-risk group) and 5 to 7 (high-risk group) dosage-altered genes experienced 49.24 and 79.56 deaths per 100 person-years (hazard ratio [HR], 1.63; 95% confidence interval [CI], 1.10-2.40; Cox regression model P=.02), respectively. These associations with survival are validated using gene expression data in 3 independent glioma studies, comprising 76 (global log-rank P=.003; 47.89 vs 15.13 deaths per 100 person-years for high risk vs low risk; Cox model HR, 3.04; 95% CI, 1.49-6.20, P=.002) and 70 (global log-rank P=.008; 83.23 vs 34.16 deaths per 100 person-years for high risk vs low risk; HR, 3.86; 95% CI, 1.59-9.35; P=.003) high-grade gliomas and 191 glioblastomas (global log-rank P=.002; 83.23 vs 34.16 deaths per 100 person-years for high risk vs low risk; HR, 2.27; 95% CI, 1.44-3.58; P<.001).

Conclusions The alteration of multiple networking genes by recurrent chromosomal aberrations in gliomas deregulates critical signaling pathways through multiple, cooperative mechanisms. These mutations, which are likely due to nonrandom selection of a distinct genetic landscape during gliomagenesis, are associated with patient prognosis.

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The goal is to reduce the complexity and functional interactions among genes.

Using more than 500 glioma samples—220 glioblastomas, 38 anaplastic astrocytomas, and 188 and 207 glioblastomas, respectively, were obtained from the Open Access and Controlled-Access Data Browser (http://smd.stanford.edu/). Agilent Human Genome CGH Microarray 244A gene dosage data, Affymetrix HT Human Genome U133A Array Plate Set gene expression data, and clinical data for 219, 188, and 207 glioblastomas, respectively, were obtained from the Open-Access and Controlled-Access Data Tiers Portal (http://tcga-data.nci.nih.gov/).
Gene dosage and gene expression data, 175 samples with gene expression and clinical data, 175 samples with gene dosage and gene expression data, and 172 samples with all 3 data types. Agilent Human Genome CGH Microarray 244A data from TCGA were background corrected using subtraction then replacement of negative or zero values with half the minimum of the positive corrected values on the array, then were LOWESS normalized using the “limma” package for R. Data from TCGA were similarly processed as described above using CBS via the “snapCGH package” for R.

Genomic Data Processing
Gene dosage data deposited into the SMD were background subtracted and included for downstream analysis if the reference channel intensity was at least 2.5 times the observed background and Pearson correlation was greater than 0.6 for the sample and reference channels. Locally weighted least squares (LOWESS) normalization was performed using SNOMAD (http://pervnerlab.kennedykrieger.org/snomad.htm) and the Institute for Genomic Research TM4 Microarray Data Analysis System (TIGR MIDAS) (http://www.tm4.org/midas.html). The GoldenPath Human Genome Assembly (http://genome.ucsc.edu/) was used to map fluorescence ratios of the arrayed human cDNAs to chromosomal positions. Circular binary segmentation (CBS)13 from the R package “DNAcopy” (http://www.r-project.org/) was used to estimate segmented regions of equal dosage along each chromosome using 33,245 clones that passed initial data normalization and filtering. To increase the number of estimated segmented regions obtained by CBS and thereby the sensitivity of the algorithm, the \( \alpha \) level was set to .05 rather than the default of .01. Segmented gene dosage values were deemed changed compared with normal human reference DNA if they fell outside of the \( \pm 3 \)-SD range of all segmented log₂ values of control (glioma and normal brain) self-to-self hybridizations, resulting in classification as copy number “gain” for gene dosage values \( >0.2135 \) and “loss” for gene dosage values \( <-0.2135 \). Agilent Human Genome CGH Microarray 244A data from TCGA were background corrected using subtraction then replacement of negative or zero values with half the minimum of the positive corrected values on the array, then were LOWESS normalized using the “limma” package for R. Data from TCGA were similarly processed as described above using CBS via the “snapCGH package” for R.

Gene expression data obtained from TCGA,2 UCLA,13 and Lee et al14 were normalized as previously described. Preprocessed gene expression data obtained from TCGA,2 UCLA,13 and Lee et al14 were normalized as previously described.

