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**PDGFRA amplification is common in pediatric and adult high-grade astrocytomas and identifies a poor prognostic group in IDH1 mutant glioblastoma**

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**CONFLICT OF INTEREST**

The authors declare they have no conflict of interest.
Abstract

High-grade astrocytomas (HGAs), corresponding to WHO grades III (AA) and IV (GBM), are biologically aggressive and their molecular classification is increasingly relevant to clinical management. PDGFRA amplification is common in HGAs, although its prognostic significance remains unclear. Using fluorescence in situ hybridization (FISH), the most sensitive technique for detecting PDGFRA copy number gains, we determined PDGFRA amplification status in 123 pediatric and 263 adult HGAs. A range of PDGFRA FISH patterns were identified and cases were scored as non-amplified (normal and polysomy) or amplified (low-level and high-level). PDGFRA amplification was frequent in pediatric (29.3%) and adult (20.9%) tumors. Amplification was not prognostic in pediatric HGAs. In adult tumors diagnosed initially as GBM, the presence of combined PDGFRA amplification and IDH1R132H mutation was a significant independent prognostic factor (p=0.01). In HGAs, PDGFRA amplification is common and can manifest as high-level and focal or low-level amplifications. Our data indicate that the latter is more prevalent than previously reported with copy number averaging techniques. To our knowledge, this is the largest survey of PDGFRA status in adult and pediatric HGAs and suggests PDGFRA amplification increases with grade and is associated with a less favorable prognosis in IDH1 mutant de novo GBMs.

Keywords

PDGFRA; astrocytoma; FISH; IDH1; isocitrate dehydrogenase 1; prognosis

INTRODUCTION

High-grade astrocytomas (HGAs), including anaplastic astrocytoma (AA), WHO Grade III, and glioblastoma (GBM), WHO Grade IV, occur in both adults and children and are among the deadliest forms of cancer. Current practice stratifies HGAs based upon clinical, histopathologic, and limited molecular features. As our understanding of gliomagenesis and tumor response to therapy improves, this stratification will likely undergo multiple revisions with incorporation of additional molecular markers. As such, the continued development and validation of robust methods to assess molecular alterations in HGA is critical.

HGAs are characterized by alterations in receptor tyrosine kinase (RTK) signaling, and abnormal platelet-derived growth factor (PDGF) signaling has been demonstrated in a significant subset of both adult and pediatric tumors. In adult HGAs, PDGF receptor alpha (PDGFRα) is the second most commonly altered RTK receptor after epidermal growth factor receptor (EGFR), with amplification of the PDGFRα locus being the most common mechanism (7, 9, 18, 25, 27, 30). While estimates vary, in a large study using array-based comparative genomic hybridization, amplification of PDGFRα was identified in 11% of patients (27). Increased PDGF pathway activity, however, has been reported in up to 33% of adult GBM (3). Indeed, PDGF signaling pathway alterations are a characteristic feature of many tumors designated as “proneural” based on genomic, transcriptomal, and proteomic...
features (3, 21, 30). However, while tumors with a proneural phenotype may have an improved overall survival (21), other studies have suggested that PDGFRA copy number gain/amplification may be associated with worse overall survival in astrocytoma (1, 29).

In children, increased PDGF signaling is also thought to be an important driver of HGAs and PDGFRA amplification is similarly considered a common mechanism, with frequencies ranging from 3.4% to 12% (16, 20, 23). In specific clinical subsets of HGA, such as diffuse intrinsic pontine glioma, up to 25% of tumors may have amplification (32), potentially corresponding to its distinctive biologic properties (22). Despite the relatively high frequency of PDGFRA amplification in both pediatric and adult HGAs, the prognostic significance of this alteration remains largely unclear.

Mutations in isocitrate dehydrogenase 1 (IDH1) are common in adult AA and in subsets of adult GBM, including GBM that have progressed from a lower grade astrocytoma (secondary GBM); they are also found in a small subset of tumors diagnosed initially as GBM (de novo GBM) (5, 8, 19, 31). IDH1 mutant tumors, with R132H being the most common mutation, exhibit unique spatial, temporal, and biologic characteristics, including enhanced overall survival relative to IDH1 wild type tumors of a similar grade (15, 17). While infrequent in de novo GBM, IDH1 mutations are enriched in the proneural subtype, which is also characterized by alterations in PDGFRA signaling (21, 30).

Currently there are no established criteria for the assessment of PDGFRA copy number gain/amplification in clinical samples. While many studies have relied on copy number averaging techniques, such as PCR and SNP arrays, these methods may underestimate the frequency of PDGFRA amplifications when only scattered cells are amplified or the degree of amplification is low level. This is particularly true given the tremendous intratumoral heterogeneity of HGAs as recently illustrated for EGFR and PDGFR (24, 25). As such, the simplest and most sensitive technique for detecting copy number gains in routinely processed pathology specimens is fluorescence in situ hybridization (FISH). Using this technique, we assess a large series of HGAs, provide practical interpretive guidelines for the clinical assessment of PDGFRA copy numbers, and examine the prognostic significance of amplification in both adult and pediatric cohorts.

