Mutations in the \textit{TERT} promoter represent the genetic underpinnings of tumor cell immortality. Beyond the two most common point mutations, which selectively recruit the ETS factor GABP to activate \textit{TERT}, the significance of other variants is unknown. In seven cancer types, we identify duplications of wildtype sequence within the core promoter region of \textit{TERT} that have strikingly similar features including an ETS motif, the duplication length and insertion site. The duplications recruit a GABP tetramer by virtue of the native ETS motif and its precisely spaced duplicated counterpart, activate the promoter and are clonal in a \textit{TERT} expressing multifocal glioblastoma. We conclude that recurrent \textit{TERT} promoter duplications are functionally and mechanistically equivalent to the hotspot mutations that confer tumor cell immortality. The shared mechanism of these divergent somatic genetic alterations suggests a strong selective pressure for recruitment of the GABP tetramer to activate \textit{TERT}.

Limitless replicative potential is one of the hallmarks of cancer cells\textsuperscript{1}, enabling malignant transformation. This cellular immortalization can be achieved by reactivating telomerase, an enzyme that maintains telomere length during cell division. Genetic alterations that lead to activation of expression in its subunit \textit{Telomerase Reverse Transcriptase (TERT)} are often found in cancer\textsuperscript{2–8}. In fact, \textit{TERT} promoter (\textit{TERTp}) mutations are the most common noncoding mutations in human cancers\textsuperscript{9} including the majority of glioblastomas\textsuperscript{9}. The two \textit{TERT} hotspot mutations G228A and G250A in the core promoter region are annotated by their genomic coordinate, chr5:1,295,228 and chr5:1,295,250 (human genome build, hg19), respectively. Lower frequency recurrent \textit{TERTp} mutations consist of T161G, G228T, GG228A and GG242A\textsuperscript{4,11}. \textit{TERTp} mutations can also be annotated by their position relative to \textit{TERT} ATG or translation start, for example -124C > T (G228A). The G228A and G250A \textit{TERT} mutations create new E\textsubscript{26} transformation-specific (ETS) binding sites (CCGGAA) and recruit the GA-binding protein (GABP) transcription factor (TF) complex to activate \textit{TERT}\textsuperscript{12–15}. The GABP TF complex is an obligate multimer consisting of the DNA binding subunit GABPA and the transactivation subunit GABPB1\textsuperscript{11}. Depending on the isoform of the GABPB1 subunit,
the complex either forms a heterotetramer containing GABPA-GABPB1L or heterodimer consisting of GABPA-GABPBP1S. In the case of the mutant TERTp, the generation of a new ETS motif is positioned so that it is in helical phase with the native ETS motif and the distance between the two motifs is ideal for GABP heterotetramer binding24.

The TERTp is subject to other alterations in cancer such as insertion/deletion mutations, amplifications and epigenetic modifications25-27. The significance of many of these alterations, and their potential mechanism of telomerase reactivation is unknown. In this regard, two recent case reports observed an intriguing 22 bp duplication of unknown function within the TERTp in two glioblastomas and a thyroid cancer22,23.

Here, we identified 21 tumor samples of 7 different cancer types with 21–25 bp duplications of wildtype sequence within the core promoter. This sizeable cross-cancer cohort allowed us to discover that the sequence, length and insertion position of the duplication are remarkably conserved, identical in many cases, strongly suggesting a pathogenic role. Furthermore, these observations raised a hypothesis which we test to explain why the features are conserved and mechanistically how they compare to hotspot mutations. Together our results point to the apparent positive selection of wildtype sequence duplications that, like hotspot mutations, recruit the GABP tetramer to activate TERT.

Results

Duplications in the core TERT promoter occur in multiple cancer types and have conserved features

Recurrent point mutations in the TERTp that generate de novo ETS binding sites recruit the GABP heterotetramer TF complex (GABPB1L and GABPA) to activate telomerase expression (Fig. 1a, b)28-30. In contrast, in cells with the wildtype TERTp GAB binding is not detectable and GABP knockdown does not affect TERT expression. In our cancer panel sequencing of glioblastomas, we identified a tumor lacking either of the hotspot point mutations but instead harboring a duplication of wildtype sequence with the core TERTp. Although the duplication was deemed a variant of unknown significance, we found intriguing similarity to three duplications reported in two case studies22,23. To begin to understand the frequency and features of ETS motif containing TERTp duplications in cancer, we surveyed the literature along with AACR Project Genomics Evidence Neoplasia Information Exchange (GENIE) and UCFS500 Pan-Cancer datasets. Most of the AACR GENIE TERTp sequencing data were generated by the MSK-Impact targeted sequencing panel. Although the MSK-Impact publication captured variants in the TERTp, hotspot mutations were the main focus. Subsequently, additional tumors have been profiled by MSK-IMPACT and are included in the GENIE cohort, v101,31. The UCFS500 Pan-Cancer dataset is available to UCSC affiliates and was generated by the UCFS500 targeted sequencing panel, which covers the TERTp. Table 1 includes 21 duplications from 18 cases we identified that are contained within the core region of the TERTp, including six samples with duplications from four patients in our cohort. Comparing the studies in Table 1, TERTp duplication variants are represented by cancer types with the highest frequencies of the hotspot TERTp mutations, which include bladder cancer, glioblastoma, squamous cell carcinoma of the oropharynx, hepatocellular carcinoma and thyroid cancer. TERTp duplications are also present in rare forms of breast and intestinal cancer. Comparing patient demographics, TERTp duplications are represented by different age groups, sex, and races (Table 1). Together these data show that TERTp duplications are not restricted to a specific patient demographic (age or sex) or tumor type (primary site, metastasis or recurrent tumor), but they do tend to occur in the same cancer types as those with frequent TERTp hotspot mutations suggesting they may serve a similar function. Although TERTp duplications and mutations occur in similar cancer types, broadly, TERTp duplications occur at a lower frequency (Supplementary Fig. 1a, Supplementary Table 1).

We next carefully compared the TERTp duplication variants in Table 1 to identify any unifying features that might give clues to their function and made two striking observations. Irrespective of cancer type, duplications have an approximate size of 22 bp and contain an exact copy of one of the native ETS motifs, specifically ETS 200 (Table 1). To determine if the ETS containing TERTp duplications in Table 1 had unique features as compared to non-ETS duplications in the TERTp, we mapped all MSK-Impact duplication variants within a 4 kb region upstream of TERTp (chr5:1295160-1299000) and observed a broad spectrum of insert sizes with no obvious bias for their insert position (Supplementary Fig. 1b). However, when specifically mapping these variants by presence or absence of the GGAA motif, 13 of the 14 GGAA containing insertions were localized within the core promoter adjacent to the native ETS motif (chr5:1295180-1295199) and approximately 22 bp in length. However, the 19 non-GGAA containing insertions were highly variable in length and dispersed throughout the TERTp, which suggests that ETS containing duplications have a selective pressure for their length and insertion point (Supplementary Fig. 1b). Interestingly, the one GGAA containing duplication variant that did not map directly adjacent to the native ETS (c.125-106) is different than the others as it is further upstream in the TERTp and the ETS motif is actually a copy of the G228A hotspot mutation (Table 1; Supplementary Fig. 1b, c). In agreement, this patient was also diagnosed with a G228A mutation. We identified one additional tumor that showed both TERTp G228A hotspot mutation and TERTp duplication containing the wildtype sequence. Although G228A and G250A hotspot mutations are considered mutually exclusive, rare cases with both mutations have been reported as well.