Gene Dosage Data Analysis
We developed a new permutation-based approach, which calculates the probabilistic fit for the coincidence of distinct chromosomal alterations. Gene-by-gene CBS dosage data were averaged for each of 802 chromosomal cytogenetic bands (Box) corresponding to standard chromosomal high-resolution banding, based on an International System for Human Cytogenetic Nomenclature (ISCN) 850 chromosome ideogram.16 For each tumor \( i \) and cytoband \( j \), measurements \( x[i,j] \) were scored +1 for gain, −1 for loss, and 0 otherwise (ie, unchanged). Scores were mapped into a data matrix \( x \) with tumors and cytobands ordered as rows and columns, respectively. The association of alterations within each possible combination of 2 cytobands \( (i,j) \) was assessed based on computing the score \( S[i,j] \) as follows:

\[
S[i,j] = \frac{\sum x[i,1] \times x[i,j]}{n[i,j]},
\]

where \( n[i,j] \) indicated the number of tumors with both \( i \) and \( j \) altered, and \( x[i,1] \times x[i,j] \) scores +1 if cytobands \( i,j \) for tumor \( t \) are both amplified or both deleted, −1 if one is amplified and the other deleted, and 0 otherwise (ie, both unchanged or only one cytoband altered). One thousand permutations were applied within the columns of the data matrix to estimate the false-discovery rate (FDR). Associations of pairs of cytobands with an FDR <0.05 were considered significant. The same approach was applied to the gene dosage data of individual candidate genes.

Gene Dosage and Expression Integration
For the Stanford samples, gene dosage and gene expression data (expression in >80% of tumors) were available for 15,608 clones that mapped to 10,351 genes, 7509 of which had either losses or gains in at least 2 tumors. For the latter, signal-to-noise (s2n) ratios were computed to assess the influence of dosage alteration for each gene on its transcript,17-19 defined by the difference of the means \( m \) of expression levels in the groups of gene dosage–altered \( m_1 \) and –unaltered \( m_2 \) samples, divided by the sum of standard deviations \( s \) of expression levels in both groups \( s_1 \) and \( s_2 \), respectively. The above data matrix \( x \) was transformed such that gene dosage change \( c \) was assigned g for gain, l for loss, and u otherwise. The significance of the s2n ratio for each gene was estimated by randomly permuting the gene alteration labeling vector 10,000 times. We further determined corresponding FDR-adjusted
q values using the “qvalue” package for R (default parameters), allowing for a more straightforward interpretation of statistical significance in the context of multiple hypothesis testing. False-discovery rate estimation was performed by setting fixed cutoffs and estimating the FDR by permutations as previously described. These computations were done separately for l vs n and g vs n. For clones with alterations in both directions, s2n ratio and corresponding q value for the more recurrent change were chosen. Similar s2n calculations and permutation-based P value computations were conducted to test and validate associations between gene dosage and gene expression for selected candidate genes using TCGA gene dosage and expression data. Due to the intentional selection of these genes for hypothesis testing in TCGA data, FDR was controlled using the method of Benjamini and Hochberg, a limiting case of q value correction.

Gene Set Enrichment Analyses and Modeling of Gene Interactions
Gene set enrichment analyses, using hypergeometric testing for the set of genes with significant gene dosage–gene expression relationships, were executed using Ingenuity Pathways Analysis (IPA) (Ingenuity Systems, Mountain View, California) to identify overrepresented biological functions and signaling pathways from the IPA database for this gene set with right-tailed \( P < .05 \). Ingenuity Pathways Analysis was also used to identify direct and indirect physical (protein–protein) and functional interactions between genes with significant gene dosage–gene expression relationships. An initial gene set was first overlaid onto the set of all cataloged IPA interactions and “focus genes” required in the IPA algorithm were identified as the subset having direct interactions with other genes in the initial set. The specificity of a gene’s interactions was calculated as the percentage of its interactions with other significant genes in the initial set. Genes with the highest specificity of connections were used for the initiation and growth of biological networks with a maximum of 35 genes. Networks of highly interconnected genes were identified by statistical likelihood as follows:

\[
\text{Score} = -\log_{10}\left(1 - \frac{\binom{C(G, i)}{C(N, s)} \binom{N - G, s - 1}{N - s}}{\binom{N}{s}}\right)
\]

where \( N \) is the number of genes in the genomic network, of which \( G \) are focus genes, for a network of \( s \) genes, \( f \) of which are focus genes; and \( C(N, s) \) is the binomial coefficient. The score reflects the negative logarithm of the \( P \) value that signifies the likelihood of the focus genes in a network being found together as a result of random chance. We merged the top-scoring networks using a “force-direct layout algorithm” to form a composite network representing the underlying biology of the process. Gene set enrichment analyses for the composite network were computed as above. For purposes of further investigation, genes with connectivity (ie, number of interactions with other genes) above the 95% quantile of the overall distribution of gene connectivity in the composite network were deemed network “hub” genes. Genes with tumor-related biological functions and interactions with hub genes were deemed “hub-interacting” genes. For the interaction of the hub genes with each other and with the hub-interacting genes, the mode of functional interaction was integrated with the direction of gene dosage change of each gene constituent of a gene pair to determine the presence of a cooperatively tumorigenic relationship. The BioGRID protein interaction repository was used to confirm high connectivity of candidate genes yielded by the IPA analyses by providing an experimentally based data source for physical protein interactions. The entire set of interactions observed in humans (release 2.0.51; March 26, 2009) was downloaded and the number of interactions (or connectivity) observed for all proteins represented in the data set was compared with the number of interactions observed for the candidate genes yielded by the IPA analyses using a Wilcoxon rank-sum test.