MATERIALS AND METHODS

Cohort

Formalin fixed, paraffin-embedded (FFPE) tumor tissue from a total of 123 pediatric HGAs, 103 adult AAs, 172 adult de novo GBMs (i.e. tumors initially diagnosed as GBM), and 17 IDH1 mutant secondary GBMs (i.e. GBM with documented progression from a lower grade astrocytoma) were obtained from ten institutions: UCSF Brain Tumor Research Center (BTRC) Tissue Bank; Department of Pathology, Newcastle General Hospital; Department of Pathology & Laboratory Medicine, University of Kentucky College of Medicine; Department of Pathology and Laboratory Medicine, Emory University; Department of Pathology, Gemelli Hospital, University of Sacred Heart, Rome Italy; Department of Pathology and Laboratory Medicine, The Children’s Hospital of Philadelphia, Philadelphia, PA; Department of Pathology and Laboratory Medicine, University of Pennsylvania; and Brain Tumor Translational Resource, Department of Pathology and Laboratory Medicine, University of California Los Angeles. These included both whole tissue and tissue microarray (TMA) sections, the latter obtained from ten previously generated HGA TMAs. Clinical characteristics are summarized in Table 1. Pediatric (0.1 – 20 years of age) and adult (25.7–83 years of age) astrocytoma patients were diagnosed with either AA, WHO grade III, or
GBM, WHO grade IV. Clinical and molecular characteristic of the tumor were obtained when available from the respective institutions and included survival from time of initial surgery, age at initial diagnosis, sex, and IDH1 mutation status (IDH1 R132H) using IDH1(R132H) immunohistochemistry (H09, Dianova GmbH, Hamburg, Germany) or sequencing (12).

Fluorescence in situ hybridization (PDGFRA)

Dual-color FISH analysis was performed on 5-micron thick FFPE whole and TMA sections as previously described (12, 14). Briefly, sections were deparaffinized, digested with pepsin, heat denatured and allowed to hybridize with probe sets overnight 37°C in a humidified oven. A SpectrumOrange (SO)-labeled home brew probe for PDGFRA (BAC clone RP11-231C18, CHORI BACPAC Resources Center, Oakland, CA, USA; previously reported in ref (2)) diluted 1:10 in DenHybe (Insitus, Albuquerque, NM) was paired with SpectrumGreen (SG)-labeled centromere enumerating probe (CEP4) 4p11-q11 (reference probe) (Abbott, Downers Grove, IL). Following washes to remove excess unbound probe, the nuclei were counterstained with 10 µl DAPI (Insitus, Albuquerque, NM) and slides were coverslipped. The fluorescent signals were enumerated under an Olympus BX41 fluorescent microscope with appropriate filters (Olympus; Melville, NY). For each hybridization, green and orange signals were enumerated in 100 non-overlapping nuclei. Slides were scanned for regional variability and were considered abnormal regardless of whether the alteration appeared focal or diffuse. Hybridizations were considered non-informative if the FISH signals were either lacking or too weak to interpret.

Statistical Analysis

A two-tailed t-test was used to compare mean values except where noted. P<0.05 was considered statistically significant. For Kaplan-Meier survival analysis, groups were compared using the log-rank (Mantel-Cox) test. Overall survival was truncated at 750 days for pediatric AAs, as there was a single death after 750 days, and at 4500 days for adult AAs, as there was a single censored patient after 4500 days. Contingency analysis was performed using Fisher’s exact test, two-sided. Multivariate Cox proportional hazard regression was used to model survival; while, 10-fold cross-validation and an integrated Brier score (6, 13) were used to compare the predictive error scores. All statistical analyses were carried out using GraphPad software (GraphPad Software Inc., La Jolla, CA) and R (26).

Study approval

All procedures were performed according to protocols approved by the University of California Committee on Research (San Francisco, California, USA). De-identified FFPE sections of human high-grade astrocytomas and TMAs were obtained from the participating institutions including the UCSF Brain Tumor Research Center Tissue Bank, Department of Pathology, Newcastle General Hospital; Department of Pathology, Children’s Hospital, Los Angeles; Department of Lab Medicine and Pathobiology, University of Toronto; Department of Pathology & Laboratory Medicine, University of Kentucky; Department of Pathology and Laboratory Medicine, Emory University; Department of Pathology, Gemelli Hospital, University of Sacred Heart, Rome Italy; The Children’s Hospital of Philadelphia; Department of Pathology and Laboratory Medicine, University of Pennsylvania; and Brain Tumor Translational Resource, Department of Pathology and Laboratory Medicine, University of California Los Angeles.
RESULTS

