that a meiosis I-specific factor from budding yeast, monopolar, generates kinetochores with more microtubule-binding elements and greater strength. These findings provide direct evidence that sister kinetochore fusion underlies the cosegregation of sister chromatids during meiosis I.

REFERENCES AND NOTES


Spatial and temporal diversity in genomic instability processes defines lung cancer evolution

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Spatial and temporal dissection of the genomic changes occurring during the evolution of human non–small-cell lung cancer (NSCLC) may help elucidate the basis for its dismal prognosis. We sequenced 25 spatially distinct regions from seven operable NSCLCs and found evidence of branched evolution, with driver mutations arising before and after subclonal diversification. There was pronounced intratumor heterogeneity in copy number alterations, translocations, and mutations associated with APOBEC cytidine deaminase activity. Despite maintained carcinogen exposure, tumors from smokers showed a relative decrease in smoking-related mutations over time, accompanied by an increase in APOBEC-associated mutations. In tumors from former smokers, genome-doubling occurred within a smoking-signature context before subclonal diversification, which suggested that a long period of tumor latency had preceded clinical detection. The regionally separated driver mutations, coupled with the relentless and heterogeneous nature of the genome instability processes, are likely to confound treatment success in NSCLC.

Lung cancer is the leading cause of cancer-related mortality (1, 2). Understanding the pathogenesis and evolution of lung cancer may lead to greater insight into tumor initiation and maintenance and may guide therapeutic interventions. Previous work characterizing the genome of non–small-cell lung cancer (NSCLC) has demonstrated that NSCLC genomes exhibit hundreds of nonsilent mutations together with copy number aberrations and genome doublings (3–9). Although subclonal populations have been identified within single biopsies (9), the extent of genomic diversity within primary NSCLCs remains unclear. Moreover, although both exogenous mutagenic processes, such as smoking (10–12), and endogenous processes, such as up-regulation of APOBEC cytidine deaminases (13–15), have been found to contribute to the large mutational burden in NSCLC, the temporal dynamics of these processes and their contribution to driver somatic aberrations over time remain unknown.

To investigate lung cancer evolution, we performed multiregion whole-exome and/or whole-genome sequencing (M-seq WES/WGS) on a total of 25 tumor regions, collected from seven NSCLC patients who underwent surgical resection before receiving adjuvant therapy. The major NSCLC histological subtypes, including adenocarcinoma (LUAD) and squamous cell carcinoma (LUSC), were represented (table S1). Sequencing of tumor and normal DNA to mean coverage depths of 107× and 54× for M-seq WES and M-seq WGS, respectively (table S2), identified 1884 nonsilent and 76,129 silent mutations (16).

To evaluate the intratumor heterogeneity of nonsilent mutations, we classified each mutation as ubiquitous (present in all tumor regions) or heterogeneous (present in at least one, but not all, regions). Spatial intratumor heterogeneity was identified in all seven NSCLCs, with a median of 30% heterogeneous mutations (range 4–90%); all these tumors also showed regions of ubiquitous mutations (table S3). The median numbers of nonsilent mutations were 22,061 (range, 14,991 to 35,856) in the 25 regions from seven NSCLCs.

When the nonsilent mutations were plotted spatially (fig. S3), we observed extensive spatial diversity in the distribution of nonsilent mutations within the tumors. This diversity was restricted to individual regions, with similar nonsilent mutation patterns seen in each sample from the same tumor but with different mutation patterns across the seven different tumors. These results demonstrate that the nonsilent mutations identified in these tumors reflect the spatial diversity of the tumors and are not due to technical or sequencing artifacts.
To resolve the extent of genomic diversity in NSCLC and to infer the ancestral relations between tumor regions, we estimated the fraction of tumor cells within each region harboring each mutation. Almost all ubiquitous mutations (>99%) were classified as fully clonal within each region. Moreover, in most regions, the majority of heterogeneous mutations was clonal and, thus, present in all cells within the region (Fig. 1B and fig. S3). However, certain regions displayed considerable subclonal diversity. For example, >75% of heterogeneous mutations present in L004 R5 were subclonal (Fig. 1B), and this region consisted of two distinct subclonal populations. The subclonal structure of each tumor region was then used to construct phylogenetic trees, by using both maximum parsimony and unweighted pair-group methods. We also took into account regional copy number losses that resulted in shared truncal mutations becoming heterogeneous later in tumor evolution (16), such as a segment of chromosome 6 in LS01 (fig. S4) and the PAX7 mutation in the lymph node of L001 (Fig. 1A). Notably, all seven NSCLCs showed evidence of branched tumor evolution (fig. S5).