Survival Analysis and Multigene Predictor
Survival curves were estimated by the Kaplan-Meier product-limit method, and survival distributions between groups were compared using the log-rank test. Univariate and multivariate Cox proportional hazards regression analyses were performed with overall survival as the dependent variable, and the proportional hazards assumption was tested using interactions with time. To create a multigene predictor model, we first selected predictor genes based on their statistical significance at \( P < .05 \) in univariate Cox models relating continuous gene dosage data to survival in the TCGA data set. To combine data for the 7 detected genes, we used a schema according to the presence of alterations of these genes in individual tumors. Based on the number of alterations we assigned tumors to a high-risk group (≥5 of 7 genes altered in the direction associated with gliomagenesis) and a low-risk group (≤2 genes altered). The remainder of the tumors were considered moderate-risk. For the data sets that only included gene expression data without gene dosage data, we used transcript levels above or below their respective medians within a data set to reflect possible chromosomal gain or loss. For genes whose deletion or amplification demonstrated association with survival data at the genetic level in the TCGA analysis, we assigned an “alteration call” to denote potential chromosomal loss or gain if the transcript level was below or above the median. We created low-, moderate-, and high-risk groups for the transcript data based on the number of alteration calls using the same criteria as for the genetic data.

Statistical Calculations and Data Visualization
Unless otherwise specified, all statistical calculations were performed using R software, version 2.8.1, and pack-
ages from Bioconductor, release 2.3 (http://www.bioconductor.org/). Paired and unpaired \( t \) test, Wilcoxon rank-sum test, and 2-way contingency table analysis based on Pearson \( \chi^2 \) test were used as appropriate. Odds ratios in the 2-way contingency table analysis were computed according to the equation \((a/b)/(c/d)\), where \( a \) and \( b \) indicate the number of landscape genes (Box)—genes mapping to the nonrandom genetic landscape—with and without significant gene-dosage effect, respectively; and \( c \) and \( d \) indicate the number of nonlandscape genes with and without significant gene dosage effect, respectively. Corresponding 95% confidence intervals (CIs) for the estimated parameters were computed on “constant \( \chi^2 \) boundaries” as detailed elsewhere.\textsuperscript{23} Unsupervised hierarchical clustering was done in Cluster (http://rana.lbl.gov/EisenSoftware.htm),\textsuperscript{24} and 2-way complete linkage clustering based on Pearson correlation as the distance metric was applied. Two-class, unpaired Significance Analysis of Microarrays (SAM; version 3.02; http://www-stat.stanford.edu/~tibs/SAM)\textsuperscript{25} was used to assess the weight of Microarrays (SAM; version 3.0-477.0) weeks. A sample population of 76 high-grade glioma patients (25 females and 51 males) treated at MDA,\textsuperscript{2} whose data were made public on March 15, 2006, constituted a first clinical validation set. Patients included 55 with glioblastomas and 21 with anaplastic astrocytomas. Mean patient age was 45.8 (SD, 12.9) years. Sixty-two patients were dead and 14 were alive at last follow-up. Median follow-up was 93.0 (range, 3.0-477.0) weeks. A sample population of 70 high-grade glioma patients (43 females and 27 males) treated at UCLA,\textsuperscript{3} made public on March 8, 2006, constituted a second clinical validation set. Patients included 47 glioblastomas, 8 anaplastic astrocytomas, 9 anaplastic oligodendrogliomas, and 6 anaplastic oligoastrocytomas. Mean patient age was 45.6 (SD, 15.0) years. Forty-four patients were dead and 26 were alive at last follow-up. Median follow-up was 63.2 (range, 1.0-359.0) weeks. A unified sample population of 191 glioblastoma patients (74 females and 117 males) from multiple institutions,\textsuperscript{14} made public on October 10, 2008, constituted a third clinical validation set. Mean patient age was 53.8 (SD, 13.6) years. One hundred seventy-six patients were dead and 15 were alive at last follow-up. Median duration of follow-up was 55.6 (range, 1.0-479.0) weeks.