Patient characteristics

The study cohort included 123 pediatric and 263 adult HGAs whose clinical characteristics are summarized in Table 1. For pediatric HGAs, a total of 71 AAs, mean age 9.06 years (range 0.2 – 20), and 57 GBMs mean age 10.1 years (range 0.1 – 19) were analyzed. For adult HGAs, 103 AAs, mean age 50.6 years (range 25.7 – 80) and 160 de novo GBMs, mean age 55.1 years (range 27 – 79) were analyzed. Based on these data, an additional 27 de novo GBM with mutations in IDH1 and a mean age of 46 years (range 21 – 83) and 17 IDH1 mutant secondary GBM, with documented progression from a lower grade astrocytoma, with a mean age of 39.9 years (range 26 – 55) were analyzed.

Detection of PDGFRA copy number gain/amplification by FISH

A wide range of FISH patterns was identified, reflecting both inter- and intra-tumoral heterogeneity. Based on this heterogeneity we devised a scoring system that reflected both the relative number of tumor cells with copy number gain and the magnitude of copy number gain (Figure 1). Cases were scored as: normal (no increase or <10% cells with <6 PDGFRA signals); polysomy (>10% cells with 2–6 signals); low-level amplification (<10% of cells with >12 or innumerable signals or >40% cells with 6–12 signals); or high-level amplification (>10% cells with >12 or innumerable signals). Both low-level and high-level amplification were considered as “PDGFRA amplified.” These definitions were based on similar definitions for amplification of other oncogenes using clinical FISH assays (28). For instance, the definition of low-level amplification with >40% cells containing ≥6 signals is nearly identical to that utilized for EGFR amplification in lung cancer (10). Although biologically, this is probably more accurately described as a “high-level polysomy” rather than true gene amplification, the definition nonetheless correlates strongly with clinical outcome, including therapeutic response to tyrosine kinase inhibitors (10, 28). It was similarly felt that the more common pattern of gene amplification by FISH (>12 or innumerable signals) found in <10% of cells would have a roughly equivalent overall increase in dosage to that of a lower level of gain in larger numbers of cells (i.e., low-level amplification). Often, these two patterns of low-level amplification could be seen together in the same tumor. The decision not to use the PDGFRA/CEP4 ratio as part of the definition was based on the finding of several cases in which co-amplification of the centromeric region was found (Figure 1F) wherein a ratio near 1.0 would falsely exclude an interpretation of gene amplification.

Pediatric HGAs

FISH analysis of 123 pediatric HGAs demonstrated that 36 of 123 (29.3%) tumors had PDGFRA amplification, including 19 (15.4%) high-level and 17 (13.8%) low-level examples. A high frequency of PDGFRA amplification has been reported in a subset of diffuse intrinsic pontine gliomas (DIPGs) (22, 32). In our cohort we did not observe an increase in PDGFRA amplification in the brainstem/cerebellum versus other brain regions (p=0.53); however, our cohort included only 12 HGAs that involved the brainstem or cerebellum. To examine potential clinical differences between patients with PDGFRA amplified versus non-amplified tumors, we analyzed GBM and AA separately.

In pediatric GBM, FISH analysis demonstrated a striking 22 of 57 (38.6%) tumors with amplification of PDGFRA, including 11 (19.3%) with low-level and 11 (19.3%) with high-level amplification. Stratification of clinical variables based on PDGFRA amplification is shown in Table 2. Patients with PDGFRA amplified tumors tended to be older than patients without amplification; however, this difference was not significant (p=0.13). There was no
difference in overall survival between patients with and without PDGFRA copy number gain/amplification (Figure 2A).

In pediatric AA, we identified a slightly lower percentage of PDGFRA amplified cases than in GBM (Table 3). A total of 14 of 66 (21.2%) tumors had amplification, including 6 (9.09%) with high-level and 8 (12.1%) with low-level amplification. Although not statistically significant, the age of patients with PDGFRA amplified tumors tended to be older (p=0.08). PDGFRA amplified tumors also tended to be more common in females than males (p=0.18). Similar trends in age and sex in the PDGFRA amplified tumors was noted in the pediatric GBM cohort. There was no significant difference in overall survival between PDGFRA amplified and non-amplified tumors, although there was a trend towards worse survival with amplification (log-rank, p=0.21) (Figure 2B).

**Adult HGAs**

FISH analysis in adult HGAs demonstrated 55 of 263 (20.9%) PDGFRA amplified tumors, including 25 (9.50%) with high-level and 30 (11.4%) with low-level amplification. Similar to pediatric HGAs, PDGFRA amplification was more frequent in de novo GBM than AA. In adults, 36 of 160 (22.5%) GBM had PDGFRA amplification, including 17 (10.6%) high-level and 19 (11.9%) low-level (Table 4). While the patient’s age, sex, and overall survival were similar between PDGFRA amplified and non-amplified tumors, there was a significant association between PDGFRA amplification and mutations in IDH1 (p=0.028) (Table 4 and Figure 3A).

