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A mathematical framework to determine the temporal sequence of somatic genetic events in cancer

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Human cancer is caused by the accumulation of genetic alterations in cells. Of special importance are changes that occur early during malignant transformation because they may result in oncogene addiction and represent promising targets for therapeutic intervention. Here we describe a computational approach, called Retracing the Evolutionary Steps in Cancer (RESIC), to deduce the temporal sequence of genetic events during tumorigenesis from cross-sectional genomic data of tumors at their fully transformed stage. When applied to a dataset of 70 advanced colorectal cancers, our algorithm accurately predicts the sequence of *APC*, *KRAS*, and *TP53* mutations previously defined by analyzing tumors at different stages of colon cancer formation. We further validate the method with glioblastoma and leukemia sample data and then apply it to complex integrated genomics databases, finding that high-level *EGFR* amplification appears to be a late event in primary glioblastomas. RESIC represents the first evolutionary mathematical approach to identify the temporal sequence of mutations driving tumorigenesis and may be useful to guide the validation of candidate genes emerging from cancer genome surveys.

mathematical modeling | stochastic framework | optimization algorithm

Recent technological advances have empowered researchers to examine the cancer genome at unprecedented throughput and resolution (1–3). Computational algorithms designed to filter random genetic events have begun to uncover mutational patterns that are typical for a particular cancer type and highly consistent between sample sets (1, 2, 4, 5). Further functional validation of these recurrent genetic events in nontransformed primary cells and mouse models of human cancer is hampered by the lack of knowledge of the sequence in which these alterations occur during human tumorigenesis. This temporal order can guide the generation of the correct genomic context in animal models of human cancer and can prioritize the validation of potential drug targets because those changes that occur early during malignant transformation may result in rewiring of the signaling circuitry or confer a state of addiction to the new signal. Here we describe a unique computational approach, called Retracing the Evolutionary Steps in Cancer (RESIC), to determine the sequence of genetic events using cross-sectional genomic data from a large number of tumors at their fully transformed stage (Fig. 1).

Model

RESIC is based on the principles of population genetics—the mathematical study of the dynamics of genetic variation within populations (6). Consider a population of N cells at risk of accumulating the genetic changes leading to cancer (Fig. 2*A*). Cells proliferate according to a stochastic process (7): At each time step, a cell is chosen proportional to fitness to produce a possibly

mutated daughter cell. Subsequently another cell is chosen at random to die and is replaced by the newly produced cell to maintain homeostasis. A mutated cell can take over the population (i.e., reach fixation) or go extinct due to stochastic fluctuations (Fig. 2*A*, *Inset*). Depending on the order of appearance of particular mutations, the population of cells follows different evolutionary paths toward the fully mutated state (Fig. 2*A*). We developed a mathematical model describing the evolutionary dynamics of this system (see *SI Text*).

We assume that cancers originate from a single population of cells per person and study the evolutionary dynamics of individuals accumulating the mutations leading to cancer (Fig. 2*B*). We consider the dynamics of patients in steady state: There is a constant influx into the unmutated state, representing diagnosis of disease, and a constant outflux from the fully mutated state, accounting for deaths of patients or their cure. At steady state, the population is distributed across all possible states; this steady-state distribution can be compared to the numbers of clinical samples with the corresponding genotypes, where the total number of patients in a dataset is equal to the sum of patients in all states. This mapping is used to optimize a subset of parameters in the mathematical model (i.e., the fitness values of cell types) by minimizing the difference between the prediction and the observed frequencies in the dataset (see *SI Text*). Other parameters, such as cellular population size, mutation rate and influx rate, are estimated from experimental results (8, 9) and tested for robustness over several orders of magnitude (see *SI Text*). The output of RESIC is given as percent of the flux through the network via each particular path.