Regarding the unique features of duplication variants (Table 1), we previously reported the favored features of the GABP heterotetramer TF complex binding for TERTp activation as having two ETS binding sites recruit the GABP tetramer to activate transcription. In support, two of the 22 bp TERTp duplications previously reported and included in Table 1 increase activity of the TERTp in a promoter-reporter assay. Based on sequence motif analysis of the duplication, over one hundred potential function and made two striking observations. Irrespective of cancer type, duplications have an approximate size of 22 bp and contain an exact copy of one of the native ETS motifs, specifically ETS 200 (Table 1). To determine if the ETS containing TERTp duplications in Table 1 had unique features as compared to non-ETS duplications in the TERTp, we mapped all MSK-Impact duplication variants within a 4 kb region upstream of TERTp (chr5:1295160-1299000) and observed a broad spectrum of insert sizes with no obvious bias for their insert position (Supplementary Fig. 1b). However, when specifically mapping these variants by presence or absence of the GGAA motif, 13 of the 14 GGAA containing insertions were localized within the core promoter adjacent to the native ETS motif (chr5:1295180-1295199) and approximately 22 bp in length. However, the 19 non-GGAA containing insertions were highly variable in length and dispersed throughout the TERTp, which suggests that ETS containing duplications have a selective pressure for their length and insertion point (Supplementary Fig. 1b). Interestingly, the one GGAA containing duplication variant that did not map directly adjacent to the native ETS (c.125-106) is different than the others as it is further upstream in the TERTp and the ETS motif is actually a copy of the G228A hotspot mutation (Table 1; Supplementary Fig. 1b, c). In agreement, this patient was also diagnosed with a G228A mutation. We identified one additional tumor that showed both TERTp G228A hotspot mutation and TERTp duplication containing the wildtype sequence. Although G228A and G250A hotspot mutations are considered mutually exclusive, rare cases with both mutations have been reported as well.

Regarding the unique features of duplication variants (Table 1), we previously reported the favored features of the GABP heterotetramer TF complex binding for TERTp activation as having two ETS binding sites in phase such that their spacing has a periodicity of 10.5 bp (full helical turns), a preference we also observed genome wide12. The distance of 22 bp between the duplicated de novo ETS motif and native ETS motif is approximately 2 full helical turns, similar to the distance from the native ETS to either hotspot mutation, putting these two motifs in phase for potentially optimal GABP heterotetramer binding (Fig. 1a, c). Supplementary Fig. 1c).

TERT promoter activation requires the ETS containing duplication and GABP

The wildtype TERT promoter is incapable of significant GABP tetramer recruitment, and yet the duplicated sequences are wildtype, leading to uncertainty about how the duplications might activate TERT in vivo, and indeed whether they do. However, the precise positioning of the newly acquired ETS motif and its position suggested to us that TERTp duplications might function like TERTp hotspot mutations to recruit the GABP tetramer to activate transcription. In support, two of the 22 bp TERTp duplications previously reported and included in Table 1 increase activity of the TERTp in a promoter-reporter assay. Based on sequence motif analysis of the duplication, over one hundred potential TF were predicted that could play a role in TERTp activation22,23. From the conserved features in our relatively large number of cases and in our prior studies, we hypothesized that any 22 bp TERTp duplication that specifically includes the native ETS motif could increase promoter activity and the increase would be dependent on insertion point and GABP. To test this hypothesis, we generated twenty luciferase reporter constructs covering all possible 22 bp TERTp duplication variants containing the native ETS motif and performed luciferase assays using bladder cancer (UMUC3) and glioblastoma (LN229) cell lines (Fig. 1). We observed that all duplications of the native ETS motif were sufficient for increased TERTp activity, at a level very similar to G228A and G250A hotspot mutations (Fig. 1d, e). Interestingly, only duplications containing a copy of native ETS 200, not ETS 195, were sufficient for increased promoter activity. Specifically, duplications
containing only a copy of native ETS 200 (-115_94 to -117_96) showed increased reporter activity when compared to the TERT wildtype promoter while duplications containing only a copy of native ETS 195 (-94_73 to -99_78) did not (Fig. 1d–f). These data agree with Table 1 where ETS 200 is present in all duplicated sequences while ETS (-94_-73 to -98_-77) did not (Fig. 1d).

**Table 1**

| Duplicated Region | Duplication Insertion | De Novo ETS 195 200 | Native ETS 195 200 | Reporter Activity |
|-------------------|-----------------------|---------------------|-------------------|------------------|
| Wild-Type         | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -98_94            | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -99_98            | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -100_99           | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -101_100          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -102_101          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -103_102          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -104_103          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -105_104          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -106_105          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -107_106          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -108_107          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -109_108          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -110_109          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -111_110          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -112_111          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -113_112          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -114_113          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -115_114          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -116_115          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -117_116          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -118_117          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -119_118          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -120_119          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |
| -121_120          | GCCGGAGGAGGAGGCCGCGGGGC| No change            | No change         | Increased         |

**Figures**

- **Fig. 1a**: TERTp Wild-Type promoter structure.
- **Fig. 1b**: TERTp Hot Spot Mutations.
- **Fig. 1c**: TERTp Duplications.
- **Fig. 1d**: Firefly/Renilla reporter activity.
- **Fig. 1e**: Firefly/Renilla reporter activity.

**Supplementary Figures**

Supplementary Fig. 1c). To determine if this distance of 22 bp between the duplicated de novo and native ETS motif is essential for promoter activity, insertion and deletion mutations were performed to alter the spacing of the c.-100_79 duplication variant (Supplementary Fig. 2a). We performed luciferase assays using bladder cancer and glioblastoma cell lines and observed that TERTp activity was similar between c.-100_79 and the G228A hotspot mutant (Supplementary Fig. 2b, c). However, promoter activity was abolished as bases were either inserted or deleted from this duplication sequence (Supplementary Fig. 2b, c).
Interestingly, **TERTp** activity increased again as bases were inserted and the distance between de novo and native ETS motifs was around 30 base pairs or approximately 3 helical turns, which put the ETS binding sites in phase with each other again (Supplementary Fig. 2b, c). Although these motifs were in phase, the activity never reached that of c-100–79, which suggests the distance of 2 helical turns may be ideal for **TERTp** duplications. The results across this large collection of engineered variants of the duplicated promoter were consistent between the two cancer cell lines with substantially different mutational profiles, transcription factor expression patterns and epigenetic states.

