Topographic correlates of driver mutations and endogenous gene expression in pediatric diffuse midline gliomas and hemispheric high-grade gliomas

Eve Kazarian1, Asher Marks2, Jin Cui1, Armine Darbinyan3, Elizabeth Tong4, Sabine Mueller5,6,7, Soonmee Cha4 & Mariam S. Aboian1,8

We evaluate the topographic distribution of diffuse midline gliomas and hemispheric high-grade gliomas in children with respect to their normal gene expression patterns and pathologic driver mutation patterns. We identified 19 pediatric patients with diffuse midline or high-grade glioma with preoperative MRI from tumor board review. 7 of these had 500 gene panel mutation testing, 11 patients had 50 gene panel mutation testing and one 343 gene panel testing from a separate institution were included as validation set. Tumor imaging features and gene expression patterns were analyzed using Allen Brain Atlas. Twelve patients had diffuse midline gliomas and seven had hemispheric high-grade gliomas. Three diffuse midline gliomas had the K27M mutation in the tail of histone H3 protein. All patients undergoing 500 gene panel testing had additional mutations, the most common being in ACVR1, PPM1D, and p53. Hemispheric high-grade gliomas had either TP53 or IDH1 mutation and diffuse midline gliomas had H3 K27M-mutation. Gene expression analysis in normal brains demonstrated that genes mutated in diffuse midline gliomas had higher expression along midline structures as compared to the cerebral hemispheres. Our study suggests that topographic location of pediatric diffuse midline gliomas and hemispheric high-grade gliomas correlates with driver mutations of tumor to the endogenous gene expression in that location. This correlation suggests that cellular state that is required for increased gene expression predisposes that location to mutations and defines the driver mutations within tumors that arise from that region.

Pediatric high-grade gliomas are aggressive tumors with different molecular characteristics than adult gliomas1–3. There is a distinct difference between the driver mutations found in pediatric diffuse midline gliomas and hemispheric high-grade gliomas. Diffuse midline gliomas represent 10–15% of all childhood brain tumors and have median survival of 9–12 months4. MR imaging plays a critical role in diagnosis and surgical biopsy may not always be performed or feasible for diffuse midline gliomas in eloquent location. Radiation therapy is the standard of care due to their location within critical structures along the midline of the brain, yet chemotherapy and convection enhanced delivery approaches have been shown to play a role in treatment5,6. Many of these tumors contain somatic mutations in genes encoding histone tails; in 2016, the World Health Organization introduced the new classification of the diffuse midline gliomas, H3 K27M-mutant7. Diffuse midline gliomas, H3 K27M-mutant, frequently appear along with mutations in signal transduction (ACVR1), chromatin remodeling (ATRX), and cell cycle regulation (TP53, PPM1D)8. Recent genomic analyses have led to the development of subtype-specific therapies8,9. On the other hand, hemispheric high-grade gliomas in children commonly have

1Department of Radiology & Biomedical Imaging, Yale School of Medicine, New Haven, CT, USA. 2Department of Pediatric Hematology & Oncology, Yale School of Medicine, New Haven, CT, USA. 3Department of Neuropathology, Yale School of Medicine, New Haven, CT, USA. 4Department of Radiology, University of California, San Francisco, San Francisco, USA. 5Division of Pediatric Hematology & Oncology, Department of Pediatrics, University of California, San Francisco, San Francisco, USA. 6Department of Neurological Surgery, University of California, San Francisco, San Francisco, USA. 7Department of Neurology, University of California, San Francisco, San Francisco, USA. 8email: mariam.aboian@yale.edu
hypermutated molecular signatures including TP53, NF1, SETD2, ATRX, PTEN, ARID1A, and POLE\(^{14,9}\). The survival rates for these tumors are poor, with 5-year survival rates of less than 20%. Maximum safe resection and radiation therapy in combination contribute to improved survival in patients with hemispheric high-grade gliomas as compared to those with diffuse midline gliomas\(^{10,11}\).