We next evaluated the regional heterogeneity of potential NSCLC driver mutations, classified into three categories on the basis of current evidence supporting driver mutation status (16). Every tumor showed evidence for ubiquitous, as well as heterogeneous, driver mutations, many of which were clonally dominant in a subset of tumor regions and entirely absent in others (Fig. 1B, fig. S3, and table S3). Note that the probability of missing a category 1 “high-confidence” driver gene by analyzing a single region for each

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**Fig. 1. Intratumor heterogeneity of somatic mutations in human NSCLC.** (A) Heat maps show the regional distribution of all nonsilent mutations; presence (blue) or absence (gray) of each mutation is indicated for every tumor region. Cartoons depict the location of each tumor. Column next to heat map shows the intratumor heterogeneity; mutation present in all regions (blue), in more than one but not all (yellow), or in one region (red). Mutations are ordered on tumor driver category with categories 1 to 3 indicated in the right column in black, dark gray, and light gray, respectively (details in table S3). Total number of nonsilent mutations is provided below each tumor with percentage of heterogeneous mutations in brackets. In L001, the mutation marked by an asterisk (*) is additional to the germline MEN1 mutation. LN, lymph node; R, region. (B) Two-dimensional Dirichlet plots show the cancer cell fraction (CCF) of the mutations in all regions of tumors L004; increasing intensity of red indicates the location of a high posterior probability of a cluster. In region R5, the majority of heterogeneous mutations were subclonal, and a cluster of mutations with a CCF below 1 can be observed.
tumor was on average 42% (range 0 to 67%) and 83% (range 67 to 100%) for all potential driver gene categories (1 to 3), which highlights the potential limitations of assessing single tumor regions. Nevertheless, category 1 and 2 driver mutations were significantly more often truncal compared with mutations in nondriver genes in our M-seq analysis ($P = 0.04$). Consistent with these data, in The Cancer Genome Atlas (TCGA) cohort, previously reported driver genes (5, 19, 20) were significantly enriched for clonal mutations ($P < 0.0001$) (fig. S6). These data indicate that, in NSCLC, most known driver mutations occur early in tumor evolution.

To determine the intratumor heterogeneity of copy number aberrations, we estimated integer DNA copy numbers for each tumor region (7, 16, 21, 22). A large fraction of the genome had undergone alterations in all tumors, and genomic profiles were more similar within tumors than between different tumors (fig. S7). To evaluate the spatial heterogeneity of potential tumor driver copy number aberrations, we explored the regional distribution of chromosomal segments identified as recurrently gained or lost in TCGA LUAD or LUSC tumors. Most segments were identified as aberrant in at least one tumor region, and many recurrent gains and losses were identified as aberrant in at least one tumor region (Fig. 2A).

For example, in L001, a focal EGFR amplification (chr7p11.2), as well as deletions of chromosomal segments harboring CDKN2A (chr9p21.3) and PTEN (chr10q23.31), was observed in all regions, whereas, in L008, we observed heterogeneous copy number losses involving CDKN2A and PTEN. In support of copy number aberrations occurring later in tumor development, we also identified subclonal copy number aberrations within tumor regions. For instance, more than 15% of the genome in region R1 of L008 was subject to subclonal copy number alterations (fig. S8). Consistent with evidence of subclonal copy number aberrations, centromeric fluorescence in situ hybridization analyses confirmed numerical chromosomal diversity within individual tumor regions (fig. S9), which suggested that chromosomal instability may provide a substrate for subclonal competition.