The shared features between the **TERTp** duplication and hotspot mutations suggested to us that GABP may play a functional role in the activation of the **TERTp** duplication. To determine if the **TERTp** duplication is dependent on GABP for the increase in promoter activity, we created UMC3 and LN229 cells with control and GABPA siRNAs to deplete the GABP TF complex then performed luciferase assays with these **TERTp** duplication sequences. Similar to our observations in Fig. 1d, e, all ETS containing duplications showed increased reporter activity when compared to the **TERT** wildtype promoter. In stark contrast, upon knockdown of GABPA, we observed a dramatic reduction to wildtype **TERTp** levels for all ETS containing duplications in both cell lines, and this level of reduction was similar to G228A and G250A hotspot mutations (Fig. 2a, b). Next, we investigated whether GABP can bind to the fragment of the **TERTp** with the 22 bp insertion in an electromobility shift assay. Our results show GABP binds to sequences with duplicated native ETS sites (Fig. 3). Together, these data show that GABP is an essential regulator of the **TERTp** duplication, mimicking the hotspot mutations in activation strength and mechanism.

**Clonality of the TERT promoter duplication in a multifocal glioblastoma**

**TERTp** mutations are sufficient for the diagnosis of glioblastoma in IDH-wildtype diffuse astrocytic gliomas according to the latest 2021 WHO Classification of Central Nervous System Tumors. In contrast, the diagnostic value of **TERTp** duplications is unknown. We identified a patient with a **TERTp** duplication whose clinical history and tumor histological and molecular profile were consistent with an integrated diagnosis of glioblastoma, IDH-wildtype and for which intratumoral samples were available for clonality analysis (Fig. 4a).

The patient presented with two months of disorientation, double vision, right-sided vision loss and neglect, as well as difficulty with fine motor movements. Imaging analysis showed a multifocal enhancing mass in the left parieto-occipital lobes (Fig. 4b, c). Histology revealed a diffuse astrocytic glioma with pleomorphic tumor cells, brisk mitotic activity, microvascular proliferation, and palisading necrosis (Fig. 4d, e). The patient was treated with radiation and temozolomide, and experienced tumor recurrence after 13 months (Supplementary Fig. 3a, d).

From a total of 13 independent punches from five FFPE tissue blocks of the two portions of the newly diagnosed tumor, we purified genomic DNA and total RNA. We first performed PCR on the DNA to determine if the **TERTp** duplication is present in a subset of these samples suggesting sub-clonality, or if it is detectable in all available samples, indicative of an evolutionarily early and clonal event. The duplicated allele resolved as a higher molecular band on the agarose gel as compared to the wildtype allele (Fig. 4f). The detection in all thirteen DNA samples further highlights the early evolutionary origin and clonality of the **TERTp** duplication in this case. We also detected a small amount of the duplication PCR product in the "normal" tissue sample, which was brain parenchyma immediately adjacent to the tumor that likely contained a small population of infiltrating tumor cells.

To determine when the **TERTp** duplication was acquired during tumor evolution relative to other alterations, we performed exome sequencing on three different locations of the occipital and two different locations of the parietal primary tumor, as well as patient-matched blood cell DNA. We determined the SNV, copy number and tumor purity estimates and then used the SNV to generate a sample-oriented phylogenetic reconstruction and a complementary tumor clone-based analysis using the PyClone algorithm. Gain of chromosome 7, loss of chromosome 10 and homozygous deletion of the **CDKN2A** locus were confirmed in all analyzed tumor parts, as was the **TERTp** duplication and an **NFI** frameshift mutation (Fig. 4g, Supplementary Figs. 3b, 5a–e). The **TERTp** duplication is present throughout the parietal and occipital portions of the tumor that were analyzed, similar to the distribution of the characteristically clonal alterations (chromosome 10 loss and chromosome 7 gain). In contrast to the clonal events that form the trunk of the phylogenetic tree, sub-clonal driver events were also readily apparent and underlie the tree branches. For example, only the occipital tumor region showed amplification of the **EGFR** locus, whereas a **PTPN11** missense mutation and **ERRF1** homozygous deletion were exclusively in the parietal tumor region (Fig. 4g, Supplementary Figs. 3b, 5a–e). PyClone analysis allowed us to infer the clonal population structure and revealed different gene mutation clusters in different parts of the multifocal tumor, except for cluster 1 which is present in all parts representing fully clonal, or "truncal" mutations that we infer to have arisen very early in the evolution of this multifocal tumor. Cluster 1 from the PyClone analysis and the trunk of the phylogenetic tree contain a shared set of 17 mutations which, along with the clonal copy number changes, may represent the ancestral clone that seeded the occipital and parietal portions of this tumor. Cluster 2 includes the **PTPN11** mutation and was present only in the parietal tumor focus while cluster 3 which includes **EGFR** amplification was only detected in the occipital tumor focus (Fig. 4h). Like cluster 3, clusters 4 and 5 were also confined to the occipital lobe but appeared to be less abundant. Panel sequencing of a low tumor purity sample of the recurrent tumor revealed a newly acquired **PTEN** tandem duplication in exon 4.
Table 1 | Tumor samples with an ETS binding site duplication in the TERT promoter

| Duplicated Region1 | Duplicated Sequence | Insert Size (bp) | Insert Position2 | Other Acquired ETS Variants | Age | Race | Sex | Cancer Type | Tumor Type | Dataset |
|-------------------|---------------------|------------------|------------------|-----------------------------|-----|------|-----|-------------|------------|---------|
| c.-100_-79        | GGCCGGCGCGGGAAAAGGAG | 22               | 1295182 1295183  | NA                          | NA | NA  | NA  | Glioblastoma Multiforme | NA         | Pierini et al. |
| c.-100_-792       | GGCCGGCGCGGGAAAAGGAG | 22               | 1295182 1295183  | None                        | 54 | NA  | Male | Glioblastoma Multiforme | NA         | UCSF500  |
| c.-100_-79        | GGCCGGCGCGGGAAAAGGAG | 22               | 1295182 1295183  | G228A                       | 29 | NA  | Female | Squamous Cell Carcinoma | Metastasis | UCSF500  |
| c.-100_-79        | GGCCGGCGCGGGAAAAGGAG | 22               | 1295182 1295183  | None                        | 68 | White | Male | Duodenal Adenocarcinoma | Metastasis | GENIE   |
| c.-104_-83        | GGCCGGCGGGAAAGGAGGGA | 22               | 1295186 1295187  | None                        | 42 | Asian | Male | Oral Cavity Squamous Cell Carcinoma | Metastasis | GENIE   |
| c.-104_-83        | GGCCGGCGGGAAAGGAGGGA | 22               | 1295186 1295187  | None                        | 50 | White | Male | Bladder Urothelial Carcinoma | Primary | GENIE   |
| c.-104_-83        | GGCCGGCGGGAAAGGAGGGA | 22               | 1295186 1295187  | NA                          | 68 | Black | Male | Hepatocellular Carcinoma | Primary | GENIE   |
| c.-110_-89        | GGAAAGAAGGAGGGGCT    | 22               | 1295192 1295193  | None                        | 48 | Asian | Female | Malignant Phyllodes Tumor of the Breast | Primary | GENIE   |
| c.-110_-89        | GGAAAGAAGGAGGGGCT    | 22               | 1295192 1295193  | None                        | 64 | White | Male | Glioblastoma Multiforme | Primary | GENIE   |
| c.-110_-89        | GGAAAGAAGGAGGGGCT    | 22               | 1295192 1295193  | None                        | 65 | NA   | Male | Glioblastoma Multiforme | Primary | GENIE   |
| c.-110_-894       | GGAAAGAAGGAGGGGCT    | 23               | 1295192 1295193  | None                        | 69 | White | Female | Papillary Thyroid Cancer | Metastasis | GENIE   |
| c.-110_-89        | GGAAAGAAGGAGGGGCT    | 23               | 1295192 1295193  | None                        | 72 | White | Female | Poorly Differentiated Thyroid Cancer | Metastasis | GENIE   |
| c.-110_-89        | GGAAAGAAGGAGGGGCT    | 23               | 1295192 1295193  | None                        | 72 | White | Female | Poorly Differentiated Thyroid Cancer | Metastasis | GENIE   |
| c.-116_-96        | GGAGCGGGAGGGCTGGGAGG | 21               | 1295199 1295200  | NA                          | 64 | White | Male | Hepatocellular Carcinoma | Primary | GENIE   |
| c.-120_-96        | GGAGCGGGAGGGCTGGGAGG | 25               | 1295199 1295200  | None                        | 80 | NA   | Female | Poorly Differentiated Thyroid Cancer | NA         | UCSF500  |
| c.-125_-106       | GGAGCGGGAGGGCTGGGAGG | 25               | 1295199 1295200  | G228A                       | 68 | White | Male | Bladder Urothelial Carcinoma | Primary | GENIE   |

