



# Clinicopathological characteristics of circumscribed high-grade astrocytomas with an unusual combination of *BRAF* V600E, *ATRX*, and *CDKN2A/B* alternations

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## Abstract

We report four cases of high-grade astrocytoma with a *BRAF* V600E mutation, *ATRX* inactivation, and *CDKN2A/B* homozygous deletion. Children to young adults aged 3–46 presented with a well demarcated contrast-enhancing mass in the supratentorial area. Pathological examination revealed packed growth of short spindle to round polygonal cells including some pleomorphic cells. The tumors had less ability to infiltrate into the adjacent brain parenchyma and presented a circumscribed growth pattern. Mitosis was readily found, accompanied by focal necrosis and/or microvascular proliferation. Tumors were histologically similar in part to pleomorphic xanthoastrocytoma (PXA) or anaplastic PXA, but did not fit criteria for either neoplasm. A *BRAF* V600E mutation and homozygous deletion of *CDKN2A/B* were observed, which is similar to the genetic features of PXA or epithelioid glioblastoma, but the additional loss of *ATRX* nuclear immunoreactivity and absence of *TERT* promoter mutation were unusual findings, indicating a novel genetic profile. Despite their malignant histological features, all patients had a favorable clinical course and remained alive for 6 months to 28 years under standard medical treatment for malignant glioma. In summary, high grade astrocytomas with *BRAF* V600E, *ATRX*, and *CDKN2A/B* alternations had unique clinicopathological features and may be a novel subset of high grade glioma.

**Keywords** High-grade astrocytoma · Circumscribed astrocytoma · *BRAF* V600E · *ATRX* · *CDKN2A/B*

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## Introduction

Astrocytic tumors can be separated by their growth pattern, either diffuse or localized. Diffuse astrocytic tumors include diffuse astrocytoma, anaplastic astrocytoma, and glioblastoma. These tumors are mainly located in the supratentorial white matter and are characterized by diffuse infiltration. Localized astrocytomas, such as pilocytic astrocytoma and pleomorphic xanthoastrocytoma (PXA), have less ability to infiltrate into the adjacent brain parenchyma and tend to show a solid growth pattern in the subarachnoid space. Diffuse astrocytic tumors have an *IDH1/2* mutation, frequent inactivation of *ATRX*, and mutation of *TP53*, whereas localized astrocytomas tend to have mutations in the MAPK pathway such as *BRAF*.

PXA with increased mitotic activity ( $\geq 5$  mitoses per 10 high-powered fields) was recognized by the WHO classification criteria in 2016 as anaplastic PXA (A-PXA). A-PXAs have typical histological features of PXA, such as

proliferation of spindle-shaped cells with intermingled pleomorphic cells or xanthomatous cells, abundant eosinophilic granular bodies, and rich pericellular reticulin fibers [1]. PXA and A-PXA commonly have a *BRAF* V600E mutation and a homozygous deletion of *CDKN2A/B*. Moreover, *TERT* promoter mutations occasionally occur in PXA (20.5% in PXA, 23–47% in A-PXA) [2, 3], but *ATRX* inactivation is rare.

Here, we report four cases of high grade astrocytomas with an unusual combination of a *BRAF* V600E mutation, *CDKN2A/B* homozygous deletion, and *ATRX* inactivation. We discuss the histopathological differences between these tumors and other astrocytic tumors including PXA.

## Materials and methods

### Tumor samples

Four cases of high-grade astrocytoma, which had *ATRX* inactivation and *BRAF* V600E mutation by immunohistochemistry and DNA sequence, were recruited. Two cases were from the pathology archives of the Department of Pathology, Gunma University Hospital. One was reported previously (Case 2) [4]. The other two cases were from the pathology archives of the Tokyo Medical University Hospital and the University of Ryukyus Graduate School of Medicine. This research was approved by the ethics committee of Gunma University.

### Clinical data

Clinical data and images were obtained from the charts and clinicians of each hospital.