The R132H mutation in IDH1 was present in 12 of 160 (7.5%) adult de novo GBM. Based on the enrichment of PDGFRA amplification in IDH1 mutant tumors, we performed a subset analysis focused only on those tumors with mutant IDH1 (IDH1R132H). To increase the number of tumors available for analysis, we obtained an additional 27 IDH1 mutant de novo GBM (the clinical characteristics of all 39 tumors are summarized in Table 5). As expected, patients with de novo GBM with IDH1 mutation had significantly better overall survival than patients without IDH1 mutations (median survival 1927 days (n=38) versus 424 days (n=137), respectively, log-rank, p<0.0001). Stratification of these IDH1 mutant tumors by PDGFRA amplification status revealed a striking difference in median overall survival between patients with and without amplification (Figure 3B). Overall median survival was 480 days (n=16) for patients with IDH1 mutant de novo GBM with PDGFRA amplification versus 2179 days (n=22) without PDGFRA amplification (log-rank, p=0.023). Other clinical characteristics including age and sex were not significantly different between PDGFRA amplified and non-amplified tumors.

In a multivariate analysis of all de novo GBMs, we examined PDGFRA status and IDH1 mutation status. While PDGFRA status alone was not a significant prognostic factor, the interaction of PDGFRA amplification and IDH1 mutation status (i.e. the group of tumors which are both PDGFRA amplified and IDH1 mutated) was found to be a significant prognostic factor (p=0.01) (Table 6) and remained significant when age was included in the model (p=0.049).

Cross-validation is a technique used for model selection as well as to assess if the model will be useful in an independent data set. Using 10-fold cross-validation and the integrated Brier score, a measure of the strength of the model, we compared the predictive error scores between several models: baseline (no variables in the model); main effect models with IDH1 mutation and/or PDGFRA amplification status; and, an interaction model which included the two main effects as well as an interaction term that signifies both IDH1 mutation and PDGFRA amplification. The interaction model had the lowest prediction error and was
considered to be the best model, resulting in a marked reduction in error over baseline (16%) and a marginal but consistent reduction in error over IDH1 mutation alone (2%).

In adult AAs the frequency of PDGFRA amplification was less than in GBMs, with 19 of 103 (18.4%) tumors amplified, including 8 (7.8%) high-level and 11 (10.7%) low-level (Table 7). PDGFRA amplified tumors tended to be located in the frontal lobes, 10 of 15 (66.7%), as compared to the non-amplified tumors, 29 of 69 (42.0%), although this was not statistically significant (p=0.15). PDGFRA amplification was not prognostic for overall survival (Figure 3C). Interestingly, PDGFRA amplification was not increased in IDH1 mutant AAs, mutations seen in 61.2% of amplified tumors and 62.8% of non-amplified tumors. We next analyzed the subset of 60 IDH1 mutant AA to determine whether PDGFRA had prognostic benefit in IDH1 mutant adult AA (Table 8). Unlike de novo GBM in which the frequency of PDGFRA amplification was highly enriched in IDH1 mutant tumors (17/39; 43.6%) versus IDH1 non-mutant tumors (30/151; 19.9%), in AA the frequency of PDGFRA amplification was similar in IDH1 mutant tumors (11/60; 18.3%) and in IDH1 non-mutant tumors (7/36; 19.5%). Furthermore, in IDH1 mutant AA, there was no statistically significant difference in overall survival between PDGFRA amplified and non-amplified tumors (median survival 960 days (n=9) versus 3300 (n=45) days, respectively, p=0.52). While PDGFRA amplified tumors were more common in the frontal lobes with 8 of 9 (88.9%) amplified tumors located in the frontal lobes as compared to 22 of 47 (46.8%) of the non-amplified tumors, this difference was not statistically significant (p=0.26).

IDH1 mutant de novo GBM share many features with secondary GBM, derived from the progression of a lower grade astrocytoma (15). Similar to IDH1 mutant de novo GBM, PDGFRA amplification was frequent in IDH1 mutant secondary GBMs, 7 of 17 (41.2%) tumors. Unlike in de novo GBM however, PDGFRA amplification was not associated with shorter overall survival (Table 9).

DISCUSSION

HGAs are a heterogeneous group of tumors and improvements in therapy will likely require stratification of patients based upon clinical, histopathologic, and molecular characteristics. Using FISH, we define a set of criteria to evaluate PDGFRA copy number alterations and determine the frequency of amplification in a large cohort of pediatric and adult HGAs. Our data suggest that PDGFRA amplification is higher than previously estimated in both pediatric (29.3%) and adult (20.7%) HGAs. To our knowledge this represents the largest reported number of pediatric and adult HGAs studied by FISH, and these data support the notion that abnormal PDGFRA signaling is important in HGA. In adults, PDGFRA amplification was associated with significantly worse overall survival in IDH1 mutant de novo GBM. Indeed, in a multivariate analysis of all adult de novo GBMs, the combination of PDGFRA amplification and IDH1 mutation status was identified as a significant prognostic factor. While additional studies in larger patient cohorts are required, our data suggest that IDH1 mutant de novo GBM may be a more heterogeneous group than previously thought.