The high-coverage M-seq WGS (mean 96×) for L002 and L008 enabled us to investigate the regional separation of large-scale genomic events in these samples. For the adenocarcinoma tumor L002, we identified 30 structural variants, most of which were found either in the LUAD region R1 or the LUSC region R3, but not both (table S4), which suggested that they occurred after subclonal diversification (Fig. 2B). By contrast, for L008, 48 of the 52 identified structural variants were shared between the two tumor regions from different lobes of the lung, which suggested that the majority of these variants occurred before tumor metastasis to the other lobe (Fig. 2B). Notably, in L008, “chains” of translocations with highly clustered breakpoints were found between chromosomes 14 and 17, as well as chromosomes 17 and 19 (fig. S10 and table S4), which disrupted the FANCM and NF1 tumor suppressor genes. Breakpoint homology profiling suggests involvement of either nonhomologous or alternative end-joining (23, 24), indicative of double-strand break events. This lesion pattern is consistent with chromoanagenesis (25) and indicates a punctuated evolution pattern where multiple oncogenic events may have occurred simultaneously (26).

Four tumors displayed evidence for whole-genome-doubling events (16). In three tumors (L001, L004, and L008), the genome-doubling event was shared across every tumor region; it occurred before diversification, with the majority of truncal mutations (84 to 88%) present at ploidy ≥2, indicative of a large mutational burden before genome doubling. In one tumor, L002, the majority of heterogeneous mutations were also present at ploidy ≥2, indicative of two independent genome-doubling events: one in the LUAD region and one in the LUSC region (fig. S11). Notably, every truncal driver mutation likely occurred before genome doubling.

To further explore the dynamics of the mutational processes shaping lung cancer genomes over time, the spectra of point mutations in each tumor were temporally dissected. Early (truncal) mutations likely reflect processes involved before and during tumor initiation and early development, whereas late (branched) mutations...
reveal mutational processes shaping the genome during tumor maintenance and progression, including those contributing to intratumor heterogeneity. For L002, we analyzed regions R1 and R3 separately to allow comparisons of LUAD and LUSC histologies within the same tumor.

In all tumors, we observed statistically significant shifts in the mutation spectra over time ($P < 0.05$ all cases) (Fig. 3A). Furthermore, every tumor exhibited a statistically significant decrease in the proportion of C>A transversions in late compared with early mutations ($P < 0.05$) (Fig. 3A), although this was more pronounced in the LUAD cases [mean odds ratio: LUAD 3.13 (range 2.07 to 5.55) and LUSC 1.34 (range 1.21 to 1.46)]. Because C>A transversions are associated with the mutagenic effects of tobacco smoke (12), a decrease in the proportion of C>A transversions indicates a relative decrease in the mutational burden attributable to smoking during LUAD development, in both former smokers and current smokers.

To validate these observations in a larger NSCLC cohort, mutations in TCGA LUAD and LUSC samples were temporally dissected (16). Consistent with our M-seq analyses, both TCGA LUAD and LUSC smokers and former smokers exhibited a decrease in the proportion of C>A transversions in late mutations (LUAD current smokers, $P < 0.0001$; former smokers, $P < 0.0001$; never-smokers, $P = 0.147$; LUSC current smokers, $P = 0.003$; former smokers, $P < 0.0001$; and never-smokers, $P = 0.073$) (Fig. 3B). Similarly, the least-pronounced decrease was observed in LUSC current smokers; 25% of LUSC displayed no decrease in C>A transversions, compared with less than 10% in LUAD. The mutational footprint of smoking exhibits a strand bias with C>A transversions accumulating preferentially on the transcribed strand (10, 12).

The mutational footprint of smoking reveals a strand bias with C>A transversions accumulating preferentially on the transcribed strand (10, 12). Both LUAD and LUSC former smokers revealed a statistically significant decrease in strand bias in late, compared with early, C>A transversions.