### Conventional histological analysis

Tissue specimens were fixed in 10% formalin and embedded in paraffin. Three-micrometer-thick tissue sections were stained with hematoxylin and eosin. For immunohistochemistry, the following primary antibodies were used: Glial fibrillary acidic protein (GFAP; polyclonal, 1:5,000; our own), Olig2 (polyclonal, 1:5000; IBL, Takasaki, Japan), S-100 protein (polyclonal, 1:10,000; our own), Nestin (monoclonal, 1:200; Merck Millipore, Billerica, MA, USA), CD34 (monoclonal, 1:200; Nichirei, Tokyo, Japan), p53 protein (monoclonal, 1:50; Leica Microsystems, Wetzlar, Germany), *ATRX* (polyclonal, 1:500; Sigma, St. Louis, MO, USA), *BRAF* V600E epitope (monoclonal, 1:50; Spring Bioscience, Pleasanton, CA, USA), Isocitrate dehydrogenase 1 (IDH1)-R132H (monoclonal, 1:50; Dianova, Hamburg, Germany), and Ki-67 (MIB-1; monoclonal, 1:100; Dako). For GFAP, S-100P, Nestin, Olig2, EMA, CD34, *ATRX*,

and *BRAF* V600E, the intensity of the staining was graded as negative, weak, moderate or strong, and the extent was scored as follows: (–) totally negative (1+) < 10% of tumor cells were positive (2+) 10–50% of tumor cells were positive, and (3+) more than 50% of tumor cells were positive. The Ki-67 labeling index (LI) was measured by Gunma-LI (Version 017; Japan Brain Tumor Reference Center, Gunma, Japan; available at [https://www.jbtrc.com/05\\_mib04.php](https://www.jbtrc.com/05_mib04.php)).

### Direct DNA sequencing for *BRAF*, *IDH1/2*, *TP53*, and *TERT* promoter mutations

DNA was extracted from paraffin sections as previously described [5] and was amplified and sequenced using the primers described previously [6–9] (Supplementary material, Table 1). PCR products were sequenced on a 3130xl Genetic Analyser (Applied Biosystems, Foster City, CA, USA) with the Big Dye Terminator v.1.1 Cycle Sequencing kit (Applied Biosystems) following standard procedures.

### Multiplex ligation-dependent probe amplification (MLPA) analysis

Copy number changes in the *CDKN2A/B* genes and chromosomes 1 and 19 were analyzed by multiplex ligation-dependent probe amplification (MLPA) as described previously [10].

## Results

### Clinical findings

The clinical features of each case are shown in Table 1. There were three females and one male. Their age of onset ranged from 3 to 41 years. The symptoms at presentation were headache, epilepsy, impaired consciousness, and facial spasm. Imaging features revealed similar findings in all cases. The tumors were located within the cerebral hemispheres; cases 1, 2, and 4 in the temporal lobe, and case 3 in the frontal lobe. The examination of gadolinium-enhanced T1-weighted magnetic resonance imaging (MRI) revealed well demarcated, nodular, contrast-enhancing lesions (Fig. 1a, e, g, j). In the recurrent tumor of case 2, heterogeneous enhancement was observed (Fig. 1e), whereas other cases were uniformly enhanced (Fig. 1a, g, j). Mild to-moderate edema surrounding the tumor was observed by T2-weighted MRI (Fig. 1b, f, h, k). No calcification was detected by computed tomography (Fig. 1c, i, l). Gross total resection and adjuvant chemoradiotherapy was performed in all patients. The follow-up period from the first operation ranged from 6 months to 28 years. Case 1 and 2 recurred

**Table 1** Clinical features

| Case    | Age | Sex | Location            | Images  | Symptoms   | Treatment  | Prognosis   |
|---------|-----|-----|---------------------|---|--|--|---|
| 1       | 46  | F   | Left temporal lobe  | Well demarcated mass on MR (CE) with focal edema; 6×4 cm                      | Epilepsy (41y.o.–), worsening headache (45y.o.–) | GTR, TMZ+ radio-chemotherapy                           | Recurrence (50y.o.), additional TMZ+ bevacizumab, alive for 7 years after first surgery |
| 2       | 3   | M   | Right temporal lobe | Hemorrhagic mass with heterogeneous enhancement (only CT scan)                | Impaired consciousness                           | GTR, radiotherapy (33Gy) + platinum-based chemotherapy | Recurrence (26y.o.)   |
| 2 (rec) | 26  |     | Right temporal lobe | Heterogeneously enhanced mass on MR (CE) with focal surrounding edema; 7×5 cm | Worsening headache                               | Resection TMZ+ bevacizumab                             | Alive for 5 years after second surgery  |
| 3       | 35  | F   | Right frontal lobe  | Well demarcated mass on MR (CE) with focal edema                              | Left facial spasm, headache                      | GTR+ radiotherapy+ TMZ                                 | Alive without recurrence for a year   |
| 4       | 32  | F   | Left temporal lobe  | Well demarcated mass on MR (CE) with focal edema                              | Headache   | GTR+ radiotherapy+ TMZ                                 | Alive without recurrence for 6 months   |

(rec) recurrence, M male, F female, y.o. years old, MR (CE) magnetic resonance with contrast enhancement, GTR gross total resection, TMZ temozolomide

23 years and 4 years after surgery, respectively, and all patients remained alive.