1Duplicated region of the TERTp relative to TERT translation start site (ATG).
2Primary and recurrent tumors for the same patient, SF12747 and SF13241.
3Two metastatic tumors for the same patient.
4Human reference genome hg19, chromosome 5; De novo native ETS 195 motif is in bold while de novo ETS 200 motif is underlined, and de novo ETS motif is in italic. Bold and italic nucleotides in the duplication sequence indicate a mismatch when compared to the duplication template sequence. NA Not available.
5De novo native ETS 195 motif is in bold while de novo ETS 200 motif is underlined, and de novo ETS motif is in italic. Bold and italic nucleotides in the duplication sequence indicate a mismatch when compared to the duplication template sequence. NA Not available.
6De novo native ETS 195 motif is in bold while de novo ETS 200 motif is underlined, and de novo ETS motif is in italic. Bold and italic nucleotides in the duplication sequence indicate a mismatch when compared to the duplication template sequence. NA Not available.
7De novo native ETS 195 motif is in bold while de novo ETS 200 motif is underlined, and de novo ETS motif is in italic. Bold and italic nucleotides in the duplication sequence indicate a mismatch when compared to the duplication template sequence. NA Not available.
and retention of the 22 bp TERTp duplication (Supplementary Fig. 3d). These somatic genetic data show a common clonal origin of the spatially divergent tumor, with the TERTp duplication being among the earliest alterations and being retained during tumor recurrence.

We identified a second glioblastoma case from the UCSF500 cohort with TERTp duplication, gain of chromosome 10 and homozygous deletion of CDKN2A as well as an NFI frameshift mutation (Supplementary Fig. 4). We did not have sufficient material to perform further analyses on this tumor sample.

Similar strength of TERT activation by duplication and hotspot mutations

TERTp hotspot mutations recruit the GABP tetramer complex to activate transcription. Whether the TERTp duplications are passenger alterations or can drive TERT reactivation in tumors is unknown. Based on the increased promoter activity we observed for ETS containing TERTp duplications (Figs. 1, 2), we hypothesized that TERTp duplications activate TERT expression in vivo. We were unable to identify a human cell line with a TERTp duplication, therefore, to determine the association between TERTp duplication and TERT expression, we used the TERTp duplication (c.-100_-79) glioblastoma sample presented in Fig. 4. TERT expression is limited to stem cells and cancer cells, therefore, to selectively measure TERT expression in tumor cells from this TERTp duplicated glioblastoma, we performed TERT RNA in situ hybridization (RNAscope) on FFPE slides. Quantification of the RNAscope staining for these TERTp duplicated glioblastoma cells broadly shows low level nuclear expression of TERT mRNA in individual cells that is comparable to hotspot G228A mutation glioblastoma tissue (Fig. 5a, b, d). To prove specificity of the assay, we assayed an IDH-mutant astrocytoma with wildtype TERTp and detected no TERT mRNA expression (Fig. 5c, d). TERT mRNA positive cells were observed in a second glioblastoma case with a TERTp duplication (Supplementary Fig. 6).

Next, we measured TERT mRNA by RT-qPCR and like the RNAscope result, TERT was expressed in the tumor tissue but not in the adjacent “normal” FFPE punches and greater than or equal to FFPE punches obtained from a separate glioblastoma case with the hotspot mutation G228A (Fig. 5e). We also measured the mRNA expression of the GABP TF complex subunits, GABPB1 and GABPA, which shows detectable and robust expression in all available tumor regions (Fig. 5f, g). These data show that the TERTp duplication associates with transcriptional reactivation of TERT, and in this glioblastoma which also has uniform expression of the mutant TERTp transcriptional regulator, GABP. Together, these data suggest that the duplications are likely drivers of TERTp reactivation and TERT expression in vitro and in vivo.

Discussion

TERTp hotspot mutations are the most common non-coding mutations in human cancers and selectively recruit the GABP TF complex. In contrast, one case report of duplication in the TERTp predicted greater than one hundred candidate transcription factors solely based on sequence motifs in the duplication. Considering only ETS factors as candidates, the matter is also complicated by the largely shared binding site preference among the 29 family members. Through a pan-cancer data analysis and our tumor samples, we discovered key shared features of the duplications that allowed insight into their mechanism of promoter activation and likely pathogenic function, leading us to the GABP tetramer. The duplications have similar and even identical length and insertion site across at least 7 cancer types, including duodenal adenocarcinoma in which TERTp mutations are rare. The duplications mimic pathogenic hotspot mutations in their transcriptional activation of the TERTp, the rate limiting step in reactivating
telomerase activity and cellular immortalization, and in the mechanism by which they do so. Although the hotspot mutations primarily recruit GABP, further studies will be needed to assess exclusivity of GABP recruitment by the duplications. On the other hand, knockdown of GABP alone is sufficient to rapidly reduce TERT expression from the promoter with either hotspot mutation or from each of the many different promoters with an ETS-containing duplication. The location and size of GGAA containing duplications in the TERT p suggested to us that they create an additional binding site to enable GABP tetramer binding, representing another genetic mechanism by which the GABP TF complex is recruited to the TERT p. In fact, we showed that the GGAA motif and GABP TF complex are both essential for increased TERT p duplication activity. The length of the TERT p duplication puts the two ETS motifs in phase and at a distance that is most ideal for tetramer binding (Fig. 6). We observed that as the distance between the ETS motifs increases, the activation of the TERT p duplication is decreased. The requirement for two ETS motifs in helical phase further supports GABP as a prime regulator of the promoter duplication because GABP, specifically the tetramer, is the only member of the ETS family that requires two binding motifs.