Fig. 3. Temporal and spatial dissection of mutation spectra in LUAD and LUSC samples. (A) Fraction of early mutations (trunk) and late mutations (branch) accounted for by each of the six mutation types in all M-seq samples. (B) Beeswarm plots showing the fraction of early mutations and late mutations accounted for by each of the six mutation types in every TCGA former smoker or current smoker with both early and late mutations. Significance is indicated. (C) APOBEC mutation enrichment odds ratio for early (trunk, blue bars) and late (branch, red bars) mutations for M-seq samples. The APOBEC signature encompasses C>T and C>G mutations in a TpC context (16). The 95% confidence intervals for Fisher’s exact test are indicated. (D and E) Three mutation types (C>A; C>G and C>T) at all 16 possible trinucleotide contexts for L002 (D) and L008 (E). For both samples, trunk mutations as well as branch mutations from two regions are depicted.
In the majority of M-seq tumors, the decreased proportion of C>A mutations was accompanied by an increase in C>T and C>G mutations at TpC sites, indicative of APOBEC-cytidine deaminase activity (13–15). Mutations consistent with APOBEC-mediated mutagenesis were more pronounced on the branches than the trunk in four out of five LUAD M-seq samples (Fig. 3C). On average, 31% (8 to 41%) of nons silent branch mutations occurred in an APOBEC-mutation context compared with 11% (7 to 16%) of trunk nonsilent mutations. Branched driver genes PIK3CA, EP300, TGFBR1, PTPRD, and AKA9 harbored mutations in an APOBEC context, which indicated a possible functional impact of APOBEC activity on subclonal expansion. Likewise, TCGA LUAD tumors with detectable APOBEC mutational signatures showed significant enrichment in late, compared with early, APOBEC mutations (P < 0.001) (fig. S12), and 20% of subclonal driver mutations were found to occur in an APOBEC context, compared with 11% of clonal driver mutations. However, for TCGA LUSC tumors with detectable APOBEC mutational signatures, temporal dissection of APOBEC mutations did not reveal such a clear trend (fig. S12), which indicated potential differences in the temporal dynamics of APOBEC-mediated mutagenesis between histological subtypes. In addition to temporal heterogeneity, spatial heterogeneity in both the proportion of APOBEC-associated mutations (Fig. 3, D and E) and APOBEC mRNA expression was observed in the M-seq tumors (fig. S13).

To gain a deeper understanding of NSCLC evolution, we focused on the two tumors with high-coverage M-seq WGS and temporally placed the genomic instability processes relative to the emergence of the most recent common ancestor (Fig. 4). In patient L002, a current smoker, tobacco carcinogens played a significant role early in tumor development, with C>A transversions representing 39% of truncal mutations (Fig. 4A). Early mutations included multiple driver genes, such as TP53 and CHD8. Upon diversification into a LUAD subclone and a LUSC subclone, copy number alterations (fig. S7) and driver mutations were acquired independently in both subclones, such as a stop-gain mutation in the tumor suppressor gene FAT1 on the LUAD branch and mutations affecting TGFBR1, ZFHX4, ARHGAP35, and PTEN in the LUAD region. APOBEC-associated mutations were elevated specifically in the LUAD region, which included the driver mutations in TGFBR1 and PTEN, and the highest APOBEC3B mRNA expression was detected in this region (fig. S13).

The tumors from patient L008 also displayed truncal C>A transversions and spatial heterogeneity in APOBEC enrichment, with a more pronounced APOBEC signature in the tumor of the middle lobe compared with the upper lobe (Fig. 4B). In L008, we gained further temporal resolution by exploring the mutations before and after the truncal genome-doubling event. All truncal driver mutations were found to occur before genome doubling. However, a tobacco smoke signature of C>A transversions was observed in more than 30% of truncal mutations both before and after doubling, and in 21% and 9% of heterogeneous mutations in the two regions R1 and R3 from separate lobes of the lung. Because L008 ceased smoking more than 20 years before surgery (table S1), these data suggest that the genome-doubling event and truncal driver mutations occurred within a smoking carcinogenic context more than 20 years ago. Similarly, the genome-doubling event and truncal driver mutations in former smoker L001 also appeared to occur before smoking cessation more than 20 years before surgery (fig. S14). These data suggest a prolonged tumor latency period after genome doubling and before clinical detection in NSCLC.