## Histological findings

A summary of the histological features is presented in Table 2. The ratio of cellular elements was assayed by visual estimation. All tumors had a solid growth pattern, with case 1, 2, and 3 showing a slight degree of infiltration to the adjacent brain parenchyma (Fig. 2a). In case 1, tumor extension along the parenchymal vessels and subpial accumulation of tumor cells was found (Fig. 2b). Medium-sized muscular arteries were entrapped in all cases, which suggests exophytic tumor growth into the subpial space.

All tumors were mainly composed of short spindle cells with round-to-oval nuclei and thick processes arranged in a fascicular pattern (Fig. 2c, d). Tumor cells with round cytoplasm and eccentric nuclei resembling epithelioid cells were occasionally seen (Fig. 2e). Large mono- or multi-nucleated pleomorphic cells were focally observed (Fig. 2f). A xanthomatous change was seen in case 2, but not conspicuous. Unlike conventional PXAs, eosinophilic granular bodies were found only in case 4 (Fig. 2g), and not apparent in the other cases. Reticulin fibers surrounding tumor cells were not observed by silver staining (Fig. 2h). Thick hyalinization of the vessel wall was found in case 2. Invasion of tumor cells into the vascular wall was seen in cases 3 and 4. Concentric fibrosis around blood vessels was observed in case 4. An increase of small

vessels was observed in all cases, and a small number of microvascular proliferations was detected in cases 1, 4, and the recurrent tumor of case 2. Relatively high levels of mitotic activity were seen with brisk mitotic figures ranging from 4 to 20 per 10 high-powered fields. Palisading necrosis was identified in cases 1 and 2 (Fig. 2i).