**RESULTS**

**Patients**

Forty-five patients treated at Stanford University between April 5, 2001, and April 19, 2004, constituted the initial molecular discovery set. Diagnoses included 26 glioblastomas, 5 astrocytic gliomas (grades I-III), 8 oligodendrogliomas (including 3 anaplastic oligodendrogliomas), and 6 anaplastic oligoastrocytomas according to World Health Organization classification.\textsuperscript{26} Two hundred nineteen glioblastoma samples collected between July 26, 1989, and November 23, 2007, and pro-

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**Gene Dosage Reflected in Gene Expression**

We observed a significant \((q < .10)\) gene dosage–gene transcript relationship for 1562 (15.1%) of 10 351 expressed genes in the 45 Stanford tumors, including genes implicated in gliomagenesis such as \( \text{EGFR}, \text{MYC}, \text{MDM2}, \text{GBAS}, \text{PTEN}, \text{CDK6}, \text{MTAP}, \text{IGF1R}, \text{MXII}, \text{WDR11}, \text{and FGR2.} \) Gene set enrichment analyses—which test for high representation of distinct biological processes and signaling pathways—revealed a significant \((P < .05)\) by right-tailed Fisher exact test) enrichment of those genes for various tumor-promoting (and developmental) functions and pathways (eFigure 2).
We noted a widespread association of changes in gene dosage and expression across the glioma genome (eFigure 3). However, there was a greater likelihood for a gene dosage effect on the expression of genes mapping to the genetic landscape than on the expression of genes mapping to random genetic alterations \((P < .001\) by Pearson \(\chi^2\) test; odds ratio, 7.7; 95% CI, 6.9-8.6) (eFigures 3 and 4). Within the genetic landscape, we noted a greater propensity for alterations to be hypomorphic, ie, to reduce gene expression, rather than hypermorphic, ie, to increase gene expression (for hypomorphic vs hypermorphic, odds ratio, 8.4; 95% CI, 7.3-9.6 vs 3.5; 95% CI, 2.8-4.2; \(P < .001\) for both) (eFigures 3 and 4).

**Highly Interacting Landscape Genes**

Genes with a role in gliomagenesis are more likely to function as part of a cooperative group or network rather than individual units.\(^\text{10,27,28}\) Our hypothesis that the characterized common genetic landscape alters biological networks to promote glioma formation prompted network modeling to identify the interaction of the genes affected by dosage effects in the glioma genome. This yielded a “node-and-edge” graph that maps 214 genes (represented as nodes) into a network of various physical (protein-protein) or functional interactions (represented as edges connecting the nodes) (Figure 4). The modeled interactions are both directed (well-defined information flow; eg, from a transcription factor to the gene it regulates) and undirected (eg, mutual binding relationships). Gene set enrichment analyses revealed a greater probability for over-representation of tumor-promoting (83.1% of genes) and developmental (29.5%) functions and pathways in the network than for all 1562 gene dosage-driven genes (eFigure 2).

The organization of genes and proteins into functional networks likely reflects cellular function.\(^\text{27}\) We studied the genes with high connectivity (a large number of incident edges), as their alteration may affect many other genes in the network; such highly influential genes should be preferred therapeutic targets. We identified 11 highly connected hub genes (Figure 4 and eFigure 5A), all of which have a tumor-promoting function and notably included genes with a known biological role in gliomagenesis, such as \(\text{EGFR, MYC, PTEN, and BAX}\). Genes that interact functionally with the hub genes are also likely to fundamentally influence the network. Analysis of the reciprocal functional relationship of the 11 hub genes with each other and with 92 hub-interacting genes with a tumor-related function disclosed 125 cooperatively tumorigenic gene-gene relationships (Figure 4). A matched-pair analysis of the number of putatively cooperative vs noncooperative interactions for those genes revealed a significant prevalence (125 [71.8%] of all 174 interactions; \(P = .003\) by paired \(t\) test) of cooperatively tumorigenic relationships in the network (eFigure 6).

We found that 6 (54.5%) of 11 hub genes and 52 (56.5%) of 92...
tumor-related hub-interacting genes mapped to the TCGA-validated genetic landscape. We tested the nonrandomness of coalteration of the individual hub and hub-interacting genes mapping to the genetic landscape (the landscape hub and hub-interacting landscape genes, constituting together the landscape genes) by applying the same approach used initially to model a genome-wide genetic landscape. We then integrated the association data with the interaction data of our network model (Figure 4) to identify significant interactions for the landscape genes; that is, physical (protein-protein) and functional interactions between genes whose gene dosage alterations are significantly associated (FDR < 0.05) (eFigure 5, B and C). This analysis revealed 37 significant interactions involving all 6 (100%) landscape hub genes—CDC42, EGFR, MYC, PTEN, BAX, and EP300—and 31 (59.6%) of the 52 hub-interacting landscape genes (eFigure 5, B and C). Thirty-one (83.7%) of these 37 significant interactions demonstrated a cooperatively tumorigenic relationship (FIGURE 5).