TERT p hotspot mutations are considered one of the early events in glioblastoma formation. Likewise, we found TERT p duplication in all five spatially independent regions and in a total of 13 punches from the five FFPE blocks of the newly diagnosed tumor and also at recurrence. We infer the TERT p duplication is fully clonal in this case and is retained during tumor progression. Therefore, like the hotspot

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**Fig. 3 | GABPBI1 containing GABP complexes bind the TERT promoter duplication.** a, b Schematic of the GABP heterotetramer TF complex binding to the de novo and native ETS motifs of the TERTp with the G228A mutation or c−100−79 duplication, respectively. The figures were created with BioRender. Representative gels (c, e, g) and quantification (d, f, h) of EMSAs comparing binding affinity of GABPA-B1L heterodimers to the TERTp c−100−79 duplication, TERTp wildtype (native ETS motif only) and G228A mutant (native and de novo ETS motifs) DNA fragments. Abbreviations: TERTp, TERT promoter. Source data are provided as a Source Data file. Values are mean and SD. N = 3 technical replicates.
mutations, TERTp duplication may also be an early event in tumorigenesis, adding to their remarkable similarities despite being fundamentally different somatic genetic alterations.

Interestingly, we identified two cases (one squamous cell and one bladder urothelial carcinoma) with TERTp duplication and a TERTp hotspot mutation. As there is limited access to these sequencing data, it remains unclear whether the duplication and hotspot mutation were on different alleles within the same cell or in different cells within the tumor. We identified one tumor with both TERTp hotspot mutation and duplication of the mutant sequence, suggesting that the duplication may have provided additional selective advantage in this highly proliferative tumor type, though further experimentation is needed. For the second tumor, the TERTp duplication harbored the wildtype sequence. As reads were low for both the TERTp duplication and TERTp hotspot mutation in this case these could potentially be two concurrent subclonal events that formed independently from each other.

**Fig. 4 | Clonal evolution of a multifocal glioblastoma with the TERTp duplication.**

- **a** Clinical timeline. Multiple tumor tissue samples were obtained from the IDH-wildtype and TERTp duplicated glioblastoma at diagnosis (SF12747) and one sample was obtained at recurrence (SF13241).
- **b** MRI of enhancing masses in the left parieto-occipital lobes with surrounding edema and mass effect.
- **c** H&E staining of an FFPE section from the (d) occipital lobe tumor focus and the (e) parietal lobe tumor focus. Scale bar denotes 50 µm. f TERTp PCR spanning the native ETS motif and duplication for the DNA isolated from FFPE punches from multiple regions of the newly diagnosed tumor, and a no-template control (NTC). Lane 1 includes the DNA ladder for reference of base pair (bp) size. C# and D# indicate the block, # indicate the tumor and N the tumor-adjacent normal brain punch, respectively. The experiment was performed once because of the limitation of patient material.
- **g** SNV-based phylogenetic tree from the exome sequencing of DNA from five regions of the tumor and patient-matched normal peripheral blood. Terminal nodes (bold text) represent samples, where N indicates the Normal germline (peripheral blood); all other terminal nodes are tumor samples. Edges (lines) are proportional to the genetic distance between samples within each patient, with the number of mutations corresponding to each major edge indicated. h PyClone analysis of the mutated gene clusters from the exome sequencing of DNA from five regions of the tumor relative to patient-matched normal peripheral blood. N = 1 biologically independent samples for each region. Error bars indicate the mean s.d. (using 10,000 post-'burn-in' samples) of Markov chain Monte Carlo–derived cellular prevalence estimates over all mutations in a cluster as defined by PyClone algorithm. TERTp TERT promoter, HD Homozygous deletion, amp Amplification, Dup Duplication. Source data are provided as a Source Data file.
examined over 3 independent experiments for TERT promoter wt. (a–c) TERT, GABPB1 and GABPA RT-qPCR results normalized to GUSB and performed on RNA isolated from the multifocal TERTp duplicated tumor (T) (SF12747), adjacent normal (N) (SF12747) and TERTp G228A (SF12180) FFPE punches. Values are mean and SD. Multiple comparisons were performed using 1-way ANOVA, and post hoc analyses were based on Tukey’s test. **p < 0.005, ***p < 0.0001. C/D# indicates the FFPE tissue block and T/N# indicates the punch. Abbreviations: TERTp, TERT promoter. Source data are provided as a Source Data file. N = 3 technical replicates.

in different cells. Similarly, rare cases of co-occurring G228A and G250A have been reported. Nevertheless, most tumors have only one hotspot mutation or the core promoter duplication.

Our results are also important for clinical diagnosis and tumor classification. For diagnostic purposes, discovering new TERTp mutations and distinguishing pathogenic mutations from variants of unknown significance is important since the 2021 WHO Classification of Central Nervous System Tumors specifies histologically low grade diffuse astrocytic gliomas without IDH-mutation but with TERTp mutations shall be diagnosed as CNS WHO grade 4 glioblastomas. There are many more examples in other tumor types where TERT promoter mutations can change grading or diagnosis. These examples include TERT promoter status in meningiomas, ovarian clear cell carcinoma, small cell carcinoma of the bladder, high-grade adenocarcinoma of the prostate, conjunctival melanoma and thyroid carcinoma. To date, only TERTp hotspot mutations are considered pathogenic. Our data indicates that TERTp duplications should also be considered pathogenic for clinical evaluation and tumor classification purposes. This could be implemented immediately, as clinical panel sequencing in use at many academic medical centers and commercial laboratories can routinely detect the duplications. One potential limitation would be for medical centers that use only Sanger sequencing for TERTp mutation detection, although even these simple assays could be modified slightly to detect the duplication, as we show in Fig. 4f. For panel sequencing, several of the algorithms used do not detect insertions or deletions greater than 5 bp, and thus the 22 bp...
TERTp duplications may be missed. This potential issue could be further exacerbated by the consistent observation that promoter regions with high GC-content such as the TERTp yield lower sequencing depth. Our findings in Supplementary Table 1 might not be representative, as TERTp mutations were only detected in one third of all glioblastomas in the GENIE dataset whereas multiple independent studies (including our own experience as part of the UCSF Glioblastoma Precision Medicine Program) suggest the frequency is over 80%. It is therefore possible that TERTp duplications are under-reported in cancer.

Our results show that TERTp duplications are recurrent, pathogenic variants across multiple cancer types, and begin to define their frequency, size, insertion site bias and functional significance. GABP is the common denominator between TERTp duplications and hotspot mutations, supporting a key role in cellular immortality. The remarkable parallel between genetically distinct hotspot mutations and core promoter duplications together suggest a selective pressure specifically for GABP tetramer recruitment to reawaken one allele of the epigenetically silenced TERTp. Understanding the characteristics of the GABP tetramer and its binding partners may shed light on this apparently unique capability.

**Methods**

This study complies with all relevant ethical regulations and was approved by the Committee on Human Research of the University of California, San Francisco. All patients provided informed written consent prior to sample acquisition. Patients were not monetarily compensated.