PDGFRA signaling is an important driver of glioma development and progression, and based on whole-genome technologies, PDGFRA copy number gain/amplification is the second most common genetic alteration in RTKs in adult GBM, commonly estimated at 11% (27). This reported frequency is nearly identical to that of the high-level amplification pattern that we noted in the current series; however, our techniques also allowed us to uncover a large number of low-level amplifications. Using FISH on routinely processed pathology specimens we identified a high frequency of PDGFRA amplification in adult HGA, including 22.5% in GBM and 18.4% in AA. In contrast to EGFR FISH, where widespread
high-level amplification is the rule, focal or low-level PDGFRA amplifications were even more common. As such, this alteration may be particularly susceptible to underestimation by dose averaging techniques, such as PCR and array CGH. To reflect the diversity of PDGFRA signals observed by FISH, our scoring system included both low-level and high-level amplification. With the clinical variables available we did not observe a significant survival difference between tumors with low- and high-level amplification; thus we considered both “positive” for amplification (data not shown). In future studies, the level of PDGFRA amplification may have unique and unexpected prognostic associations as has been seen with EGFR (11).

In adult de novo GBM with IDH1 mutations, the frequency of PDGFRA amplification was striking with nearly half of tumors positive for amplification. Interestingly, this was not true in AAs. In IDH1 mutant AA, the frequency of PDGFRA amplification was only 18.3%. While the number of IDH1 mutant GBM and AA was relatively small (39 GBM and 60 AA) this difference was significant (43.6% vs. 18.3%, p=0.011 (OR 3.442; CI 1.385 – 8.554)). These data suggest that PDGFRA amplification may be an important event in the transition from AA to GBM in IDH1 mutant tumors. In support of this idea, analysis of a small number of IDH1 mutant secondary GBMs revealed a high frequency of PDGFRA amplification. Furthermore, the percent of cells with PDGFRA amplification tended to be less in IDH1 mutant de novo GBM as compared to IDH1 wild type GBM, as suggested by the number of cases with low-level amplification (12/17; 71%) versus (20/36; 56%), respectively. Gene amplification may be particularly important in the progression of IDH1 mutant astrocytoma as Lai et al. (15) identified EGFR amplification in a smaller percentage of cells in IDH1 mutant versus IDH1 wild type tumors. Interestingly, while survival tended to be shorter in PDGFRA amplified, IDH1 mutant AA and secondary GBM this did not reach statistical significance. The difference in survival between IDH1 mutant de novo GBM versus AA and secondary GBM may reflect an insufficient sample size and high number of censored subjects for the latter or it may suggest potential biologic differences between clinically defined de novo GBM and IDH1 mutant HGAs that progress from a lower grade tumor.

Reports on the frequency of PDGFRA amplification in pediatric HGA vary (range 3.4% to 12%) with up to 50% reported in irradiation-induced HGAs (16, 20, 23). In a large study of pediatric HGAs, PDGFR amplification as detected by array CGH was identified in 12% of HGAs overall and 17% of GBM (20). In DIPG, PDGFRA amplification may be even more common than in other HGAs, with reported estimates of 29% (22, 32). Using FISH we identified PDGFRA amplification in 29.3% of HGAs, 38.6% in GBM and 21.2% in AA. While we did not observe a significant association between brainstem location and PDGFRA amplification, we did not specifically target this group of tumors for analysis and only 12 HGAs involved the brainstem or cerebellum. PDGFRA status was not associated with overall survival in our cohort of pediatric HGAs.

This large multi-institutional study included a broad cross-section of cases from ten major medical centers; however, it was a retrospective study and has inherent biases including potential case selection bias. In addition, we had access to only limited clinical and molecular data. Indeed, genetic information such as PDGFRA mutation status, shown to be common in amplified tumors (18), and co-amplification of EGFR was not assessed (24, 25). Due to several cases with co-amplification of PDGFRA and the centromeric region for which a ratio near 1.0 would falsely exclude PDGFRA amplification, we did not use the PDGFR/CEP4 ratio to define gene amplification. While this allowed us to identify all cases with PDGFRA amplification we were not able to assess whether PDGFRA was selectively amplified or was co-amplified with other potential oncogenic genes on chromosome 4, particularly KIT and KDR (VEGFR2), given that specific probes for these
other genes were not applied. In addition, while our M:F ratio for adult GBM (1.6) was similar to the reported ratio of 1.58, our adult AAs had a very high M:F ratio (8.1) relative to the reported ratio of 1.39 (4). As PDGFRA amplification tended to be more common in female patients, we may have underestimated the overall frequency of PDGFRA amplification in this population.