Mapping of these relationships to a human chromosome ideogram reveals that the involved genes are broadly dispersed throughout the TCGA-validated landscape (Figure 5).

To validate a significant gene-transcript correlation for the 31 landscape genes with cooperatively tumorigenic relationships, we integrated corresponding gene dosage and expression profiles in TCGA using a statistical approach similar to that above. Combined gene dosage and gene expression data were available for 30 of the 31 genes in TCGA’s data set. This analysis confirmed significant gene dosage effects on expression for 27 (90.0%) of the 30 genes, excluding MGAT3, MYBL2, and PCNA (Benjamini-
Hochberg FDR-corrected \( P < 0.05 \) (Figure 5).

Finally, we confirmed high connectivity for 25 of the 27 validated landscape genes (MFN2 and ABCC4 did not have interactions reported at the time of analysis) using BioGRID, a general repository for protein-protein interactions. For 8445 available genes (the entire set of human protein interactions) with at least 1 interaction reported in humans, there were 30,386 gene-gene pairs with at least 1 reported interaction between them. The median number of reported interactions for the entire data set was 3 (range, 1-202). For the 25 validated landscape genes available for analysis, the median number of reported interactions was 12 (range, 1-108). A Wilcoxon rank-sum test comparing the number of observed interactions yielded \( P < 0.001 \), confirming the significantly higher interconnectivity of the TCGA-validated landscape genes than of the entire set of human protein interactions.

Landscape Gene Alterations Associated With Patient Outcome

Unsupervised hierarchical clustering—which assigns objects (herein, patient and genes) into groups (called clusters) so that objects from the same cluster are more similar to each other than are objects from different clusters—of 189 glioblastomas with gene dosage and clinical data in TCGA based on the 27 TCGA-validated landscape genes identified 2 major subgroups with a significant difference in their mean number of landscape gene alterations (\( P < 0.001 \) by unpaired \( t \)-test): one consisting of 153 tumors showing substantial alterations of the landscape genes (mean, 54.7% [SD, 16.1%]) and one consisting of 36 tumors showing far fewer alterations (mean, 24.3% [SD, 15.5%]) (Figure 6, A and B). Supervised group predictor analysis revealed highest group-predictive power for altered genes on chromosomes 7 and 10.

We hypothesized that the group of tumors with extensive landscape gene alterations might be those that are biologically more aggressive and, hence, have a worse prognosis. A 2-class model confirmed a significant difference in

**Figure 3.** Model of a Nonrandom Genetic Landscape in Gliomas: Confirmation by Validation Set

A Co-occurrence of gene dosage alteration in validation set (219 glioblastomas, 802 cytobands)

B Significant associations in the discovery set that were confirmed in the validation set (21 associations)

Permutation-based approach described in Figure 2 was applied to the validation set from The Cancer Genome Atlas Pilot Project (TCGA). The association map (A) indicates chromosomal regions (territories) with significant co-occurrence of gene dosage alteration (false-discovery rate [FDR]–corrected \( P < 0.05 \)). Red scores denote the significant association of 2 chromosomal gains or 2 chromosomal losses; blue scores, the significant association of a gain and a loss event. The color gradation reflects a score that indicates the significance of cytoband-cytoband associations. The association map shows a consistent pattern of significant associations, which denotes a nonrandom genetic landscape in gliomas. B, Significant associations identified in the discovery set (Figure 2B) that were confirmed in the TCGA validation set. The blue bars and corresponding blue labels indicate chromosome territories with TCGA-confirmed significant associations to other territories (TCGA-validated nonrandom genetic landscape). (See interactive Figure 1 showing significant associations in the discovery and validation sets at http://www.jama.com.)