**AACR project GENIE dataset**

The tenth dataset of AACR Project Genomics Evidence Neoplasic Information Exchange (GENIE), Cohort v10.0-public, was accessed from cBioPortal and queried for TERT promoter mutations. This dataset was filtered for TERTp mutations and insertions containing GCAA. De-identified clinical data such as cancer diagnosis, sex, age, race and the anatomic site of the tumor sequenced were also obtained for the associated cases.

**UCSF500 pan-cancer dataset**

The UCSF500 NGS Panel is a panel sequencing assay of 479 genes that are involved in human cancer. The UCSF500 NGS Panel cancer dataset was accessed from a private cBioPortal instance that is available to UCSF clinicians, researchers, or other staff. The UCSF500 cBioPortal instance currently has de-identified UCSF500 NGS Panel data as well as select de-identified clinical data elements such as diagnosis, sex, and the anatomic site of the tumor sequenced with consent to publish this information. Race was self-defined. This dataset was filtered for TERTp mutations and insertions containing GCAA.

**UCSF500 glioblastoma patient cohort and tumor samples**

Two patients with glioblastomas harboring duplications within the upstream regulatory region of the TERT gene as identified by capture-based next-generation DNA sequencing (UCSF500) performed on a prospective clinical basis as part of the UCSF Glioblastoma Precision Medicine Program were included in this study. All tumor specimens were fixed in 10% neutral-buffered formalin and embedded in paraffin. Tumor tissue was selectively scraped from unstained slides or punched from formalin-fixed, paraffin-embedded blocks using 2.0 mm disposable biopsy punches (Integra Miltex Instruments, #33-31-P/25) to enrich for maximal tumor content. Genomic DNA was extracted from this macro-dissected formalin-fixed, paraffin-embedded tumor tissue using the QIAamp DNA FFPE Tissue Kit (Qiagen, #56404).

**Targeted next-generation DNA sequencing**

Capture-based next-generation DNA sequencing was performed using a custom cancer gene panel, UCSF500, and an Illumina HiSeq 2500 instrument. This assay targets all coding exons of 479 cancer-related genes, selected introns and upstream regulatory regions of 47 genes. DNA regions captured were also chosen to enable detection of structural variants including gene fusions, and DNA segments at regular intervals along each chromosome to enable genome-wide copy
number and zygosity analysis. The total sequencing footprint is 2.8 Mb. 250 ng of DNA was used for multiplex library preparation using the KAPA Hyper Prep Kit (Roche, #07962347001) according to the manufacturer’s specifications. Hybrid capture of pooled libraries was performed using a custom oligonucleotide library (Nimblegen SeqCap EZ Choice). Captured libraries were sequenced as paired-end 100 bp reads. GRCh37 (hg19) was used as the reference genome, and sequenced reads were aligned using the Burrows-Wheeler aligner (BWA). Recalibration and deduplication were performed using the Genome Analysis Toolkit (GATK). Coverage and sequencing statistics were performed using Picard CalculateHsMetrics and Picard CollectInsertSizeMetrics. FreeBayes, Unified Genotyper, and Pindel were used to call single nucleotide variant and small insertion/deletion mutations. Delly was used for large insertion/deletion and structural alteration calling. Variant annotation was performed with Annovar. 

**Exome sequencing**

Exome capture was performed using xGen Exome Research Panel v2. Tumor cell purity and chromosomal copy number was estimated using FACETS from the exome sequencing data. Exome libraries were sequenced on an Illumina NovaSeq 6000 instrument. The reference human genome build used for all sequencing data alignment in this study was GRCh37 (hg19).

**Phylogenetic tree construction and PyClone analysis**

SNVs were called from the exome sequencing of tumor and matching normal genomic DNA by MuTect to generate sample-oriented SNV-based phylogenetic trees. To generate a phylogenetic tree, an Ordinary least squares minimum evolution from ape R package was implemented using a distance matrix for all samples from the patient. The Manhattan distance was computed for the binary call matrix, and fasteme.ols from R package ape was used to construct a rooted binary tree with 0.01. PyClone (version PyClone-0.13.1) was also used to perform the clonal frequency analysis by grouping SNVs into clonal clusters.

**RNAseq**

FFPE sections were evaluated by RNAscope chromogenic in situ hybridization (CISH) assay for the expression of TERT in individual cells using Advanced Cell Diagnostics (ACD) probes (Newark, CA) specific for the TERT promoter PCR and genotyping. RNAscope probe/cell from each image was counted, paraffin sections were evaluated by RNAscope chromogenic in situ hybridization (CISH) assay for the expression of TERT in individual cells using Advanced Cell Diagnostics (ACD) probes (Newark, CA) specific for the TERT promoter PCR and genotyping.

**TERT promoter PCR and genotyping**

Genomic DNA was purified from UMC3 and LN229 cell lines using the Quick DNA Plus Kit (Zymo Research, #D4069). 100 ng of genomic DNA was used for PCR with Phusion Polymerase (Thermo Fisher Scientific, #F5305) containing GC buffer and 5% DMSO and standard desalted primers. PCR primer sequences were designed with tails incorporating M3 primer sequences: 5′_TERTp_PCR_F 5′-GTAACACGACGGCCA GACGGAGGAGGACTG-3′ and 5′_TERTp_PCR_R 5′-CAGGAA CACGCTATGACGGCCAGTCCACGTCG-3′. Cycling conditions were: 98 °C for 1 min, 69 °C for 15 s and 72 °C for 30 s for a total of 32 cycles with a final extension at 72 °C for 5 min. A fraction of the PCR product was resolved on a 2% TBE agarose gel containing 1X SYBR Safe DNA Gel Stain (Thermo Fisher Scientific). Internal control and positive control DNA were obtained from the DSMZ, German Collection of Microorganisms and Cell Cultures, Braunschweig. PyClone (version PyClone-0.13.1) was also used to perform the clonal frequency analysis by grouping SNVs into clonal clusters.