Pediatric high-grade gliomas can be distinguished by their location of origin: hemispheric high-grade gliomas originate in the cerebral hemispheres and diffuse midline gliomas originate in the midline structures, commonly in the pons, thalamus, or spinal cord\(^{8,10,12-15}\). In this study, we characterized the genetic mutations in 19 pediatric high-grade gliomas and correlated the location of tumors to endogenous gene expression within the region of tumor origin. Knowing the exact locations of the tumors and their genetic mutations allowed us to form a hypothesis that normal gene expression signature within particular part of the brain can drive the mutation of tumors that arise from that anatomic location. A driver mutation is an alteration that gives a cancer cell a fundamental growth advantage for its neoplastic transformation. It differs from passenger mutations in that these do not necessarily determine the development of the cancer\(^{16}\). Previous studies have shown that chromatin structure affects distribution and types of mutations in cancer\(^{16}\). To test this hypothesis, we analyzed normal gene expression profile within different parts of the brain using microarray data from publicly available Allen Brain Atlas and correlated expression patterns of genes that were characterized to be driver mutations in high-grade gliomas that originated from these locations. The purpose of our study was to characterize the endogenous gene expression profile in anatomic locations where pediatric high-grade gliomas occur and correlate the gene expression level to susceptibility for developing a mutation, i.e., presence of tumors within this location with driver mutations within this gene. The Allen Brain Atlas provides data for normal intracranial gene expression in human adult and developmental brain (www.brain-map.org). This correlation would suggest that presence of chromatin microenvironment that is required for increased gene expression predisposes that location to mutations and defines the drive mutations within a specific region of the brain to tumor formation. Confirmation of gene expression patterns within specific regions of the brain associated with the presence of high-grade gliomas will lead to a better understanding of key genetic drivers involved in brain tumorigenesis.

**Methods**

**Patient characteristics.** A total of nineteen patients were either originally treated or seen for a second opinion regarding treatment options at two tertiary hospitals\(^6\). Informed consent was obtained prior to genetic sequencing for use of patient samples. Informed consent was obtained for all the subjects who are under 18 from a parent and/or legal guardian. This study was reviewed and approved by the Yale University and University of San Francisco institutional review boards. All methods were carried out in accordance with relevant guidelines and regulations.

**Characterization of tumor molecular profile.** Genomic DNA was extracted from peripheral blood and tumor tissue micro-dissected from formalin-fixed, paraffin-embedded blocks. Capture-based next-generation sequencing (NGS) was performed at the Clinical Cancer Genomics Laboratory within our institution, using an assay targeting the coding regions of 510 cancer-related genes, TERT promoter, select introns from 40 genes (for detection of gene fusions and other structural variants), and intergenic regions at regular intervals along each chromosome (for chromosomal copy number assessment), altogether with a total sequencing footprint of 2.8 Mb\(^6\).

Sequencing libraries were prepared from genomic DNA with target enrichment performed by hybrid capture using a custom oligonucleotide library. Sequencing of the 500 gene panel assay was performed on an Illumina HiSeq 2500. Duplicate sequencing reads were removed computationally to allow for accurate allele frequency determination and copy number estimates. The analysis was based on the human reference sequence build hg19 (NCBI build 37), using the following software packages: BWA, Samtools, Picard tools, GATK, CNVkit, Pindel, SATK, Annovar, Freebayes, Delly, and Nexus Copy Number. Single nucleotide variants and small insertions/deletions (indels) were visualized and verified using Integrated Genome Viewer\(^8\). For samples with at least 25% tumor, where > 200 × coverage for the tumor sample and > 100 × coverage for the normal sample, the fully clonal single nucleotide variant sensitivity is 99% and small indel sensitivity is 83% while the specificity is 98% and 71%, respectively. Large insertions, deletions, and gene rearrangements may be detected but have only been individually validated for select examples. 50 gene panel testing was performed at Tumor Profiling Laboratory of Yale-New Haven Hospital under regulations of Clinical Laboratory Improvement Amendments of 1988 (CLIA). 343 gene panel testing was performed by Foundation One (Foundation Medicine Inc).

Molecular pathologists with specialization in neuropathology and brain tumor genetics organized results into formal reports, which detailed somatic and germline alterations as well as association with any known tumor predisposition syndromes, diagnostic or prognostic implications, and potential targeted therapies. Results were discussed at weekly multidisciplinary molecular tumor boards that included surgical and molecular pathologists together with oncologists, surgeons, and radiation oncologists from a wide variety of specialties\(^8\).