In this study we define a set of criteria to assess PDGFR amplification in routine clinical samples and provide an estimate of the frequency of PDGRA amplification in a large set of pediatric and adult HGAs. In our cohort, PDGFRA amplification did not have prognostic significance in pediatric HGA. In adults, we identified PDGFRA amplification as an independent prognostic factor in IDH1 mutant de novo GBM. These data have important potential implications regarding tumor biology and prognosis and additional studies in a larger number of IDH1 mutant de novo GBM are required.

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REFERENCES


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Figure 1. Determination of PDGFRA copy number gain/amplification by FISH
Fluorescent images illustrating the different patterns of PDGFRA amplification in HGAs. (A) Normal, no increase in PDGFRA signals. (B) Polysomy, >10% of cells with >2 but <6 signals for both PDGFRA and CEP4. (C) The most frequent pattern of low-level amplification, innumerable PDGFRA signals in <10% of cells. (D) Another pattern of low-level amplification, >40% cells with ≥6 signals (a few signals are beyond the plane of focus) (E) High-level amplification, >10% cells with >12 or innumerable PDGFRA signals. (F) Rare tumors demonstrated high-level co-amplification of PDGFRA and CEP4. Amp.
denotes \textit{PDGFR}A amplification. PDGFR\textit{A} probe (\textit{red}) and CEP4 (\textit{green}), magnification x1000.
Figure 2. Pediatric HGA patients have similar overall survival with and without PDGFRA amplification

(A) Pediatric patients with GBM with (n=13) and without (n=16) PDGFRA amplification have no significant difference in overall survival based on Kaplan-Meier survival analysis, p=0.97. (B) Kaplan-Meier survival analysis examining overall survival for pediatric patients with anaplastic astrocytoma with (n=9) and without (n=25) amplification of PDGFRA, p=0.21
Figure 3. PDGFRA amplification is associated with worse overall survival in adult IDH1<sup>R132H</sup> mutant GBM

(A) Kaplan-Meier survival analysis patients with de novo GBM demonstrates no significant difference in overall survival for patients with (red, n=34) and without (blue, n=114) PDGFRA amplification, p=0.45. (B) Comparison of overall survival in larger cohort of patients stratified for IDH1 mutation (IDH1<sup>R132H</sup>) demonstrating decreased survival in IDH1 mutant GBM with PDGFRA amplification (red, n=16) as compared to those without PDGFRA amplification (blue, n=22) by Kaplan-Meier survival analysis, p=0.023. Survival curves are also shown for IDH1 non-mutant GBM with (green, n=29) and without (black, n=108) PDGFRA amplification. A similar analysis in adult patients with AA demonstrated
overall survival was similar regardless of PDGFRA amplification in (C) all patients (amplified, red (n=11); not amplified, blue (n=58)) and in (D) patients stratified for IDH1 mutation status (IDH1 mutant and amplified, red (n=9); IDH1 mutant and not amplified, blue (n=45); IDH1 non-mutant and amplified, green (n=2); IDH1 non-mutant and not amplified, black (n=21)) by Kaplan-Meier survival analysis, p=0.41 or 0.52, respectively.
### Table 1

Clinical and molecular characteristics of 123 pediatric and 263 adult HGAs.

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<tr>
<td>Mean age (years ± SD)</td>
<td>9.06 ± 5.16</td>
<td>10.1 ± 5.0</td>
<td>50.6 ± 13.8</td>
<td>55.1 ± 13.1</td>
</tr>
<tr>
<td>Sex ratio (M:F)</td>
<td>1.0</td>
<td>1.0</td>
<td>8.1</td>
<td>1.6</td>
</tr>
<tr>
<td>Locationb (supra- vs. infra-tentorial) (%)</td>
<td>86.0%</td>
<td>72.3%</td>
<td>96.7%</td>
<td>N/A</td>
</tr>
<tr>
<td>Location (Frontal vs. Other)</td>
<td>14.0%</td>
<td>6.38%</td>
<td>47.2%</td>
<td>N/A</td>
</tr>
<tr>
<td>IDH1 mutant</td>
<td>N/A</td>
<td>N/A</td>
<td>62.5% (n=96)</td>
<td>7.50% (n=160)</td>
</tr>
<tr>
<td>Median survival (days)</td>
<td>490 (n=34)</td>
<td>596 (n=29)</td>
<td>2070 (n=69)</td>
<td>450 (n=148)</td>
</tr>
</tbody>
</table>

a n = Number analyzed.

b Data on tumor location was available for 43 pediatric AA, 47 pediatric GBM, and 91 adult AA.

c N/A = Data is not available.
Table 2
Clinical characteristics of 57 pediatric patients with GBM.