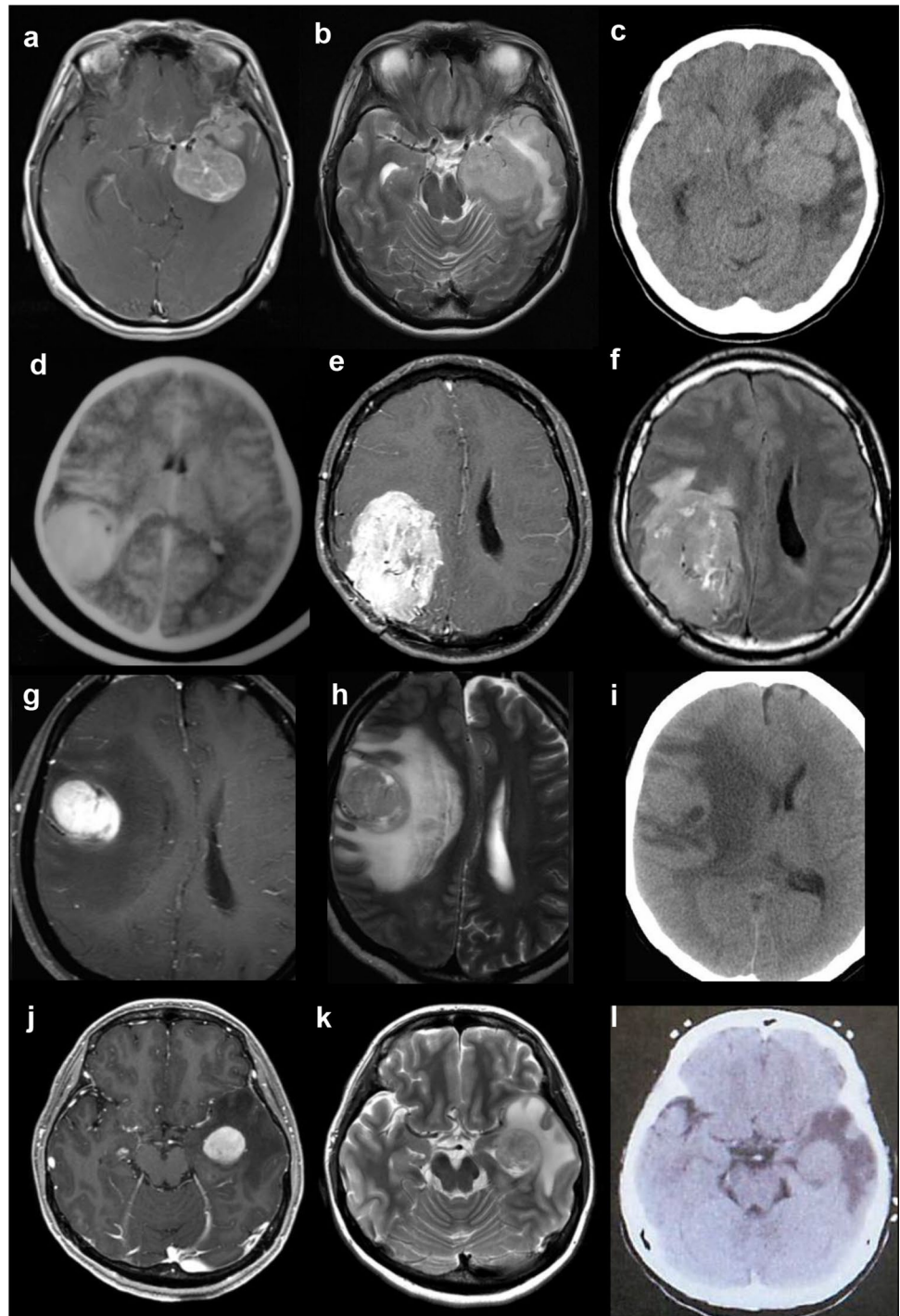
## Immunohistochemistry

The immunohistochemical findings are summarized in Table 2. All tumors were diffusely positive for S-100 protein, Olig2, and Nestin (Fig. 3a, b). Spindle-shaped cells were diffusely-to-sparsely positive for GFAP (Fig. 3c). CD34 immunoreactivity was not detected in any tumors. Loss of ATRX nuclear expression (Fig. 3d) and BRAF V600E immunoreactivity were observed in all cases (Fig. 3e). The Ki-67 labeling index ranged from 6.1 to 20.1% (Gunma-LI) (Fig. 3f).

## Genetic findings

A *BRAF V600E* mutation (Fig. 4a) and *CDKN2A/B* homozygous deletion (Fig. 4b) were detected in all cases by direct DNA sequencing or MLPA analysis. The *TERT* promoter and *IDH1/2* were intact. Direct sequencing of *TP53* revealed an R248Q mutation in case 3 (data not shown).

**Fig. 1** Radiological images of all four cases. Case 1 (a–c). **a** Gadolinium-enhanced T1-weighted image (Gd-T1WI) shows a well demarcated mass by a homogeneous enhancement. **b** A Fluid attenuated inversion recovery (FLAIR) image shows peritumoral edema. **c** Calcification was not detected by computed tomography (CT). Case 2 (d–f). **d** Contrast-enhanced CT image of the initial onset when the patient was 3 years old. A circumscribed mass with hemorrhage was located within the right temporal lobe. **e, f** The recurrent tumor at age 26. **e** Gd-T1WI shows a heterogeneously enhanced tumor in the same region. **f** A FLAIR image shows a limited edema around the tumor. Case 3 (g–i). **g** Gd-T1WI shows a homogeneously enhanced circumscribed tumor in the right frontal lobe. **h** Surrounding edema was observed in the FLAIR image. **i** Calcification was not obvious in the CT image. Case 4 (j–l). **j** Gd-T1WI shows a well demarcated mass with a homogeneous contrast enhancement within the left temporal lobe. **k** A FLAIR image shows surrounding edema. **l** No calcification was detected in the CT image



## Discussion

Alteration of *ATRX* in combination with *BRAF* V600E mutation and homozygous deletion of *CDKN2A/B* is very rare in gliomas. Among 681 grade I–IV gliomas in recent reports analyzing all three genetic mutations (82 adult glioblastomas, 148 grade II and III diffuse gliomas, 368 pediatric low- and high-grade gliomas, 24 PXAs and A-PXAs,

21 PAs and PAs with anaplasia, and 40 gangliogliomas) [11–23], only one case (histologically defined as anaplastic PXA) had all three mutations [11]. Five cases (three adult glioblastomas, one grade II diffuse glioma, and one pediatric GBM) had both *BRAF* V600E mutation and *ATRX* alteration [17, 21, 22]. However, detailed histology was not reported in these studies. Moreover, we searched 2395 grade I–IV gliomas in a public database of The Cancer Genome