**Gene expression profile analysis.** Gene expression profiles for PPM1D, ASXL1, BCO1L1, ACVR1, PIK3CA, TP53, SETD2, BRCA2, PTEN, POLE, BLM, ARID1A, ATRX, CDKN2A, MSH6, PDGFA were obtained from the publicly available microarray data of gene expression in the human brain from the Allen Brain Sciences Institute (www.brain-map.org). The gene expression z scores were determined based on microarray analysis of 6 adult brains that were micro-dissected into anatomically distinct brain regions (500), representing 93% of known genes that are represented by at least 2 probes. For each gene, gene expression profiles were downloaded for each available probe.
For example, one of the patients had histone H3 K27M mutation within a highly conserved arginine location within the DNA binding domain to a hydrophobic leucine. Mutations in these genes suggest that disruption of chromatin structure and gene expression mediated by histone H3 tails, combined with disruptions in the PI3K pathway and p38/MAPK pathway are critical for tumorigenesis. This 3-year-old patient presented with a solid tumor that originated within the pons with extension into the middle cerebellar peduncle. The tumor was infiltrative in appearance, had evidence of central necrosis, mild enhancement, and no reduced diffusion. On six months follow up, the tumor developed significant progression with local extension into the cerebral peduncles, temporal stem, thalamus, corpus callosum, and to the subependymal surface of the lateral ventricles. Overall survival was 6.6 months.

The second patient with diffuse midline glioma was found to have mutations in histone H3 K27M mutation, additional driver mutations, such as ACVR1 and PPM1D were noted in all of the patients that were analyzed with 500 gene panel testing (Table 1, Supplementary Table 1). Five hundred gene panel testing was performed on tumor tissue of seven pediatric patients with diffuse midline glioma (n = 4) or hemispheric high-grade glioma (n = 3). Average age at surgery was 3.3 ± 1.9 years with overall survival of 9.8 ± 6.8 months. Three of the diffuse midline gliomas had the K27M mutation in the tail of histone H3 protein. One patient with diffuse midline glioma had wildtype histone H3.1 and H3.3 proteins. In addition to three hemispheric gliomas characterized in current study, tumor locations and described mutations were obtained from previously published six patients (Table 1). All of the patients were found to have additional mutations in genes that play a critical role in tumorigenesis. The gene mutations were different between the patients with diffuse midline gliomas and hemispheric high-grade gliomas. While patients with diffuse midline gliomas predominantly had the histone H3 K27M mutation, additional driver mutations, such as PIK3CA and PPM1D were noted in all of the patients that were analyzed with 500 gene panel testing (Table 1, Supplementary Table 1). For example, one of the patients had histone H3 K27M mutation, PIK3CA mutation within a highly conserved helical domain, and PPM1D truncation mutation (Supporting Fig. S1). Mutations in these genes suggest that disruption of chromatin structure and gene expression mediated by histone H3 tails, combined with disruptions in the PI3K pathway and p38/MAPK pathway are critical for tumorigenesis. This 3-year-old patient presented with a solid tumor that originated within the pons with extension into the middle cerebellar peduncle. The tumor was infiltrative in appearance, had evidence of central necrosis, mild enhancement, and no reduced diffusion. On six months follow up, the tumor developed significant progression with local extension into the cerebral peduncles, temporal stem, thalamus, corpus callosum, and to the subependymal surface of the lateral ventricles. Overall survival was 6.6 months.

Table 1. Mutations identified within diffuse midline gliomas and hemispheric gliomas and their effect on protein structure.