Results

The established sequence of genetic events arising during the multistep process of colorectal carcinogenesis (10) provides a unique example to test the ability of RESIC to recover this sequence from a cross-sectional dataset. We gathered genomic data from 70 advanced colorectal tumors (1, 2) (Table S1) and used RESIC to predict the temporal relationship between alterations in the *APC*, *KRAS*, and *TP53* genes. RESIC predicts that the most likely

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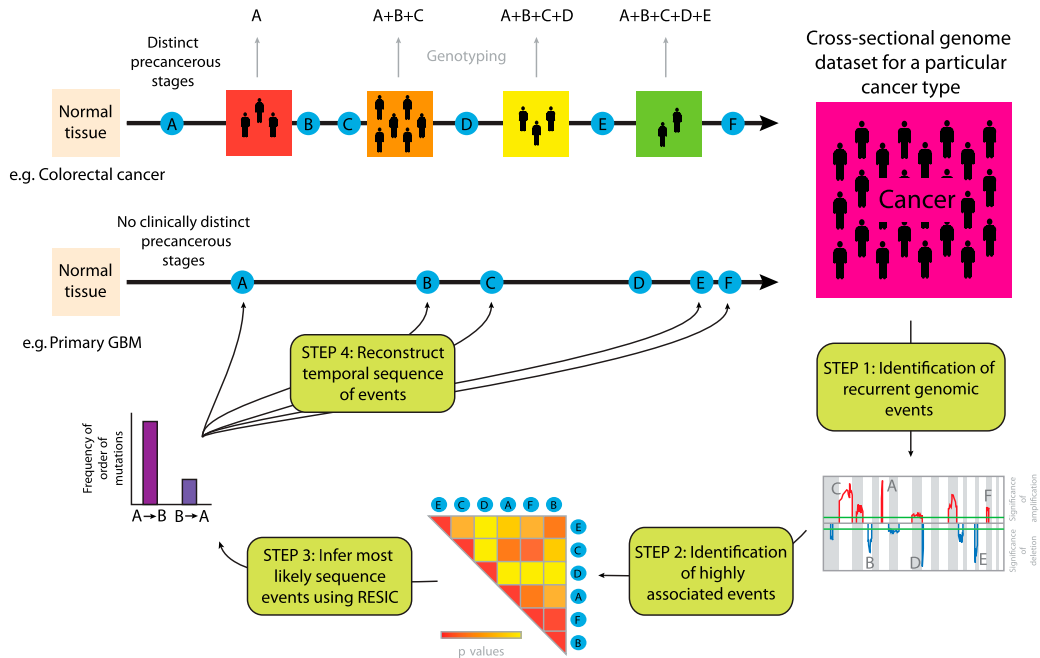


Fig. 1. Schematic diagram of RESIC. For cancer types with clinicopathologically defined stages (e.g., colorectal cancer), the temporal sequence in which genetic alterations arise during tumorigenesis can be identified by genotyping samples from patients at different stages of disease progression. For cancer types that are diagnosed de novo without detectable precursor lesions (e.g., primary GBM), the order of alterations cannot be identified with a similar approach. We present an evolutionary computational algorithm (RESIC) to identify the temporal sequence of events arising during tumorigenesis utilizing genomic data from a large number of samples (one per patient) of a particular histological type. In step 1, we use an algorithm such as GISTIC (4) to identify recurrent genetic aberrations in the genomics dataset. In step 2, these aberrations are ranked according to their pairwise association (statistically significant correlation, e.g., Fisher's exact test). In step 3, the most likely sequence of these associated events is identified using RESIC. The results generated by RESIC are used to reconstruct the order in which alterations arise during development of a particular cancer type (step 4). Our methodology is applicable to large-scale datasets and can be used to identify the temporal sequence of many genetic alterations.