<table>
<thead>
<tr>
<th>Patient characteristics</th>
<th>No amplification</th>
<th>PDGFRA amplification</th>
<th>Odds ratio (95% CI)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number (%)</td>
<td>35 (61.4%)</td>
<td>22 (38.6%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mean age (years ± SD)</td>
<td>9.23 ± 5.49</td>
<td>11.40 ± 3.82</td>
<td></td>
<td>0.13</td>
</tr>
<tr>
<td>Sex ratio (M:F)</td>
<td>1.1</td>
<td>1.0</td>
<td>1.067 (0.3464 – 3.285)</td>
<td>1.0</td>
</tr>
<tr>
<td>Median survival (days)</td>
<td>596</td>
<td>494</td>
<td></td>
<td>0.97</td>
</tr>
<tr>
<td>Location: BS/CB² vs. other</td>
<td>24.1% (n=29)</td>
<td>11.2% (n=18)</td>
<td>2.545 (0.4656 – 13.92)</td>
<td>0.45</td>
</tr>
</tbody>
</table>

²BS/CB = Brainstem/Cerebellum
Table 3
Clinical characteristics of 66 pediatric patients with AA.

<table>
<thead>
<tr>
<th>Patient characteristics</th>
<th>No amplification</th>
<th>PDGFRA amplification</th>
<th>Odds ratio (95% CI)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number (%)</td>
<td>52 (78.8%)</td>
<td>14 (21.2%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mean age (years ± SD)</td>
<td>8.16 ± 5.49</td>
<td>11.2 ± 3.76</td>
<td>0.08</td>
<td></td>
</tr>
<tr>
<td>Sex ratio (M:F)</td>
<td>1.38</td>
<td>0.45</td>
<td>3.115 (0.7859 – 12.35)</td>
<td>0.18</td>
</tr>
<tr>
<td>Median survival (days)</td>
<td>513</td>
<td>403</td>
<td>0.8571 (0.0707 – 10.38)</td>
<td>1.0</td>
</tr>
<tr>
<td>Location: BS/CB² vs. other</td>
<td>6.67% (n=30)</td>
<td>7.69% (n=13)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

²BS/CB = Brainstem/Cerebellum
Table 4
Clinical and molecular characteristics of 160 adult GBM\(^a\) patients.

<table>
<thead>
<tr>
<th>Patient and tumor characteristics</th>
<th>No amplification</th>
<th>PDGFRA amplification</th>
<th>Odds ratio (95% CI)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number (%)</td>
<td>124 (77.5%)</td>
<td>36 (22.5%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mean age (years ± SD)</td>
<td>55.7 ± 13.3</td>
<td>53.6 ± 12.7</td>
<td>0.56</td>
<td></td>
</tr>
<tr>
<td>Sex ratio (M:F)</td>
<td>1.9</td>
<td>1.1</td>
<td>0.5806 (0.2091 – 1.612)</td>
<td>0.31</td>
</tr>
<tr>
<td>% IDH1 mutanta</td>
<td>5.08% (n=124)</td>
<td>16.7% (n=36)</td>
<td>3.933 (1.184 – 13.07)</td>
<td>0.028</td>
</tr>
<tr>
<td>Median survival (days)</td>
<td>450 (n=114)</td>
<td>455 (n=34)</td>
<td>0.45</td>
<td></td>
</tr>
</tbody>
</table>

\(^a\) All tumors were de novo GBM and diagnosed at initial presentation as GBM.

\(^b\) IDH1 mutation=IDH1\(^R132H\)
Table 5

Clinical characteristics of 39 adult patients with IDH1 mutant GBM\textsuperscript{a,b}.

<table>
<thead>
<tr>
<th>Patient characteristics</th>
<th>No amplification</th>
<th>( PDGFRA ) amplification</th>
<th>Odds ratio (95% CI)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number (%)</td>
<td>22 (56.4%)</td>
<td>17 (43.6%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mean age (years ± SD)</td>
<td>44.7± 12.8</td>
<td>46.8 ± 16.8</td>
<td>0.67</td>
<td></td>
</tr>
<tr>
<td>Sex ratio (M:F)</td>
<td>1.8</td>
<td>2</td>
<td>0.9167 (0.2121 – 3.963)</td>
<td>1.0</td>
</tr>
<tr>
<td>Median survival (days)</td>
<td>2179 (n=22)</td>
<td>480 (n=16)</td>
<td>0.023</td>
<td></td>
</tr>
</tbody>
</table>

\textsuperscript{a}All tumors were de novo GBM and diagnosed at initial presentation as GBM.