Through sequencing multiple surgically resected tumor regions, we were able to unravel both the extent of genomic heterogeneity and the evolutionary history of seven NSCLCs. In contrast to the situation in clear cell renal cell carcinoma (cRCC) (27, 28), known driver mutations typically occurred early in NSCLC development, and the majority of high-confidence driver events were fully clonal. Conceivably, this explains the progression-free survival benefits associated with NSCLC oncogenic driver targeting (29). However, like cRCC (27, 28), all NSCLCs exhibited heterogeneous driver mutations and/or recurrent copy number aberrations and many heterogeneous mutations gave the “illusion of clonality,” as they are present in all cells from certain regions but undetectable within other regions. Notably, although our multiregional sampling approach allowed us to evaluate spatial heterogeneity, only a small part of the entire tumor was sampled (on average <5%), which indicates that we might be underestimating the full extent of heterogeneity in these tumors.

Conceivably, intratumor heterogeneity may compromise the ability of a single biopsy to define all driver events comprehensively for optimal tumor control. For instance, L008 presented with an activating BRAF (G469A) mutation (30) in all regions and an activating PIK3CA (E542K) mutation (31) only in region R3. Thus, a biopsy taken from R3 might suggest treatment with an inhibitor of the phosphatidylinositol 3-kinase–mammalian target of rapamycin (PI3K/mTOR) signaling axis and combination therapy. Conversely,
a single biopsy from any other region would suggest treatment with a BRAF inhibitor, for which the tumor cells from R3 might be resistant because of the PI3KCA mutation (32).

Our study also sheds light on the divergent genomic instability processes involved in NSCLC evolution and their dynamics over time. Evidence for spatial diversity in genomic instability processes suggests that opportunities to exploit such mechanisms therapeutically may be limited in this disease (33). In three tumors, we detected genome-doubling events occurring before subclonal diversification but after acquisition of driver mutations, consistent with findings in colorectal cancer that genome doubling may accelerate cancer genome evolution (34). The relation of chromosomal instability with drug resistance and early tumor recurrence (35, 36) suggests that targeting truncal driver events may accelerate cancer genome evolution. A large proportion of subclonal additional genomic instability process (APOBEC-driven) may cause of the mutations, as well as the selective advantage of differences in mutation spectra over time and space may reflect the activity of the process generating the mutations, as well as the selective advantage of the acquired mutations.

The presence of subclonal, regionally separated driver events, coupled with the relentless and dynamic nature of genomic instability processes observed in this study, highlight the therapeutic challenges associated with NSCLC. Engaging an adaptable immune system may present a tractable approach to manage the dynamic complexity in NSCLC (37). Longitudinal studies will be required to decipher drivers of subclonal expansion, identify the origins of subclones contributing to metastatic recurrence, and resolve the evolutionary principles that underpin the dismal outcome associated with this disease.

REFERENCES AND NOTES


16. Detailed information on methods is available on Science online.

LUNG CANCER EVOLUTION

Intratumor heterogeneity in localized lung adenocarcinomas delineated by multiregion sequencing

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Cancers are composed of populations of cells with distinct molecular and phenotypic features, a phenomenon termed intratumor heterogeneity (ITH). ITH in lung cancers has not been well studied. We applied multiregion whole-exome sequencing (WES) on 11 localized lung adenocarcinomas. All tumors showed clear evidence of ITH. On average, 76% of all mutations and 20 out of 21 known cancer gene mutations were identified in all regions of individual tumors, which suggested that single-region sequencing may be adequate to identify the majority of known cancer gene mutations in localized lung adenocarcinomas. With a median follow-up of 21 months after surgery, three patients have relapsed, and all three patients had significantly larger fractions of subclonal mutations in their primary tumors than patients without relapse. These data indicate that a larger subclonal mutation fraction may be associated with increased likelihood of postsurgical relapse in patients with localized lung adenocarcinomas.

Intratumor heterogeneity (ITH) may have impacts on tumor biopsy strategy, characterization of actionable targets, treatment planning, and drug resistance (1–6). ITH has recently been elucidated in substantial detail in several cancer types with the use of next-generation sequencing (NGS) approaches (7–14). Recent evidence supports a model of branched evolution leading to variable ITH in different tumors (9, 13, 15, 16). Studies in clear cell renal carcinoma

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SUPPLEMENTARY MATERIALS

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Materials and Methods

Figs. S1 to S14

Tables S1 to S4

References (38–64)

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