| Gene     | Mutation | Result                                                                 |
|----------|----------|------------------------------------------------------------------------|
| PIK3CA   | Q524K    | Substitution of glutamine for lysine, interferes with PI3Ka_1 domain responsible for substrate presentation |
| PIK3CA   | Q546K    | Substitution of glutamine for lysine, interferes with PI3Ka_1 domain responsible for substrate presentation |
| PIK3CA   | pC420R   | Protein substitution of cysteine for arginine, C2_PI3K_class_1_alpha domain involved in cell processes such as cell growth, differentiation, proliferation, and motility |
| PPM1D    | L450*    | Truncation mutation, PP2C domain conserved                             |
| BCO RL1  | Q1043*   | Truncation mutation, leads to loss of ankyrin repeat domains and PUF6_like_1 domain (PCGF Ub-like fold discriminator of BCO-likelike 1) |
| TP53     | R280L    | Substitution of arginine for leucine, interferes with tumor suppressor p53 |
| TP53     | pR273C   | Protein substitution of arginine for cysteine, interferes with tumor suppressor function of p53 |
| TP53     | Q192H    | Substitution of glutamine for histidine, interferes with tumor suppressor p53 |
| TP53     | 673-1G>A | Substitution of glycine for alanine, domains conserved                  |
| TP53     | R158C    | Substitution of arginine for cysteine, interferes with tumor suppressor p53 |
| TP53     | N239D    | Substitution of asparagine for aspartic acid, interferes with tumor suppressor p53 |
| ACVR1    | G328E    | Substitution of glycine for glutamic acid, interferes with STKc_ACVR1_ALK1 includes ATP binding and substrate binding sites |
| ASXL1    | pE566*   | On protein truncation mutation, domains conserved                       |
| NF1      | cT395-1G>T | On cDNA substitution of glycine for threonine, domains conserved         |
| NF1      | 2851-1G>T | Substitution of glycine for threonine, domains conserved                 |
| NF1      | R1362*   | Truncation mutation, interferes with RasGAP domain                      |
| SETD2    | T305fs   | Frame shift mutation, domains conserved                                |
| SETD2    | W1306*   | Truncation mutation, domains conserved                                 |
| SMARCB1  | Deletion | 22 Mb deletion on chromosome 22                                        |
| NF2      | Deletion | 22 Mb deletion on chromosome 22                                        |
| POLE     | P286L    | Substitution of proline for leucine, interferes with domain involved in catalytic subunit of DNA polymerase |
| POLE     | 286-1G>T | Substitution of glycine for threonine, interferes with domain involved in catalytic subunit of DNA polymerase |
| PTEN     | K128N    | Substitution of lysine for asparagine, interferes with PTEN_C2 domain responsible for membrane binding |
| PTEN     | T277I    | Substitution of threonine for isoleucine, interferes with PTEN_C2 domain responsible for membrane binding |

**Results**

**Imaging features of pediatric diffuse midline gliomas.** Five hundred gene panel testing was performed on tumor tissue of seven pediatric patients with diffuse midline glioma (n = 4) or hemispheric high-grade glioma (n = 3). Average age at surgery was 3.3 ± 1.9 years with overall survival of 9.8 ± 6.8 months. Three of the diffuse midline gliomas had the K27M mutation in the tail of histone H3 protein. One patient with diffuse midline glioma had wildtype histone H3.1 and H3.3 proteins. In addition to three hemispheric gliomas characterized in current study, tumor locations and described mutations were obtained from previously published six patients. All of the patients were found to have additional mutations in genes that play a critical role in tumorigenesis. The gene mutations were different between the patients with diffuse midline gliomas and hemispheric high-grade gliomas.

While patients with diffuse midline gliomas predominantly had the histone H3 K27M mutation, additional driver mutations, such as ACVR1 and PPM1D were noted in all of the patients that were analyzed with 500 gene panel testing (Table 1, Supplementary Table 1). For example, one of the patients had histone H3 K27M mutation, PIK3CA mutation within a highly conserved helical domain, and PPM1D truncation mutation (Supporting Fig. S1). Mutations in these genes suggest that disruption of chromatin structure and gene expression mediated by histone H3 tails, combined with disruptions in the PI3K pathway and p38/MAPK pathway are critical for tumorigenesis. This 3-year-old patient presented with a solid tumor that originated within the pons with extension into the middle cerebellar peduncle. The tumor was infiltrative in appearance, had evidence of central necrosis, mild enhancement, and no reduced diffusion. On six months follow up, the tumor developed significant progression with local extension into the cerebral peduncles, temporal stem, thalamus, corpus callosum, and to the subependymal surface of the lateral ventricles. Overall survival was 6.6 months.