\textsuperscript{b}IDH1 mutation=IDH1\textsuperscript{R132H}
Table 6
Hazard ratios from a multivariate survival analysis in adult patients with de novo GBM$^a$

<table>
<thead>
<tr>
<th></th>
<th>No Amplification</th>
<th>PDGFRA amplification</th>
</tr>
</thead>
<tbody>
<tr>
<td>IDH1 wt</td>
<td>1.0</td>
<td>0.739</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(0.477–1.145)$^b$</td>
</tr>
<tr>
<td></td>
<td></td>
<td>p=0.18</td>
</tr>
<tr>
<td>IDH1 mutant$^c$</td>
<td>0.140</td>
<td>0.416$^d$</td>
</tr>
<tr>
<td></td>
<td>(0.0639–0.3055)$^b$</td>
<td>(0.2131–0.8105)$^b$</td>
</tr>
<tr>
<td></td>
<td>p=8.2x10$^{-7}$</td>
<td>p=0.01</td>
</tr>
</tbody>
</table>

$^a$GBM=Complete information was available for 136 of the 178 patients, including 38 IDH1 mutant tumors.

$^b$CI=Confidence interval

$^c$IDH1 mutation=IDH1R132H

$^d$Both IDH1 is mutated and PDGFRA is amplified
Table 7
Clinical and molecular characteristics of 103 adult AA patients.

<table>
<thead>
<tr>
<th>Patient and tumor characteristics</th>
<th>No amplification</th>
<th>PDGFRA amplification</th>
<th>Odds ratio (95% CI)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number</td>
<td>84 (81.6%)</td>
<td>19 (18.4%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mean age (years ± SD)</td>
<td>50.5 ± 13.2</td>
<td>50.9 ± 16.8</td>
<td>0.91</td>
<td></td>
</tr>
<tr>
<td>Sex ratio (M:F)</td>
<td>9.2</td>
<td>5.5</td>
<td>1.673 (0.2857 – 9.794)</td>
<td>0.62</td>
</tr>
<tr>
<td>IDH1 mutant&lt;sup&gt;a&lt;/sup&gt;</td>
<td>62.8% (n=78)</td>
<td>61.2% (n=18)</td>
<td>1.075 (0.3750 – 3.083)</td>
<td>1.00</td>
</tr>
<tr>
<td>Median survival (days)</td>
<td>2070 (n=58)</td>
<td>960 (n=11)</td>
<td>0.41</td>
<td></td>
</tr>
<tr>
<td>Location: Frontal vs. other</td>
<td>42.0% (n=69)</td>
<td>66.7% (n=15)</td>
<td>0.3718 (0.1146 – 1.206)</td>
<td>0.15</td>
</tr>
</tbody>
</table>

<sup>a</sup>IDH1 mutation=IDH1<sup>R132H</sup>
Table 8
Clinical characteristics of 60 adult IDH1 mutant AA<sup>a</sup>.

<table>
<thead>
<tr>
<th>Patient characteristics</th>
<th>No amplification</th>
<th>PDGFRA amplification</th>
<th>Odds ratio (95% CI)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number</td>
<td>49 (81.7%)</td>
<td>11 (18.3%)</td>
<td></td>
<td>0.29</td>
</tr>
<tr>
<td>Mean age (years ± SD)</td>
<td>45.1 ± 10.9</td>
<td>41.3 ± 7.30</td>
<td></td>
<td>0.29</td>
</tr>
<tr>
<td>Sex ratio (M:F)</td>
<td>10</td>
<td>2.5</td>
<td>4.000 (0.5281 – 30.30)</td>
<td>0.20</td>
</tr>
<tr>
<td>Median survival (days)</td>
<td>3300 (n=45)</td>
<td>960 (n=9)</td>
<td></td>
<td>0.52</td>
</tr>
<tr>
<td>Location: Frontal vs. other</td>
<td>46.8% (n=47)</td>
<td>88.9% (n=9)</td>
<td>0.5266 (0.1790 – 1.549)</td>
<td>0.26</td>
</tr>
</tbody>
</table>

<sup>a</sup>IDH1 mutation=IDH<sub>1</sub><sup>R132H</sup>
Table 9
Clinical characteristics of 17 adult IDH1 mutant secondary GBM<sup>a</sup>.

<table>
<thead>
<tr>
<th>Patient characteristics</th>
<th>No amplification</th>
<th>PDGFRA amplification</th>
<th>Odds ratio (95% CI)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number</td>
<td>10 (58.8%)</td>
<td>7 (41.2%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mean age (years ± SD)</td>
<td>41.4 ± 10.2</td>
<td>39.5 ± 7.41</td>
<td></td>
<td>0.89</td>
</tr>
<tr>
<td>Sex ratio (M:F)</td>
<td>3</td>
<td>6</td>
<td>0.5000 (0.0415 – 6.021)</td>
<td>1.0</td>
</tr>
<tr>
<td>Median survival (days)</td>
<td>540 (n=10)</td>
<td>317.5 (n=7)</td>
<td></td>
<td>0.15</td>
</tr>
</tbody>
</table>

<sup>a</sup>Secondary GBM = GBM with documented progression from a lower grade astrocytoma.