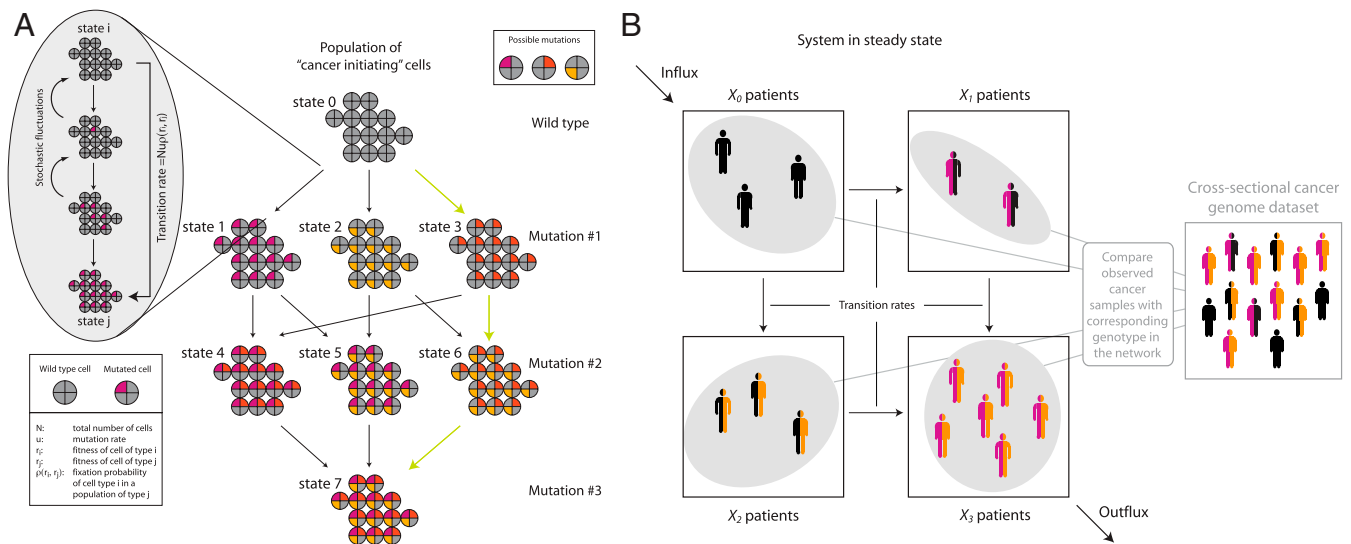


Fig. 2. Evolutionary dynamics of genetic alterations leading to cancer. (A) Transition between mutational states and schematic representation of different evolutionary trajectories toward cancer. Initially, the population consists of N cells with genotype i and fitness (i.e., growth rate) r_i (detail). During each time step, a cell is chosen at random proportional to fitness to divide, and its daughter cell replaces another randomly chosen cell. During each cell division, a mutation arises with probability u . The mutated daughter has genotype j and fitness r_j . If $r_j > r_i$, the mutated daughter cell is advantageous as compared to the mother cell; if $r_j < r_i$, it is disadvantageous, and if $r_j = r_i$, it is selectively neutral. The probability that a mutated cell takes over the population is given by its fixation probability, $\rho(r_i, r_j) = [1 - 1/(r_j/r_i)]/[1 - 1/(r_j/r_i)^N]$. If $r_j = r_i$, then $\rho(r_i, r_j) = 1/N$. The transition rate between states i and j is given by $Nu\rho(r_i, r_j)$ in small populations. A population of wild-type cells may accumulate mutations in different orders; an example path from the unmutated population to a state with three mutations is highlighted in green. (B) Population dynamics. The dynamics of patients accumulating mutations is represented in this network where nodes (i.e., mutational states) are populated according to the transition rates from one mutational state to the next. In the example shown here, cells can accumulate two mutations. The number of patients harboring cells with no mutations are denoted by X_0 , whereas those harboring mutations are denoted by X_1 , X_2 , and X_3 . There is a constant influx of cases into the initial node. Cells in these patients accumulate mutations and populate the mutational states. The outflow from the fully mutated state eventually drives the system into steady state. An optimization algorithm is used to identify the transition rates for which the number of patients in each node at steady state coincides with the observations in a cross-sectional genomics dataset. The optimized parameter values of the evolutionary process serve to identify the most likely trajectory through the network.

sequence of events is homozygous inactivation of *APC* occurring before alterations of *KRAS* (Fig. 3A and Table S2). Similarly, we found that both *APC* alleles are likely mutated before *TP53* is homozygously inactivated, and that at least one *KRAS* allele is likely mutated before inactivation of *TP53* (Fig. 3A). Although the small number of samples prevents us from investigating the complete network of *APC*, *KRAS*, and *TP53* mutations in a single computational analysis, the results of the separate smaller analyses can be combined into a sequence of events (see also Figs. S1 and S2) that coincides with the multistep model of colorectal cancer (10, 11). The results are very robust with regard to sampling stochasticity and variations of the population size, mutation rates, and influx values (Figs. S3–S6 and Table S3).