The second patient with diffuse midline glioma was found to have mutations in histone H3.3 subtype (K27M), BCO RL1 truncation mutation, and TP53 mutation within a highly conserved arginine location within the DNA binding domain to a hydrophobic leucine. Mutations in these genes suggest that disruptions in gene expression mediated by modifications of histone H3 tails, the TP53 pathway, and a related pathway that regulates gene expression in a histone de-acetylase related manner is required to drive tumorigenesis (Supplementary Table 1). This patient presented at 4.7 years of age with a large heterogeneously enhancing tumor within the thalamus that demonstrated minimal reduced diffusion on DWI and was associated with mass effect on the third ventricle that resulted in non-communicating hydrocephalus (Supplementary Fig. 2). On three months follow up, the tumor progressed locally to the adjacent corona radiata, subependymal surface of the ventricles, and demonstrated...
The overall survival was 9.7 months. (Supplementary Fig. 4). Within 9 months of presentation, the tumor demonstrated extensive local metastatic evidence of vascular compression. There was mass effect on the fourth ventricle, but no hydrocephalus was noted. The pons was expanded with near circumferential surrounding of the basilar artery without FLAIR hyperintense infiltrating pontine glioma that demonstrated minimal enhancement and did not show pathway, and chromatin modification mediated gene expression driven by ASXL1 are driving the tumorigenesis of pediatric high-grade gliomas. Three of the patients in our cohort had hemispheric high-grade gliomas. One of these patients was diagnosed with neurofibromatosis 1 and developed three separate glioblastomas over the course of a year and a half (Supplementary Fig. 5). Two of the patient's separately appearing glioblastomas within different locations (corpus callosum and temporal lobe) were sequenced and hypermutated phenotype was shown with mutations in genes TP53, SETD2, ATRX, PTEN, POLE, BLM, ARID1A, APC, KDM6A, PTPN11, and BRCA2, which are predominantly involved in DNA repair. Both of the tumors contained mutations in TP53 and NF1, but additional mutations were different. The tumors arose in separate locations and at separate times from a background of extensive FLAIR hyperintense changes within the supratentorial and infratentorial brain parenchyma. The first tumor arose from the corpus callosum at 1.6 years of age, demonstrating FLAIR hyperintense signal and peripheral enhancement. Gross total resection of the tumor was achieved and there was no local recurrence of the tumor within the resection bed. Sixteen months after resection of the original tumor, a separate independent tumor arose from the contralateral temporal lobe and demonstrated rapid growth into a heterogeneous FLAIR hyperintense enhancing mass. Subtotal resection of the new tumor was achieved. During treatment of the temporal lobe tumor, additional tumor within a cerebellar hemisphere was also noted to arise and demonstrated rapid growth. The patient survived 23 months after diagnosis of the first tumor.

The second patient with hemispheric high-grade glioma demonstrated inactivating mutations in TP53 and PIK3CA. These mutations were similar to what is typically seen in diffuse midline gliomas but lacked a mutation in a chromatin modification related pathway. This patient was diagnosed at 2.8 years of age and was found to have a left temporal FLAIR hyperintense expansile mass that showed minimal initial enhancement and heterogeneous diffusion characteristics with regions of low and high ADC values (Supplementary Fig. 6). This tumor did not undergo gross total resection and demonstrated significant local progression within 6 months of diagnosis. On six months follow up, the tumor demonstrated heterogeneous enhancement and local spread into the adjacent basal ganglia, corpus callosum, and cerebral peduncles. The patient lived for 7 months after diagnostic biopsy.

The third patient with hemispheric high-grade glioma had wildtype histone H3 and demonstrated a hypermutated genotype with mutations in ATRX, TP53, CDKN2A, SETD2, MSH6, and PDGFRA. These genes are involved in the DNA repair pathway, regulation of cell cycle progression, and signaling via ERK pathway (Supplementary Tables 1, 2). This tumor was centered in the coronal radiata of the parietal lobe and demonstrated avid enhancement, central necrosis, and prominent peripheral reduced diffusion. The tumor was encircled by FLAIR hyperintense signal suggesting prominent edema with microscopic infiltration of tumor into adjacent brain parenchyma. There were no follow up imaging for this patient to determine the overall survival.