**Table 2** Histological and immunohistochemical findings

| Case   | Spindle cell | Pleomorphic cell | Xanthic cell | Epithelioid cell | Piloid cell | Eosinophilic granular bodies | Necrosis    | MVP                  | Mitosis (/10HPF)     | Other findings   | Initial pathological diagnosis   |
|--------|--------------|------------------|--------------|------------------|-------------|------------------------------|-------------|----------------------|----------------------|--|----------------------------------|
| 1      | 90           | 5                | 0            | 5                | 0           | -                            | +           | +                    | 20                   | Accumulation of tumor cell to the subpial space and perivascular space | Glioblastoma                     |
| 2      | 60           | 10               | A few        | 5                | 25          | -                            | +           | -                    | 4                    | Microcalcification, Hyalinization of blood vessels                     | Anaplastic astrocytoma           |
| 2(rec) | 75           | 15               | A few        | 5                | 5           | -                            | +           | ±                    | 6                    | Hyalinization of blood vessels   | Localized high grade astrocytoma |
| 3      | 75           | 5                | 0            | 20               | 0           | ±                            | -           | -                    | 6                    | Localized high grade astrocytoma                                       | Localized high grade astrocytoma |
| 4      | 85           | A few            | 0            | 15               | 0           | +                            | -           | ±                    | 8                    | Prominent fibrosis around blood vessels                                | Localized high grade astrocytoma |
| Case   | GFAP         | S-100P           | Olig2        | nestin           | EMA         | CD34                         | mIDH1 R132H | ATRX                 | BRAFV600E            | p53  | Ki-67 LI (%)                     |
| 1      | 2+(s)        | 3+(s)            | 3+(s)        | 3+(m)            | 2+(m)       | -                            | -           | -                    | 3+(w)                | 1+(m)  | 20.1                             |
| 2      | 3+(s)        | 3+(s)            | 3+(m)        | 3+(s)            | ND          | -                            | ND          | Sample deterioration | Sample deterioration | Sample deterioration   | Sample deterioration             |
| 2(rec) | 2+(s)        | 3+(s)            | 3+(m)        | 3+(w)            | -           | -                            | ND          | -                    | 3+(m)                | 2+(m)  | 20                               |
| 3      | 1+(s)        | 2+(m)            | 3+(s)        | 3+(w)            | ND          | -                            | -           | -                    | 2+(w)                | ND   | 13                               |
| 4      | 3+(s)        | ND               | 3+(s)        | ND               | ND          | ND                           | -           | -                    | 3+(w)                | ND   | 6.1                              |