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Graphical representation of the interactions and networking of 214 glioma genes. The 214 genes represent the subset of all 1562 genes with significant gene dosage–transcript relationships (false-discovery rate–adjusted $q < 10^{-3}$) that map to the top-scoring subnetworks identified in a network modeling approach using Ingenuity Pathway Analysis. These subnetworks were merged using a force-directed layout algorithm to form a composite network representing the underlying biology of the process. Genes are represented as nodes using various shapes that represent the functional class of the gene product. The interactions of genes with tumor-related biological functions and fulfilling the criterion of a network hub (11 genes) or interacting with hub genes (hub-interacting gene; 92 genes) are highlighted. Hub gene refers to a gene that shows a number of interactions with other genes (the hub-interacting genes or other hub genes) that is above the 95% quantile of the overall distribution of the number of interactions of all genes in the network (eFigure 5A). Cooperatively tumorigenic interactions are interactions for which integration of the mode of interaction for a hub-hub (solid red lines) or hub–hub-interacting gene pair (blue lines) with the direction of gene dosage–gene expression change (gain or loss) reveals an effect on this interaction that synergistically promotes tumorigenesis. Paired $t$ testing comparing the number of cooperative vs noncooperative interactions for 11 hub genes with each other and 92 hub-interacting genes with tumor-related functions reveals a significant prevalence ($P = 0.003$) of cooperatively tumorigenic interactions (125 [71.8%] of all 174 interactions) (eFigure 4). (See interactive Figure 2 of the complete network at http://www.jama.com.)
overall survival between the 2 groups (median duration of survival of 51 vs 72 weeks), such that tumors with extensive alterations demonstrated a comparably unfavorable outcome (hazard ratio [HR], 1.75; 95% CI, 1.19-2.59; log-rank $P = .004$) (Figure 6C). Median duration of follow-up was 73.1 (range, 10.6-503.4) weeks for the group with modest alterations and 47.0 (range, 1.1-307.4) weeks for the group with extensive alterations. Tumors with wide

Figure 5. Networking of 31 Hub and Hub-Interacting Genes With Cooperatively Tumorigenic Relationships That Map to the TCGA Validated Landscape

Human chromosome ideogram representation of the networking of 31 hub and hub-interacting genes that map to The Cancer Genome Atlas Pilot Project (TCGA)–validated genetic landscape and possess cooperatively tumorigenic relationships. These cooperatively tumorigenic relationships emerge through the mode of interaction between the genes and their significant coalteration. Gene dosage–gene expression integration for 30 of the 31 genes with combined availability of gene dosage and expression data in TCGA confirmed significant gene dosage effects on transcription for 27 (90.0%) of 30 genes (Benjamini-Hochberg false-discovery rate–corrected $P < .05$). Labels indicate the cytogenetic band mapping of the genes.

a Genes with a significant ($P < .05$) relationship between gene dosage and duration of patient survival in the TCGA data set based on Cox proportional hazards regression analysis.

b Genes without confirmed significant gene dosage–gene expression relationship in the TCGA data set.

c TIMP3 is embedded within an intron of the SYN3 gene. Gene dosage information for TIMP3 was only indirectly available through a probe mapping to SYN3 in TCGA. Integration of gene dosage information using this probe and TIMP3 transcript information revealed a significant ($P < .05$) gene dosage–transcript relationship.
spread landscape gene alterations had a 2-year survival rate of 12.9% and 82.34 deaths per 100 person-years vs a 2-year survival rate of 32.0% and 48.31 deaths per 100 person-years for tumors with modest alterations.

**Multigene Predictor Model**

To determine the potential utility of a concise molecular profile in estimat-
ing the prognosis of an individual glioblastoma patient, we assessed the survival association of gene dosage for each of the 27 validated landscape genes in TCGA glioblastomas. Cox proportional hazard regression indicated statistically significant survival associations for 7 landscape genes, including the DNA polymerase delta 2, small subunit (POLD2) (\(P = .01\)) and cytochrome C, somatic (CYCS) (\(P = .01\)) genes on chromosome 7, the MYC oncogene on chromosome 8 (\(P = .03\)), and the aldo-keto reductase family 1, member C3 (AKR1C3) (\(P = .002\)), YME1-like 1 (YME1L1) (\(P = .006\)), annexin A7 (ANXA7) (\(P = .008\)), and programmed cell death 4 (neoplastic transformation inhibitor) (PDCD4) (\(P = .01\)) genes on chromosome 10. Each of the 7 genes has a biological function whose alteration might contribute to tumorigenesis. The avian myelocytomatosis homologue (MYC) is a major oncogene\(^3\); ANXA7 is a candidate tumor suppressor gene in breast and prostate cancer\(^28,30\) and its protein product is a phospholyation target of the epidermal growth factor receptor (EGFR) oncoprotein\(^34\); AKR1C3 and PDCD4 have been implicated in colorectal cancer\(^35,36\) and their expression is regulated by phosphatase and tensin homolog (PTEN)\(^37\); YME1L1 has been related to cancer development and progression\(^38\) and has been identified as an MYC-responsive gene,\(^39\) as has CYCS.\(^40\)

The protein encoded by POLD2 is a subunit of the DNA polymerase delta complex, which also contains the protein product of the landscape gene PCNA; it is involved in DNA replication and repair and its expression is regulated by PTEN.\(^37\)