**Mammalian cell culture**

All cell lines were maintained at 37 °C in a humidified incubator with 5% CO₂. Cell culture medium was changed every 3–5 days depending on cell density. For routine passage, cells were split at a ratio of 1:3-10 when they reached 80% to 90% confluence. LN229 human glioblastoma cells (ATCC, #CRL-2611) were cultured in Dulbecco’s Modified Eagle Medium/Nutrient Mixture F-12 (DMEM/F-12; Corning, #10-090-CV) supplemented with 10% fetal bovine serum (FBS; Gibco) and 100 U/mL Penicillin-Streptomycin (Gibco). UMC3 human bladder cancer cells (ATCC, #CRL-T2759) were cultured in Minimum Essential Medium (MEM; Corning, #10-009-CV) supplemented with 10% FBS (Gibco) and 100 U/mL Penicillin-Streptomycin (Gibco). Cell lines were authenticated by short tandem repeat (STR) analysis at the University of California Berkeley Sequencing Facility and confirmed to be Mycoplasma free by PCR followed by TBE agarose gel electrophoresis using a previously published method. For mycoplasma testing, cell lines were grown for a period of 72 h in the absence of Penicillin-Streptomycin then 1 mL of culture supernatant was collected and centrifuged at 13,000 x g for 5 min at room temperature. Supernatant was aspirated then the resulting pellet was lysed with 50 μL of QuickExtract (Lucigen) and incubated on a thermal cycler at 65 °C for 15 min, 68 °C for 15 min and 98 °C for 10 min. A total of 2 μL QuickExtract lysate was used as template in 20 μL PCR reaction containing Phusion Polymerase (Thermo Fisher Scientific) and standard desalted primers. PCR primer sequences: 5′_Myco_F1CGCCTGAGTAGTACGT WGC; 5′_Myco_F 2′_TGCTGRTGATGACATTGC; 5′_Myco_F3CRCTGATGATGACTG CTG; 5′_Myco_F4CGCCTGGTATGATCCTGC; 5′_Myco_R1CGGGTG TGTACARACCCGA; and 5′_Myco_R2GGGGTGTGTCACAAAAACCGA. Cycling conditions were: 1 cycle at 98 °C for 30 s, 34 cycles at 98 °C for 10 s, 65 °C for 10 s and 72 °C for 20 s, and a final extension at 72 °C for 5 min. PCR reactions were resolved on a 1.3% TBE agarose gel containing 0.5 mg/mL SYBR Safe DNA Gel Stain (Thermo Fisher Scientific). Internal control and positive control DNA were obtained from the DSMZ, German Collection of Microorganisms and Cell Cultures, Braunschweig. Internal control and positive control DNA were obtained from the DSMZ, German Collection of Microorganisms and Cell Cultures, Braunschweig.
Molecular cloning of luciferase reporter plasmids

**TERT** promoter reporter variants were generated by PCR amplification from single stranded DNA Ultramer oligos (IDT) using Phusion Polymerase (Thermo Fisher Scientific, #F5305) and the following standard desalted primers (IDT): Xhol_dup_F 5′-CTAGCTCGAGTCTGCCCC CTTACCTTCCAG-3′ and HindIII_dup_R 5′-TTGCAAGCTTGGCG CGGAGGGAGATCAGGCTGCTGAAA-3′. PCR products were gel purified using Zymoclean Gel DNA Recovery Kit (Zymo Research, #D4001), digested with Xhol and EcoRI (Thermo Fisher Scientific, #FD0695, #FD0275) then column purified using DNA Clean & Concentrator-5 (Zymo Research, #D4040). Digested fragments were then ligated into Xhol and HindIII (Thermo Fisher Scientific, #FD0695, #FD0505) linearized pGL4.9-TERT WT (Addgene, #84924) plasmid. All PGL4 TERTp duplication variants were verified by Sanger sequencing (GENEWIZ).

Luciferase promoter reporter assay

Transfection of TERTp reporter plasmids was carried out with ViaFect transfection reagent (Promega, #E4981). Briefly, cells were seeded at a density of 3000 or 6000 cells per well in a 96-well clear bottom white polystyrene microplate (Corning) in antibiotic free media. 24 h post-seeding, cells were transfected with 90 ng PGL4.10 plasmid (experimental promoter with Firefly luciferase, Promega), 9 ng PGL4.74 plasmid (control promoter with Renilla luciferase, Promega), and 0.3 µL of ViaFect transfection reagent (Promega, #E4981) in 10 µL of Opti-MEM serum free media (Gibco). Non-targeting (#D-001206-13-20) and GABPA (#M-011662-01) directed siGENOME SMART siRNA (Dharmacon) were used for knockdown experiments. For knockdown experiments, cells were seeded at a density of 2,000 or 5,000 cells per well in a 96-well plate in antibiotic free media. 24 h post-seeding, cells were transfected with 50 nM of siRNA and 0.3 µL of DharmaFECT reagent (Dharmacon, #T-2001-02) in 10 µL of Opti-MEM serum free media (Gibco). At 48 h post-transfection, cells were transfected with reporter plasmids as described above. 24 h post transfection of reporter plasmids, Firefly luciferase activity was measured by using the Dual-Luciferase Reporter assay system (Promega, #E1960) on the Promega GloMax 96 well microplate luminometer and normalized against Renilla luciferase activity then presented as a ratio to Firefly to Renilla using arbitrary units.

Protein purification

Genes for GABP β1L and GABP α were cloned to pET-Duet-1 vector using Gibson assembly. GABP β1L was inserted after the first ribosome binding site and GABP α after the second ribosome binding site. GABP β1L has a N-terminal His-Tag separated from the subunit by TEV cleavage sequence.

Subunits were expressed in Rosetta DE3 cell line. A single colony was used to inoculate an overnight starter culture in Luria-Bertani media (LB) with 100 µg/ml Ampicillin. Next, starter culture was used to inoculate large Terrific Broth (TB) media cultures with 100 µg/ml Ampicillin. These cultures were grown at 37 °C until the OD600 reached 0.8-1 at which point they were chilled on ice and induced with 0.5 mM IPTG. Protein expression was conducted at 16 °C overnight. Cells were harvested via centrifugation, resuspended in 50 mM HEPES pH 7, 500 mM NaCl, 1 mM TCEP, 5% (v/v) glycerol and flash frozen. For protein purification pellets were thawed and mixed with 0.5 mM PMSF and 20 µL of 10X purification buffer in 5 mL G25 (GE) desalting column. For annealing, puriﬁed ETS sites were cloned to pET-Duet-1 vector (Promega) and duplication mutant creating a new de novo site (5′-AGGGCGGGGCCGGAGGGGCGGGGGCCG-3′) were obtained from IDT as PAGE puriﬁed. Then one of the strands of each sequence was labeled with 32P and annealed to its complementary unlabeled strand in 20 mM Tris pH 7.5, 100 mM LiCl, 1 mM MgCl2 buffer and used for the EMSA assay.

To introduce 3p label, Duplexes of single stranded DNA were incubated with 5 Units of T4 PNK (NEB) and 10Ci of 32P-γ-ATP (Perkin-Elmer) in PNK buffer in total volume of 25 µL for 30 min at 37 °C and heat inactivated at 65 °C for 20 min. To remove excess of ATP, reaction was diluted by addition of 25 µL of DEPC-treated water and applied to a G25 (GE) desalting column. For annealing, puriﬁed oligos were further diluted by addition of 50 µL of DEPC-treated water.

To perform EMSA assays, GABP proteins were thawed and diluted to starting stock concentration of 2 µM and then further serial dilutions were performed in Superdex 200 buffer to create a series of sixteen total protein stocks. Annealed DNA oligonucleotides were diluted to 1 nM stock concentration in annealing buffer. Reactions were performed by mixing DEPC-treated water, 5x reaction buffer and DNA to create a master stock, which was then equally distributed between reaction tubes and supplemented with protein from sixteen stocks. The final reaction component concentrations were as follows: 0.2 nM 60 bp DNA, GABPA/B heterodimer (0 to -1000 nM) with ﬁnal concentration of reaction buffer 20 mM Tris pH 7.5, 50 mM KCl, 5.2 mM MgCl2, 1 mM TCEP, 5% (v/v) glycerol in 15 µL ﬁnal reaction volume. Reactions were incubated at room temperature for 1 h and then 3 µL were applied to a 5% (v/v) native 0.5X TBE PAGE gel with 5 mM MgCl2, which was prerun at 8 W for -1 h at 4 °C. The gel was then run in 0.5X TBE buffer with 5 mM MgCl2 for -2.5 h at 8 W at 4 °C. The gel was dried and phosphorous screen was exposed to the gel overnight. Imaging was performed using a Typhoon phosphorimager and quantiﬁcation with ImageQuant software. The fraction of promoter sequence bound was calculated by dividing the signal of bound DNA fraction by the sum of bound and unbound signals in each lane.