Gene expression levels based on Allen Brain Atlas gene expression atlas. Allen Brain Atlas microarray gene expression data for human brain was used to determine the normal gene expression levels of genes that were found to be mutated in diffuse midline gliomas and hemispheric high-grade gliomas. The microarray gene expression data represents gene expression in 500 anatomic locations that were micro-dissected from 6 adult normal human brains and 4 midgestational prenatal brains. H3F3A, HIST1H3B, ACRV1, PPM1D, BCO1L1, and ASXL1 genes were found to be mutated within our diffuse midline gliomas that were sequenced with 500 gene panel testing and agreed with published literature. Microarray gene expression data from normal brain demonstrated that these genes are preferentially expressed along the midline including the brainstem, cerebellum, and thalamus (Figs. 1, 2, Supplementary Fig. 7). On the contrary, these genes demonstrated lower overall expression within the cerebral hemispheres (Figs. 2, 3, and 4). Two of the
genes (BCORL1 and ASXL1) demonstrate increased expression along both the midline and within the cerebral hemispheres. BCORL1 gene was mutated in thalamic diffuse midline glioma in our cohort and its expression is normally elevated within the ventral thalamus, claustrum, and cingulum bundle (Fig. 1, Supplementary Fig. 7A). PPM1D gene was mutated in diffuse midline glioma that originated in superior and middle cerebellar peduncle, cerebral peduncle, and pons. Normally elevated expression of PPM1D gene is found within the thalamus, oculomotor nuclear complex of the midbrain, which is the location where tumor originated, globose nucleus, nucleus suberuless, superior olivary complex, and central medullary reticular group (Figs. 2, 3, and 4, Supplementary Fig. 7B). ASXL1 gene was mutated in diffuse midline glioma that originated within the pons. Normally elevated expression of ASXL1 gene was found within superior olivary complex of the pons, medullà's central glial substance and cuneate nucleus, rostral group of the interliminar nuclei of the thalamus, and additional nuclei within the thalamus (Figs. 2, 3, and 4, Supplementary Fig. 7C). ACVR1 gene mutation was found within diffuse midline glioma that originated within the cervical spine but can also be seen in other midline gliomas. There was increased expression of ACVR1 in multiple midline locations that included the thalamus and pons (Figs. 2, 3, and 4, Supplementary Fig. 7D). PIK3CA gene was mutated in tumors found within superior and middle cerebellar peduncle, cerebral peduncle, pons, cervical spine, temporal lobe, and insula. Elevated gene expression was also found within these anatomic regions including midbrain, pons, temporal lobe, and claustrum (Figs. 2, 3, and 4, Supplementary Fig. 7E). TP53 gene was mutated in tumors that were found in pons, thalamus, corpus callosum, temporal lobe, and insula. Elevated gene expression was also found within these anatomic regions including thalamus, pons, temporal lobe, and claustrum (Figs. 2, 3, and 4, Supplementary Fig. 7F).

Figure 1. Location based human gene expression levels within the cerebral hemispheres and along the midline structures for H3F3A, H3F3B, H3F3C, HIST1H3B, ACVR1, PPM1D, BCORL1, and ASXL1. Distribution of gene expression z scores within the brain with green indicating low expression and red indicating high expression.
In patients who had hemispheric high-grade gliomas, there were several characteristic imaging features. These tumors were primarily aggressive locally with prominent infiltrative spread best appreciated on FLAIR MR imaging. One of the patients had gross total resection of one of the tumors with distant recurrence. These tumors were predominantly hypermutated, with one of the patients demonstrating mutations in up to 12 genes. Some of these genes were mutated at different locations and recurred in the second tumor that grew at a separate location. When looking at gene expression of these genes in normal brains, they were predominantly expressed in the cerebral hemispheres with only some of the genes demonstrating strong expression along the midline structures and along the hemispheres together (Figs. 2, 3, and 4, Supplementary Fig. 8A). SETD2 gene was mutated in tumors originating from the corpus callosum and temporal lobe. Normal elevated expression of

**Figure 2.** Expression profile of genes mutated in midline and hemispheric high-grade gliomas.
SETD2 was found in corpus callosum and cingulate gyrus. There was decreased expression of SETD2 within the temporal lobe (Supplementary Fig. 8A). PTEN gene was mutated in tumor originating within the temporal lobe and in none of the diffuse midline gliomas. There were elevated expression levels of PTEN within the temporal lobe (Figs. 2, 3, and 4, Supplementary Fig. 8B). PTPN11 gene was mutated within a tumor that originated within corpus callosum and had highest gene expression in the corpus callosum and cingulum bundle (Figs. 2, 3, and 4, Supplementary Fig. 8C). KDM6A gene was mutated within a tumor that originated within corpus callosum with similar gene expression pattern being elevated within the midline structures of corpus callosum, cingulum bundle, and thalamus (Figs. 2, 3, and 4, Supplementary Fig. 8D). POLE, BLM, ARID1A and APC genes were mutated within a tumor that originated within the temporal lobe, but for this gene the highest expression was in the midline structures as opposed to temporal lobe (Figs. 2, 3, and 4, Supplementary Fig. 8E–H). ATRX gene was mutated within a tumor that originated within the white matter of the parietal lobe, but its expression within the parietal lobe was not elevated (Supplementary Fig. 8I). CDKN2A gene was mutated within a tumor that originated within the white matter of the parietal lobe, with its expression being highest within the cingulum bundle, claustrum, thalamus, and pons (Supplementary Fig. 8J). MSH6 gene was mutated within a tumor that originated within the white matter of the parietal lobe with highest gene expression of being seen in corpus callosum, cingulum bundle, and thalamus (Supplementary Fig. 8K). PDGFRA gene was mutated within a tumor that originated within the white matter of the parietal lobe with the predominant normal gene expression being found in pons, thalamus, and claustrum (Supplementary Fig. 8L). These findings suggest that not all of the genes that are mutated in the hemispheric tumors strictly have highest expression within the hemispheric structures, but when looking at overall expression patterns they still have higher expression along hemispheric structures (Figs. 2, 3, and 4).