In the TCGA training set of 189 glioblastomas, the combined 7-gene predictor model assigned 119 tumors to a high-risk group (\(\geq 5\) of 7 genes altered) with a median duration of follow-up of 49.3 (range, 2.4-307.4) weeks, 39 to a low-risk group (\(\leq 2\) genes altered) with a median follow-up of 62.9 (range, 10.6-503.4) weeks, and 31 to a moderate-risk group (all others) with a median follow-up of 51.0 (range, 1.1-183.1) weeks. There was a significant difference in overall survival for the 3 groups (log-rank \(P = .02\); Cox model HR, 1.63; 95% CI, 1.10-2.40 [\(P = .02\)] comparing high-risk vs low-risk groups and HR, 1.87; 95% CI, 1.13-3.10 [\(P = .01\)] comparing moderate-risk vs low-risk groups; 49.24, 92.66, and 79.56 deaths per 100 person-years for low, moderate, and high-risk groups, respectively (Figure 7A). Similar multigene models using gene expression data revealed significant differences in survival for the MDA and UCLA validation sets (Figure 7B and C). In the MDA set, 23, 33, and 20 patients were assigned to the low-, moderate-, and high-risk groups and had median follow-up times of 175 (range, 33-477) weeks, 70 (range, 12-467) weeks, and 62 (range, 3-311) weeks, respectively. The low-, moderate-, and high-risk groups demonstrated 15.13, 41.10, and 47.89 deaths per 100 person-years, respectively. A log-rank test for all 3 curves yielded \(P = .003\) and Cox modeling yielded an HR of 3.04 (95% CI, 1.49-6.20; \(P = .002\)) comparing high-risk vs low-risk groups and an HR of 2.52 (95% CI, 1.33-4.78; \(P = .005\)) comparing moderate-risk vs low-risk groups (Figure 7B). In the UCLA set, 20, 34, and 16 patients were assigned to the low-, moderate-, and high-risk groups and had median follow-up times of 128.5 (range, 6-359) weeks, 58.5 (range, 8-320) weeks, and 49 (range, 1-147) weeks, respectively. The low-, moderate-, and high-risk groups demonstrated 16.14, 38.41, and 83.43 deaths per 100 person-years, respectively. A log-rank test for all 3 curves yielded \(P = .003\) and Cox modeling yielded an HR of 3.86 (95% CI, 1.59-9.35; \(P = .003\)) comparing high-risk vs low-risk groups and an HR of 2.12 (95% CI, 0.97-4.63; \(P = .06\)) comparing moderate-risk vs low-risk groups, respectively (Figure 7C). In the multi-institutional validation set of 191 glioblastomas, 43, 92, and 56 patients were assigned to the low-, moderate-, and high-risk groups and had median follow-up times of 73 (range, 1-479), 57.5 (range, 8-313), and 47 (range, 1-242) weeks, respectively. The low-, moderate-, and high-risk groups demonstrated 34.16, 63.49, and 83.23 deaths per 100 person-years, respectively. A log-rank test for all 3 curves yielded \(P = .002\) and Cox modeling yielded an HR of 2.27 (95% CI, 1.44-3.58; \(P < .001\)) comparing high-risk vs low-risk groups and an HR of 1.72 (95% CI, 1.14-2.59; \(P = .01\)) comparing moderate-risk vs low-risk groups (Figure 7D), confirming the robustness of the multigene predictor model in glioblastoma at the expression level.

Although gene dosage was prognostic in TCGA tumors, we did not find an association between mRNA expression and survival in the TCGA data set. Heterogeneity of patient characteristics and treatment (tumor samples collected over nearly 2 decades at varying institutions) and the fact that gene transcription is more dynamic and fluctuant than gene dosage may explain the disparity in findings in the TCGA group at the genetic and transcript levels.

**COMMENT**

The pathobiology of human gliomas emerges through the actions of multiple genes and their interactions with each other. As the classic theories regarding glioma origin and gliomagenesis are currently being reappraised,\(^41\) we used a systems biology approach without regard to morphological subtypes to generate a model of a nonrandom genetic landscape in gliomas. Herein, nonrandomness describes a significant association of distinct chromosomal alterations. Our model proposes that this genetic landscape involves multiple reciprocal gene alterations, which cooperatively promote gliomagenesis and contribute to unfavorable patient prognosis, thus providing the selection pressure for its conservation.