The temporal relationship between *TP53* inactivation and mutations in the *RAS* pathway may not be the same in all cancer types. For example, genetic inactivation of neurofibromin-1 (*NF1*), a negative regulator of *RAS* activity (12), induces senescence of human astrocytes in the absence of *TP53* mutations (13); furthermore, mice develop high-grade gliomas only if *TP53* is inactivated prior to or simultaneously with *NF1* (14). To investigate the temporal relationship between *NF1* and *TP53* inactivation in human primary glioblastoma (GBM), we applied RESIC to a genomic dataset containing sequence and copy number information regarding these two genes for 91 primary human GBM samples (3) (Table S1). We found that *TP53* is likely inactivated before *NF1* is lost (Fig. 3B). Because several GBMs in this particular sample set were collected after treatment with radiation and/or chemotherapy—therapies that might increase mutation rates (3), we reran RESIC using only GBMs that had not received prior therapy ($n = 72$). We again found that *TP53* is likely mutated first (Fig. 3B). Our predictions are thus consistent with experimental data and suggest that *TP53* inactivation may not only be an early genetic event in *TP53*-mutant secondary glioblastomas (15) but also in *TP53*-mutant primary glioblastomas.

We next applied RESIC to a leukemia dataset to test whether it also accurately predicts the order of mutations arising in “liquid” tumors. We investigated the order of mutations in the *JAK2* and *TET2* genes in a set of 57 secondary acute myelogenous leukemia (AML) samples transformed from a preexisting myeloproliferative neoplasm (MPN) (Fig. 4A). The classical MPNs are clonal disorders of hematopoiesis characterized by the presence of the *JAK2V617F* mutation in most patients as well as an increased likelihood of transformation to AML (16). Clonality studies suggest that acquired *JAK2V617F* mutations may not represent the earliest genetic event in MPN pathogenesis and are not required for transformation to AML (17, 18). Recent studies have identified mutations in the putative tumor suppressor gene *TET2* in MPNs (17, 19–23), and clonal analysis suggested that *TET2* mutations precede the acquisition of *JAK2V617F* mutations in MPN pathogenesis (19). However, these studies were done on a small number of patients. We therefore analyzed a larger set of patients with leukemic transformation from a preceding MPN, including samples from two different disease states (MPN and post-MPN AML) from 14 different patients (19, 24). When applying RESIC to the set of secondary AML samples for which *JAK2* and *TET2* mutational status was known, we found that *JAK2* mutations likely preceded mutation in *TET2* (Fig. 4B). Notably, analysis of 14 patients for which samples were available from the MPN and AML disease states showed that *TET2* mutations were present in the AML, but not preceding MPN sample, in 5 patients with concomitant *JAK2/TET2* mutations (Fig. 4C) at the time of leukemic transformation. These data demonstrate, contrary to previous reports of smaller patient cohorts from a single time point, that *TET2* mutations are more commonly acquired subsequent to *JAK2* in MPN/AML pathogenesis (24), as predicted by RESIC.

After validation of RESIC using examples for which the temporal order of alterations is known, we analyzed a large integrated genomics dataset of primary glioblastoma samples ($n = 594$)