Comparing endogenous gene expression z scores between genes that are commonly mutated in high grade midline gliomas versus high grade hemispheric gliomas demonstrates a difference in gene expression patterns in both adult and developmental human datasets (Figs. 2, 3, and 4). In the developmental brain dataset, the differences in hemispheric expression of PTEN and APC are not as elevated as seen in adults (Figs. 2 and 4). We noted the expression patterns of histone genes HIST1H3B and H3F3A (Supplementary Table 20, Supplementary Table 21, Supplementary Fig. 8M, Supplementary Fig. 8N). BCO1L1, PPM1D, PIK3CA, and TP53 appear to have higher expression z scores along the midline structures as compared to hemispheric structures. On the other hand, SETD2, PTEN, PTPN11, POLE, BLM, and APC appear to have higher expression z scores along the hemispheric structures as compared to midline structures.

Discussion
We report the imaging features of pediatric high-grade gliomas that are located along the midline and within the cerebral hemispheres and that had targeted next-generation gene sequencing analysis. We found that in our diffuse midline gliomas, additional driver mutations are always present, regardless of histone H3 mutation status. This agrees with published literature that multiple mutations are found in these tumors17,18. The genes that are mutated in the diffuse midline gliomas contribute to coordination of three distinct pathways. A chromatin...
modification alteration was always present in these tumors and is represented by modification of histone H3 tails or by presence of mutation in a chromatin modifying gene, such as ASXL1. In addition to alteration of chromatin modification, mutations in PI3K pathway are also found. One patient did not have PI3K mutation, but did have alteration in BCORL1, which is involved in histone deacetylation and has been shown to be involved in cross-talk with PI3K pathway19. In addition to chromatin modification and PI3K pathway alterations, mutations in TP53 pathway were also noted in our diffuse midline gliomas. In cases where TP53 was not mutated, genes involved in MAPK pathway, which is involved in crosstalk with TP53 pathway, were mutated20. This suggests that drivers of tumorigenesis in diffuse midline gliomas need to combine three distinct pathways—chromatin modification alteration, PI3K pathway, and p53 mediated MAPK cross-talk (Supplementary Fig. 9).

While the genes mutated in diffuse midline gliomas are normally expressed along the midline structures of the brain (Figs. 2, 3, and 4), in patients with hemispheric glioblastomas with wildtype histone H3, hypermutated genomes with mutations in multiple genes that play a critical role as tumor suppressors and DNA repair were seen in one of our patients and have been described in the literature (Supplementary Table 22)6,8. These genes are found to be normally expressed either along the cerebral hemisphere white matter or along both the cerebral hemispheres and midline structures (Figs. 2, 3, and 4). The differences in gene expression z-scores are very small, which is expected in normal brain tissues. The confirmation of the patterns that we see in adults with developmental brain data supports our original hypothesis.

TP53 and PIK3CA were found to be mutated in both midline and hemispheric high-grade gliomas. The exact TP53 mutations were different between the different patients that were analyzed, but involved critical function of DNA binding domains. Two of the diffuse midline gliomas and all three of the hemispheric gliomas had

Figure 4. Gene expression patterns from prenatal brain atlas data.
mutations in TP53. In addition, three of the diffuse midline gliomas and one of the hemispheric gliomas had mutation in PIK3CA, indicating a critical role of the PI3 kinase pathway in tumorigenesis. Normal gene expression analysis of these genes in humans demonstrates prominent expression in both the hemispheres and midline structures suggesting they are important in normal function. Expression of these genes within specific parts of the brain may predispose them to become mutated in the setting of open actively transcribed chromatin and may play a role in specific selection of drivers of tumorigenesis by tumors within specific locations of the brain.