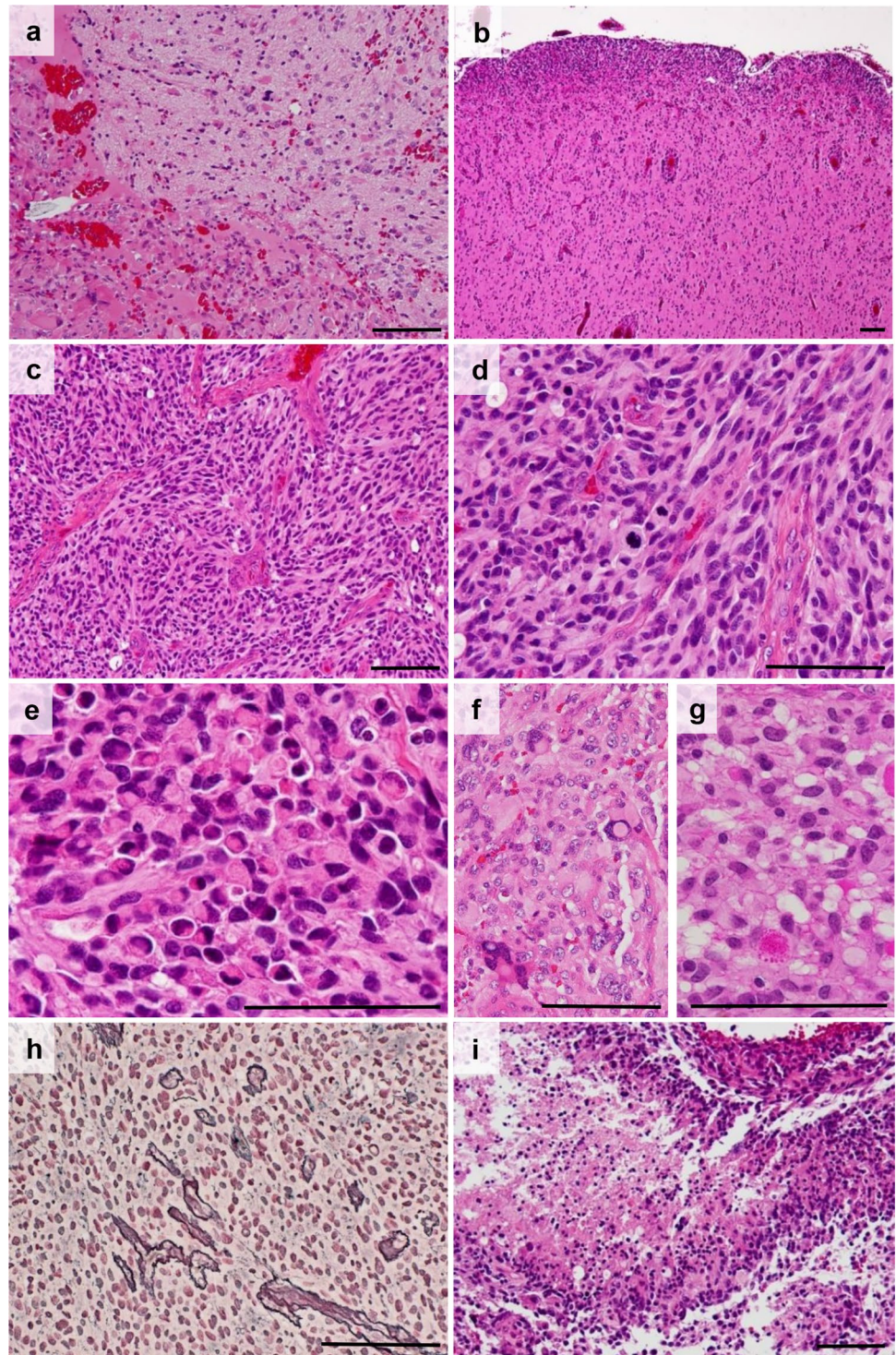
The visually estimated ratio of cellular elements (spindle, pleomorphic, xanthic, epithelioid and piloid cell) are described in percentage (%)

+ observed, ± only few observed, -not observed, /10HPF per 10 high power fields

The intensity and extent of the of immunopositive tumor cells were scored as follows: (w)—weak, (m)—moderate, (s)—strong, (-)totally negative, (1+) < 10%, (2+) 10–50%, (3+) > 50%

ND not done, GFAP glial fibrillary acidic protein, S-100P S-100 protein, S-100P S-100 protein, EMA epithelial membrane antigen, mIDH1 R132H mutated isocitrate dehydrogenase-1/2 R132Hs

**Fig. 2** Histological findings. **a** Case 2. The border between the tumor and adjacent brain parenchyma was relatively sharp. **b** Case 1. Accumulation of tumor cells in the subpial and perivascular space. **c, d** Case 1. Proliferation of spindle cells in a fascicular pattern with frequent mitosis is shown (arrow). **e** Case 1. Tumor cells with eccentric nuclei and round cytoplasm resembled epithelioid cells. **f** Case 3. Pleomorphic cells with bizarre nuclei. **g** Case 4. Eosinophilic granular bodies. **h** Case 1. Reticulin fibers were limited around blood vessels. **i** Case 2. Palisading necrosis. Scale bar = 50  $\mu$ m (a–i)