(3, 4). These samples were used together with RESIC to identify genetic alterations occurring early during malignant transformation. Such alterations may result in “rewiring” of the signaling circuitry, confer a state of “addiction” to the new signal (25), and thus represent particularly promising targets for therapeutic intervention (26). We first identified areas of statistically significant gene copy number alterations using GISTIC (4). For each locus, we distinguished gain of a single copy from high-level amplification, and hemizygous from homozygous deletions (Table S1). We then identified alterations that are significantly positively correlated (Fig. 4D), because the determination of an order of oncogenic events is meaningful only for those events that co-occur sufficiently often. Although *MDM2* and *CDK4* on chromosome 12 and *EGFR* and *MET* on chromosome 7 were the most significantly correlated mutations, these associations are likely the result of large-region deletion or amplification events; because such changes cannot be attributed to independent mutational events, we excluded them from our analyses. Low- and high-level amplification of *EGFR* and homozygous loss of *PTEN* are the most significantly correlated events on separate chromosomes (p values $< 10^{-30}$ and 10^{-24}) (Fig. 4D). Because the dataset contains copy number information for 552 GBMs (excluding samples with homozygous loss of *PTEN* due to their relative infrequency) but sequence data for only 125 GBMs and because for both genes, copy number alterations and point mutations occur frequently, we first sought to investigate the robustness of the algorithm to the exclusion of sequence information. With only copy number data, we found that the two most frequent mutational paths through this network—initiating with *EGFR* low-level amplification and homozygous *PTEN* loss, respectively (Fig. S1)—have very similar frequencies. Therefore, RESIC identifies no clear order of events for this mutational network. Using both copy number and point mutation data, we found that *EGFR* biallelic alterations likely occur before *PTEN* loss (Fig. S1). Hence the only difference between the two analyses is loss of significance of the dominant path, suggesting that analyzing copy number information only may be a viable option.

To perform a computational analysis on a large mutational network, we next determined that *p16* homozygous deletions frequently co-occur with *EGFR* and *PTEN* alterations (p value $< 10^{-8}$) in the dataset containing copy number alterations only (Fig. 4D). When studying the mutational network of *EGFR*, *PTEN*, and *p16* (Figs. S2–4), RESIC predicts that the most common early alterations are *EGFR* low-level amplification and *p16* deletion, with similar likelihood (Fig. 4E). Although there is no single most frequent path through the network (Fig. S3), the frequency of paths concluding with high-level amplification of *EGFR* is highest; the second most frequent final event is homozygous *p16* deletion (Fig. 4E). These data suggest that glial progenitor cells may tolerate full *EGFR* activation only after inactivation of *p16* or *PTEN*. This result agrees with the fact that *EGFR* overexpression is insufficient for tumorigenesis in mouse models of glioblastoma (27, 28), providing support for the temporal order of events predicted by RESIC.

Discussion

We have presented a rational methodology, RESIC, to identify the temporal order of oncogenic events during tumorigenesis. This computational pipeline uses cross-sectional genomic data as input and provides the most likely temporal order of genomic alterations as output. We validated the predictions of RESIC with events for which the temporal order is known and then applied it to a large integrated genomic dataset of primary glioblastoma samples. RESIC is based on three assumptions: (i) For each network under consideration, all cancers initiate without any of the mutations in the network. Hence all patients enter the network through the unmutated node, but might already have accumulated other mutations causing malignancy. This assump-