Identification of the importance and mechanistic interplay of such gene alterations in gliomas prompts evaluation of their potential as therapeutic targets. The network context of a gene likely affects the efficacy of therapies that target its protein. The complexity...
of our landscape model helps explain the lack of therapeutic efficacy of strategies targeting single gene products. For example, targeting the oncoprotein EGFR alone has had limited clinical effect despite the prominence of EGFR in our model as a landscape hub gene.42 This paradox is resolved by the fact that the deregulation of EGFR in glioblastomas does not exist in isolation. But rather, the functional organization of EGFR, and other gene dosage–altered genes within the landscape, in a complex network of interactions predicts that monotherapeutic approaches will fail: extensive cross-talk between interacting genes and their pathways permits resistance to therapy. The topology of our landscape model suggests that molecular targeting of multiple (hub) genes will be more effective.

Our study has limitations. First, our landscape model reduces the biological complexity of human gliomas to the

**Figure 7. Multigene Risk Scoring Model in Malignant Gliomas**

- **A** Glioblastomas from TCGA validation set (n = 189)
- **B** High-grade gliomas from MDA validation set (n = 76)
- **C** High-grade gliomas from UCLA validation set (n = 70)
- **D** Glioblastomas from unified validation set (n = 191)

Patient assignment to low-, moderate-, and high-risk groups is based on risk scores generated by the number of gene dosage alterations (0-2, 3-4, and 5-7, respectively) of 7 landscape genes (POLD2, CYCS, MYC, ARRC13, YME1L1, ANXA7, and PDCD4), each of which was individually linked to patient survival in Cox proportional hazard regression analyses in The Cancer Genome Atlas Pilot Project (TCGA). A, Kaplan-Meier estimates of overall survival for the 3 groups in 189 glioblastomas from TCGA with available survival data. Median follow-up for the low-, moderate-, and high-risk groups was 62.9 (range, 10.6-503.4), 49.3 (range, 2.4-307.4), and 47.4 (range, 1.1-183.1) weeks, respectively. B, Estimates of overall survival in the University of Texas M. D. Anderson Cancer Center (MDA) validation set of 76 high-grade gliomas. Median follow-up for the low-, moderate-, and high-risk groups was 175 (range, 33-477) weeks, 70 (range, 12-467) weeks, and 62 (range, 3-311) weeks, respectively. C, Estimates of overall survival in the University of California, Los Angeles (UCLA) validation set of 70 high-grade gliomas. Median follow-up for the low-, moderate-, and high-risk groups was 73 (range, 1-479), 57.5 (range, 8-313), and 47 (range, 1-242) weeks, respectively.

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GENETIC LANDSCAPE MODEL FOR BRAIN TUMORS

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The current work provides a network model and biological rationale for the selection of a nonrandom genetic landscape in human gliomas. Coincident genetic alterations of multiple landscape genes may evolve rapidly under positive selection to provide multiple, synergistic mechanisms of dysregulation of critical signaling pathways toward gliomagenesis. A multigene predictor model incorporating 7 landscape genes demonstrates how molecular insights emerging from our integrative multidimensional analysis could translate into relevant clinical end points affecting the future management of gliomas.

CONCLUSION

The conclusions of this study were drawn to support the proposition that multigene predictors of glioblastoma may be more amenable to predictive screens.

into account the key mechanisms contributing to glioblastoma pathogenesis. Several gene sets have been proposed as multigene predictors of glioblastoma patient survival. Many of these gene sets have used so-called metagenes—patterns of gene expression—for risk classification. Although metagenes support the notion that, in a complex glioblastoma tumor, it is not a single gene but, rather, multiple genes that drive the disease process, the risk associations of metagenes are primarily correlative. While metagenes, by definition, constitute gene subsets identified without considering underlying mechanisms, some studies have shown that such metagenes can characterize clinically relevant molecular subtypes of high-grade gliomas. Recently, a consensus gene expression profile associated with patient outcome across independent data sets and an optimized 9-gene metagene predictor have been reported for high-grade gliomas. While the primary role of multigene predictor models would be to prospectively identify glioma patients with low vs high likelihood for a durable response to standard therapy, molecular predictor models that place genes into pathways and networks allow mechanistic insights that are crucial for developing novel molecularly targeted therapies. We show that a focused set of 7 landscape genes associates with patient survival across several different study populations with discrimination similar to that of correlative multigene predictors. These observations lay the foundation for future development of a mechanistically based molecular risk estimation model in glioblastomas and high-grade gliomas. In our study, the multigene predictor model associates with survival when taking into account either genetic or transcriptional information about those genes. The genetic model might lend itself better to translation into clinical practice for 2 reasons. First, while the expression of an individual gene can vary over short periods of time in cancers, gene dosage is more static and more amenable to predictive screens.

Second, the gene expression model depends on risk groupings in which the transcript profile of an individual patient is related to the profiles of other patients, with a potential for demographic bias. In contrast, the genetic predictor allows risk estimation based on a patient’s individual biological profile by using a binary model (the gene is altered or it is not).
REFERENCES