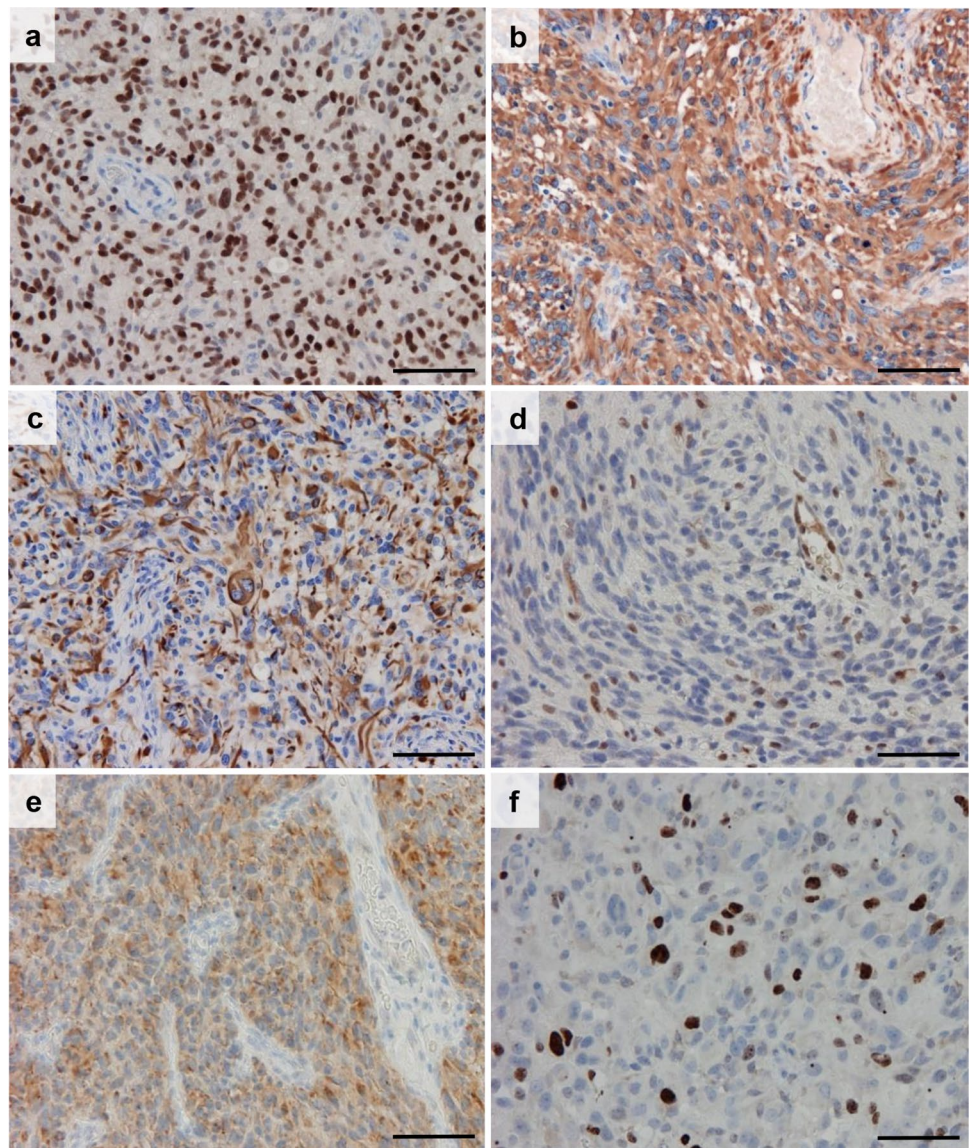
Atlas (TCGA) (cBioPortal for Cancer Genomics: <https://www.cbioportal.org>), but no tumor had all three alternations.

Reinhardt et al. recently reported a subset of histologically defined anaplastic pilocytic astrocytomas, showing a common DNA methylation profile [14]. These cases had a high frequency of shared mutations in the MAPK pathway, *CDKN2A/B*, and *ATRX*. Changes in the MAPK pathway

were mostly a *BRAF-KIAA1549* fusion or *NF1* mutation, and only one case had the *BRAF* V600E mutation. The case with the *BRAF* V600E mutation also had a *CDKN2A/B* alternation but no loss of *ATRX* immunoreactivity.

Both *BRAF* V600E mutation and *CDKN2A/B* homozygous deletion commonly occur in PXA and A-PXA.[24, 25], whereas concurrent *BRAF* V600E mutation, *CDKN2A/B*

**Fig. 3** Immunohistochemical findings. **a** Uninterrupted nuclear immunoreactivity for Olig2. **b** Cytoplasmic positivity for Nestin. **c** GFAP was positive in patches. **d** The tumor cell nuclei show loss of ATRX expression, but expression in vessels was retained. **e** Diffuse moderate positivity for BRAF V600E. **f** Relatively high Ki-67 labeling index is shown. **a, b:** case 4, **c, f:** case 3, **d, e:** case 1, Scale bar = 50  $\mu$ m (**a–f**)