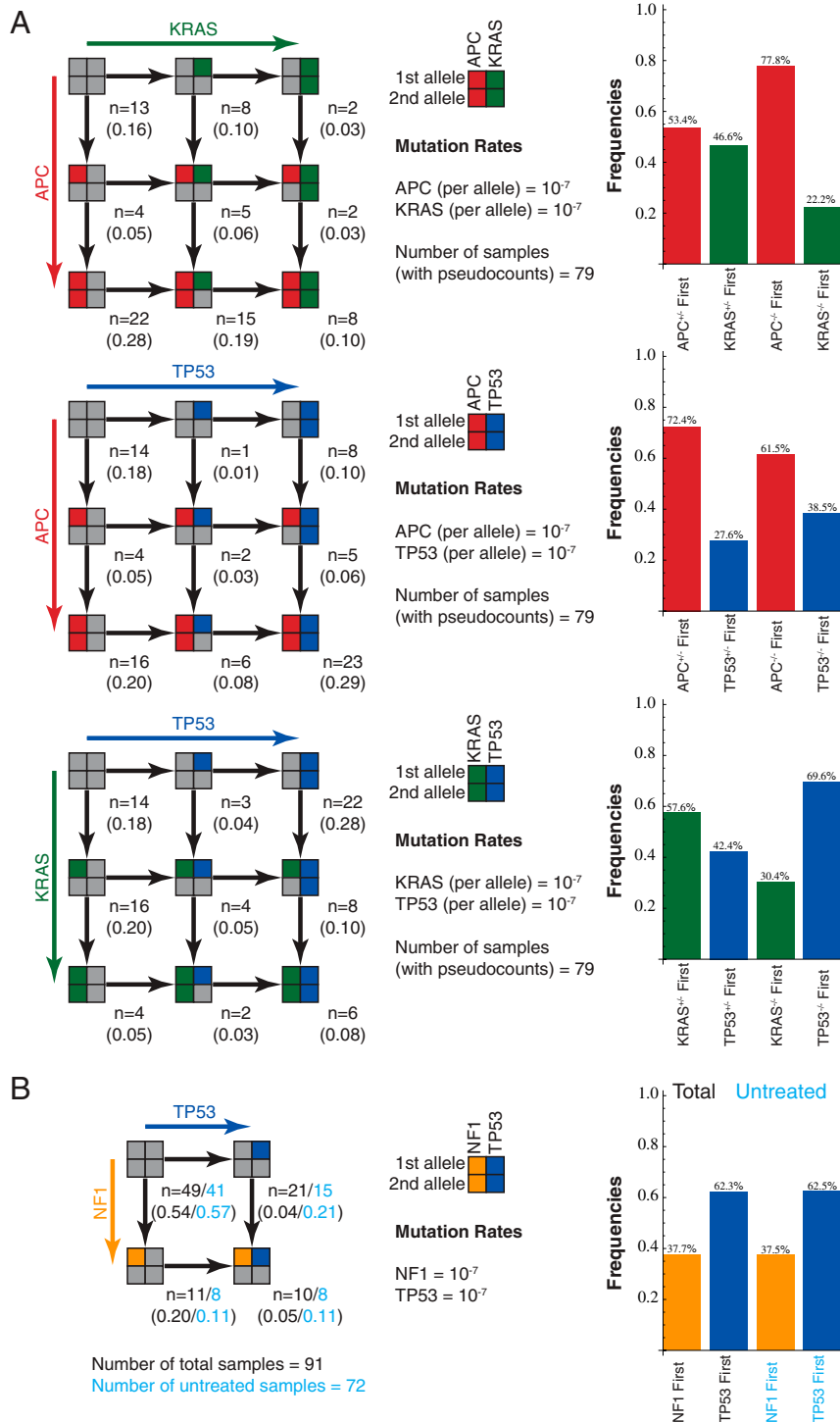


Fig. 3. Validation of RESIC utilizing colorectal cancer and glioblastoma data. We tested the predictions of RESIC in colorectal cancer and glioblastoma because the order of some events leading to those tumor types has been identified (10, 14). In the schematics of the networks, nodes represent the numbers of patients with a particular genotype, whereas black arrows represent transitions between mutational states. **(A)** The order of *APC*, *KRAS*, and *TP53* in colorectal cancer. *APC* is shown in red, *KRAS* in green, and *TP53* in blue. All mutation rates are 10^{-7} per allele per cell division. We apply a pseudocount of 1 to the entire system to prevent states with zero observations. Schematics of the networks are shown at *Left*. We display the numbers and frequencies of patients in each mutational node in the network and show the most frequent paths through the network in the histograms at *Right*. Detailed results are listed in [Table S2](#). *(First Row)* The *APC*–*KRAS* network. RESIC predicts that biallelic inactivation of *APC* likely occurs before any *KRAS* alteration. *(Second Row)* The *APC*–*TP53* network. RESIC predicts that biallelic inactivation of *APC* likely occurs before *TP53* inactivation. *(Third Row)* The *KRAS*–*TP53* network. RESIC predicts that an alteration of *KRAS* likely occurs first. **(B)** The order of *NF1* and *TP53* in glioblastoma. *NF1* is shown in orange. We study the mutational network of heterozygous alterations only since all *NF1* and most *TP53* mutations in the dataset are heterozygous ([Table S1](#)). A schematic of the network is shown at *Left*. Detailed results are listed in [Table S2](#). We show the number of samples with each genotype observed in the complete set of 91 The Cancer Genome Atlas samples (black) and in the restricted set of 72 untreated samples (blue) (3). For both the unrestricted and the restricted sets, RESIC predicts that a *TP53* point mutation likely occurs before *NF1* is altered (*Right*).