Reverse transcription quantitative PCR (RT-qPCR)

Glioblastoma tumor tissue (SF12747) was selectively punched from single stranded 60 bp oligonucleotides with wild type TERTp including native ETS sites (5′-AGGGCGGGGCCGGAGGGGCGGGGGCCG-3′) and duplication mutant creating a new de novo site (5′-AGGGCGGGGCCGGAGGGGCGGGGGCCG-3′) were obtained from IDT as PAGE puriﬁed. Then one of the strands of each sequence was labeled with 32P and annealed to its complementary unlabelized strand in 20 mM Tris pH 7.5, 100 mM LiCl, 1 mM MgCl2 buffer and used for the EMSA assay.

To introduce 3P label, Duplexes of single stranded DNA were incubated with 5 Units of T4 PNK (NEB) and 10 Ci of 32P-γ-ATP (Perkin-Elmer) in PNK buffer in total volume of 25 µL for 30 min at 37 °C and heat inactivated at 65 °C for 20 min. To remove excess of ATP, reaction was diluted by addition of 25 µL of DEPC-treated water and applied to a G25 (GE) desalting column. For annealing, puriﬁed oligos were further diluted by addition of 50 µL of DEPC-treated water.

To perform EMSA assays, GABP proteins were thawed and diluted to starting stock concentration of 2 µM and then further serial dilutions were performed in Superdex 200 buffer to create a series of sixteen total protein stocks. Annealed DNA oligonucleotides were diluted to 1 nM stock concentration in annealing buffer. Reactions were performed by mixing DEPC-treated water, 5x reaction buffer and DNA to create a master stock, which was then equally distributed between reaction tubes and supplemented with protein from sixteen stocks. The final reaction component concentrations were as follows: 0.2 nM 60 bp DNA, GABPA/B heterodimer (0 to -1000 nM) with ﬁnal concentration of reaction buffer 20 mM Tris pH 7.5, 50 mM KCl, 5.2 mM MgCl2, 1 mM TCEP, 5% (v/v) glycerol in 15 µL ﬁnal reaction volume. Reactions were incubated at room temperature for 1 h and then 3 µL were applied to a 5% (v/v) native 0.5X TBE PAGE gel with 5 mM MgCl2, which was prerun at 8 W for -1 h at 4 °C. The gel was then run in 0.5X TBE buffer with 5 mM MgCl2 for -2.5 h at 8 W at 4 °C. The gel was dried and phosphorous screen was exposed to the gel overnight. Imaging was performed using a Typhoon phosphorimager and quantiﬁcation with ImageQuant software. The fraction of promoter sequence bound was calculated by dividing the signal of bound DNA fraction by the sum of bound and unbound signals in each lane.
All samples were run in triplicate using the QuantStudio 5 (Thermo Fisher Scientific) and all gene expression data were normalized to GUSB mRNA. PCR was performed with an annealing temperature of 60 °C and a total of 45 cycles was run for all primer pairs. Dissociation curves were performed to confirm specific product amplification. RT-qPCR standards for each gene were generated from a mixture of human cell line cDNA via end-point RT-PCR then gel purification, using the appropriate primer pair. Gradient PCR reactions were performed with the C1000 Touch Thermal cycler (Bio-Rad) to determine the annealing temperature and specificity for each primer set. Primer sequences corresponding to each gene for the mRNA expression analysis were designed using NCBI Primer Blast or selected from those previously reported in the literature: TERT_RT_F 5′-TCACGGAGAC CACGTTTCAA-3′; TERT_RT_R 5′-TTCAAGTGCTGTCTATCATCAAT-3′; GABPB1_RT_F 5′-AAACGGGTTATCTGCTGTTC-3′; GABPB1_RT_R 5′- GCCACACTCGGAGTTCCTGAA-3′; GABA_RT_F 5′-AACAGGGCTTTTGATGG-3′; GABA_RT_R 5′-GTGAGGTCTATATCGGTCATGCT-3′; GUSB_RT_F 5′-CTCATTTGGAATTTTGCCGATT-3′; and GUSB_RT_R 5′- CCGAGTAGAAGATCCCTTTTTA-3′.

Statistics & reproducibility
No statistical method was used to predetermine sample size. Sample sizes were chosen based on availability of suitable material. All experiments were done in at least technical duplicates and compared using two different cell lines if suitable. All attempts at replication were successful. No data were excluded from the analyses. The experiments were not randomized. The investigators were not blinded to allocation during experiments and outcome assessment. Representative images of H&E stained and RNA scope sections from diagnostic tumor specimens are shown. Each tumor specimen was carefully examined by an expert neuropathologist and showed similar findings throughout the resected tumor tissue from which representative images were selected.

Reporting summary
Further information on research design is available in the Nature Research Reporting Summary linked to this article.

Data availability
The raw exome sequencing data generated in this study have been deposited in the European Genome-Phenome Archive database under accession code EGAD00001008768. The raw panel sequencing data generated in this study have been deposited in the European Genome-Phenome Archive database under accession code EGAD00001009286. The exome and panel sequencing data are available under restricted access for IRB requirements, access can be obtained by contacting J.F.C. (joseph.costello@ucsf.edu). The GENIE dataset can be accessed via https://genie.cbioportal.org/login.jsp after registration on the platform. The data from Panebianco et al. (ref. 22) are available from the corresponding author upon reasonable request (we used the processed and published data). Pierini et al. (ref. 23) used TERT promoter specific PCR and Sanger sequencing and thus the data (sequencing traces) and results are available in their publication. Source data are provided with this paper (including uncropped gels and blots). Source data are provided with this paper.

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Author contributions
J.F.C. supervised this work. C.J.B., A.K.S. and J.F.C. wrote the manuscript with support from all authors. C.J.B., K.M.S., A.S.W., M.Y.K. and C.H. prepared and processed the analyses. S.M.C., J.J.P. and D.A.S. provided samples and clinical data. C.J.B., A.K.S., K.M.S. J.A.D., J.J.P., D.A.S. and J.F.C. performed data analysis and interpretation.

Competing interests
J.A.D. is a co-founder of Caribou Biosciences, Editas Medicine, Intellia Therapeutics, Scribe Therapeutics, and Mammoth Biosciences. J.A.D. is a scientific advisory board member of Caribou Biosciences, Intellia Therapeutics, effector Therapeutics, Scribe Therapeutics, Synthego, Metagenomi, Mammoth Biosciences, and Inari. J.A.D is a Director at Johnson & Johnson and has sponsored research projects by Pfizer, Roche Biopharma, and Biogen. J.F.C. is a co-founder of Telo Therapeutics, Inc. and has ownership interests. The other authors declare no competing interests.

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