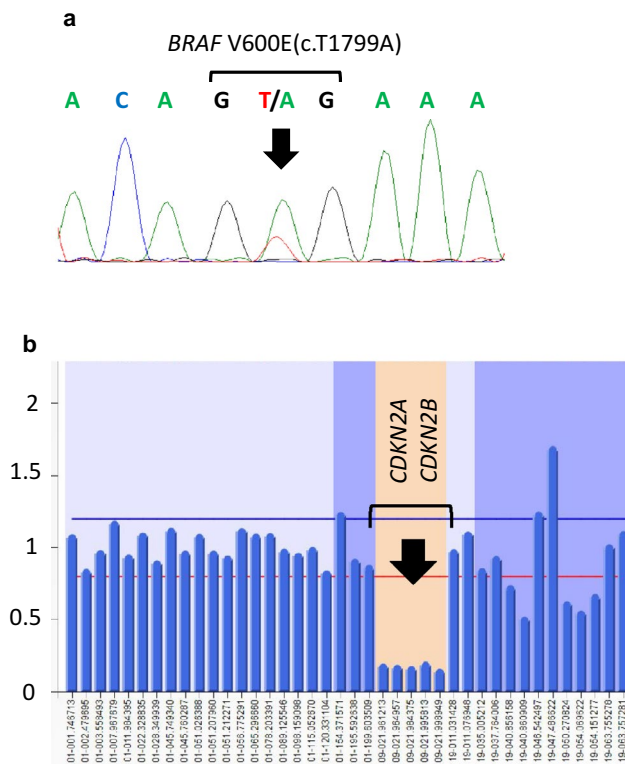


homozygous deletion, and *TERT* promoter mutation are frequent in epithelioid glioblastoma and A-PXA [10, 11]. In our current study, mutations in the *TERT* promoter were not detected, but loss of nuclear ATRX immunoreactivity was observed. This is interesting because *TERT* and *ATRX* are both telomere maintenance-related genes and are mutually exclusive in diffuse gliomas [26, 27]. Additionally, concurrent *TERT* promoter mutation, *CDKN2A/B* homozygous deletion, and *BRAF* V600E mutation are observed in a subset of melanomas and thyroid carcinomas [28, 29], but the combination of *ATRX* inactivation, *CDKN2/B* homozygous deletion, and *BRAF* V600E mutation has not been reported in other non-CNS neoplasms.

Histologically, all of our cases had a solid growth pattern, mainly composed of spindle-shaped cells with pleomorphism resembling PXA. A-PXA is sometimes misdiagnosed as glioblastoma or anaplastic astrocytoma because of

its proliferative capacity [30]. In fact, tumors histologically diagnosed as A-PXA are occasionally re-classified into glioblastoma using methylation analysis [31]. The diagnosis of A-PXA can be supported not only by a demarcated border, but also numerous eosinophilic granular bodies, pleomorphic cells, and xanthic cells. In our cases, these histological features were not prominent, which made it more difficult to differentiate glioblastoma. However, all our cases resulted in an excellent post-operative state with the longest survival of 28 years. Therefore, distinguishing these tumors from glioblastoma is important from a clinical aspect.

Previous studies of gliomas harboring *BRAF* V600E reported relatively favorable prognosis compared to those without this mutation [32–35]. Chi et al. reported five low-grade diffuse gliomas with the *BRAF* V600E mutation demonstrating a distinct phenotype [36]. Several of these cases had relatively well demarcated borders. One



**Fig. 4** Genetic findings. **a** Direct DNA sequencing of case 1. The *BRAF V600E* mutation (arrow) was detected. **b** Multiplex ligation-dependent probe amplification analysis of case 1. Homozygous deletion of *CDKN2A/B* is shown (arrow)

case had packed growth of spindle-shaped glial cells without typical characteristics of PXA, pilocytic astrocytoma, or ganglioglioma. The other case displayed proliferation of tumor cells in the subarachnoid and Virchow-Robin spaces. Although these cases were all low-grade, and the authors did not investigate mutations in *ATRX* and *CDKN2A/B*, the cases shared some clinicopathological features of our cases.

In summary, the present cases represent circumscribed high-grade astrocytomas with the unusual combination of *BRAF V600E*, *ATRX*, and *CDKN2A/B* alternations. These cases lacked typical histological characteristics of PXA and cannot be classified into the WHO classification of CNS tumors. Even with an anaplastic appearance, the prognosis was relatively favorable, and over-diagnosis of glioblastoma may be concerned. Alternations in *ATRX* and *BRAF V600E* can be readily detected by immunohistochemistry to verify the diagnosis. To understand whether these tumors are a distinct entity, or they are a subgroup of other tumors such as PXA, more cases are needed, and additional methylation analysis is worth investigating.

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## Compliance with ethical standards

**Conflict of interest** The authors declare that they have no conflict of interest.

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