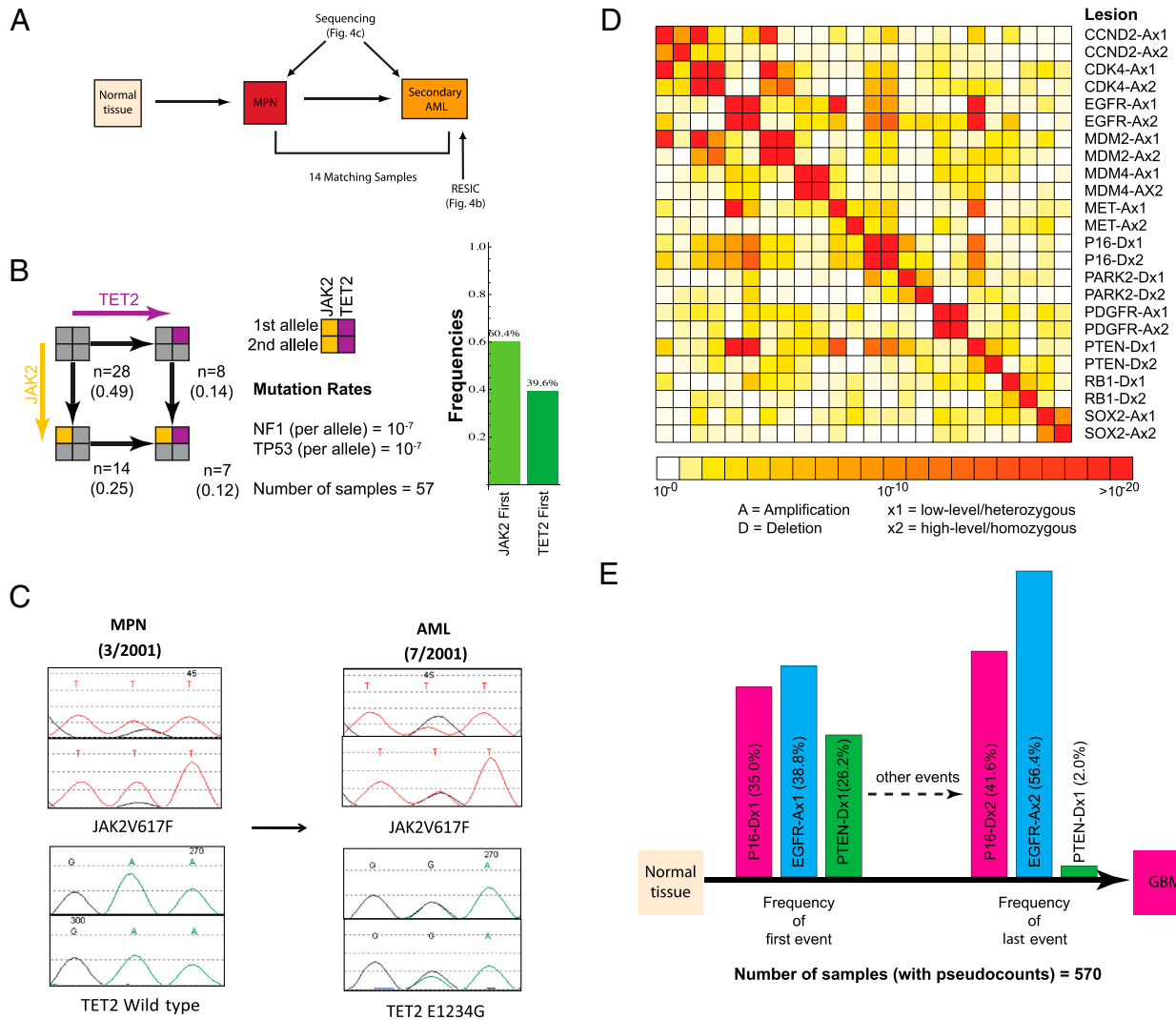


Fig. 4. Application of RESIC to secondary AML and primary GBM. (A) We analyzed a dataset of 57 patients with AML, including samples from two different disease states (MPN and post-MPN AML) from 14 different patients (19, 24). The data of AML patients were analyzed with RESIC (see B), whereas data from both MPN and AML patients were sequenced for *JAK2* and *TET2* (see C). (B) When applying RESIC to a set of secondary AML samples for which *JAK2* and *TET2* mutational status was known for both AML and MPN disease states, we found that *JAK2* mutations likely precede *TET2* mutations in this sample set. (C) Analysis of 14 patients for which samples were available from the MPN and AML disease states showed that *TET2* mutations were present in the AML, but not preceding MPN sample, in 5 patients. (D) Genetic alterations in primary glioblastoma. The statistical significance of correlations between genetic alterations was calculated with Fisher's exact test. Color codes range from 10^{-20} (red) to 10^0 (white). Significance after Bonferroni correction is marked at $\sim 3 \times 10^{-7}$ (orange). Note that lesions colocalized on the same chromosome have stronger correlations likely caused by large-region amplifications or deletions and thus cannot be considered as independent genetic events. (E) Prediction of RESIC for the *PTEN-p16-EGFR* network in primary glioblastoma. We show the frequencies of the initiating and final mutational events of this network. RESIC predicts that *p16* deletion or *EGFR* low-level amplification are the most common initiating events with frequencies of about 35–39% each, whereas high-level amplification of *EGFR* is the most frequent last event of this network with a frequency of 56.4%.

tion ensures that the model is consistent with the use of cross-sectional data and is important for situations in which more than one set of mutually exclusive mutations lead to cancer; in such cases, tumor samples can be subdivided into classes of different tumor subtypes, and each subtype must be analyzed individually. (ii) The likelihood of diagnosis is uniform across all states. This assumption can be partially relaxed if more biological data are included in the model, which is equivalent to changes in the number of observed samples with each combination of mutations; variations in these numbers have negligible impact on the results (Fig. S5). (iii) The order of mutations arising during tumorigenesis can be inferred from the order of mutations arising after diagnosis. This assumption links the results obtained from a database containing diagnosed cancer samples to the processes of tumorigenesis before diagnosis. Interactions between tumor cells and the microenvironment or immune system can be considered