There a number of limitations in our paper including small sample size and our cohort not including patients with mutations in IDH1/2 genes, which are common in adult hemispheric gliomas. Even though our cohort size is small, the genes that were identified on targeted next generation sequencing agree with published literature on more comprehensive datasets. In addition, gene expression profiles in humans ranging from 24 to 57 years of age were analyzed in Allen Brain Atlas Database, which we confirmed on the developing brain database within Allen Brain Atlas. Future studies should include gene expression profiles from pediatric patients as this data becomes more available. In addition, clonality of mutations within the tumors was not evaluated in the current study, which is an important area of research for future analysis. In addition to regional microdissection-based analysis of gene expression, focused single cell analysis within specific regions of the brain of the genes that are known to be mutated in pediatric brain tumors will provide critical information on etiology of tumorigenesis in midline versus hemispheric pediatric gliomas.

In conclusion, we characterize the imaging localization of tumors in pediatric patients which had targeted next-generation sequencing. We found that in diffuse midline gliomas three major pathways were mutated in every patient—chromatin modification, PI3 kinase pathway, and TP53 pathway. Among the hemispheric high-grade glioma patients, we had one patient with hypermutated glioblastomas and another patient with defects in the TP53 pathway and PI3 kinase pathway. To increase the number of our patients, we performed a multi-institutional study, which included patients with 4 additional hemispheric high-grade gliomas, 1 cerebellar hemisphere glioma, and 6 pontine diffuse midline gliomas (Table 2). Gene mutations in these patients followed similar trend, with TP53 mutations being found in hemispheric high-grade gliomas, while histone H3 K27M mutation being found in diffuse midline gliomas. The limitation of added patients from a second institution is that sequencing analysis in these tumors was limited to 50 gene panel, which does not include many of the driver mutations found in pediatric brain tumors. On analysis of temporal expression profile of normal gene expression in the brain, we found that genes that were mutated in patients with midline gliomas, were usually had normal higher expression in the anatomic locations where these tumors arise. This suggests that normal gene expression profile and brain microenvironment affects predisposition to mutagenesis and may play a role in defining driver mutations in tumors that arise from different parts of the brain.

Table 2. Mutations identified in individual pediatric brain tumors. Patients 1–8 underwent 500 gene panel mutation analysis. Patients 9–16 underwent either 50 gene panel mutation analysis. Patient 17 underwent Foundation One analysis. Patients 18 and 19 underwent standard pathologic analysis with immunohistochemistry stains for histone H3 K27M mutation. 50 gene panel analysis includes PIK3CA but not ACVR1 mutation analysis.

| Patient | Pathological diagnosis | Mutations | Tumor origin |
|---------|------------------------|-----------|-------------|
| 1       | Diffuse midline glioma | H3 K27M, PIK3CA, PPM1D | Pons        |
| 2       | Diffuse midline glioma | H3 K27M, BCORL1, TP53, PDGFRA | Thalamus    |
| 3       | Diffuse midline glioma | H3 K27M, PIK3CA, ACVR1 | Cervical spinal cord |
| 4       | Diffuse midline glioma | PIK3CA, TP53, ASXL1 | Pons        |
| 5       | Glioblastoma           | TP53, NF1, SETD2, ATRX, KDM6, PTPN11, BRCA2 | Corpus collosum |
| 6       | Glioblastoma           | TP53, NF1, PTEN, SETD2, POLE, BLM, ARID1A, APC | Temporal lobe |
| 7       | Glioblastoma           | ATRX, TP53, CDKN2A, SETD2, MSH6, PDGFRA | Parietal lobe |
| 8       | Glioblastoma           | SMARC1, NF2 | Frontal lobe |
| 9       | Glioblastoma           | TP53, EGFR | Temporal lobe |
| 10      | Anaplastic astrocytoma | IDH1, TP53 | Frontal lobe |
| 11      | Glioblastoma           | No histone H3 or IDH mutation | Cerebellar hemisphere |
| 12      | Glioblastoma           | IDH1, TP53 | Frontal lobe |
| 13      | Glioblastoma           | TP53 | Frontal lobe |
| 14      | Diffuse midline glioma | Histone H3 K27M, TP53 | Pons        |
| 15      | Diffuse midline glioma | PDGFRA, no histone H3 mutation | Pons        |
| 16      | Diffuse midline glioma | Histone H3 K27M, TP53, PTEN | Pons        |
| 17      | Diffuse midline glioma | Histone H3 K28M, TERT | Pons        |
| 18*     | Diffuse midline glioma | Histone H3 K27M | Pons        |
| 19*     | Diffuse midline glioma | Histone H3 K27M | Pons        |
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Author contributions

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Competing interests

The authors declare no competing interests.

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Correspondence and requests for materials should be addressed to M.S.A.

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