as a modulation of mutation rates and fitness values but are excluded from the current implementation of RESIC for clarity. The results of RESIC are very robust to changes in the number of samples per genotype due to variability in sampling or the rate of diagnosis (Figs. S5 and S6), the population size of cells at risk of accumulating mutations, the influx value into the unmutated state, and the mutation rate (Table S3).

The frequencies of evolutionary paths through mutational networks identified by RESIC are never 100%. This finding may suggest that there is not a unique order in which these alterations occur—a hypothesis proposing that the genetic model of colorectal cancer (10) describes the order of mutations of only a subset of colorectal tumors (29). Alternatively, it may be attributed to the alteration of signaling pathways through alterations of mutually exclusive but functionally equivalent genes; it could also be the result of distinct mutational classes within the same

cancer subtype. The latter issue is addressed by studying highly correlated mutations because strong correlation may reflect functional associations.

We have restricted our analysis to only a subset of mutations implicated in tumorigenesis because RESIC requires correlated alterations as input; the question of the sequence of alterations is meaningful only if those alterations co-occur sufficiently frequently. Even the largest currently available dataset, the The Cancer Genome Atlas data, combined with separate glioblastoma studies ($n = 594$), contains only a small number of associated lesions (Fig. 4D). These correlated lesions also reflect individual molecular subtypes of glioblastoma. Based upon our analyses, we expect that of the order of 100 samples are sufficient to analyze mutational networks containing several loci. However, RESIC can be applied to any tumor type by analyzing significantly correlated genetic alterations in separate computational analyses (see Fig. 3). Furthermore, RESIC can be applied to individual subtypes of cancers after these subtypes have been identified utilizing gene expression or other data.

Based on the examples presented here, we anticipate that our algorithm will provide the research community with a tool for the identification of tumor-initiating events using the emerging cross-sectional cancer genome datasets and will help with

the generation of hypotheses about carcinogenesis that can be tested using modern mouse models of human cancer (30). The identification of the order of genetic alterations in specific cancer (sub)types may lead to important insight into cancer biology and should inspire studies aimed at elucidating how specific genes cooperate (or in a different order may prevent) tumorigenic processes.

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